



National Library  
of Canada

Bibliothèque nationale  
du Canada

Canadian Theses Service

Services des thèses canadiennes

Ottawa, Canada  
K1A 0N4

## CANADIAN THESES

## THÈSES CANADIENNES

### NOTICE

The quality of this microfiche is heavily dependent upon the quality of the original thesis submitted for microfilming. Every effort has been made to ensure the highest quality of reproduction possible.

If pages are missing, contact the university which granted the degree.

Some pages may have indistinct print especially if the original pages were typed with a poor typewriter ribbon or if the university sent us an inferior photocopy.

Previously copyrighted materials (journal articles, published tests, etc.) are not filmed.

Reproduction in full or in part of this film is governed by the Canadian Copyright Act, R.S.C. 1970, c. C-30.

**THIS DISSERTATION  
HAS BEEN MICROFILMED  
EXACTLY AS RECEIVED**

### AVIS

La qualité de cette microfiche dépend grandement de la qualité de la thèse soumise au microfilmage. Nous avons tout fait pour assurer une qualité supérieure de reproduction.

S'il manque des pages, veuillez communiquer avec l'université qui a conféré le grade.

La qualité d'impression de certaines pages peut laisser à désirer, surtout si les pages originales ont été dactylographiées à l'aide d'un ruban usé ou si l'université nous a fait parvenir une photocopie de qualité inférieure.

Les documents qui font déjà l'objet d'un droit d'auteur (articles de revue, examens publiés, etc.) ne sont pas microfilmés.

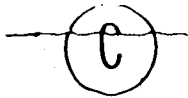
La reproduction, même partielle, de ce microfilm est soumise à la Loi canadienne sur le droit d'auteur, SRC 1970, c. C-30.

**LA THÈSE A ÉTÉ  
MICROFILMÉE TELLE QUE  
NOUS L'AVONS REÇUE**

THE UNIVERSITY OF ALBERTA

Systematics, phylogeny and biogeography of New World weevils  
traditionally of the tribe Cleonini (Coleoptera:  
Curculionidae; Cleoninae)

by



Robert Samuel Anderson

A THESIS

SUBMITTED TO THE FACULTY OF GRADUATE STUDIES AND RESEARCH  
IN PARTIAL FULFILMENT OF THE REQUIREMENTS FOR THE DEGREE  
OF Doctor of Philosophy

Department of Entomology

EDMONTON, ALBERTA

Spring 1987

Permission has been granted to the National Library of Canada to microfilm this thesis and to lend or sell copies of the film.

The author (copyright owner) has reserved other publication rights, and neither the thesis nor extensive extracts from it may be printed or otherwise reproduced without his/her written permission.

L'autorisation a été accordée à la Bibliothèque nationale du Canada de microfilmer cette thèse et de prêter ou de vendre des exemplaires du film.

L'auteur (titulaire du droit d'auteur) se réserve les autres droits de publication; ni la thèse ni de longs extraits de celle-ci ne doivent être imprimés ou autrement reproduits sans son autorisation écrite.

ISBN 0-315-37601-5

THE UNIVERSITY OF ALBERTA

RELEASE FORM

NAME OF AUTHOR Robert Samuel Anderson  
TITLE OF THESIS Systematics, phylogeny and biogeography  
of New World weevils traditionally of  
the tribe Cleonini (Coleoptera:  
Curculionidae; Cleoninae)  
DEGREE FOR WHICH THESIS WAS PRESENTED Doctor of Philosophy  
YEAR THIS DEGREE GRANTED Spring 1987

Permission is hereby granted to THE UNIVERSITY OF ALBERTA LIBRARY to reproduce single copies of this thesis and to lend or sell such copies for private, scholarly or scientific research purposes only.

The author reserves other publication rights, and neither the thesis nor extensive extracts from it may be printed or otherwise reproduced without the author's written permission.

(SIGNED) *Robert S. Anderson*

PERMANENT ADDRESS:  
*DEPARTMENT OF ENTOMOLOGY*  
*TEXAS A&M UNIVERSITY*  
*COLLEGE STATION, TX, USA*

DATED *December 1* ..... 1986

THE UNIVERSITY OF ALBERTA  
FACULTY OF GRADUATE STUDIES AND RESEARCH

The undersigned certify that they have read, and recommend to the Faculty of Graduate Studies and Research, for acceptance, a thesis entitled Systematics, phylogeny and biogeography of New World weevils traditionally of the tribe Cleonini (Coleoptera: Curculionidae; Cleoninae) submitted by Robert Samuel Anderson in partial fulfilment of the requirements for the degree of Doctor of Philosophy.

..... *Le. P. Hill* .....

Supervisor

..... *M. A. Wilson* .....

..... *D. A. Long* .....

..... *Ruth A. Street* .....

..... *W. H. ...* .....

External Examiner

Date..... *November 21, 1986* .....

## Abstract

New World species of Cleoninae traditionally placed in Cleonini are revised. Natural history and nomenclatural history of Cleoninae are briefly reviewed, a key to New World genera of Cleoninae is presented, and diagnostic characters of the subfamily are briefly discussed. Four genera are represented in the New World. These are: *Cleonis* Dejean (introduced; 1 species); *Stephanocleonus* Motschulsky (Holarctic, northern; 6 species); *Apleurus* Chevrolat (southwestern North America, arid lands; 8 species); and, *Cleonidius* Casey (eastern and western North America, semi-arid and arid lands; 19 species).

The following information is given in detail for each of *Cleonis*, *Stephanocleonus*, *Apleurus* (*Gibbostethus*) n. subg. (type species, *Dinocleus hystrix* Fall by monotypy), *Apleurus* (*Apleurus*), and *Cleonidius*: synonymic list, notes about synonymy, diagnosis, description, list of included species, and a discussion of phylogenetic relationships. Keys to species of North American *Stephanocleonus* and *Apleurus*, and world *Cleonidius*, are presented. The following information is given in detail for each species considered: synonymic list, notes about synonymy, problems in recognition, description, distribution, natural history information, and chorological relationships.

Only for species of *Apleurus* and *Cleonidius* are reconstructed phylogenies presented based on analyses of 31

Both *Apleurus* and *Cleonidius* are hypothesized to have entered North America in savanna or grassland habitats (or their precursors) across Beringia during late Eocene time. In North America, primary diversification has taken place in the arid lands of the southwestern United States although two species of *Cleonidius* are found in coastal regions of eastern North America. In both genera there appears to be an initial vicariance of lineages, on Compositae, in grassland or mesquite-grassland habitats between California, central Mexico, and the southern Rocky Mountains. Further speciation events in *Apleurus* appear due primarily to shifts to new, apotypic habitat types; host plant shifts, primarily between Leguminosae-Rosaceae, and within-habitat-type allopatric speciation appear to predominate as the causes of further speciation in *Cleonidius*.

Evolutionary trends in *Apleurus* and *Cleonidius* are discussed and patterns and predictions of biogeographic or evolutionary interest summarized and presented for examination for general applicability.

## Acknowledgments

First and foremost I wish to thank my supervisor, George E. Ball, for his criticism, advice and support during the course of this study. The academic environment for systematic studies at the University of Alberta, created largely by his enthusiasm and dedication, was a pleasure to work in. In addition, financial aid from Natural Sciences and Engineering Research Council grant A-1399, held by G.E. Ball, is also gratefully acknowledged.

For advice in selecting a thesis topic and for their assistance and guidance at various stages and with various aspects of the study, I thank D.R. Whitehead, C.W. O'Brien, E.L. Sleeper, H.R. Burke, W.E. Clark, and A.T. Howden; they were always available for consultation. R.T. Thompson was especially helpful in finding specimens and literature during my visit to the British Museum (Natural History). His help in solving nomenclatural problems and in demonstrating his novel preparatory techniques is also very gratefully acknowledged.

I also thank members of my advisory committee (M.V.H. Wilson, D.A. Craig, R.A. Stockey) and external thesis examiner (W.E. Clark) for their critical review of the penultimate draft of the thesis. For help in finding collecting localities in Arizona and for companionship in the field, I thank Scott McCleve of Douglas, Arizona. For permission to collect in Big Bend National Park, Texas, I thank C.M. Fleming of that same facility.



Finally, I thank all my fellow students at the University of Alberta (D.C. Currie, J.M. Cumming, G.A.P. Gibson, J.E. O'Hara, D.R. Maddison, J.S. Ashe, R.E. Roughley, J.-F. Landry, J.H. Acorn) for their friendship at home, companionship in the field, for their constant encouragement, but most of all, for expressing a keen interest in problems in weevil systematics and biogeography when called upon for their help and comments. Thanks are also due A. Sen for preparation of the photographic plates.

Additional financial support during the course of this study came from The University of Alberta, (Dissertation Fellowship, 1985-1986), and the Government of the Province of Alberta (Province of Alberta Scholarships, 1983-1984, 1984-1985).

I offer a last word of fond appreciation to my parents Robert Hamilton and Daisy, and to my wife Catherine; you were there when I needed you in ways you didn't realize and I can't explain.

## Table of Contents

Chapter	Page
Abstract .....	iv
Acknowledgments .....	vi
List of Tables .....	xvii
List of Figures .....	xxv
1. Introduction .....	1
2. Materials and Methods .....	3
2.1 Materials .....	3
2.2 Methods .....	6
2.2.1 Collection of specimens .....	6
2.2.2 Specimen preparation and examination .....	7
2.2.3 Structural terms .....	11
2.2.4 Illustrations .....	14
2.2.5 Measurements and descriptive statistics .....	16
2.2.6 Descriptive format .....	20
2.2.7 Criteria for recognition of species-group taxa .....	24
2.2.8 Criteria for recognition of genus-group taxa with notes on classification .....	26
3. Natural history of Cleoninae .....	30
4. Supraspecific taxa and included species .....	32
4.1 Subfamily Cleoninae .....	32
4.1.1 Historical review .....	32
4.1.2 Recognition of Cleoninae .....	33
4.1.3 Sex differentiation .....	34
4.1.4 Key to genera of New World Cleoninae .....	34
4.2 Genus <i>Cleonis</i> Dejean .....	39

4.2.1	Species accounts .....	47
4.2.1.1	<i>Cleonis pigra</i> (Scopoli) .....	47
4.3	Genus <i>Stephanocleonus</i> Motschulsky .....	52
4.3.1	Key to species of adult <i>Stephanocleonus</i> .....	60
4.3.2	Species accounts .....	63
4.3.2.1	<i>Stephanocleonus confusus</i> Anderson, new species .....	64
4.3.2.2	<i>Stephanocleonus cristaticollis</i> Csiki .....	75
4.3.2.3	<i>Stephanocleonus immaculatus</i> Anderson, new species .....	82
4.3.2.4	<i>Stephanocleonus parshus</i> Anderson, new species .....	91
4.3.2.5	<i>Stephanocleonus plumbeus</i> LeConte .....	102
4.3.2.6	<i>Stephanocleonus stenothorax</i> Anderson, new species .....	111
4.4	Genus <i>Apleurus</i> Chevrolat .....	119
4.4.1	Subgenus <i>Gibbostethus</i> Anderson, new subgenus .....	125
4.4.1.1	Species accounts .....	126
4.4.1.1.1	<i>Apleurus (Gibbostethus) hystrix</i> (Fall), new combination .....	126
4.4.2	Subgenus <i>Apleurus</i> Chevrolat .....	135
4.4.2.1	Key to species of adult <i>Apleurus (Apleurus)</i> .....	141
4.4.2.2	Species accounts .....	145
4.4.2.2.1	<i>Apleurus (Apleurus) aztecus</i> (Champion), new combination .....	145
4.4.2.2.2	<i>Apleurus (Apleurus) lutulentus</i> (LeConte), new combination .....	153
4.4.2.2.3	<i>Apleurus (Apleurus) porosus</i> (LeConte), new combination .....	166
4.4.2.2.4	<i>Apleurus (Apleurus) angularis</i> (LeConte), new combination .....	178

4.4.2.2.5	<i>Apleurus (Apleurus) jacobinus</i> (Casey), new combination .....	193
4.4.2.2.6	<i>Apleurus (Apleurus) albovestitus</i> (Casey), new combination .....	203
4.4.2.2.7	<i>Apleurus (Apleurus) saginatus</i> (Casey), new combination .....	222
4.5	Genus <i>Cleonidius</i> Casey .....	231
4.5.1	Key to species of adult <i>Cleonidius</i> .....	242
4.5.2	Species accounts .....	252
4.5.2.1	<i>Cleonidius poricollis</i> species group ....	252
4.5.2.1.1	<i>Cleonidius poricollis</i> (Mannerheim) ..	253
4.5.2.1.2	<i>Cleonidius eustictorrhinus</i> Anderson, new species .....	265
4.5.2.1.3	<i>Cleonidius modestus</i> (Mannerheim) ...	276
4.5.2.1.4	<i>Cleonidius subcylindricus</i> Casey ....	286
4.5.2.1.5	<i>Cleonidius longinasus</i> Anderson, new name .....	296
4.5.2.1.6	<i>Cleonidius texanus</i> (LeConte) .....	307
4.5.2.2	<i>Cleonidius americanus</i> species group ....	319
4.5.2.2.1	<i>Cleonidius americanus</i> Csiki .....	320
4.5.2.2.2	<i>Cleonidius frontalis</i> (LeConte) .....	331
4.5.2.2.3	<i>Cleonidius canescens</i> (LeConte) .....	343
4.5.2.2.4	<i>Cleonidius infrequens</i> Anderson, new species .....	355
4.5.2.2.5	<i>Cleonidius puberulus</i> (LeConte) .....	365
4.5.2.2.6	<i>Cleonidius collaris</i> (LeConte) .....	376
4.5.2.2.7	<i>Cleonidius notolomus</i> Anderson, new species .....	385
4.5.2.3	<i>Cleonidius lobigerinus</i> species group ...	396
4.5.2.3.1	<i>Cleonidius lobigerinus</i> Casey .....	397

4.5.2.3.2	<i>Cleonidius calandroides</i> (Randall)	..413
4.5.2.4	<i>Cleonidius boucardi</i> species group	.....423
4.5.2.4.1	<i>Cleonidius boucardi</i> (Chevrolat)	....424
4.5.2.4.2	<i>Cleonidius trivittatus</i> (Say)	.....436
4.5.2.4.3	<i>Cleonidius placidus</i> Csiki	.....449
4.5.2.4.4	<i>Cleonidius quadrilineatus</i> (Chevrolat)	.....460
4.5.2.5	<i>Incertae sedis</i>	.....471
4.5.2.5.1	<i>Cleonidius vibex</i> (Pallas), new combination	.....472
4.6	Adventitious taxa	.....478
4.7	Extant taxa erroneously placed in Cleoninae	.....478
4.8	Fossil material	.....479
5.	Evolutionary analysis	.....481
5.1	Introductory comments	.....481
5.2	Phylogeny	.....483
5.2.1	Methods of phylogeny reconstruction	.....483
5.2.2	Phylogenetic position of Cleoninae	.....486
5.2.3	Phylogenetic relationships among tribes of Cleoninae	.....489
5.2.4	Phylogenetic relationships of genera of New World Cleonini	.....490
5.2.4.1	Genus <i>Apleurus</i> Chevrolat	.....498
5.2.4.1.1	Character evolution	.....498
5.2.4.1.2	Phylogenetic relationships of included species	.....512
5.2.4.2	Genus <i>Cleonidius</i> Casey	.....515
5.2.4.2.1	Character evolution	.....515
5.2.4.2.2	Phylogenetic relationships of included species	.....526

5.3 Biogeography .....	530
5.3.1 Methods of biogeographic analysis .....	530
5.3.2 Genus <i>Apleurus</i> Chevrolat .....	535
5.3.2.1 Descriptive biogeography .....	535
5.3.2.1.1 Diversity patterns .....	535
5.3.2.1.2 Habitat associations .....	536
5.3.2.1.3 Plant associations .....	538
5.3.2.1.4 Altitudinal zonation .....	538
5.3.2.1.5 Seasonal occurrence .....	539
5.3.2.1.6 Co-existence .....	540
5.3.2.1.7 Wing length polymorphism .....	540
5.3.3 Genus <i>Cleonidius</i> Chevrolat .....	541
5.3.3.1 Descriptive biogeography .....	541
5.3.3.1.1 Diversity patterns .....	541
5.3.3.1.2 Habitat associations .....	543
5.3.3.1.3 Plant associations .....	544
5.3.3.1.4 Altitudinal zonation .....	545
5.3.3.1.5 Seasonal occurrence .....	546
5.3.3.1.6 Co-existence .....	546
5.3.3.1.7 Wing length polymorphism .....	547
5.3.4 Historical biogeography of New World <i>Apleurus</i> and <i>Cleonidius</i> .....	548
5.3.4.0.1 Introductory Comments--A Warning ...	548
5.3.4.0.2 Historical Explanations .....	550
5.4 Evolutionary trends and adaptations .....	572
5.4.1 Adaptive trends to arid conditions .....	572
5.4.2 Other evolutionary trends .....	575

5.5 Patterns and predictions .....	579
6. Conclusions and future research .....	582
7. Literature cited .....	585
8. Appendix 1 .....	604
9. Appendix 2 .....	612
10. Appendix 3 .....	738

LIST OF TABLES

Table	Page
1. Descriptive statistics for <i>S. confusus</i> based on eight males from Medicine Hat, Alberta.....	73
2. Descriptive statistics for <i>S. confusus</i> based on eight females from Medicine Hat, Alberta.....	74
3. Descriptive statistics for <i>S. cristaticollis</i> based on one male from Monarch, Alberta.....	80
4. Descriptive statistics for <i>S. cristaticollis</i> based on one female from Medicine Hat, Alberta.. .....	81
5. Descriptive statistics for <i>S. immaculatus</i> based on eight males from throughout the species range.....	89
6. Descriptive statistics for <i>S. immaculatus</i> based on eight females from throughout the species range.....	90
7. Descriptive statistics for <i>S. parshus</i> based on eight males from southern Manitoba and Ontario.. .....	100
8. Descriptive statistics for <i>S. parshus</i> based on eight females from southern Manitoba and Ontario.....	101
9. Descriptive statistics for <i>S. plumbeus</i> based on eight males from Rivière-au-Tonnerre, Québec.... .....	109



10.	Descriptive statistics for <i>S. plumbeus</i> based on eight females from Rivière-au-Tonherre, Québec.. .....	110
11.	Descriptive statistics for <i>S. stenothorax</i> based on eight males from Bluefish Caves Archaeological Site, Yukon Territory.....	117
12.	Descriptive statistics for <i>S. stenothorax</i> based on eight females from Bluefish Caves Archaeological Site, Yukon Territory.....	118
13.	Descriptive statistics for <i>A. hystrix</i> based on eight males from the El Segundo Sand Dunes, Los Angeles County, California.....	133
14.	Descriptive statistics for <i>A. hystrix</i> based on eight females from the El Segundo Sand Dunes, Los Angeles County, California.....	134
15.	Descriptive statistics for <i>A. aztecus</i> based on 13 males from throughout the species range...152	152
16.	Descriptive statistics for <i>A. aztecus</i> based on six females from throughout the species range... .....	153
17.	Descriptive statistics for <i>A. lutulentus</i> based on eight males from 2 mi. NE. Portal, Arizona... .....	165
18.	Descriptive statistics for <i>A. lutulentus</i> based on eight females from 2 mi. NE. Portal, Arizona. .....	166

19. Descriptive statistics for *A. porosus* based on  
seven males from 68.0 km. W. Ejido  
Viscaino, Baja California Sur, Mexico.....176

20. Descriptive statistics for *A. porosus* based on  
eight females from 68.0 km. W. Ejido  
Viscaino, Baja California Sur, Mexico.....177

21. Descriptive statistics for *A. angularis* based on  
15 males from Mercury, Nevada.....191

22. Descriptive statistics for *A. angularis* based on  
10 females from Mercury, Nevada.....192

23. Descriptive statistics for *A. jacobinus* based on  
eight males from Turlock, California.....202

24. Descriptive statistics for *A. jacobinus* based on  
eight females from Turlock, California.....203

25. Descriptive statistics for *A. albovestitus* based  
on eight males from Portal, Arizona.....220

26. Descriptive statistics for *A. albovestitus* based  
on eight females from Portal, Arizona.....221

27. Descriptive statistics for *A. saginatus* based on  
eight males from 5 mi. SW. Patagonia, Arizona...  
.....229

28. Descriptive statistics for *A. saginatus* based on  
eight females from 5 mi. SW. Patagonia, Arizona.  
.....230

29. Descriptive statistics for *C. porlcollis* based  
on eight males from Redondo, California.....263

30.	Descriptive statistics for <i>C. porlcollis</i> based on eight females from Redondo, California....	264
31.	Descriptive statistics for <i>C. eustictorrhinus</i> based on 13 males from throughout the species range.....	274
32.	Descriptive statistics for <i>C. eustictorrhinus</i> based on 14 females from throughout the species range.....	275
33.	Descriptive statistics for <i>C. modestus</i> based on eight males from Palm Springs, California....	284
34.	Descriptive statistics for <i>C. modestus</i> based on eight females from Palm Springs, California.....	285
35.	Descriptive statistics for <i>C. subcylindricus</i> based on seven males from throughout the species range.....	294
36.	Descriptive statistics for <i>C. subcylindricus</i> based on 17 females from throughout the species range.....	295
37.	Descriptive statistics for <i>C. longinasus</i> based on 10 males from Hemet Reservoir, Herkey Creek, and Idyllwild, California.....	305
38.	Descriptive statistics for <i>C. longinasus</i> based on 10 females from Hemet Reservoir, Herkey Creek, and Idyllwild, California.....	306

39. Descriptive statistics for *C. texanus* based on  
eight males from Mile 19, Hitchcock  
Highway, Santa Catalina Mountains, Arizona...317

40. Descriptive statistics for *C. texanus* based on  
eight females from Mile 19, Hitchcock  
Highway, Santa Catalina Mountains, Arizona...318

41. Descriptive statistics for *C. americanus* based  
on eight males from Mercy Hot Springs,  
California.....329

42. Descriptive statistics for *C. americanus* based  
on eight females from Mercy Hot Springs,  
California.....330

43. Descriptive statistics for *C. frontalis* based on  
eight males from Medicine Hat, Alberta.....341

44. Descriptive statistics for *C. frontalis* based on  
eight females from Medicine Hat, Alberta.....342

45. Descriptive statistics for *C. canescens* based on  
eight males from Denver, Colorado.....353

46. Descriptive statistics for *C. canescens* based on  
eight females from Denver, Colorado.....354

47. Descriptive statistics for *C. infrequens* based  
on two males from throughout the species range..  
.....363

48. Descriptive statistics for *C. infrequens* based  
on six females from throughout the species  
range.....364

49.	Descriptive statistics for <i>C. puberulus</i> based on six males from Colorado Springs, Colorado....	374
50.	Descriptive statistics for <i>C. puberulus</i> based on 10 females from Colorado Springs, Colorado...	375
51.	Descriptive statistics for <i>C. collaris</i> based on seven males from throughout the species range... .....	383
52.	Descriptive statistics for <i>C. collaris</i> based on six females from throughout the species range... .....	384
53.	Descriptive statistics for <i>C. notolomus</i> based on 11 males from throughout the species range...	394
54.	Descriptive statistics for <i>C. notolomus</i> based on 13 females from throughout the species range.... .....	395
55.	Descriptive statistics for <i>C. lobigerinus</i> based on eight males from 1mi. N. Little Lake, California.....	411
56.	Descriptive statistics for <i>C. lobigerinus</i> based on eight females from 1mi. N. Little Lake, California.....	412
57.	Descriptive statistics for <i>C. calandroides</i> based on eight males from Hampton, New Hampshire...	421
58.	Descriptive statistics for <i>C. calandroides</i> based --on eight females from Hampton, New Hampshire.... .....	422

59. Descriptive statistics for *C. boucardi* based on eight males from north of Tepatitlan, Jalisco, Mexico.....434

60. Descriptive statistics for *C. boucardi* based on eight females from 9 mi. W. Tepatitlan, Jalisco, Mexico.....435

61. Descriptive statistics for *C. trivittatus* based on eight males from Albuquerque, New Mexico.....447

62. Descriptive statistics for *C. trivittatus* based on eight females from Albuquerque, New Mexico...448

63. Descriptive statistics for *C. placidus* based on 10 males from Panoche, California.....458

64. Descriptive statistics for *C. placidus* based on eight females from Panoche, California.....459

65. Descriptive statistics for *C. quadrilineatus* based on eight males from southern Arizona localities.....469

66. Descriptive statistics for *C. quadrilineatus* based on eight females from southern Arizona localities.....470

67. Chorological relationships of *Cleonidius* species.....604

68. Character states in *Apleurus* species and related taxa.....605

69.	Character states in <i>Cleonidius</i> species and related taxa.....	606
70.	Plesiotypic and apotypic states of characters used in phylogenetic analysis of <i>Apleurus</i> species and related taxa.....	604
71.	Plesiotypic and apotypic states of characters used in phylogenetic analysis of <i>Cleonidius</i> species and related taxa.....	607

Figure	Page
1. Lateral view of head of <i>Microlarinus</i> species.....	611
2. Lateral view of head of <i>Rhinocyllus conicus</i> .....	611
3. Lateral view of head of <i>Larinus planus</i> .....	611
4. Lateral view of head of <i>Lixus scrobicollis</i> .....	611
5. Apex of tibia of <i>Cleonidius infrequens</i> .....	611
6. Apex of tibia of <i>Apleurus lutulentus</i> .....	611
7. Ventral view of prosternum of <i>Apleurus</i> species....	611
8. Ventral view of prosternum of <i>Cleonidius</i> species.....	611
.....	611
9. Lateral view of head of <i>Cleonis pigra</i> .....	613
10. Dorsal view of head of <i>Cleonis pigra</i> .....	613
11. Lateral view of internal sac of male <i>Cleonis pigra</i> ...	613
.....	613
12. Dorsal view of internal sac of male <i>Cleonis pigra</i> ....	613
.....	613
13. Lateral view of aedeagus of male <i>Cleonis pigra</i> ....	613
14. Ventral view of apical sclerite complex of	
internal sac of male <i>Cleonis pigra</i> .....	613
15. Lateral view of gonocoxite <sup>II</sup> and stylus of	
female <i>Cleonis pigra</i> .....	613
16. Spermatheca of female <i>Cleonis pigra</i> .....	613
17. Ventral view of sternum VIII of female <i>Cleonis</i>	
<i>pigra</i> .....	613
18. Ventral view of metatarsus of <i>Cleonis pigra</i> .....	613



19.	Dorsal habitus of <i>Cleonis pigra</i> .....	615
20.	Dorsal habitus of <i>Stephanocleonus confusus</i> .....	615
21.	Dorsal habitus of <i>Stephanocleonus immaculatus</i> .....	615
22.	Dorsal habitus of <i>Stephanocleonus plumbeus</i> .....	615
23.	Dorsal habitus of <i>Stephanocleonus stenothorax</i> .....	615
24.	Dorsal habitus of <i>Apleurus (Gibbostethus) hystrix</i> .....	615
25.	Head of <i>Stephanocleonus confusus</i> .....	617
	a. Lateral view.....	617
	b. Dorsal view.....	617
26.	Head of <i>Stephanocleonus cristaticeps</i> .....	617
	a. Lateral view.....	617
	b. Dorsal view.....	617
27.	Head of <i>Stephanocleonus immaculatus</i> .....	617
	a. Lateral view.....	617
	b. Dorsal view.....	617
28.	Head of <i>Stephanocleonus parshus</i> .....	617
	a. Lateral view.....	617
	b. Dorsal view.....	617
29.	Head of <i>Stephanocleonus plumbeus</i> .....	617
	a. Lateral view.....	617
	b. Dorsal view.....	617
30.	Head of <i>Stephanocleonus stenothorax</i> .....	617
	a. Lateral view.....	617
	b. Dorsal view.....	617
31.	Male genitalia of <i>Stephanocleonus confusus</i> .....	619
	a. Lateral view of aedeagus.....	619

	b. Ventral view of apex of aedeagus.....	619
	c. Ventral view of sternite of sternum VII.....	619
32.	Male genitalia of <i>Stephanocleonus cristaticollis</i> .....	
	.....	619
	a. Lateral view of aedeagus.....	619
	b. Ventral view of apex of aedeagus.....	619
	c. Ventral view of sternite of sternum VII.....	619
33.	Male genitalia of <i>Stephanocleonus immaculatus</i> .....	619
	a. Lateral view of aedeagus.....	619
	b. Ventral view of apex of aedeagus.....	619
	c. Ventral view of sternite of sternum VII.....	619
34.	Male genitalia of <i>Stephanocleonus parshus</i> .....	619
	a. Lateral view of aedeagus.....	619
	b. Ventral view of apex of aedeagus.....	619
	c. Ventral view of sternite of sternum VII.....	619
35.	Male genitalia of <i>Stephanocleonus plumbeus</i> .....	619
	a. Lateral view of aedeagus.....	619
	b. Ventral view of apex of aedeagus.....	619
	c. Ventral view of sternite of sternum VII.....	619
36.	Male genitalia of <i>Stephanocleonus stenothorax</i> .....	619
	a. Lateral view of aedeagus.....	619
	b. Ventral view of apex of aedeagus.....	619
	c. Ventral view of sternite of sternum VII.....	619
37.	Ventral view of sternum VIII of female	
	<i>Stephanocleonus confusus</i> .....	619
38.	Ventral view of sternum VIII of female	
	<i>Stephanocleonus cristaticollis</i> .....	619

39.	Ventral view of sternum VIII of female	
	<i>Stephanocleonus immaculatus</i> .....	619
40.	Ventral view of sternum VIII of female	
	<i>Stephanocleonus parshus</i> .....	619
41.	Ventral view of sternum VIII of female	
	<i>Stephanocleonus plumbeus</i> .....	619
42.	Ventral view of sternum VIII of female	
	<i>Stephanocleonus stenothorax</i> .....	619
43.	Internal sac of male <i>Stephanocleonus confusus</i> .....	621
	a. Lateral view.....	621
	b. Dorsal view.....	621
44.	Internal sac of male <i>Stephanocleonus cristaticeps</i> ..	
	.....	621
	a. Lateral view.....	621
	b. Dorsal view.....	621
45.	Internal sac of male <i>Stephanocleonus immaculatus</i> .....	
	.....	621
	a. Lateral view.....	621
	b. Dorsal view.....	621
46.	Internal sac of male <i>Stephanocleonus parshus</i> .....	621
	a. Lateral view.....	621
	b. Dorsal view.....	621
47.	Internal sac of male <i>Stephanocleonus plumbeus</i> .....	621
	a. Lateral view.....	621
	b. Dorsal view.....	621
48.	Internal sac of male <i>Stephanocleonus stenothorax</i> .....	
	.....	621

a.	Lateral view.....	621
b.	Dorsal view.....	621
49.	Hubbs-Hubbs diagram illustrating variation among specimens of <i>Stephanocleonus</i> species; length of elytra (LE1): a, males; b, females....	623
50.	Hubbs-Hubbs diagram illustrating variation among specimens of <i>Stephanocleonus</i> species; form of elytra (WE1M/LE1): a, males; b, females...	625
51.	Hubbs-Hubbs diagram illustrating variation among specimens of <i>Stephanocleonus</i> species; relative length of pronotum compared to elytra (LP/LE1): a, males; b, females.....	625
52.	Hubbs-Hubbs diagram illustrating variation among specimens of <i>Stephanocleonus</i> species; form of rostrum (WRA/LR): a, males; b, females....	627
53.	Hubbs-Hubbs diagram illustrating variation among specimens of <i>Stephanocleonus</i> species; relative width of frons compared to rostrum (WRA/WF): a, males; b, females.....	629
54.	Lateral view of gonocoxite II and stylus of female <i>Stephanocleonus</i> .....	631
55.	Spermatheca of female <i>Stephanocleonus</i> .....	631
56.	Genitalia of <i>Chromoderus fasciatus</i> .....	631
a.	Lateral view of aedeagus of male.....	631
b.	Lateral view of internal sac of male.....	631
c.	Dorsal view of internal sac of male.....	631

d.	Ventral view of apical sclerite complex of internal sac of male.....	631
e.	Ventral view of sternum VIII of female.....	631
57.	Genitalia of <i>Cnemodontus llimpidus</i> .....	631
a.	Lateral view of aedeagus of male.....	631
b.	Lateral view of internal sac of male.....	631
c.	Dorsal view of internal sac of male.....	631
d.	Ventral view of apical sclerite complex of internal sac of male.....	631
e.	Ventral view of sternum VIII of female.....	631
58.	Lateral view of mesosternum of <i>Apleurus</i> ( <i>Gibbostethus</i> ) <i>hystrix</i> .....	631
59.	Lateral view of mesosternum of <i>Apleurus</i> ( <i>Apleurus</i> ) <i>lutulentus</i> .....	<del>631</del>
60.	Spermatheca of female <i>Apleurus</i> .....	631
61.	Ventral view of ligula of <i>Apleurus</i> ( <i>Gibbostethus</i> ) <i>hystrix</i> .....	631
62.	Ventral view of ligula of <i>Apleurus</i> ( <i>Apleurus</i> ) <i>lutulentus</i> .....	631
63.	Ventral view of ligula of <i>Apleurus</i> ( <i>Apleurus</i> ) <i>saginata</i> .....	631
64.	Ventral view of maxilla of <i>Apleurus</i> ( <i>Apleurus</i> ) <i>saginata</i> .....	631
65.	Ventral view of maxilla of <i>Apleurus</i> ( <i>Gibbostethus</i> ) <i>hystrix</i> .....	631
66.	Tarsal claws of <i>Apleurus angularis</i> .....	<del>631</del>
67.	Tarsal claws of <i>Apleurus jacobinus</i> .....	631

68.	Dorsal habitus of <i>Apleurus aztecus</i> .....	633
69.	Dorsal habitus of <i>Apleurus lutulentus</i> .....	633
70.	Dorsal habitus of <i>Apleurus porosus</i> .....	633
71.	Dorsal habitus of <i>Apleurus jacobinus</i> .....	633
72.	Dorsal habitus of <i>Apleurus angularis</i> (3 mi. N Baker, Nevada).....	633
73.	Dorsal habitus of <i>Apleurus angularis</i> (Colorado Springs, Colorado).....	633
74.	Dorsal habitus of <i>Apleurus angularis</i> (Baboquivaria Mountains, Arizona).....	633
75.	Dorsal habitus of <i>Apleurus albovestitus</i> (18 mi. SW Mendota, California).....	635
76.	Dorsal habitus of <i>Apleurus albovestitus</i> (16 mi. S. Vidal, California).....	635
77.	Dorsal habitus of <i>Apleurus albovestitus</i> (Virgin, Utah).....	635
78.	Dorsal habitus of <i>Apleurus albovestitus</i> (Phoenix, Arizona).....	635
79.	Dorsal habitus of <i>Apleurus albovestitus</i> (Yuma, Arizona).....	635
80.	Dorsal habitus of <i>Apleurus saginatus</i> .....	635
81.	Head of <i>Apleurus hystrix</i> .....	637
	a. Lateral view.....	637
	b. Dorsal view.....	637
82.	Head of <i>Apleurus aztecus</i> .....	637
	a. Lateral view.....	637
	b. Dorsal view.....	637

83.	Head of <i>Apleurus lutulentus</i> .....	637
	a. Lateral view.....	637
	b. Dorsal view.....	637
84.	Head of <i>Apleurus porosus</i> .....	637
	a. Lateral view.....	637
	b. Dorsal view.....	637
85.	Head of <i>Apleurus angularis</i> .....	637
	a. Lateral view.....	637
	b. Dorsal view.....	637
86.	Head of <i>Apleurus jacobinus</i> .....	637
	a. Lateral view.....	637
	b. Dorsal view.....	637
87.	Head of <i>Apleurus saginatus</i> .....	637
	a. Lateral view.....	637
	b. Dorsal view.....	637
88.	Head of <i>Apleurus albovestitus</i> .....	637
	a. Lateral view.....	637
	b. Dorsal view.....	637
89.	Female genitalia of <i>Apleurus hystrix</i> .....	639
	a. Ventral view of sternum VIII.....	639
	b. Lateral view of gonocoxite II and stylus.....	639
90.	Female genitalia of <i>Apleurus aztecus</i> .....	639
	a. Ventral view of sternum VIII.....	639
	b. Lateral view of gonocoxite II and stylus.....	639
91.	Female genitalia of <i>Apleurus lutulentus</i> .....	639
	a. Ventral view of sternum VIII.....	639
	b. Lateral view of gonocoxite II and stylus.....	639

92.	Female genitalia of <i>Apleurus porosus</i> .....	639
	a. Ventral view of sternum VIII.....	639
	b. Lateral view of gonocoxite II and stylus.....	639
93.	Female genitalia of <i>Apleurus angularis</i> .....	639
	a. Ventral view of sternum VIII.....	639
	b. Lateral view of gonocoxite II and stylus.....	639
94.	Female genitalia of <i>Apleurus jacobinus</i> .....	639
	a. Ventral view of sternum VIII.....	639
	b. Lateral view of gonocoxite II and stylus.....	639
95.	Female genitalia of <i>Apleurus albovestitus</i> .....	639
	a. Ventral view of sternum VIII.....	639
	b. Lateral view of gonocoxite II and stylus.....	639
96.	Female genitalia of <i>Apleurus saginatus</i> .....	639
	a. Ventral view of sternum VIII.....	639
	b. Lateral view of gonocoxite II and stylus.....	639
97.	Male genitalia of <i>Apleurus hystrix</i> .....	641
	a. Lateral view of internal sac.....	641
	b. Dorsal view of internal sac.....	641
	c. Ventral view of apical sclerite complex of internal sac.....	641
	d. Lateral view of aedeagus.....	641
	e. Ventral view of aedeagus.....	641
	f. Ventral view of sternite of sternum VII.....	641
98.	Male genitalia of <i>Apleurus aztecus</i> .....	641
	a. Lateral view of internal sac.....	641
	b. Dorsal view of internal sac.....	641
	c. Lateral view of aedeagus.....	641



	d. Ventral view of apical sclerite complex of internal sac.....	641
99.	Male genitalia of <i>Apleurus lutulentus</i> .....	641
	a. Lateral view of internal sac.....	641
	b. Dorsal view of internal sac.....	641
	c. Lateral view of aedeagus.....	641
	d. Ventral view of apical sclerite complex of internal sac.....	641
100.	Male genitalia of <i>Apleurus angularis</i> .....	643
	a. Lateral view of internal sac.....	643
	b. Dorsal view of internal sac.....	643
	c. Lateral view of aedeagus.....	643
	d. Ventral view of apical sclerite complex of internal sac.....	643
101.	Male genitalia of <i>Apleurus jacobinus</i> .....	643
	a. Lateral view of internal sac.....	643
	b. Dorsal view of internal sac.....	643
	c. Lateral view of aedeagus.....	643
	d. Ventral view of apical sclerite complex of internal sac.....	643
	e. Ventral view of sternite of sternum VII.....	643
102.	Male genitalia of <i>Apleurus porosus</i> .....	643
	a. Lateral view of internal sac.....	643
	b. Dorsal view of internal sac.....	643
	c. Lateral view of aedeagus.....	643
103.	Male genitalia of <i>Apleurus albovestitus</i> .....	645
	a. Lateral view of internal sac.....	645

	b. Dorsal view of internal sac.....	645
	c. Lateral view of aedeagus.....	645
	d. Ventral view of apical sclerite complex of internal sac.....	645
	e. Ventral view of sternite of sternum VII.....	645
104.	Male genitalia of <i>Apleurus saghnatus</i> .....	645
	a. Lateral view of internal sac.....	645
	b. Dorsal view of internal sac.....	645
	c. Lateral view of aedeagus.....	645
	d. Ventral view of apical sclerite complex of internal sac.....	645
105.	Hubbs-Hubbs diagram illustrating variation among specimens of <i>Apleurus</i> species; length of elytra (LE1): a, males; b, females.....	647
106.	Hubbs-Hubbs diagram illustrating variation among specimens of <i>Apleurus</i> species; form of rostrum (WRA/LR): a, males; b, females.....	649
107.	Hubbs-Hubbs diagram illustrating variation among specimens of <i>Apleurus</i> species; form of elytra (WE1M/LE1): a, males; b, females.....	651
108.	Hubbs-Hubbs diagram illustrating variation among specimens of <i>Apleurus</i> species; width of frons compared to width at apex of rostrum (WRA/WF): a, males; b, females.....	651
109.	Hubbs-Hubbs diagram illustrating variation among specimens of <i>Apleurus</i> species; form of pronotum (WPT/WPB): a, males; b, females.....	653

110.	Hubbs-Hubbs diagram illustrating variation among specimens of <i>Apleurus</i> species; length of pronotum compared to length elytra (LP/LE1); a, males; b, females.....	653
111.	Male genitalia of <i>Cylindropterus luxeri</i> .....	655
	a. Lateral view of internal sac.....	655
	b. Dorsal view of internal sac.....	655
112.	Male genitalia of <i>Lixus (Lixoglyptus) spartii</i> .....	655
	a. Lateral view of internal sac.....	655
	b. Dorsal view of internal sac.....	655
113.	Ventral view of sternum VIII of female <i>Cylindropterus luxeri</i> .....	655
114.	Ventral view of sternum VIII of female <i>Lixus (Lixoglyptus) spartii</i> .....	655
115.	Internal sac of male <i>Cleonidius</i> .....	655
	a. Lateral view.....	655
	b. Dorsal view.....	655
116.	Spermatheca of female <i>Cleonidius</i> .....	655
117.	Lateral view of gonocoxite PI and stylus of <i>Cleonidius</i> .....	655
118.	Dorsal habitus of <i>Cleonidius poricollis</i> .....	657
119.	Dorsal habitus of <i>Cleonidius eustictorrhinus</i> .....	657
120.	Dorsal habitus of <i>Cleonidius modestus</i> .....	657
121.	Dorsal habitus of <i>Cleonidius subcylindricus</i> .....	657
122.	Dorsal habitus of <i>Cleonidius longinasus</i> .....	657
123.	Dorsal habitus of <i>Cleonidius texanus</i> .....	657
124.	Dorsal habitus of <i>Cleonidius americanus</i> .....	659

125.	Dorsal habitus of <i>Cleonidius frontalis</i> .....	659
126.	Dorsal habitus of <i>Cleonidius canescens</i> .....	659
127.	Dorsal habitus of <i>Cleonidius infrequens</i> .....	659
128.	Dorsal habitus of <i>Cleonidius puberulus</i> .....	659
129.	Dorsal habitus of <i>Cleonidius collaris</i> .....	659
130.	Dorsal habitus of <i>Cleonidius notolomus</i> .....	659
131.	Dorsal habitus of <i>Cleonidius lobigerinus</i> .....	661
132.	Dorsal habitus of <i>Cleonidius calandroides</i> .....	661
133.	Dorsal habitus of <i>Cleonidius boucardi</i> .....	661
134.	Dorsal habitus of <i>Cleonidius trivittatus</i> .....	661
135.	Dorsal habitus of <i>Cleonidius placidus</i> .....	661
136.	Dorsal habitus of <i>Cleonidius quadrilineatus</i> .....	661
137.	<i>Cleonidius porlcollis</i> .....	663
	a. Lateral view of head.....	663
	b. Dorsal view of head.....	663
138.	<i>Cleonidius eustictorrhinus</i> .....	663
	a. Lateral view of head.....	663
	b. Dorsal view of head.....	663
139.	<i>Cleonidius modestus</i> .....	663
	a. Lateral view of head.....	663
	b. Dorsal view of head.....	663
140.	<i>Cleonidius subcylindricus</i> .....	663
	a. Lateral view of head.....	663
	b. Dorsal view of head.....	663
141.	<i>Cleonidius longinasus</i> .....	663
	a. Lateral view of heads of male and female.....	663
	b. Dorsal view of head of female.....	663

142.	<i>Cleonidius texanus</i> .....	663
	a. Lateral view of head.....	663
	b. Dorsal view of head.....	663
143.	<i>Cleonidius americanus</i> .....	665
	a. Lateral view of head.....	665
	b. Dorsal view of head.....	665
	c. Lateral view of pro-tibia of female.....	665
144.	<i>Cleonidius frontalis</i> .....	665
	a. Lateral view of head.....	665
	b. Dorsal view of head.....	665
	c. Lateral view of pro-tibia of female.....	665
145.	<i>Cleonidius canescens</i> .....	665
	a. Lateral view of head.....	665
	b. Dorsal view of head.....	665
146.	<i>Cleonidius infrequens</i> .....	665
	a. Lateral view of head.....	665
	b. Dorsal view of head.....	665
147.	<i>Cleonidius puberulus</i> .....	665
	a. Lateral view of head.....	665
	b. Dorsal view of head.....	665
148.	<i>Cleonidius collaris</i> .....	665
	a. Lateral view of head.....	665
	b. Dorsal view of head.....	665
149.	<i>Cleonidius notolomus</i> .....	665
	a. Lateral view of head.....	665
	b. Dorsal view of head.....	665
150.	<i>Cleonidius lobigerinus</i> .....	667

	a. Lateral view of head.....	667
	b. Dorsal view of head.....	667
151.	<i>Cleonidius calandroides</i> ./.....	667
	a. Lateral view of head.....	667
	b. Dorsal view of head.....	667
152.	<i>Cleonidius boucardi</i> .....	667
	a. Lateral view of head.....	667
	b. Dorsal view of head.....	667
153.	<i>Cleonidius trivittatus</i> .....	667
	a. Lateral view of head.....	667
	b. Dorsal view of head.....	667
154.	<i>Cleonidius placidus</i> .....	667
	a. Lateral view of head.....	667
	b. Dorsal view of head.....	667
155.	<i>Cleonidius quadrilineatus</i> .....	667
	a. Lateral view of head.....	667
	b. Dorsal view of head.....	667
156.	<i>Cleonidius vibex</i> .....	667
	a. Lateral view of head.....	667
	b. Dorsal view of head.....	667
157.	Ventral view of sternum VIII of female	
	<i>Cleonidius poricollis</i> .....	669
158.	Ventral view of sternum VIII of female	
	<i>Cleonidius eustictorrhinus</i> .....	669
159.	Ventral view of sternum VIII of female	
	<i>Cleonidius modestus</i> .....	669

160. Ventral view of sternum VIII of female  
*Cleonidius subcylindricus*.....669

161. Ventral view of sternum VIII of female  
*Cleonidius longinasus*.....669

162. Ventral view of sternum VIII of female  
*Cleonidius texanus*.....669

163. Ventral view of sternum VIII of female  
*Cleonidius americanus*.....669

164. Ventral view of sternum VIII of female  
*Cleonidius frontalis*.....669

165. Ventral view of sternum VIII of female  
*Cleonidius canescens*.....669

166. Ventral view of sternum VIII of female  
*Cleonidius infrequens*.....669

167. Ventral view of sternum VIII of female  
*Cleonidius puberulus*.....669

168. Ventral view of sternum VIII of female  
*Cleonidius notolomus*.....669

169. Ventral view of sternum VIII of female  
*Cleonidius collaris*.....669

170. Ventral view of sternum VIII of female  
*Cleonidius lobigerinus*.....669

171. Ventral view of sternum VIII of female  
*Cleonidius calandroides*.....669

172. Ventral view of sternum VIII of female  
*Cleonidius boucardi*.....669

173.	Ventral view of sternum VIII of female	
	<i>Cleonidius trivittatus</i> .....	669
174.	Ventral view of sternum VIII of female	
	<i>Cleonidius placidus</i> .....	669
175.	Ventral view of sternum VIII of female	
	<i>Cleonidius quadrilineatus</i> .....	669
176.	Ventral view of sternum VIII of female	
	<i>Cleonidius vibex</i> .....	669
177.	<i>Cleonidius porticollis</i> .....	671
	a. Lateral view of internal sac of male.....	671
	b. Dorsal view of internal sac of male.....	671
	c. Lateral view of aedeagus of male.....	671
178.	<i>Cleonidius eustictorrhinus</i> .....	671
	a. Lateral view of internal sac of male.....	671
	b. Dorsal view of internal sac of male.....	671
	c. Lateral view of aedeagus of male.....	671
179.	<i>Cleonidius modestus</i> .....	671
	a. Lateral view of internal sac of male.....	671
	b. Dorsal view of internal sac of male.....	671
	c. Lateral view of aedeagus of male.....	671
180.	<i>Cleonidius subcylindricus</i> .....	671
	a. Lateral view of internal sac of male.....	671
	b. Dorsal view of internal sac of male.....	671
	c. Lateral view of aedeagus of male.....	671
181.	<i>Cleonidius longinusus</i> .....	671
	a. Lateral view of internal sac of male.....	671
	b. Dorsal view of internal sac of male.....	671



	c. Lateral view of aedeagus of male.....	671
182.	<i>Cleonidius texanus</i> .....	671
	a. Lateral view of internal sac of male.....	671
	b. Dorsal view of internal sac of male.....	671
	c. Lateral view of aedeagus of male.....	671
183.	<i>Cleonidius americanus</i> .....	673
	a. Lateral view of internal sac of male.....	673
	b. Dorsal view of internal sac of male.....	673
	c. Lateral view of aedeagus of male.....	673
184.	<i>Cleonidius frontalis</i> .....	673
	a. Lateral view of internal sac of male.....	673
	b. Dorsal view of internal sac of male.....	673
	c. Lateral view of aedeagus of male.....	673
185.	<i>Cleonidius canescens</i> .....	673
	a. Lateral view of internal sac of male.....	673
	b. Dorsal view of internal sac of male.....	673
	c. Lateral view of aedeagus of male.....	673
186.	<i>Cleonidius infrequens</i> .....	673
	a. Lateral view of internal sac of male.....	673
	b. Dorsal view of internal sac of male.....	673
	c. Lateral view of aedeagus of male.....	673
187.	<i>Cleonidius puberulus</i> .....	673
	a. Lateral view of internal sac of male.....	673
	b. Dorsal view of internal sac of male.....	673
	c. Lateral view of aedeagus of male.....	673
188.	<i>Cleonidius notolomus</i> .....	673
	a. Lateral view of internal sac of male.....	673

	c. Lateral view of aedeagus of male.....	673
189.	<i>Cleonidius collaris</i> .....	673
	a. Lateral view of internal sac of male.....	673
	b. Dorsal view of internal sac of male.....	673
	c. Lateral view of aedeagus of male.....	673
190.	<i>Cleonidius lobigerinus</i> .....	675
	a. Lateral view of internal sac of male.....	675
	b. Dorsal view of internal sac of male.....	675
	c. Lateral view of aedeagus of male.....	675
191.	<i>Cleonidius calandroides</i> .....	675
	a. Lateral view of internal sac of male.....	675
	b. Dorsal view of internal sac of male.....	675
	c. Lateral view of aedeagus of male.....	675
192.	<i>Cleonidius boucardi</i> .....	675
	a. Lateral view of internal sac of male.....	675
	b. Dorsal view of internal sac of male.....	675
	c. Lateral view of aedeagus of male.....	675
193.	<i>Cleonidius trivittatus</i> .....	675
	a. Lateral view of internal sac of male.....	675
	b. Dorsal view of internal sac of male.....	675
	c. Lateral view of aedeagus of male.....	675
194.	<i>Cleonidius placidus</i> .....	675
	a. Lateral view of internal sac of male.....	675
	b. Dorsal view of internal sac of male.....	675
195.	<i>Cleonidius quadrilineatus</i> .....	675
	a. Lateral view of internal sac of male.....	675

	c. Lateral view of aedeagus of male.....	675
196.	<i>Cleonidius vibex</i> .....	675
	a. Lateral view of internal sac of male.....	675
	b. Dorsal view of internal sac of male.....	675
	c. Lateral view of aedeagus of male.....	675
197.	Hubbs-Hubbs diagram illustrating variation among specimens of <i>Cleonidius</i> species; length of elytra (LE1): a, males; b, females.....	677
198.	Hubbs-Hubbs diagram illustrating variation among specimens of <i>Cleonidius</i> species; form of rostrum (WRA/LR): a, males; b, females.....	679
199.	Hubbs-Hubbs diagram illustrating variation among specimens of <i>Cleonidius</i> species; width of frons compared to width at apex of rostrum (WRA/WF): a, males; b, females.....	681
200.	Hubbs-Hubbs diagram illustrating variation among specimens of <i>Cleonidius</i> species; length of pronotum compared to length of elytra (LP/LE1): a, males; b, females.....	683
201.	Hubbs-Hubbs diagram illustrating variation among specimens of <i>Cleonidius</i> species; form of elytra (WELM/LE1): a, males; b, females.....	683
202.	Collection localities for <i>Stephanocleonus</i> <i>immaculatus</i> .....	685
203.	Collection localities for <i>Stephanocleonus parshus</i> .... .....	685

	<i>confusus</i> (dots) and <i>Stephanocleonus</i>	
	<i>stenothorax</i> (stars).....	687
205.	Collection localities for <i>Stephanocleonus</i>	
	<i>plumbeus</i> (dots) and <i>Stephanocleonus</i>	
	<i>cristicollis</i> (stars).....	687
206.	Collection localities for <i>Apleurus saginatus</i> .....	689
207.	Collection localities for <i>Apleurus lutulentus</i> .....	689
208.	Collection localities for <i>Apleurus jacobinus</i> .....	689
209.	Collection localities for <i>Apleurus albovestitus</i> ...	689
210.	Collection localities for <i>Apleurus aztecus</i>	
	(dots) and <i>Apleurus porosus</i> (stars).....	691
211.	Collection localities for <i>Apleurus angularis</i> .....	691
212.	Collection localities for <i>Apleurus hystrix</i> .....	691
213.	Collection localities for <i>Cleonis pigra</i> .....	691
214.	Collection localities for <i>Cleonidius longinatus</i> ...	693
215.	Collection localities for <i>Cleonidius subcylindricus</i> ..	
	.....	693
216.	Collection localities for <i>Cleonidius eustictorrhinus</i> .	
	.....	693
217.	Collection localities for <i>Cleonidius modestus</i> .....	693
218.	Collection localities for <i>Cleonidius texanus</i> .....	693
219.	Collection localities for <i>Cleonidius poricollis</i> ...	693
220.	Collection localities for <i>Cleonidius trivittatus</i> ...	
	.....	695
221.	Collection localities for <i>Cleonidius boucardi</i> .....	695

	(dots) and <i>Cleonidius infrequens</i> (stars).....	695
223.	Collection localities for <i>Cleonidius americanus</i> (dots) and <i>Cleonidius collaris</i> (stars).....	695
224.	Collection localities for <i>Cleonidius quadrilineatus</i> .....	695
225.	Collection localities for <i>Cleonidius notolomus</i> (dots) and <i>Cleonidius puberulus</i> (stars).....	695
226.	Collection localities for <i>Cleonidius frontalis</i> ....	697
227.	Collection localities for <i>Cleonidius canescens</i> ....	697
228.	Collection localities for <i>Cleonidius lobigerinus</i> .....	697
229.	Collection localities for <i>Cleonidius calandroides</i> ....	697
230.	Numbers of species of <i>Apleurus</i> in 5° intervals....	699
231.	Numbers of species of <i>Cleonidius</i> in 5° intervals.....	701
232.	Hubbs-Hubbs diagram illustrating variation among specimens of <i>Apleurus</i> species; elevation.....	703
233.	Hubbs-Hubbs diagram illustrating variation among specimens of <i>Cleonidius</i> species; elevation...	703
234.	Reconstructed phylogeny based on primary characters illustrating hypothesized phylogenetic relationships among <i>Apleurus</i> species and related out-groups.....	705

secondary characters illustrating  
hypothesized phylogenetic relationships  
among *Apleurus* species and related out-groups...  
.....707

236. Reconstructed phylogeny based on primary  
characters illustrating hypothesized  
phylogenetic relationships among *Cleonidius*  
species and related out-groups.....709

237. Reconstructed phylogeny based on primary and  
secondary characters illustrating  
hypothesized phylogenetic relationships  
among *Cleonidius* species and related out-groups.  
.....711

238. Habitat associations of *Apleurus* species.....713

239. Plant associations of *Apleurus* species.....715

240. Habitat associations of *Cleonidius* species.....717

241. Plant associations of *Cleonidius* species.....719

242. Origin and initial diversification of *Apleurus*  
in North America: *Apleurus hystrix*, *A.*  
*aztecus*, and *A. lutulentus*-*A. porosus*-*A.*  
*angularis*-*A. jacobinus*-*A. albovestitus*-*A.*  
*saginitus* lineage.....721

243. Diversification of *Apleurus* in North America:  
*Apleurus lutulentus* and *A. porosus*-*A.*  
*angularis*-*A. jacobinus*-*A. albovestitus*-*A.*  
*saginitus* lineage.....723

	<i>Apleurus angularis</i> , <i>A. jacobinus</i> and <i>A. porosus</i> .	723
245.	Diversification of <i>Apleurus</i> in North America: <i>Apleurus albovestitus</i> and <i>A. saginatus</i> .....	723
246.	Origin and initial diversification of <i>Cleonidius</i> in North America: <i>Cleonidius poricollis</i> group, <i>C. americanus</i> group, <i>C. lobigerinus</i> group, and <i>C. boucardi</i> group.....	725
247.	Diversification of <i>Cleonidius</i> in North America: <i>Cleonidius poricollis</i> - <i>C. eustictorrhinus</i> , <i>C. modestus</i> , and <i>C. subcylindricus</i> - <i>C.</i> <i>longinusus</i> - <i>C. texanus</i> lineages.....	727
248.	Diversification of <i>Cleonidius</i> in North America: <i>Cleonidius poricollis</i> and <i>C. eustictorrhinus</i> ....	727
249.	Diversification of <i>Cleonidius</i> in North America: <i>Cleonidius subcylindricus</i> , <i>C. longinusus</i> and <i>C. texanus</i> .....	727
250.	Diversification of <i>Cleonidius</i> in North America: <i>Cleonidius americanus</i> - <i>C. frontalis</i> , <i>C.</i> <i>canescens</i> - <i>C. infrequens</i> - <i>C. puberulus</i> - <i>C.</i> <i>collaris</i> - <i>C. notolomus</i> lineages.....	729
251.	Diversification of <i>Cleonidius</i> in North America: <i>Cleonidius americanus</i> and <i>C. frontalis</i> .....	729
252.	Diversification of <i>Cleonidius</i> in North America: <i>Cleonidius puberulus</i> and <i>C. infrequens</i> .....	729

	) <i>Cleonidius collaris</i> and <i>C. notolomus</i> .....	729
254.	Diversification of <i>Cleonidius</i> in North America: <i>Cleonidius boucardi</i> and <i>C. trivittatus</i> - <i>C.</i> <i>placidus</i> - <i>C. quadrilineatus</i> lineage.....	731
255.	Diversification of <i>Cleonidius</i> in North America: <i>Cleonidius trivittatus</i> and <i>C. placidus</i> - <i>C.</i> <i>quadrilineatus</i> lineage.....	731
256.	Diversification of <i>Cleonidius</i> in North America: <i>Cleonidius placidus</i> and <i>C. quadrilineatus</i> .....	731
257.	Geographic ranges of species in the <i>Cleonidius</i> <i>poricollis</i> species group.....	733
258.	Geographic ranges of species in the <i>Cleonidius</i> <i>americanus</i> species group.....	733
259.	Geographic ranges of species in the <i>Cleonidius</i> <i>boucardi</i> species group.....	735



J.B.S. Haldane, in an oft repeated quote, once stated that the creator must have held a special fondness for beetles for he made so many of them. To this I would hasten to add that he must also have had a special predilection towards weevils, for the Curculionidae are the largest single family of organisms known. More than 44883 species had been described as of 1971 (O'Brien and Wibmer 1978) and countless others no doubt await discovery and formal scientific description. In so far as numerous species await description, some clearer understanding has yet to be achieved of those taxa already formally recognized. Crowson (1955), commenting on this very problem, noted that "A satisfactory resolution of the Curculionidae into subfamilies and tribes is probably the largest and most important outstanding problem in the higher classification of Coleoptera particularly as a great number of the species are more or less seriously injurious to economically valuable crops."

But to begin to make progress towards achieving this goal, we must start at some lower level, come to some understanding of a subfamily, tribe or genus, and then proceed to the larger and more complex problem. This I have chosen to do by conducting a study of the North American species that have traditionally been placed in the tribe Cleonini of the Cleoninae. The group had not been revised since LeConte and Horn (1876) and subsequently Casey (1891),

systematists as problematical and in need of work. Not only were species very difficult to reliably identify, but classification of the North American fauna in a world perspective had not been attempted. Use of the group in applied problems (e.g. as biological control agents of pest weeds), or to address theoretical problems of general evolutionary or biogeographic importance, was not yet possible.

This study will alleviate those problems of identification and provide more detailed and reliable definitions of species and species-group taxa than those previously presented. It will also address the classification of the North American fauna in a world perspective and represent a start at a consistent scheme for use by scientists in all faunal regions. Phylogenetic and biogeographic patterns can then be recognized and resultant process-hypotheses formulated so as to be available for general consideration and applicability by students from a wide array of disciplines, not just those interested in Curculionoidea.

## 2.1 Materials

This study was based on examination of 9498 adult specimens of *Cleonidius* (3586); *Apleurus* (5627); North American *Stephanocleonus* (224), and *Cleonis pligra* (61), borrowed from various collections, both institutional and private, and including material I personally collected in the western United States and Canada. The following abbreviations represent these collections, both individual and institutional, and their respective curators.

AMNH American Museum of Natural History, Central Park West at 79th Street, New York, New York, 10024, U.S.A.; L.H. Herman.

ASUT Arizona State University, Tempe, Arizona, 85281, U.S.A.; F.F. Hasbrouck.

BMNH British Museum (Natural History), Cromwell Road, London, SW7 5BD, England; R.T. Thompson and C.H.C. Lyal.

CASC California Academy of Sciences, Department of Entomology, Golden Gate Park, San Francisco, California, 94118, U.S.A.; D.H. Kavanaugh.

CDFA California Department of Food and Agriculture, 1220 N Street, Sacramento, California, 95814, U.S.A.; T.N. Seeno.

CNCI Canadian National Collection of Insects, Arachnids and Nematodes, Biosystematics Research Institute, Research Branch, Ottawa, Ontario, K1A 0C6, Canada; D.E. Bright.

CUIC Cornell University Insect Collections, Department of Entomology, Cornell University, Ithaca, New York, 14853, U.S.A.; J.K. Liebherr.

University, Indianapolis, Indiana, 46207, U.S.A.; C. O'Brien.

- DLCC Douglas L. Caldwell Collection, Chemical Lawn R & D Center, P.O. Box 395, Milford Center, Ohio, 43045, U.S.A.; D.L. Caldwell.
- ELSC Elbert L. Sleeper Collection, Department of Biology, California State University at Long Beach, Long Beach, California, 90801, U.S.A.; E.L. Sleeper.
- FMNH Field Museum of Natural History, Roosevelt Road and Lake Shore Drive, Chicago, Illinois, 60605, U.S.A.; L.E. Watrous.
- HAHC Henry F. and Anne T. Howden Collection, Department of Biology, Carleton University, Ottawa, Ontario, K1S 5B6, Canada; A.T. Howden.
- JLCC John L. and Alberta F. Carr Collection, 23 Dalrymple Green NW, Calgary, Alberta, T3A 1Y2, Canada; J.L. Carr.
- JVMC John V. Matthews Collection, Energy, Mines, and Resources Canada, 601 Booth Street, Ottawa, Ontario, K1A 0E8, Canada; J.V. Matthews.
- LACM Los Angeles County Museum of Natural History, 900 Exposition Boulevard, Los Angeles, California, 90007, U.S.A.; R.R. Snelling.
- MCZC Museum of Comparative Zoology, Harvard University, Cambridge, Massachusetts, 02138, U.S.A.; A.F. Newton, Jr.
- MNHP Musée Nationale d'Histoire Naturelle, Entomologie, 45 Rue Buffon, Paris, France; N. Berti.
- MSUC Montana State University, Department of Biology, Bozeman, Montana, 59717, U.S.A.; S. Rose.
- NMSU New Mexico State University, Department of Biology, Las Cruces, New Mexico, 88003, U.S.A.; J.R. Zimmerman.
- NRS Naturhistoriska Riksmuseet, Sektionen för entomologi, S-104 05, Stockholm, Sweden; P.I. Persson.
- NSDA Nevada State Department of Agriculture, 350 Capitol Hill Avenue, P.O. Box 11100, Reno, Nevada, 89510, U.S.A.; R.C. Bechtel.
- OSDA Oregon State Department of Agriculture, 635 Capitol Street N.E., Salem, Oregon, 97310, U.S.A.; R.L.

- OSUC Oregon State University, Department of Entomology,  
Corvallis, Oregon, 97331, U.S.A.; G.L. Peters.
- PUL Purdue University, Department of Entomology, West  
Lafayette, Indiana, 47907, U.S.A.; A. Provonsha.
- RSAC denotes specimens in my own personal collection.
- SBMN Santa Barbara Museum of Natural History, 2559 Puesta  
del Sol Road, Santa Barbara, California, 93105, U.S.A.;  
S. Miller and L. Marx.
- SMC Scott McCleve Collection, 2210 13th Street, Douglas,  
Arizona, 85607, U.S.A.; S. McCleve.
- TAMU Texas Agricultural and Mechanical University,  
Department of Entomology, College Station, Texas,  
77843, U.S.A.; H.R. Burke.
- UASM University of Alberta, Strickland Museum, Department of  
Entomology, Edmonton, Alberta, T6G 2E3, Canada; G.E.  
Ball.
- UAT University of Arizona, Department of Entomology, Tucson,  
Arizona, 85721, U.S.A.; F.G. Werner.
- UCBC University of California, Division of Entomology and —  
Parasitology, Berkeley, California, 94720, U.S.A.; J.A.  
Chemsak.
- UCMC University of Colorado Museum, Campus Box 218, Boulder,  
Colorado; U.N. Lanham.
- UIM University of Idaho, Department of Entomology, Moscow,  
Idaho, 83843, U.S.A.; W.F. Barr.
- UNSM University of Nebraska State Museum, Research and  
Systematics Collections, W-436 Nebraska Hall, Lincoln,  
Nebraska, 68588, U.S.A.; B.C. Ratcliffe.
- USNM United States National Museum (Natural History),  
Washington D.C., 20560, U.S.A.; D.R. Whitehead.
- WECC Wayne E. Clark Collection, Department of Entomology,  
Auburn University, Auburn, Alabama, 36849, U.S.A.; W.E.  
Clark.
- WSU Washington State University, Department of Entomology,  
Pullman, Washington, 99164, U.S.A.; R.S. Zack.

perspective required knowledge of structural diversity of taxa from other regions of the world. In view of this, representatives of type-species (where designated) of 50 of a total of 77 described genus-group names of the tribe Cleonini (not including New World taxa or junior homonyms where replacement names are available) from the British Museum (Natural History) were studied. In instances where a type-species has not been designated for a given genus-group name, representatives of a species initially placed in that taxon (and thus available for subsequent designation as type-species) were selected for study. These taxa are listed in Appendix 3.

## 2.2 Methods

### 2.2.1 Collection of specimens

Most specimens of Cleoninae collected by me were acquired simply by placing a standard entomological beating sheet underneath foliage of a particular plant and then striking the plant sharply with a sturdy stick in a downward motion over the sheet. Weevils then on the plant will feign death and drop downward onto the white sheet where they are easily seen and captured. Specimens were killed in vials containing fine ethyl acetate-soaked wood chips. A few

were handpicked from bare soil. Deciduous shrubs in oak-pinyon-juniper woodlands and annual or perennial woody shrubs along dry stream beds, desert washes, or various lotic waterways were found to be the most productive places to concentrate collecting efforts.

### 2.2.2 Specimen preparation and examination

A Wild M5 stereo dissecting microscope was used for routine examination of external structural features of dried adult specimens. Dissections were made of mouthparts and internal genitalia and these were examined with both a Wild M5 dissecting microscope and a Leitz SM-Lux compound brightfield microscope. Special preparation procedures for specific characters examined in detail are given following. For such characters, note is made in the text of the number of specimens that were examined in detail.

Genitalia.-- For each species, a variable number of male and female individuals were selected from various localities for study of internal genitalic structure. Individuals of both sexes were first softened by immersion in hot distilled water, to which a few drops of liquid soap had been added. Dissection consisted of insertion of the apices of a pair of fine forceps into the sternal suture separating the base of the abdomen from the metasternum. Careful application of pressure resulted in separation of

tissues. Repeated examinations were made of the abdomen during this process to determine the suitable degree of treatment. The abdomen was then transferred to a 5-10% acetic acid-alcohol solution to neutralize KOH activity, and then to distilled water. Genitalia were then removed from the abdomen and placed in 95% ethanol and then into glycerine for subsequent examination. The abdomen was dried and mounted on a card placed on the specimen pin.

No special techniques were used in the examination of female genitalia; however, for the internal sac of the aedeagus of males to be adequately examined, it was necessary that it be everted and inflated. This was best accomplished using an unpublished technique devised by Mr. Richard T. Thompson of the British Museum (Natural History).

Most practical for large specimens, this technique consists of eversion of the internal sac following softening and treatment with KOH by careful insertion of a hooked minuten pin mounted on the end of a wooden probe into the apex of the aedeagus. Slow and numerous repeated tugs on the sac, which is visible through the wall of the aedeagus, worked best and could be continued until the sac was completely everted, the apical sclerite complex exposed and apical. Attempts to extract the sac rapidly and using only one or a few insertions of the minuten pin frequently damaged the sac such that inflation was not possible.



viscosity, such as glycerine, is recommended). The base of the aedeagus was placed at the end of a narrow piece of flexible rubber tubing and the rubber adjacent to the base of the aedeagus pinched with a pair of flat-tipped forceps placed firmly in a hole in a rubber stopper attached to a sturdy metal stand. A continuous force was maintained on the forceps by placement of a small elastic band around the forceps near their apices such that they needed to be forced open using another pair of forceps. It was important that the aedeagus was held firmly in place in the apex of the tubing, but that the base of the aedeagus was not sealed by the forceps. The other end of the piece of tubing was attached to a syringe filled with the liquid in which the aedeagus was immersed. The syringe was also placed in a hole in a rubber stopper attached to a separate sturdy metal stand. Gentle pressure on the syringe forced the liquid out the apex of the tubing but ideally only through the aedeagus and thus the internal sac. As liquid passed into the sac it became inflated and remained so for short periods of time while the pressure exerted by the syringe continued to exceed the force limiting inflation of the sac. Drawings were thus made of the sac by repeated application of slight pressure on the syringe. Various views of the sac were obtained by manipulation of the aedeagus in the apex of the tubing and repetition of the entire procedure. Proper

syringe did not inflate the sac because either the forceps were sealing the base of the aedeagus or, too much liquid was passing out the tubing adjacent to the base of the aedeagus and not through it. In this latter instance the aedeagus was expelled from the apex of the tubing because it was not held firmly in place by the pressure exerted onto the tubing by the forceps. Repeated attempts and manipulation of the location of pressure of the forceps were required for good results.

Following examination, internal genitalia were placed in glycerine in a microvial attached to the specimen pin.

Mouthparts.-- Heads were removed from individuals softened as noted above, and placed in 5-10% KOH (aqueous) for a variable amount of time as required for removal of soft tissues. The head was then transferred to a 10% hydrogen peroxide-95% ethanol solution to bleach cuticle to facilitate examination of surface features with a brightfield microscope. Following removal to a 95% ethanol-distilled water solution, maxillary palpi and the prementum were carefully removed from the head by severing connections to the apex of the rostrum with a hooked minuten pin mounted on a wooden probe. These structures were then transferred to glycerine on a depression slide for detailed examination using a brightfield microscope. The head was air dried and mounted on a card placed on the specimen pin.

mouthpart examination was only carried out for species of *Apleurus*. Initial surveys of diversity of mouthpart structure in *Cleonidius* and *Stephanocleonus* species did not prove promising and were discontinued.

### 2.2.3 Structural terms

Structural terms follow those of Kissinger (1970) except as noted in Anderson (1984b). Structure of the internal sac of the aedeagus of males in Cleoninae has not been previously studied and an informal nomenclature has been developed for use (Fig. 115). Future detailed studies on other Cleoninae should result in a more formal nomenclature based on apparent homology between structures in a more comprehensive survey of structural diversity.

Ambiguity in descriptors used in association with various structural terms is a problem worthy of comment. Objectively delimited character states (e.g. presence/absence, red/black, three setae/one seta, etc.) are ideal for taxonomic and phylogenetic purposes because there is no ambiguity in assigning a state to a particular taxon. Most character states however, are subjectively expressed (e.g. elongate-narrow/robust, large/small, deep/shallow, etc.). For these latter characters to be maximally useful in a taxonomic sense as criteria for identification or

taxa. I have attempted to do this for those character states that are taxonomically important in this study but which are not augmented in the text in a quantitative manner or accompanied with figures. Characters and their states in which there is little ambiguity involved are not considered.

Vestiture.-- I term all body vestiture of apparent non-sensory primary function as scales. Body vestiture is important both phylogenetically and taxonomically. Pronotal scale patterns serve to distinguish genera, elytral scale patterns species of *Stephanocleonus*, and species groups and species of *Cleonidius*. Abdominal scale patterns are important in separating some species of *Apleurus*.

Scales vary from flat or appressed, to recumbent, sub-erect and erect. I use the terms flat or appressed for scales at an angle of more or less  $0^\circ$  relative to body surface; recumbent,  $0^\circ-45^\circ$ ; sub-erect,  $46^\circ-89^\circ$ ; and erect, more or less  $90^\circ$ .

Shape and size of the scale is important in determining macroscopic patterns of surface vestiture. Small, fine scales obscure less of the underlying dark cuticle and thus the ground color of the cuticle predominates in the pattern; larger, robust scales obscure more cuticle and impart a predominantly whitish color to the pattern. Density of the scales is not as important as size and shape in determining the macroscopic pattern.

concepts of "hair". Length of "hair-like scales" is important both taxonomically and phylogenetically. Members of some species clearly lack sub-erect or erect surface vestiture; members of most species have short sub-erect or erect vestiture that is not more than approximately twice the length of the largest appressed scales on that particular body part. Members of few species have moderately long sub-erect or erect vestiture that is between twice and three times the length of the largest appressed scale. Finally, members of few species have long vestiture, greater than three times the length of the largest appressed scale. Sub-erect or erect vestiture is abraded to various degrees on some specimens causing problems in identification.

Density of all vestiture and size of scales are indicated qualitatively with reference to other taxa.

Punctuation.-- This character is important both phylogenetically and taxonomically. Three aspects of punctuation are described; size, density and depth. States of all three aspects are part of a continuum from small, sparse or shallow, to large, dense and deep.

Density of punctuation is defined on the basis of the relationship between diameter of a puncture and the distance between it and adjacent punctures. Sparse punctures are those with diameters less than the distance between punctures; moderately dense are those with diameters more or

punctures.

Size and depth of punctation are more difficult to assess in an objective manner. Shallow punctures are those with the walls of the puncture at about a  $45^\circ$  angle or less; deep are those with walls between  $45^\circ$  and  $90^\circ$ . Moderately deep are those with the state difficult to assign as either deep or shallow. Small punctures are those where a very rough estimate of the number of punctures, if lined side by side spanning that body part, would number more than 50; large are those where the number is less than 25. Estimates between 25 and 50 are considered moderately large.

Punctation of the head, rostrum and pronotum are most important and most frequently referred to in keys and descriptions.

#### 2.2.4 Illustrations

With the specimen placed in glycerine, all drawings of genitalia and mouthparts were made on paper with the aid of a camera lucida attached to Wild M5 or Leitz SM-Lux microscopes at various magnifications and later traced, then inked, onto mylar drafting film for reproduction. Drawings of other structural features were similarly made of dried pinned specimens with the aid of a camera lucida attached to a Wild M5 microscope. Details of surface vestiture and

structures are drawn to the same scale (in most instances) and orientation to facilitate comparisons. Only taxonomically or phylogenetically valuable structures are illustrated; thus, comparisons of certain attributes of the structures for the purposes of identification may be misleading. Such attributes although illustrated for aesthetic completeness are not discussed in the text.

Distribution maps are presented for all species with the exception of *C. vibex*. Special maps and illustrations were prepared to illustrate general ecological and historical biogeographic patterns, evolutionary trends or adaptations, etc., or as accompanying visual simplifications of various discussions in the text.

Hubbs-Hubbs diagrams representing intraspecific and interspecific variation in certain mensural species attributes were prepared separately for each sex. Range, mean, 1.5 standard deviations (SD) on either side of the mean, and two standard errors (SE) on either side of the mean are plotted. Choice of 1.5 SD and 2 SE follows Whitehead (1971) as noted in the section on descriptive statistics.

Features illustrated are as follows: LE1, WRA/LR, WRA/WF, WE1M/LE1, LP/LE1 for *Stephanocleonus* (Figs. 49-53) and *Cleonidius* (Figs. 197-201); and, LE1, WRA/LR, WRA/WF, WE1M/LE1, LP/LE1, WPT/WPB for *Apleurus* (Figs. 105-110).

were also prepared to illustrate the altitudinal distributions of *Apleurus* (Fig. 232) and *Cleonidius* (Fig. 233) species. Samples used for preparation of Hubbs-Hubbs diagrams minimally include those samples used in preparation of descriptive statistics tables. In those instances where measurements were taken of individuals from more than the sample used in the descriptive statistics tables, these other samples constituted part of an examination for possible intraspecific variation in mensural features.

#### 2.2.5 Measurements and descriptive statistics

**Measurements.**--- Measurements were made on pinned specimens at various magnifications with the aid of an ocular micrometer in a Wild M5 microscope. Measurements selected for detailed study and for presentation were those that upon initial survey revealed differences between species of potential taxonomic or phylogenetic value. For this reason not all measurements or ratios presented were taken on species of all genera.

These measurements and their abbreviations are:

- LR, length of rostrum measured in lateral aspect from anterior margin of eye to point of articulation of mandible;
- WF, minimum width of frons between eyes in dorsal aspect;
- WRA, maximum width of rostrum measured across apex in dorsal



of pronotum measured across base in dorsal aspect; WPT, width of pronotum measured across apical one-third or one-quarter at tubercles (if present) or immediately posterior to sub-apical constriction in dorsal aspect; WELH, width of elytra measured across base at humeri with elytra contiguous throughout length at suture in dorsal aspect; WELM, width of elytra measured across midlength with elytra contiguous throughout length at suture in dorsal aspect; LE1, length of elytra measured along suture from apex of scutellum to apex of elytra in dorsal aspect; LEy, maximum length of eye measured in lateral aspect; WEy, maximum width of eye measured in lateral aspect.

Some of these measurements were used to obtain ratios which serve to express the form of a particular part of the body. These ratios are; WEy/LEy, shape of eye; WPB/LP, form of pronotum; WRA/LR, relative length of rostrum; WF/WRA, relative separation of eyes; WELM/LE1, form of elytra; WP/LE1, relative length of elytra; and, WPT/WPB, position of maximum width of pronotum.

Descriptive statistics.--- A comprehensive set of descriptive statistics is given for one sample of each species treated. Statistical analysis is less comprehensive for those species where there was not adequate material. Sexes are treated separately and ideally each sample includes 8 members of each sex collected at the same

or, for the purposes of more comprehensive statistical analysis, if markedly less than 8 specimens are represented, are composed of a number of specimens collected from a more extensive geographic area. Least preferred samples are composed of all specimens of the species available for study. Mensural data treated in the descriptive statistics section include only those measurements and ratios given previously. Number of specimens, geographic locality of the specimens, range, mean, 1.5 standard deviations (SD), and 2 standard errors (SE) are given where number of specimens permits. In instances where too few specimens were available for meaningful statistical treatment, values for SD and SE are not given. Number of standard deviations and standard errors on either side of the mean were chosen following Whitehead (1971). Non-overlap of two standard errors from the mean is equivalent to a t-test at 0.05 probability, and indicates samples to be statistically significantly different; non-overlap of 1.5 standard deviations from the mean indicates that 90% or more of the specimens from one sample can be distinguished from 90% or more from the other sample, and indicates samples to be taxonomically significantly different (Whitehead 1971).

For most species these mensural features serve primarily as aids in identification and as characters valuable for phylogenetic analysis. For some species,

the mensural features have been verified as reliable by further measurements taken on a larger number of specimens than that indicated in the descriptive statistics tables or on the Hubbs-Hubbs diagrams (e.g. rostrum length in *C. texanus*-*C. longinasus*, form of elytra in *C. trivittatus*-*C. quadrilineatus*).

Analysis of geographic variation.--- Where problems in species recognition were apparent in terms of geographic variation in structural features, population samples from sites throughout the species range were selected and examined to determine the taxonomic significance of this variation. If more or less extensive differences were found between samples, geographically intermediate samples (if available) were examined to infer the degree of reproductive isolation of the geographically more distant samples. Should geographically intermediate populations prove structurally intermediate between the two more distant samples, gene flow is inferred and all samples are considered conspecific. Divergence from both of the more distant samples or close similarity to any one of the two distant samples by the intermediate sample may indicate barriers to gene flow and may result in separate species status for the two most distant samples, particularly if geographically correlated with other variable structural features or aspects of natural history.

complete list of synonymy is given. Type material examined and depository are noted following each of the names of the nominal taxa. Lectotypes and neotypes are designated where appropriate; paralectotypes have not been designated. Complete label data for primary type material examined, type locality, gender, and type species are given where appropriate for species, subgenera and genera. All literature citations of names of North American species, *Cleonidius vibex* (Pallas), and all citations pertaining to *Cleonis pigra* (Scopoli) in North America are given. A concerted attempt has been made to make these listings as complete as possible. Page numbers have not been given for all citations, but only for those wherein new species are described. Published misidentifications ("misident." in synonymy list) are indicated under both the incorrect and correct names. Notes in parentheses following most literature references indicate the nature of that reference as follows: biol., biology; catal., catalog; check., checklist; desc., description; distn., distribution; and redesc., redescription. Lack of any notation indicates use of that name with little or no added information about the taxon.

For each species, subgenus or genus, a "Problems in recognition" section is presented which I feel to be more desirable than a diagnosis because the latter, in most

species that I had problems separating from, and perhaps at one time misidentified or confused with, the species in question. In this manner I point out the species with which others are likely to make errors. This section is meant to be consulted after a tentative identification has been made using the keys.

A section dealing with notes about synonymy and type material follows and deals in various detail with specific criteria for newly proposed synonymy and with other specific details about type material as required. For all newly described species, label data for holotype and allotype are given in full with data from each label enclosed in separate quotation marks starting from the top label down. Separate lines on each label are indicated by a "/". A specific type locality is designated. Paratypes are treated less rigorously and label information is given in a simple standard format of: locality, date, collector, sex, location of deposition. Specimens are ordered alphabetically according to locality. Derivation of the specific epithet is also given.

Following this section is a detailed description of the taxon including structural variation. Complementing this verbal description are various illustrations of important structural features and for all species, a set of variously comprehensive descriptive statistics from one population for

each of males and females. Only attributes that vary between species are described. Character states are described in consistent order to facilitate comparisons. Subjective terms such as "large" and "small" have been avoided as much as possible but if by necessity used, are augmented with illustrations. All structural variation is incorporated in the appropriate places in the species description.

Geographic distribution of each species is presented in the form of dot maps illustrating positions of localities of collections. State records and questionable localities of collection are noted in the text but not placed on maps. Literature records are not placed on maps nor noted in text unless they fall outside the known range of the species based on other specimens examined. Complete lists of specimens examined for each species are not presented but have been placed in the Archives of the Department of Entomology, University of Alberta.

Natural history information from publications, specimen label data, and personal collecting experiences are summarized for each species. For the purposes of biogeographic and evolutionary interpretations, habitat associations were recognized using, as appropriate, schemes by Rzedowski (1978), Brown (1982), or Livingston and Shreve (1921). Elevations are either taken directly from specimen labels (if noted), or, if site of collection is a major geographic feature, are taken from a gazetteer or one of a number of maps. Hubbs-Hubbs diagrams illustrate variation in

elevational distribution within and among species of *Cleonidius* and *Apleurus*. Seasonal data are taken from labels. Supra-generic classification of plants follows Cronquist (1968). Generic names of plants follow Kearney and Peebles (1960); species names are following Kearny and Peebles (1960), Munz and Keck (1968) and Correll and Johnston (1970).

Patterns of allopatry and sympatry for each species are summarized in the chorological relationships section. They are summarized for *Apleurus* and *Stephanocleonus* species, but presented in a table format for *Cleonidius* species.

Consultation of this section will not only reduce possible misidentifications in instances where specimen locality is known, but also indicates retention of species identity under variable conditions of species packing and species occurrence. For this reason special attention is given to chorological relationships with close phylogenetic relatives.

A section on phylogenetic relationships concludes the descriptive format and verbally states the phylogenetic relationships of the taxon and the evidence for this placement. For species of *Apleurus* and *Cleonidius*, further details are given in the "Phylogeny" section. Phylogenetic relationships have not been examined for species of *Stephanocleonus*.

For species groups of *Cleonidius*, only a short diagnosis and a statement of evidence for monophyly and

phylogenetic relationships are presented. Structural features not variable within a species group but variable within *Cleonidius* are repeated in the species group diagnosis and all species descriptions.

#### 2.2.7 Criteria for recognition of species-group taxa

As with most systematists, my primary criterion for species recognition is the biological species concept (Mayr 1963). I find the evolutionary species concept of Wiley (1981), although theoretically desirable, so vague as to be of little practical applicability. Since experimental evidence of reproductive isolation is lacking for most insect species, Cleoninae being no exception, such isolation is inferred from differences primarily in structural features, but also in features of natural history, especially host plant associations.

In instances of sympatry, individuals are regarded as belonging to separate biological species if a variable number of structural features are substantially and consistently different. Differing host plant associations alone are not evidence of separate species status, but in combination with differences in structural features are used herein to indicate separate species status. This consideration is especially important should the degree of structural difference be less than is found in sympatric species where host plant associations are broader or not



consistently different yet structural differences are relatively extensive.

For allopatric individuals the situation is more complex and difficult to resolve. Allopatric individuals are regarded as conspecific if they exhibit structural features that do not differ, or if differing, do so to a lesser degree than differences noted between separate sympatric species and in a geographically based clinal manner. If the allopatric forms differ in various features of structure and/or natural history, and these features cannot be shown to have a consistent geographically determined pattern of continuous variation, the forms are considered separate species. Obviously, the more characters that differ in this manner, the more marked the evidence for separate species status. It is very important to note that marked phenetic differences are not the primary consideration, rather it is the pattern of geographic variation (assumed indicative of degree of gene flow) that determines the taxonomic status of the various populations. Phenetic differences may simply be the result of locally different selection pressures and not reproductive isolation.

These are my general criteria. Especially complicated and/or problematical patterns of structural variation must be considered individually and as such are discussed in detail in the text where appropriate.

The subspecific category has not been used. As expressed elsewhere (Anderson and Peck 1986), I feel that

this category should only be used for allopatric populations where lack of intermediate populations precludes analysis of geographic variation and inference of degree of reproductive isolation. In no instances in this study was such a consideration warranted; geographically intermediate populations and natural history information were consistently available to permit assessment of extent of gene flow and recognition of status at the species level.

#### 2.2.8 Criteria for recognition of genus-group taxa with notes on classification

Some consideration must be here given to criteria for recognition and ranking of genus-group taxa for this study is concerned with a regional fauna and classification of its constituents in terms of the world fauna. All genus-group taxa should preferably be monophyletic (holophyletic) although paraphyletic taxa are acceptable in certain special situations; polyphyletic taxa are not acceptable.

Genus-group taxa should contain an assemblage of related species sharing structural features and/or features of natural history, and separated from other assemblages by a gap in such features. Aside from this requirement, naming and ranking of taxa are always debatable because there are no objective criteria for formal recognition of taxa, and if recognized, at what hierarchical level they should be placed. Although this has been attempted (Hennig 1966), it

has not met with widespread acceptance or success.

All decisions about naming and ranking must be made with reference to related taxa with genealogical and phenetic relationships of the taxa as the prime considerations. Genealogical relationships (including some consideration of confidence in them), should take priority over phenetic relationships because only classifications based on evolutionary parameters can logically contain more information, due to inheritance, than was initially used in their creation (Whitehead 1972). Classifications of this nature may be maximally predictive and highest in information content, but may prove difficult to employ as information retrieval systems depending upon their complexity and intricacy. Many recent books (e.g. Eldredge and Cracraft 1980; Wiley 1981; and references cited therein) have dealt with this problem at length and simple compromise between information content and facility for use is perhaps the most important general consideration in constructing classifications.

Four genera are recognized in this study as occurring in North America. *Cleonis* is monotypic; *Stephanocleonus* is represented in North America by only 6 species but in the Palearctic region by numerous species, so no intrageneric classification has been attempted (species in North America are herein arranged alphabetically); *Cleonidius* is represented by 20 species; 19 Nearctic, 1 Palearctic (at least); *Apieurus* is represented by 8 Nearctic species.

In *Cleonidius* only informal species groups, including the *Incertae sedis* category, are recognized primarily because of low confidence in genealogical relationships and thus possible instability of the resultant classification, and little variation in structural features. By using only species groups, the possible predictive value of the classification is enhanced and can be tested without the problems associated with increased nomenclatural complexity, not desirable for such a small number of species of a relatively restricted geographic distribution.

In *Apleurus*, two subgenera have been recognized rather than species groups. Although *Apleurus* contains only 8 species, the genealogically most primitive member of *Apleurus* differs from other species of *Apleurus* in a number of structural features, some not yet found elsewhere in Cleoninae. Confidence in this primitive phylogenetic position is high and structural differences are marked, with remaining *Apleurus* species forming a structurally cohesive grouping. For these reasons, recognition of subgenera was deemed desirable. Furthermore, desire for recognition of *Gibbostethus* as a subgenus is enhanced by the largely unknown natural history, restricted distribution, and possible endangered status of this phylogenetically most primitive species of *Apleurus*.

Classification above the generic level is not attempted, although some statements concerning relatives of genera occurring in North America and their shared character

states are made, forming part of the basis for a future reclassification of the subfamily.

### 3. Natural history of Cleoninae

Members of the subfamily Cleoninae are found from mesic to very arid habitats. Taxa traditionally placed in the tribe Lixini are those usually associated with more mesic habitats whereas most traditional Cleonini are found in arid habitats. Exceptions, especially of some *Lixus* species associated with arid habitats, are known.

Most species breed in herbs or shrubs of a wide variety of plant taxa but most notably Chenopodiaceae, Compositae, Leguminosae, Rosaceae, Cruciferae and Polygonaceae. Larvae of most species mine stems or roots of these plants but those of some feed in flower heads or on seeds (e.g. *Rhinocyllus conicus*, *Larinus planus*, *Microlarinus* spp.). Natural history of *Microlarinus* spp. (Kirkland and Goeden 1977, 1978a,b) and of *Rhinocyllus conicus* (Zwölfer and Harris 1984) have been particularly well studied; however, most other information on the natural history of Cleoninae, especially in North America, is brief and fragmentary (e.g. Bargagli 1884; Pierce 1907).

Adults of most species of traditional Cleonini are brachypterous and even those that are macropterous have not been observed to fly. Most traditional Lixini are macropterous.

Adults of various species have been collected as prey of *Bufo* sp. (Amphibia), *Geococcyx californicus* (Aves), and various predatory Hymenoptera. Tachinidae (Diptera) (Arnaud 1978) and various parasitic Hymenoptera (Krombein *et al.*

1979) have been recorded as parasites of members of the genus *Lixus* in North America.

Various species of Cleoninae have been employed as, or are being considered as, biological control agents of weeds including thistle (*Cirsium*, Compositae), knapweed (*Centaurea*, Compositae), and puncturevine (*Tribulus*, Zygophyllaceae) (Anderson 1984a; Kirkland and Goeden 1978a,b; Maddox 1976; Wheeler and Whitehead 1985; Zwölfer and Harris 1984). None of the species are considered to be economic pests although some are occasionally found on cultivated Chenopodiaceae or Cruciferae.

## 4. Supraspecific taxa and included species

### 4.1 Subfamily Cleoninae

#### 4.1.1 Historical review

As of 1978, 1445 species of Cleoninae had been described (O'Brien and Wibmer 1978). However, the number of genera stated is perhaps on the low side for many taxa given subgeneric rank under *Cleonus* likely require separate generic status. The subfamily name Cleoninae (as "*Cleonides*") and generic name *Cleonus* were first proposed by Schoenherr (1826) but three years earlier, Schoenherr (1823) had also proposed the subfamily name Geomorinae (as "*Geomorides*") and the generic name *Geomorus* for the same taxa. Earlier still, Dejean (1821) had also proposed the name *Cleonis* to include species, some of which were subsequently placed by Schoenherr in *Geomorus* and *Cleonus*. Under Article 23(d) of the International Code of Zoological Nomenclature, Geomorinae has priority over Cleoninae, although the latter is in current general use. *Cleonis* Dejean likewise has priority over both *Geomorus* Schoenherr and *Cleonus* Schoenherr. The name Geomorinae has however, not subsequently been used whereas the name Cleoninae has been in widespread and constant use since its introduction (e.g.



Schoenherr 1834; Chevrolat 1873; Faust 1904; Csiki 1934; O'Brien and Wibmer 1982). A proposal has therefore recently been submitted to the International Commission of Zoological Nomenclature requesting suppression of Geomorinae in favour of Cleoninae (Anderson in press). Pending a decision, the latter name will be used.

North American species have traditionally been included in the tribe Cleonini (Casey 1891; LeConte 1876a; LeConte and Horn 1883) which is comparable to Cleoninae of other authors (e.g. Csiki 1934). There has been little confusion regarding included world taxa with the exception of the Holarctic genus *Lepyryus* now generally regarded not to be a cleonine.

#### 4.1.2 Recognition of Cleoninae

Adults of the subfamily Cleoninae are recognized by two features not known in other Curculionidae. These are: 1), short, telescoping, ventrally situated, three-articled labial palpi; and 2), presence of paired symbiont sacs attached to the vagina near the base of gonocoxite II in females. A more detailed diagnosis describing other features of the subfamily is not presented, pending examination of further taxa.

#### 4.1.3 Sex differentiation

Sexes are differentiated in Cleoninae as in most other weevils. Males are easily distinguished by flat to concave abdominal sterna III and IV and by possession of two clearly recognizable sclerotized terga at the abdominal apex, usually only visible upon displacement of the elytral apices. Females possess more evenly convex abdominal sterna III and IV and possess only a single sclerotized tergum at the abdominal apex. In addition, females tend to have a longer rostrum than do males (especially in *Cleonidius*) and are generally larger in body size.

#### 4.1.4 Key to genera of New World Cleoninae

1. Rostrum, in dorsal view, from apex of epistoma to anterior margin of eye, more or less as long as greatest width (Figs. 1-2).....2
- 1'. Rostrum, in dorsal view, from apex of epistoma to anterior margin of eye, much longer than greatest width (Figs. 3-4, 10, 25-30, 81-88, 137-156).....3
- 2(1). Elytron with intervals each with row of conspicuous erect setae. Anterolateral margin of pronotum straight, with long postocular vibrissae immediately behind eye (Fig. 1). Eye more or less round (Fig. 1).....*Microlarinus* Hochhuth  
(2 species, deliberately introduced)

2'. Elytron with intervals without row of erect setae.  
Anterolateral margin of pronotum with rounded  
postocular lobe and short postocular vibrissae  
behind eye (Fig. 2). Eye distinctly elongate-oval  
(Fig. 2).....*Rhinocyllus* Germar  
(1 species, deliberately introduced)

3(1'). Pronotum dorsally and laterally with numerous shiny  
glabrous tubercles, lacking distinct punctures.  
Rostrum dorsomedially sulcate (Fig. 10).....  
.....*Cleonis* Dejean  
(1 species, adventive)

3'. Pronotum dorsally and laterally distinctly  
punctate, with at most the outer margins of  
punctures of lateral margins swollen, glabrous and  
shiny. Rostrum either dorsomedially carinate to  
various degree (Figs. 25-30, 84-86) or not, not  
dorsomedially sulcate.....4

4(3'). Mesosternum with mesosternal process markedly  
tumescent (Fig. 58). Male with aedeagus markedly  
expanded laterally from midlength to apical  
one-third (Fig. 97e).....  
.....*Apleurus* subgenus *Gibbostethus* Anderson  
(1 species, native)

4'. Mesosternum with mesosternal process flat or at  
most only slightly convex, not tumescent (Fig. 59).  
Male with aedeagus more or less uniform in width  
throughout median portion of length.....5

only humerus and very base of interval 3 swollen and convex. Pronotum with disk with scale pattern various. Prosternum with or without swellings; swellings, if present, situated immediately anterior to each procoxal cavity (Fig. 8).....6

5'. Tibia with corbel ridge sharp (Fig. 6). Elytron with all intervals equally flat or with humerus and variously, sutural interval and intervals 3, 5, 7, and 9 elevated and convex throughout the greater part of their lengths. Pronotum with disk with large white scales present in a lateral stripe of various width, small and fine in moderately broad to very broad apically narrowed median stripe; median area largely black in color, underlying dark cuticle not obscured by overlying scales.

Prosternum with or without swellings; swellings, if present, situated immediately anterior to each prosternal impression (Fig. 7).....8

6(5). Antenna with article 2 of funiculus distinctly longer than wide, distinctly longer than each of articles 3 to 6, slightly shorter than to distinctly longer than article 1 (Fig. 4). Pronotum with anterolateral margins straight, slightly sinuate, or with at most variously developed (usually small) acute postocular projection;

one-half width of eye) behind base of eye (Fig. 4).

Ventral surface of femur dentate or not. (Some South American species of *Lixus* have slightly developed rounded postocular lobes and moderately long postocular vibrissae of more or less equal length, but have dentate femora).....\*

.....*Lixus* Fabricius  
(numerous species, native; worldwide)

6'. Antenna with article 2 of funiculus more or less as long as wide, more or less subequal in length to each of articles 3 to 6, shorter than article 1 (Figs. 3, 137-156). Pronotum with anterolateral margins straight or with slightly to well-developed, rounded postocular lobe; postocular vibrissae uniformly short (less than one-half width of eye in length) to unequal in length, greatest length (more or less equal to or greater than one-half width of eye) behind base of eye (Figs. 3, 137-156). Femur not dentate.....7

7(6'). Elytra elongate-narrow (width at midlength less than 0.65 times length) (Figs. 118-136). Pronotal disk with distinct white scales of various sizes (Figs. 118-136). Elytra with white scales, various in size, but more or less obscuring view of underlying cuticle over large part of elytral

7'. Elytra more robust (width at midlength greater than 0.65 times length). Pronotal disk with at most only very short indistinct setae, distinct scales absent. Elytra with scattered patches of elongate fine scales in addition to very short setae; underlying cuticle not obscured by scale cover.....

.....*Larinus* Schoenherr  
(1 species, introduced)

8(5'). Epistoma with anterior margin emarginate (Figs 25-30). Pronotum with well-developed, rounded postocular lobes; postocular vibrissae indistinct, uniformly short (Figs. 25-30). Eye elongate "kidney shaped" (Figs. 25-30); flat, or only slightly convex in dorsal view...*Stephanocleonus* Motschulsky

(6 species, native; also in Palearctic Region)

8'. Epistoma with anterior margin rounded (Fig. 82-88). Pronotum with anterior margin straight behind eyes or with small acute postocular projection immediately behind base of eye; postocular vibrissae distinct and long, longest immediately behind base of eye (Figs. 82-88). Eye "teardrop shaped" (Figs. 82-88); protruding and moderately to markedly convex in dorsal view.....

.....*Apleurus* subgenus *Apleurus* Chevrolat  
(7 species, native)

a separate genus, states that "it is scarcely distinct from *Lixus*". Accordingly, individuals of species that are assigned to *Ileomus* by O'Brien and Wibmer (1982) key out in the above key to *Lixus*. Detailed study of phylogenetic relationships of the various groupings of *Lixus* species is needed to assess their validity as nominal subgeneric or generic taxa.

#### 4.2 Genus *Cleonis* Dejean

*Curculio*; Scopoli 1763 (in part; sp. desc.). Linnaeus 1767 (in part; sp. desc.). Goeze 1777 (in part; sp. desc.). Knoch 1781 (in part; sp. desc.). Villers 1789 (in part; sp. desc.). Gmelin 1790 (in part; sp. desc.).

*Cleonis* Dejean 1821:96. Gender, feminine. Type species *Curculio sulcirostris* Linnaeus by subsequent designation (O'Brien and Wibmer 1982:73). Silfverberg 1979 (in part; check.). O'Brien and Wibmer 1982 (in part; catal., distn., misident. *Apleurus*, *Cleonidius*, *Stephanocleonus*). O'Brien and Wibmer 1984 (in part; misident. *Apleurus*, *Cleonidius*, *Stephanocleonus*).

*Geomorus* Schoenherr 1823:column 1141. Gender, masculine. Type species *Curculio sulcirostris* Linnaeus by original designation.

*Cleonus* Schoenherr 1826:145. Unjustified replacement name

International Code of Zoological Nomenclature. Say 1831  
(misident. *Cleonidius*). Kirby 1837 (misident.  
*Cleonidius*). LeConte 1850 (misident. *Stephanocleonus*).  
Melsheimer 1853 (misident. *Cleonidius*). LeConte 1858  
(misident. *Apleurus*). LeConte 1859 (misident.  
*Apleurus*). Gemminger and von Harold 1871 (in part;  
misident. *Apleurus*, *Cleonidius*). LeConte 1876a  
(misident. *Cleonidius*). Henshaw 1881-1882 (misident.  
*Cleonidius*). LeConte and Horn 1883 (misident.  
*Cleonidius*). Henshaw 1885 (misident. *Cleonidius*).  
Wickham 1889 (misident. *Cleonidius*). Wickham 1896  
(misident. *Cleonidius*). Fall 1897 (misident.  
*Cleonidius*). Fall 1901 (misident. *Cleonidius*). Wickham  
1902 (misident. *Cleonidius*). Fletcher 1906 (misident.  
*Cleonidius*). Fall and Cockerell 1907 (misident.  
*Cleonidius*). Pierce 1907 (misident. *Cleonidius*). Ely  
1913 (misident. *Cleonidius*). Gibson 1914 (misident.  
*Cleonidius*). Anderson 1914 (misident. *Cleonidius*).  
Blatchley and Leng 1916 (misident. *Cleonidius*). Yothers  
1916 (misident. *Cleonidius*). Leng 1920 (misident.  
*Stephanocleonus*, *Apleurus*, *Cleonidius*). Leonard 1926  
(misident. *Stephanocleonus*, *Cleonidius*). Böving 1927  
(in part; larval key). Bradley 1930 (misident.  
*Apleurus*, *Cleonidius*). Crosby and Blauvelt 1930  
(biol.). Leng and Mutchler 1933 (catal.). Wilcox et al.



*Stephanocleonus, Cleonidius, Apleurus, 1960 (misident. Cleonidius)*. Brown 1940 (distn.). Blackwelder 1947 (misident. *Apleurus, Cleonidius*). Bruhn 1947 (misident. *Cleonidius*). Hicks 1947 (distn.). Hicks 1949 (distn.). Anderson 1956 (biol., distn.). Essig 1958 (misident. *Cleonidius*). Sanders 1960 (misident. *Cleonidius*). Kissinger 1964 (in part; biol.). Tanner 1966 (misident. *Apleurus, Cleonidius*). Kingsolver 1972 (misident. *Cleonidius*). Burke and Anderson 1976 (biblio.). Kumar et al. 1976 (misident. *Cleonidius*). Krombein 1979 (misident. *Stephanocleonus, Apleurus*). Arnett et al. 1980 (misident. *Cleonidius*). Batra et al. 1981 (distn.). Peschken 1984 (biol.).

*Cleonus s.s.*; Csiki 1934 (catal.).

*Cleonus (Cleonidius)*; Arnett 1960-1962 (in part; misident. *Apleurus*).

Notes about synonymy.--- *Eplimeces* Billberg, 1820, p. 45, predates *Cleonis* Dejean, 1821 and originally included various Cleoninae (among others, *Curculio sulcirostris* Linnaeus, the type species of *Cleonis* Dejean, and, *Curculio filiformis* Fabricius, presently assigned to *Lixus* Fabricius [Csiki 1934]). The name *Eplimeces*, to my knowledge, has not subsequently been used but because no type species has been designated for the genus, the name represents a threat to

species of *Epimeces* Billberg, 1820, p. 45. *Epimeces* therefore becomes a new junior subjective synonym of *Lixus* Fabricius, 1801, p. 498, type species *Curculio paraplecticus* Linnaeus, 1758, p. 380, by subsequent designation (Latreille 1810, p. 430).

Schoenherr (1823), seeking consistency in gender of generic names, changed all feminine generic names in Curculionidae to a masculine name (not always the masculine form of the former feminine name). As a result of this, and according to Schoenherr (1823), inadequate characterization to recognize a type species, use of the name *Cleonis* Dejean, 1821 was discontinued, and the species initially placed in *Cleonis* were assigned to one of four genera by Schoenherr (1826). One of these was *Geomorus* Schoenherr, 1823. As such, *Geomorus* cannot be regarded as an express replacement name for *Cleonis*. Subsequently, Schoenherr (1826) proposed the genus *Cleonus* expressly as a replacement name (although unjustified) for *Geomorus* as indicated by his designation of *Curculio sulcirostris* Linnaeus, 1767 as the type species. This same species had already been designated by him as the type species of *Geomorus*. Schoenherr (1826) also placed all species initially included in *Geomorus* in *Cleonus*. The generic name *Cleonus* has since been in widespread use. However the name *Cleonis* was recently resurrected by Silfverberg (1979) and used by O'Brien and Wibmer (1982),

(subgenus *Cleonis* of Csiki [1934]) including only *C. pligra* (Scopoli), *C. japonicus* (Faust), and *C. sardous* (Chevrolat), the latter two species not examined by me. *Cleonis* appears closely related to *Cyphocleonus* Motschulsky and *Adosomus* Faust, and subsequent phylogenetic analysis may reveal the three to be congeneric.

**Diagnosis.**--- Adult Cleoninae with moderately robust body form (Fig. 19). Rostrum with broad low median carina longitudinally sulcate throughout length (Fig. 10). Pronotum with small glabrous shiny tubercles, not distinctly punctate; pronotal postocular lobes slightly to moderately well-developed, rounded. Tibia with corbel ridge sharp (as in Fig. 6). Elytra with strial punctures individually indistinct, dorsal surface of elytra with scattered, small, irregularly shaped (usually transverse), glabrous, shiny swellings.

**Description.**--- **Size.** Moderately large, moderately robust in form. **Mouthparts.** Prementum flat, with single large seta on each side. Maxillary palpus with palpifer and stipes each with large seta. Basal articles of labial palpi separated by distance subequal to width of a palpus. **Rostrum.** Moderately robust; with broad low median carina longitudinally moderately deeply sulcate throughout length from base of

emarginate medially. Antenna with funiculus with article 1 very slightly longer than article 2 (Fig. 9); apical three articles of club with placoidal sensillae. Head. Eye elongate-quadrate, slightly wider at top than bottom; flat (Fig. 9). Vestiture. Dorsum lacking erect or suberect vestiture, with only simple elongate-narrow appressed white scales. Procoxae, mesocoxae, metasternum and base of abdominal sternum III of both sexes with moderately long suberect hair-like scales. Prothorax. Dorsal surface of pronotum with small glabrous shiny tubercles, not distinctly punctate. Median basal area shallowly impressed. Disk with elongate, moderately large, moderately dense, white scales present in broad lateral stripe and narrow median line, with scales small and fine in pair of paramedian apically narrowed stripes; median area largely black in color, underlying dark cuticle not obscured by overlying scales (Fig. 19). Pronotum widest at base, lateral margins slightly rounded and convergent from base to apex; apical constriction very slight (Fig. 19). Pronotal postocular lobes rounded, slightly to moderately well-developed (Fig. 9). Prosternum with slight impression, anterolaterad of each procoxal cavity and with slight swelling immediately anterior to each prosternal impression. Legs. Tarsi broad, articles 2 and 3 more or less as broad as long, subequal in length; article 1 only slightly longer than articles 2 or 3;

3 (Fig. 18). Claws connate in basal one-third, not to slightly divergent. Foretibia with small to moderately well-developed second spur; inner margin with moderately large denticles throughout greater portion of length. Tibia with corbel ridge sharp. Metatibia of male with uncus with ventral margin slightly sinuate. Wings. Present. Elytra. Base of interval 3 and humerus very slightly swollen and convex, otherwise intervals uniformly flat. Humeri distinct. Strial punctures individually indistinct, dorsal surface of elytra with scattered small irregularly shaped (usually transverse) glabrous shiny swellings. Abdomen. Ventral surface with small shiny glabrous patches, each with single small appressed scale. Genitalia. Female. Abdominal sternum VIII lacking basal arm (Fig. 17). Gonocoxite II elongate-triangular, apex not prolonged into marked lobe; stylus moderately large, apical in position (Fig. 15). Spermathecal gland slightly elongate-oval (Fig. 16). Male. Aedeagus elongate, moderately robust; in lateral view, moderately arcuate near base then only slightly arcuate throughout rest of length (Fig. 13). Apex slightly spatulate. Internal sac with median dorsal pocket high; various lobes present; apical and dorsal median pockets individually distinct; apical sclerite complex present, individual sclerites distinctly scythe-like (Figs. 11, 12, 14).

pronotum and elytra that are distinctly tuberculate and not punctate. *Adosomus* and *Cleonis* species possess moderately well-developed, rounded, postocular lobes, whereas postocular lobe are lacking from *Cyphocleonus* species. The only feature which distinguishes *Cleonis* from both of these two genera is the medially longitudinally sulcate rostrum of the former. In *Adosomus* the rostrum is medially tumescent and not carinate or sulcate, whereas in *Cyphocleonus* species, the rostrum is variously medially tumescent and carinate. *Cyphocleonus trisulcatus* (Herbst) has a low broad median carina but which is only apicomediaally sulcate.

Checklist of included species.--- Following Csiki (1934), three species are recognized. They are *C. japonicus* (Faust), *C. sardous* (Chevrolat), and *C. pigra* (Scopoli). *Cleonis pigra* is the only one of the three species that I have examined and is the only one that occurs in the Nearctic Region.

Faust (1904) distinguishes *C. pigra* and *C. japonicus* from *C. sardous* by the latter having metatarsal article 2 slightly longer than article 3 and by differences in the pronotal scale patterns. *Cleonis pigra* and *C. japonicus* are separated on the basis of extent of distribution and development of the pronotal and elytral tubercles, the distribution of the ventral abdominal glabrous patches, and

vary somewhat, and suspect that the three included species may prove conspecific.

Phylogenetic relationships.--- *Cleonis*, *Cyphocleonus* and *Adosomus* appear to represent a monophyletic group based on the presence of a tuberculate rather than punctate pronotum and elytra. Beyond this, affinities of the group are uncertain.

#### 4.2.1 Species accounts

##### 4.2.1.1 *Cleonis pigra* (Scopoli)

(Figs. 9-19, 213)

*Curculio piger* Scopoli 1763:23.

*Curculio sulcirostris* Linnaeus 1767:617.

*Curculio transversofasciatus* Goeze 1777:409.

*Curculio nebulosus* Knoch 1781:87 [not Linnaeus 1758:385].

*Curculio fasciatus* Villers 1789:216 [not Müller 1776:86].

*Curculio fasciatus* Gmelin 1790:1804.

*Cleonis sulcirostris*; Dejean 1821.

*Cleonus sulcirostris*; Gyllenhal 1834.

*Cleonus indicus* Fähræus 1842:55.

*Cleonus piger*; Everts 1903.

*Cleonis pigra*; O'Brien and Wibmer 1984.

**Notes about synonymy.**--- The synonymical list follows Csiki (1934); I have not examined type material of this species or of any of its conspecific forms. This is the species frequently referred to in previous publications as *C. sulcirostris*.

**Description.**--- **Specimens examined.** 61 unsexed. **Size.** Length, male, 7.5-13.4 mm; female, 11.4-14.8 mm. Width, male, 3.7-6.6 mm; female, 5.8-7.0 mm. **Rostrum.** Lateral margins sharp, indicated by low rounded carina from above point of antennal insertion to frons (Fig. 10). Side portion of rostrum immediately ventrad of lateral carina moderately deeply sulcate from immediately anterior to eye to point of antennal insertion; with moderately deep elongate-narrow punctures, most linearly confluent or nearly so. **Prothorax.** Surface dorsally and laterally with numerous small shiny glabrous tubercles; medially with small to large elongate-oval shiny glabrous swelling. **Prosternum** in lateral view with apical portion flat, slightly shorter than adjacent posterior steeply declivous portion. **Elytra.** Sutural interval in basal one-half, intervals 2 to 5 basally, and two obliquely posteromedially directed patches



slightly elevated shiny glabrous patches (Fig. 19). Wings. Long (greater than elytra in length). Bases of 2A joined but very lightly sclerotized. Abdomen. Ventral surface with small shiny glabrous patches distinct and dense on abdominal sterna III and IV, less distinct on sterna V to VII.

Genitalia. Female (2 specimens). Abdominal sternum VIII with lateral arms narrow, evenly and slightly inwardly arcuate throughout length, expanded apically (Fig. 17). Male (2 specimens). Aedeagus robust; in lateral view narrowed from approximately apical one-third to apex (Fig. 13). Internal sac with single, large, broad, dorsally directed, lobe; with moderately large, paired, laterally directed, lobes and small dorsally directed median lobe on dorsal surface at midlength; with small dorsolaterally directed paired lobes at midlength and midheight (Figs. 11-12). Eversible apical sclerite complex with paired scythe-like sclerites, each with median elongate-narrow projection long, visible in lateral view (Figs. 11, 14); adjacent ventral surface and basal portion of sides of apex of sac with pair of large transverse dark sclerites (Fig. 11).

Geographic distribution.--- This species was accidentally introduced into North America from Europe prior to 1919, the year of the earliest known North American record (Anderson 1956). It occurs throughout New York and southern Ontario,

west to Michigan and east to eastern Quebec and New Brunswick (Fig. 213). It is widespread in the Palearctic Region (Csiki 1934).

**Variation.**--- As part of a program investigating the potential for use of *C. pigra* as a biological control agent for the introduced *Centaurea diffusa* Lam. (diffuse knapweed) in the prairie provinces of Canada, a study of variation based on 15 mensural features was carried out on adult individuals of *C. pigra* reared from species of *Cirsium*, *Carduus* and *Centaurea* (all Compositae) to determine if structurally distinct host races exist (Anderson, 1984a). Measurements were made of 15 characters and a linear discriminant analysis was performed. Separation using the generated discriminant function proved unreliable and because absolute size was the principal component upon which the function was based, it was concluded that only a single polyphagous morphotype was present. Differences in absolute size are likely a result of structural attributes of the different larval feeding and pupation sites in the different host plants and are doubtfully heritable.

**Natural history.**--- Anderson (1956) summarizes information on the natural history of this species in North America as follows. Adults are the overwintering stage and are first found on Canada thistle, *Cirsium arvense* (L.) Scop., or bull thistle, *Cirsium vulgare* (Savi) Tenore, in early June.

Copulation and oviposition take place through mid-July. Eggs are laid singly in the lower portions of the stems in cavities chewed by the females and subsequently plugged with frass following deposition of the egg. Larvae bore downward into the primary root where they feed for approximately 30 days. Pupation takes place in the root. Adults emerge in August or September and overwinter under ground debris.

Although a wide range of Compositae serve as host plants in Europe (Anderson 1956; Peschken 1984, and references cited therein), only two plants, *Cirsium arvense* and *Cirsium vulgare*, both adventive, are known to serve as hosts in North America (Anderson 1956). Native North American *Cirsium* species or other Compositae are not attacked by the weevils.

A similar range of host plant associations to those of *C. plura* in Europe is also found in another species of Cleoninae, *Rhinocyllus conicus* Froelich, which was recently shown to consist of various host specific races (Zwölfer and Preiss 1983). As such, host plant races are also suspected in *C. plura* although, as noted, an examination of the structural features of adults reared from various Compositae did not reveal differences between individuals reared from different plant taxa.

Immature stages were described by La Ferla (1939).

#### 4.3 Genus *Stephanocleonus* Motschulsky

*Cleonus*; LeConte 1850 (in part). Leng 1920 (in part; catal.). Leonard 1926 (in part; check.). Bleasdel 1937 (in part). Krombein 1979 (in part; as prey).

*Stephanocleonus* Motschulsky 1860:540 (in key). Gender, masculine. Type species *Curculio flaviceps* Palliser by original designation. LeConte 1876a (key, sp. desc.). Henshaw 1881-1882 (in part; check.). LeConte and Hörn 1883 (key). Henshaw 1885 (check.). Blatchley and Leng 1916 (key, sp. redesc.).

*Conlocleonus* Motschulsky 1860:540 (in key). New synonymy. Gender, masculine. Type species *Cleonus carlinrostris* Gyllenhal by original designation.

*Plagiographus* Chevrolat 1873:21. Gender, masculine. Type species not designated.

*Cleonus* (*Stephanocleonus*); Casey 1891 (in part; key). Fall and Cockerell 1907 (in part; check.). Csiki 1934 (in part; catal.). Arnett 1960-1962 (in part; catal., key). Kissinger 1964 (in part; key).

*Cleonis*; O'Brien and Wibmer 1982 (in part; catal., distn.). O'Brien and Wibmer 1984 (in part).

Notes about synonymy.--- The genera *Stephanocleonus* and *Conlocleonus* were both originally proposed by Motschulsky (1860). They were distinguished by the rostrum having only a single median carina in the latter, whereas in

*Stephanocleonus*, the rostrum, in addition to the median carina, also had a smaller oblique carina on each side. *Stephanocleonus* is also noted as being apterous but nothing is said of the state of the wings in *Coniocleonus*. I have examined specimens of *Stephanocleonus flaviceps* Palliser, the type species of *Stephanocleonus*, and did not see small oblique lateral carinae, but only a single uniformly steeply declivous median carina in the single male available. The two females examined have the median carina similarly steeply declivous basally but decreasingly declivous apically. A very slight oblique carina is present where this more gradual apical declivity meets the flat surface of the rostrum on each side of the median carina. These three individuals are wingless.

I have not seen specimens of *Coniocleonus carlinirostris* Gyllenhal, the type species of that genus, but I have examined a male and a female of each of *C. excoriatus* Gyllenhal and *C. glaucus* Fabricius, a female of *C. cineritius* Gyllenhal, and a male of each of *C. cinerascens* Hochhuth and *C. ferrugineus* Fähræus. Individuals of *C. excoriatus* and *C. glaucus* are macropterous and the rostrum has a single median carina which is gradually declivous throughout its length. There are no small lateral carinae evident. However, the specimens of *C. cineritius*, *C. cinerascens* and *C. ferrugineus* each have a rostrum with slightly to moderately well-developed oblique lateral carinae in the same position as in *S. flaviceps*, and have

wings of variable length.

In view of this variation and the lack of other distinguishing characters, I cannot accept other than that *Stephanocleonus* is simply an apterous *Conlocleonus*. I do not think the rostral differences between the type species warrant separate generic status. I therefore consider *Stephanocleonus* and *Conlocleonus* to be new subjective synonyms, a decision anticipated by Casey (1891:188) who stated, regarding the Nearctic species, that "It is a question whether our species should be placed in *Stephanocleonus* or *Plaglographus* [a junior synonym of *Conlocleonus* according to Faust 1904 and Csiki 1934], but this is a matter of but slight importance as the differences between the subgenera appear to be very inconsiderable".

I choose to give the name *Stephanocleonus* priority over *Conlocleonus* because the former has been the only name used in reference to the Nearctic species here placed in the genus. Furthermore, no major works have recently appeared dealing with *Conlocleonus*, however, a recent review by Ter-Minasyan (1979) of the *Stephanocleonus* of the Palearctic Region includes species that in my opinion are congeneric with species in North America.

**Diagnosis.**--- Adult Cleoninae with moderately robust to robust body form (Figs. 20-23). Eye kidney-shaped, flat (Figs. 25-30). Rostrum with well-developed sharp median carina (Figs. 25-30). Pronotum punctate; pronotal postocular

lobes well-developed, rounded; postocular vibrissae uniformly short (Figs. 25-30). Meso- and especially metatarsus elongate-narrow, with all articles markedly longer than broad; article 2 distinctly longer than article 3; article 1 markedly longer than articles 2 or 3. Tibia with corbel ridge sharp (as in Fig. 6). Procoxae, mesocoxae, metasternum, metacoxae and base of abdominal sternum III of male with moderately long to very long erect hair-like scales; hair-like scales sparse to lacking in female. Pronotal disk with elongate, white, moderately large, moderately dense scales present in lateral stripe of various width, small and fine in moderately broad to very broad, apically narrowed, median stripe, and variously small and fine to moderately large and elongate along lateral margins; median area largely black in color, underlying dark cuticle not obscured by overlying scales (Figs. 20-23). Prosternum with or without swellings, swellings, if present, immediately anterior to each prosternal impression (as in Fig. 7). Metauncus of male with ventral margin slightly to markedly sinuate. Female with abdominal sternum VIII with basal arm very long (Figs. 37-42); gonocoxite II rounded basally, with apex prolonged into a marked lobe (Fig. 54). Male with apex of aedeagus spatulate or not (Figs. 31-36); internal sac with or without large paired sclerite at midlength (Figs. 31-36).

**Description.**--- **Size.** Moderately large, moderately robust to

robust in body form (Figs. 20-23). **Mouthparts.** Prementum ventrally longitudinally carinate, slightly swollen to flat, with one or more large setae on each side. Maxillary palpus with palpifer and stipes each with at least one large seta. Bases of labial palpi separated by distance subequal to width of basal article of labial palpus. **Rostrum.** Moderately to markedly robust, not medially tumescent; with low, to high, sharp median carina (Figs. 25-30). Epistoma markedly swollen or not, with apical margin emarginate medially (Figs. 25-30). Antenna with funiculus with article 1 slightly to markedly longer than article 2 (Figs. 25-30); apical three articles of club with placoidal sensillae. **Head.** Eye kidney-shaped, flat (Figs. 25-30). Upper margin of eye rounded to sharp, frons convex to markedly concave. **Vestiture.** Dorsum lacking or with at most only very short indistinct superect or erect vestiture, with simple elongate-narrow appressed white scales of various size and density. Procoxae, mesocoxae, metasternum, metacoxae and base of abdominal sternum III of male with moderately long to very long erect hair-like scales; hair-like scales sparse to lacking from female. **Prothorax.** Dorsal surface of pronotum punctate. Pronotum with median basal area shallowly to deeply impressed, disk with or without various other impressions. Disk with elongate white moderately large moderately dense scales present in lateral stripe of various width, small and fine in moderately broad to very broad apically narrowed median stripe, and variously small and



fine to moderately large and elongate along lateral margins; median area largely black in color, underlying dark cuticle not obscured by overlying scales (Figs. 20-23). Pronotum widest at base to subequal in width from base to apical one-quarter, then constricted to various extent and convergent to apex (Figs. 20-23). Pronotal postocular lobes moderately to well-developed; postocular vibrissae uniformly short (length less than one-half maximum width of eye) (Figs.; 25-30). Prosternum with slight impression anterolaterad of each procoxal cavity; with or without slight swelling immediately anterior to each prosternal impression (as in Fig. 7). Legs. Foretarsus moderately broad, articles 2 and 3 more or less subequal in length, at most slightly longer than broad; article 1 only slightly longer than articles 2 or 3; article 3 moderately deeply bilobed. Meso- and especially metatarsus elongate-narrow; all articles markedly longer than broad, article 2 distinctly longer than article 3; article 1 markedly longer than articles 2 or 3, article three slightly bilobed. Ventral tarsal pilosity various in extent, lacking entirely on at least more basal articles in most species. Claws connate in basal one-third, not to slightly divergent. Foretibia with inner margin near apex with moderately-developed second spur, inner margin with at most only small denticles in apical one-half. Metatibia of male with uncus with ventral margin slightly to markedly sinuate. Tibia with corbel ridge sharp. Wings. Absent or present

(various in length). Elytra. Intervals flat to variously slightly swollen and convex; striae distinctly punctate. Humeri acute to rounded. Dorsal surface of most species with variously developed, posteromedially directed oblique patches of small and fine scales at each of basal one-third and apical one-third to midlength (Figs. 20, 22-23).

**Abdomen.** Ventral surface with small shiny glabrous patches, each with single small appressed scale. Sternum VII of male with apex variously medially emarginate in most species.

**Genitalia.** Female. Abdominal sternum VIII with long basal arm (Figs. 37-42). Gonocoxite II rounded basally, with apex prolonged into marked lobe; stylus moderately large, apical in position (Fig. 54). Spermathecal gland round (Fig. 48).

Male. Aedeagus robust and short to elongate-narrow, various in shape in lateral view; apex not to markedly spatulate (Figs. 31-36). Internal sac various; median pocket low, apical and dorsal median pockets individually distinct or not, various lobes present; apical sclerite complex with individual sclerites distinctly scythe-like, large paired sclerite at midlength present or absent (Figs. 43-48).

**Comparisons.**--- *Stephanocleonus* is very similar to *Pleurocleonus*, however, the latter lacks pronotal postocular lobes, has an elongate-teardrop-shaped eye, tarsal claws that are widely divergent, and a low broad longitudinally sulcate median carina. Species of *Stephanocleonus* may also be confused with *Nomimonyx* and *Epirhynchus* because of

similarities in head and rostral form, and eye shape.

*Epirhynchus* however, has a single tarsal claw, and both have short broad tarsal articles, not the markedly elongate-narrow articles of *Stephanocleonus* species.

Although unlikely, some *Connorhynchus*, *Bothynoderes* and *Chromonotus* may also be confused with *Stephanocleonus* species. These taxa however have article 2 of the antennal funiculus longer than article 1; in *Stephanocleonus* species, article 1 is longer than article 2.

**Checklist of included species.**--- In the Nearctic Region, six species are placed in *Stephanocleonus*. Numerous species occur in the Palearctic Region (Csiki 1934; Ter-Minasyan 1979) but are not listed here. The six Nearctic species are as follows:

1. *S. confusus* Anderson, n. sp.
2. *S. cristaticollis* Csiki
3. *S. immaculatus* Anderson, n. sp.
4. *S. parshus* Anderson, n. sp.
5. *S. plumbeus* (LeConte)
6. *S. stenothorax* Anderson, n. sp.

**Phylogenetic relationships.**--- Distribution of the apotypic character state of female gonocoxite II (rounded basally and with a marked apical lobe [Fig. 54]) suggests that *Stephanocleonus*, *Pleurocleonus* Motschulsky, *Menecleonus* Faust, *Xanthochelus* Chevrolat, and *Connorhynchus* Motschulsky

(=*Temnorhinus* Chevrolat) form a monophyletic group. *Menecleonus*, *Xanthochelus* and *Pleurocleonus* appear to further comprise another monophyletic group (based upon presence of an unpaired sclerite near the confluence of the apical pocket and the dorsal median pocket in the internal sac of the aedeagus of males) that is the sister-group of *Stephanocleonus*. This proposed sister-group relationship is based on article 1 of the antennal funiculus being longer than article 2. *Conorhynchus* has article 2 of the funiculus longer than article 1 and is likely the sister-group to the *Stephanocleonus-Pleurocleonus, Menecleonus, Xanthochelus* lineage. *Bothynoderes* Schoenherr, *Chromonotus* Motschulsky and also *Chromosomus* Motschulsky, have the apotypic state of elongate-narrow meso- and metatarsi and together with the above taxa likely comprise another monophyletic group based upon shared possession of this character state. They also have a very similar eye-shape to *Conorhynchus* and similarly all have article 2 of the antennal funiculus longer than article 1.

#### 4.3.1 Key to species of adult *Stephanocleonus*

1. Dorsal surface of elytra uniformly covered with small white scales, lacking distinct maculations; with or without a small shiny glabrous area at confluence of intervals 4 to 6 (Fig. 21). Elytra with humeri distinct; lateral margins more or less

straight in basal one-half (Fig. 21).....

.....*S. Immaculatus* Anderson

1'. Dorsal surface of elytra faintly to distinctly maculate, with small white scales less dense in two posteriorly directed oblique patches (one at basal one-third from intervals 2 to 6; the other at apical one-third from intervals 2 to 4), and with a small triangular shiny glabrous area at confluence of intervals 4 to 6 (Figs. 20, 22-23). Elytra with humeri less distinct or rounded; lateral margins more or less arcuate in basal one-half (Figs. 20, 22-23).....2

2(1'). Elytra fused together; metasternal wings short, approximately equal to one-half length of elytra.....3

2'. Elytra not fused together; metasternal wings long, approximately equal to or greater than length of elytra.....4

3(2). Elytra with scattered small recurved hair-like scales, most evident on declivity from sutural interval to interval 3; with greater part of length of sutural interval and intervals 3 and 5 elevated and convex. Female with sternum VII flat, lacking paired tumescence. Male with median lobe elongate; apex slightly spatulate; in lateral view narrowed from apical one-third to apex; with at most a small basal ventral tubercle (Fig. 31).....

- .....*S. confusus* Anderson
- 3'. Elytra lacking erect hair-like scales, with only the extreme base of interval 3 convex, otherwise with all intervals flat throughout their length. Female with ventral surface of sternum VII with paired tumescence. Male with median lobe less elongate; apex markedly spatulate; in lateral view, narrowed from approximately midlength to apex; with a large basal ventral tubercle (Fig. 32).....
- .....*S. cristat/collis* Csiki
- 4(2'). Pronotum with subapical constriction well-defined dorsally and laterally, with distinct moderately deep impressions laterad of median carina at apical one-quarter laterally continuous with subapical constriction; with apical margin broadly and moderately deeply emarginate at middle. Elytra with tubercle at confluence of intervals 4 to 6 markedly elevated. Rostrum robust (width at apex greater than 0.90 times length); approximately uniform in width throughout length (Fig. 30).....
- .....*S. stenothorax* Anderson
- 4'. Pronotum with subapical constriction only laterally well-defined, lacking or with at most only very shallow indistinct impressions laterad of median carina; with apical margin not or only very slightly emarginate at middle. Elytra with tubercle at confluence of intervals 4 to 6 only slightly

- elevated. Rostrum more elongate-narrow (width at apex less than 0.90 times length); slightly narrowed at midlength (Figs. 28-29).....5
- 5(4'). Pronotum with punctures large, deep and dense, the distance between punctures distinctly less than the diameter of a puncture (Fig. 28). Female metatarsal article 3 with large ventral pilose pads. Male with median lobe in ventral view with apex symmetrical (Fig. 34b); internal sac lacking median dorsal pocket and large dorsally directed paired lobes on dorsal surface immediately anterior to midlength (Fig. 46) .....*S. parshus* Anderson
- 5'. Pronotum with punctures small, shallow and sparse, the distance between punctures greater than the diameter of a puncture (Fig. 29). Female metatarsal article 3 entirely spinose, lacking ventral pilose pads. Male with median lobe in ventral view with apex asymmetrical (Fig. 35b); internal sac with median dorsal pocket and lacking dorsally directed lobes from dorsal surface (Fig. 47) .....  
.....*S. plumbeus* (LeConte)

#### 4.3.2 Species accounts

4.3.2.1 *Stephanocleonus confusus* Anderson, new species

(Figs. 20, 25, 31, 37, 43, 204)

*Stephanocleonus plumbeus*; LeConte 1876a (in part; desc., misident., mixed type series). Henshaw 1881-1882 (in part; check., misident.). Henshaw 1885 (in part; check., misident.). Wickham 1902 (check., misident.). Wickham 1909 (check., misident.). Leng 1919 (in part; check., poss. misident.). Leng 1920 (in part; catal., misident.). Danks 1981 (in part; check., poss. misident.).

*Cleonus cristatus*; Casey 1891 (key, misident.).

*Cleonus (Stephanocleonus) plumbeus*; Fall and Cockerell 1907 (in part; check., misident.).

*Stephanocleonus cristatus*; LeConte 1878 (distn., misident.). Leng 1920 (in part; catal., misident.).

*Cleonus plumbeus*; Krombein 1979 (in part; prey, poss. misident.).

*Cleonis cristatifollis*; O'Brien and Wibmer 1982 (in part; catal., distn., misident.).

*Cleonis plumbeus*; O'Brien and Wibmer 1982 (in part; catal., distn., misident.).

Type material:--- Holotype, male, labelled with a red label "HOLOTYPE", "Spearfish Canyon/ Black Hills, S.D./ 15-18:VI:1910.", inverted "*Cleonus/ cristatus/ Lec.*", "HOLOTYPE/ *Stephanocleonus/ confusus/ Anderson*" and with



abdomen on card and genitalia in microvial attached to pin (USNM). Allotype, female, labelled with a red label "ALLOTYPE", "Twp. 21 Rge. 6/ W. 4 Mer. Alberta/ 12.V.1980/ Lot 3 BF&JL Carr", "ALLOTYPE/ *Stephanocleonus/ confusus/ Anderson*" and with abdomen on card and genitalia in microvial attached to pin (CNCI). Type locality, Spearfish Canyon, Black Hills, South Dakota.

Paratypes. 36 males, 29 females. CANADA: Alberta: Township 1, Range 4, West 4 Meridian, 23.V.82, B.F. and J.L. Carr, 1M (JLCC); Township 13, Range 14, West 4 Meridian, 16.IV.82, B.F. and J.L. Carr, 1M (JLCC); Township 14, Range 25, West 4 Meridian, 15.V.82, B.F. and J.L. Carr, 1F (JLCC); Township 10, Range 2, West 5 Meridian, 11.V.82, B.F. and J.L. Carr, 1F (JLCC); Medicine Hat, 22.IV.29, F.S. Carr, 1F (UASM), 13.VI.32, F.S. Carr, 2M, 1F (UASM), 24.VI.30, F.S. Carr, 1F (UASM), 27.III.27, F.S. Carr, 1F (UASM), 14.IV.26, F.S. Carr, 1F (UASM), 13.VI.33, F.S. Carr, 1F (UASM), 2.VII.23, F.S. Carr, 1M (UASM), 1.V.32, F.S. Carr, 1M (UASM), 13.V.27, F.S. Carr, 1M (UASM), 13.V.23, F.S. Carr, 1M (UASM), 6.IX.27, F.S. Carr, 1F (UASM), 12.VII.24, F.S. Carr, 1F (UASM), 18.II.24, F.S. Carr, 1F (UASM), 8.V.65, J.L. Carr, 1F (JLCC); Township 14, Range 13, West 4 Meridian, 22.VII.79, J.L. Carr, 1M (JLCC); Ghost Dam, 2.VII.80, B.F. and J.L. Carr, 1F (JLCC); Seebe, 7.VI.73, B.F. and J.L. Carr, 1M (JLCC); Suffield, 20.IV.23, 1F (UASM); Township 6, Range 1, West 4 Meridian, 20.V.80, B.F. and J.L. Carr, 1F (JLCC); Township 10, Range 2, West 5

Meridian, 28.IV.81, B.F. and J.L. Carr, 1M (JLCC); Township 15, Range 14, West 4 Meridian, 9.V.80, B.F. and J.L. Carr, 1F (JLCC); 27mi. sw. Nordegg, 13.VI.66, R. Freitag/T.L. Erwin, 1F (CWOB); Calgary, IV.11, Criddle, 1F (CNCI); Lethbridge, 6.V.30, J.H. Pepper, 1M (CNCI). Northwest Territories: 1M (CNCI). Saskatchewan: Township 6, Range 2, West 3 Meridian, 13.VIII.73, B.F. and J.L. Carr, 1M (JLCC); Cypress Hills Provincial Park, 29.V.63, Cook et al., 1F (CNCI).

UNITED STATES OF AMERICA: Arizona: Coconino County, 2.4mi. n. Kaibab Lodge, 28.VII.71, Lawton/Willis, 2M (CWOB). Colorado: Grant/Jackson County, Rabbit Ears Pass, 2872 me., 19.VI.81, M. Kaulbars, 1M (RSAN); Top of Range below Sapello and Pecos Rivers, 11,000 ft., 1.VIII.00, H.F. Wickham, 1M (USNM); Cumbres Pass, 10,000 ft., 22.VI.35, E.C. Van Dyke, 1M (CASC); 1mi. n. Nederland, 11.VIII.73, C.W. O'Brien, 1M (CWOB); Longs Peak, 10-11,000 ft., 8.VII.26, E.C. Van Dyke, 1M (CASC); Argentine Road, H.F. Wickham, 1M (USNM); Kenosha Pass, 16.VII.38, J.W. Green, 1F (CASC); Gore Pass, 15.IX.48, O. Bryant, 1F (CASC); Leadville, 7-14.VII.96, H.F. Wickham, 1M (USNM). Montana: Helena, Hubbard and Schwartz, 2M (USNM); Flathead Lake, 12.VII.11, 1F (USNM); Gallatin County, 9.IV.32, 1M (MSU), 8.V.32, 1F (MSU), 1VI.31, 1F (MSU); Bozeman, 17.V.28, 1M (MSU), 11.V.25, 1M (MSU); 15mi. ne. Bozeman, 30.VI.72, Burleson, 1M (MSU); 6mi. e. Dell, 30.VI.54, Anderson, 1M (MSU). Nevada: White Pine County, 17.7mi. n., 3.5mi. nw. McGill, Monte Neva Hot Springs,

25.VII.71, Lawton and Willis, 1M, 1F (CWOB). New Mexico: 1F (MCZ); Cloudcroft, V.18, 1M (CASC). Utah: Alta, 29.VI.47, O. Bryant, 1F (CASC). Wyoming: 1F (USNM); Beulah, 21.VII., T.D.A. Cockerell, 1M (UASM); Park County, Beartooth Plateau, 6.VII.61, S.M. Sutton, 1M (UCM), 31.VIII.59, J.G. Edwards, 1F (CWOB); Beartooth Plateau, Hairgrass Canyon, 7.VII.59, 1F (CWOB); Bighorn Mountains, Medicine Mountain, 18.VIII.62, J.G. Edwards, 1F (CWOB); Tie Siding, VIII.25, 1F (USNM).

**Notes about synonymy.**--- An individual of this species from New Mexico was included in the original type series of *S. plumbeus* LeConte. *Stephanocleonus plumbeus* as herein recognized does not occur in New Mexico.

**Derivation of specific epithet.**--- From the Latin "*confusio*" meaning mixed or confused. This name is used in reference to the previous confusion of this species with *S. plumbeus*.

**Problems in recognition.**--- This is the only species of North American *Stephanocleonus* in which the elytra have small recurved hair-like scales on the dorsal surface and in which elytral intervals 3 and 5 are markedly elevated and convex throughout the greater part of their lengths. The internal sac of the male aedeagus is also distinctive (Fig. 43).

Description.--- Data about variation in LR, WF, WRA, LP, WPB, WEH, WELM, LEI, LEy, WEy, WEy/LEy, WPB/LP, WRA/LR, WRA/WF, WELM/LEI, and LP/LEI are presented in Tables 1 and 2. Size. Length, male, 9.2-15.0 mm; female, 10.5-14.8 mm. Width, male, 4.7-6.5 mm; female, 4.9-7.2 mm. Head. Frons with punctures large, sparse, shallow, not confluent; also with small broad appressed white scales, dense laterally, sparse medially. Some specimens with a slightly raised carina on vertex. Rostrum. Moderately robust, slightly more so in males (width at apex 0.850-0.947 times length in male; 0.810-0.909 in female) (Fig. 25). With well-developed, sharp, steeply declivous, low, median carina from above point of antennal insertion to base of rostrum; both basally and apically terminated at small, moderately deep fovea (Fig. 25). Rostrum flat from immediately laterad of median carina to lateral margins, lateral margins sharp from above point of antennal insertion to anterior margin of eye, indicated in a few specimens by a slightly raised rounded carina. Dorsal punctures large, sparse, shallow, not confluent. With scales small broad appressed white, dense laterally, sparse medially. Pronotum. Dorsal apical margin not or shallowly emarginate at middle. With low broad indistinct median carina in anterior one-half. Subapical constriction well defined laterally, not so dorsally; some specimens with shallow impressions laterad of median carina, but discontinuous with lateral portion of subapical constriction. Median basal area narrowly, moderately deeply

impressed. Dorsal punctures moderately large to large, dense and moderately deep; medially, the distance between punctures markedly less than the diameter of an individual puncture, some punctures confluent and irregularly impressed; laterally, the distance between punctures subequal to or greater than diameter of an individual puncture. Prosternum. With shallow impression anterior to each procoxal cavity, and with slightly to moderately developed transverse swelling anterior to each impression. Elytra. Robust in general form (width at midlength 0.644-0.718 times length in male; 0.624-0.708 in female) (Fig. 20). In dorsal view with lateral margins evenly arcuate from midlength to base, humerus rounded, not distinct (Fig. 20). Sutural interval and intervals 3 and 5 (in some specimens also basal one-half of interval 7) elevated and convex throughout the greater part of their lengths. Scales white small moderately dense; smaller and less dense in two posteromedially directed oblique patches, one at basal one-third from intervals 2 to 6, the other at apical one-third from intervals 2 to 4. With a slightly elevated triangular glabrous shiny area at apical one-quarter at confluence of intervals 4 to 6 (Fig. 20). With scattered very short recurved hair-like scales, most evident on declivity from sutural interval to interval 3. Wings. Short (more or less equal to one-half length of elytra). Legs. Foretibia of female with inner margin with small denticles in apical one-half; subapical tooth small to

moderately large, indistinct to distinct from apical denticles. Foretibial and mesotibial unci of both sexes moderately large, metatibial unci small. Tarsal claws connate in basal one-quarter to one-third, divergent, each with basal internal flange well-developed. Ventral tarsal pilose vestiture of foretarsus of male present as small elongate-narrow pads covering apical one-quarter on article 1, as large elongate-oval pads covering apical three-quarters on article 2, and as large elongate-oval pads covering more or less the entire ventral surface on article 3; of female, pilose vestiture lacking from article 1, lacking to present as elongate-narrow pads covering apical one-quarter to one-half on article 2, as elongate-oval pads covering apical one-half on article 3; of mesotarsus of male, lacking to as apical tufts on outer lobe on article 1, as elongate-narrow pads covering apical three-quarters on outer lobe and as apical tufts on inner lobe on article 2, as large elongate-oval pads covering apical three-quarters on article 3; of mesotarsus of female, lacking from article 1, lacking from to as elongate-narrow pads covering apical one-half on outer lobe on article 2, as small to large elongate-oval pads covering apical one-third to two-thirds on article 3; of metatarsus of male, lacking from articles 1 and 2, as elongate-oval pads covering apical one-quarter to three-quarters on article 3; of metatarsus of female, lacking from articles 1 to 3 to as elongate-oval pads covering apical one-third on article 3. Abdomen. Abdominal

sternum VII in female evenly rounded, not tumescent; in male with apical margin slightly emarginate medially. Genitalia. Female (four examined). Abdominal sternum VIII with basal arm expanded at apex; lateral arms narrow, slightly inwardly and evenly arcuate throughout length (Fig. 37). Male (12 examined). Abdominal sternum VIII with paired sclerite with inner apices truncate (Fig. 31c). Aedeagus elongate-narrow; in lateral view slightly and evenly arcuate throughout length, narrowed from approximately apical one-third to apex, apex slightly spatulate; in ventral view with apex symmetrical, medially produced into acuminate apical projection (Figs. 31a,b). Base of aedeagus with at most a small ventral tubercle. Internal sac elongate with large dorsally directed paired lobe on dorsal surface at apical one-third; median dorsal pocket high, with large laterally directed paramedial lobes at basal one-third near ventral margin (Fig. 43). In lateral view with basal paired sclerite and apical sclerite complex widely separated (Fig. 43a).

**Geographic variation.**--- None noted.

**Geographic distribution.**--- This species is known from southern Alberta and southern Saskatchewan, south to Arizona and New Mexico (Fig. 204). Most published records of *S. cristaticollis* refer to this species.

**Natural history.**--- Adults of this species have been

collected in prairie habitats in southern Alberta to alpine habitats at elevations of 3400 m in Colorado and New Mexico.

**Chorological relationships.**--- This species is broadly sympatric with *S. cristaticollis* and *S. parshus*, and narrowly sympatric with *S. immaculatus* along the western limits of the range of the latter.



Table 1. Descriptive statistics for *S. confusus* based on eight males from Medicine Hat, Alberta

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.60-2.20	1.900	0.278	0.131
WF	1.60-2.20	1.938	0.299	0.141
WRA	1.50-2.00	1.713	0.246	0.116
LP	2.40-3.40	2.913	0.477	0.225
WPB	3.00-4.30	3.725	0.698	0.329
WE1H	3.80-5.30	4.638	0.810	0.382
WE1M	4.60-6.40	5.550	0.935	0.441
LE1	6.90-9.90	8.350	1.425	0.672
LEy	1.30-1.70	1.513	0.246	0.116
WEy	0.60-0.80	0.763	0.112	0.053
B. Proportions.				
WEy/LEy	0.46-0.57	0.506	0.058	0.027
WPB/LP	1.23-1.40	1.278	0.080	0.038
WRA/LR	0.85-0.95	0.902	0.060	0.028
WRA/WF	0.81-0.94	0.886	0.059	0.027
WE1M/LE1	0.64-0.72	0.665	0.036	0.017
LP/LE1	0.33-0.36	0.349	0.017	0.008

Table 2. Descriptive statistics for *S. coronatus* based on eight females from Medicine Hat, Alberta.

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.80-2.30	2.113	0.233	0.110
WF	1.80-2.40	2.150	0.289	0.136
WRA	1.50-2.00	1.838	0.265	0.125
LP	2.70-3.50	3.213	0.396	0.186
WPB	3.30-4.60	4.113	0.682	0.321
WE1H	4.20-5.80	5.175	0.768	0.362
WE1M	5.00-7.00	6.163	0.948	0.447
LE1	7.80-10.10	9.325	1.048	0.493
LEy	1.40-1.80	1.613	0.187	0.088
WEy	0.70-0.90	0.800	0.113	0.053
B. Proportions.				
WEy/LEy	0.47-0.53	0.496	0.029	0.013
WPB/LP	1.22-1.35	1.278	0.070	0.033
WRA/LR	0.81-0.91	0.869	0.064	0.030
WRA/WF	0.81-0.90	0.855	0.042	0.020
WE1M/LE1	0.62-0.71	0.660	0.042	0.019
LP/LE1	0.33-0.36	0.344	0.013	0.006

(Figs. 26, 32, 38, 44, 205)

*Stephanocleonus cristatus* LeConte 1876a:147 [not Chevrolat 1873:98]. Holotype (examined), female, labelled "Utah", "S./ *cristatus*/ Lec", "Horn Coll/ H8514" and with my label indicating it as the holotype "*Stephanocleonus/ cristatus* LeConte/ HOLOTYPE labelled/ by Anderson" (MCZC). Type locality, Utah. LeConte 1878 (distn., misident. of *S. confusus*). Henshaw 1881-1882 (check.). Henshaw 1885 (check.). Leng 1920 (in part; catal., misident. *Stephanocleonus confusus*).

*Cleonus (Stephanocleonus) cristatus*; Casey 1891 (key, misident. *Stephanocleonus confusus*). Csiki 1934 (catal.).

*Cleonus (Stephanocleonus) cristaticollis* Csiki 1934:26. New name for *Stephanocleonus cristatus* LeConte.

*Cleonis cristatus*; O'Brien and Wibmer 1982 (catal., distn., as jr. homonym).

*Cleonis cristaticollis*; O'Brien and Wibmer 1982 (in part; catal., distn., misident. *Stephanocleonus confusus*).

Problems in recognition.--- This is one of two brachypterous species of North American *Stephanocleonus*. Unlike *S. confusus* individuals, in which elytral intervals 3 and 5 are elevated and convex throughout the greater part of their lengths, individuals of *S. cristaticollis* have only the

They also do not possess the short recurved hair-like scales on the dorsal surface of the elytra that are found in *S. confusus*. Abdominal sternum VII of females has a pair of low rounded swellings, and the shape of the male aedeagus and form of the internal sac are distinctive (Figs. 32, 44).

**Description.**--- Data about variation in LR, WF, WRA, LP, WPB, WE1H, WE1M, LE1, LEy, WEy, WEy/LEy, WPB/LP, WRA/LR, WRA/WF, WE1M/LE1, and LP/LE1 are presented in Tables 3 and 4. **Size.** Length, male, 13.8 mm; female, 13.6-14.1 mm. Width, male, 6.4 mm; female, 6.4-6.6 mm. **Head.** Frons with punctures large, sparse, shallow, indistinct; also with small elongate appressed white scales, dense laterally, sparse medially. **Rostrum.** Moderately robust (width at apex 0.857 times length in male; 0.905 in female) (Fig. 26). With well-developed, sharp, steeply declivous, low, median carina from above point of antennal insertion to base of rostrum; both basally and apically terminated at small, moderately deep fovea (Fig. 26). Rostrum with shallow trough immediately laterad of median carina, lateral margins sharp from above point of antennal insertion to anterior margin of eye, indicated by a slightly raised rounded carina. Dorsal punctures large, sparse, shallow, indistinct. With scales small, elongate, appressed, white, dense laterally, sparse medially. **Pronotum.** Dorsal apical margin shallowly emarginate at middle. With low broad indistinct median carina in anterior

one-half. Subapical constriction well defined laterally, not so dorsally; with shallow impressions laterad of median carina, but discontinuous with lateral portion of subapical constriction. Median basal area broadly, moderately deeply impressed. Dorsal punctures small, sparse and shallow; medially and laterally with the distance between punctures subequal to or greater than diameter of an individual puncture; punctures irregularly impressed and dense in median basal area. Prosternum. With only shallow impression anterior to each procoxal cavity. Elytra. Moderately robust in general form (width at midlength 0.627 times length in male; 0.647 in female). In dorsal view with lateral margins evenly arcuate from midlength to base, humerus rounded, not distinct. Base of interval 3 swollen and convex, otherwise elytral intervals uniformly flat. Scales white small moderately dense; smaller and less dense in two posteromedially directed oblique patches, one at basal one-third from intervals 2 to 6, the other at apical one-third from intervals 2 to 4. With a slightly elevated triangular glabrous shiny area at apical one-quarter at confluence of intervals 4 to 6. Suberect or erect vestiture lacking. Wings. Short (more or less equal to one-half length of elytra). Legs. Foretibia of female with inner margin with small denticles in apical one-half; subapical tooth small, distinct from apical denticles. Foretibial and mesotibial unci of both sexes moderately large, metatibial unci small. Tarsal claws connate in basal one-quarter to one-third,

divergent, each with basal internal flange well-developed. Ventral tarsal pilose vestiture of foretarsus of male present as small elongate-oval pads covering apical one-third on article 1, as large elongate-oval pads covering apical three-quarters on article 2, and as large elongate-oval pads covering more or less the entire ventral surface on article 3; of female, pilose vestiture lacking from article 1, present as apical tufts on article 2, as large elongate-oval pads covering apical three-quarters on article 3; of mesotarsus of male, as elongate-narrow pads covering apical one-quarter on outer lobe, lacking from inner lobe on article 1, as elongate-narrow pads covering apical three-quarters on article 2, as large elongate-oval pads covering more or less entire ventral surface on article 3; of mesotarsus of female, lacking from articles 1 and 2, as elongate-narrow pads covering apical three-quarters on article 3; of metatarsus of male, lacking from articles 1 and 2, as elongate-oval pads covering apical three-quarters on article 3; of metatarsus of female, lacking from articles 1 to 3. **Abdomen.** Abdominal sternum VII in female with pair of rounded swellings at midlength; in male with apical margin slightly emarginate medially. **Genitalia.** Female (one examined). Abdominal sternum VIII with basal arm slightly expanded at apex; lateral arms broad, slightly inwardly and evenly arcuate throughout length (Fig. 38). Male (two examined). Abdominal sternum VIII with paired sclerite with inner apices acuminate (Fig. 32c). Aedeagus moderately

robust; in lateral view markedly arcuate, more so in basal one-half, narrowed gradually from midlength to apex, apex markedly spatulate; in ventral view with apex symmetrical, medially produced into elongate rounded apical projection (Figs. 32a,b). Base of aedeagus with a large ventral tubercle (Fig. 32a). Internal sac globose with dorsally directed median lobe on dorsal surface; with moderately large dorsolaterally directed paired lobe at midlength near dorsal margin (Fig. 44). In lateral view with basal paired sclerite and apical sclerite complex approximate (Fig. 44a).

**Geographic variation.**--- None noted.

**Geographic distribution.**--- This species is known from only four specimens: two males and a female from southern Alberta, and the female holotype from Utah (no further locality data given) (Fig. 205). Most published records of this species refer to *S. confusus*.

**Natural history.**--- No information is available about the natural history of this species.

**Chorological relationships.**--- The distribution of the few known specimens indicates that this species is probably broadly sympatric with *S. confusus* and narrowly sympatric with western *S. parshus*.

Table 3. Descriptive statistics for *S. cristaticollis*  
based on one male from Monarch, Alberta

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	2.10	2.100	-	-
WF	2.10	2.100	-	-
WRA	1.80	1.800	-	-
LP	3.30	3.300	-	-
WPB	4.10	4.100	-	-
WE1H	5.40	5.400	-	-
WE1M	7.40	6.400	-	-
LE1	10.20	10.200	-	-
LEy	1.70	1.700	-	-
WEy	0.80	0.800	-	-
B. Proportions.				
WEy/LEy	0.47	0.471	-	-
WPB/LP	1.24	1.242	-	-
WRA/LR	0.86	0.857	-	-
WRA/WF	0.86	0.857	-	-
WE1M/LE1	0.63	0.637	-	-
LP/LE1	0.32	0.324	-	-



Table 4. Descriptive statistics for *S. cristaticollis* based on one female from Medicine Hat, Alberta

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	2.10	2.100	-	-
WF	2.20	2.200	-	-
WRA	1.90	1.900	-	-
LP	3.40	3.400	-	-
WPB	4.30	4.300	-	-
WE1H	5.60	5.600	-	-
WE1M	6.60	6.600	-	-
LE1	10.20	10.200	-	-
LEy	1.80	1.800	-	-
WEy	0.80	0.800	-	-
B. Proportions.				
WEy/LEy	0.44	0.444	-	-
WPB/LP	1.27	1.265	-	-
WRA/LR	0.91	0.905	-	-
WRA/WF	0.86	0.864	-	-
WE1M/LE1	0.65	0.647	-	-
LP/LE1	0.33	0.333	-	-

4.3.2.3 *Stephanocleonus immaculatus* Anderson, new species

(Figs. 21, 27, 33, 39, 45, 202)

*Cleonus* (*Stephanocleonus*) *plumbeus*; Fall and Cockerell 1907  
(in part; check., misident.). Csiki 1934 (in part;  
catal., misident.).

*Stephanocleonus plumbeus*; Blatchley and Leng 1916 (in part;  
key, redesc., misident.). Leng 1919 (in part; check.,  
poss. misident.). Leng 1920 (in part; catal.,  
misident.). Danks 1981 (in part; check., poss.  
misident.).

*Cleonus plumbeus*; Leonard 1926 (in part; check., poss.  
misident.). Krombein 1979 (in part; prey, poss.  
misident.).

*Cleonis plumbeus*; O'Brien and Wibmer 1982 (in part; catal.,  
distn., misident.).

Type material.--- Holotype, male, with a red label  
"HOLOTYPE", "McMurray, / Alta. V.10.53/ W.J. Brown",  
"HOLOTYPE/ *Stephanocleonus/ Immaculatus/ Anderson*" and with  
abdomen on card and genitalia in microvial attached to pin  
(CNCI). Allotype, female, with a red label "ALLOTYPE",  
"Gillam Man./, 10.VI.1949/ J.B. Wallis", "ALLOTYPE/  
*Stephanocleonus/ Immaculatus/ Anderson*" and with abdomen on  
card and genitalia in microvial attached to pin (CNCI). Type  
locality, Fort McMurray, Alberta.

Paratypes. 10 males, 16 females. CANADA: Alberta: Township 93, Range 10, West 4 Meridian, 1.VI.85, B.F. and J.L. Carr, 1F (JLCC); McMurray, 21.VI.53, W.J. Brown, 1F (CNCI), 29.VII.53, G.E. Ball, 1M (CNCI), 22.VI.53, G.E. Ball, 1F (CNCI); Lac La Biche, NE shore, 30.VI.63, L.M. Kenakin, 1F (CWOB); Township 37, Range 18, West 5 Meridian, 18.VII.73, B.F. and J.L. Carr, 1F (JLCC); Township 35, Range 18, West 5 Meridian, 18.VIII.67, B.F. and J.L. Carr, 1F (JLCC); Exshaw, 27.VI.54, B.F. and J.L. Carr, 1M, 1F (JLCC); High Prairie, 26.VII.61, A.R. Brooks, 1F (CNCI). British Columbia: Canal Flats, 13.V.82, B.F. and J.L. Carr, 1M (JLCC). Manitoba: Gillam, 16.VI.50, F. McAlpine, 1F (CNCI); Aweme, 31.V.10, N. Criddle, 1F (USNM); Beaujou [sic], 24.VI.61, H.E. Milxiron, 1M (CNCI). Northwest Territories: Fort Smith, 17.V.50, J.B. Wallis, 1F (CNCI), 16.VI.50, W.G. Helps, 1F (CNCI). Ontario: Lake Superior, Whitefish Point, 46°38'N 84°33'W, Hubbard and Schwartz, 1M, 1F (USNM). Saskatchewan: Junction Highways 2 and 165 east, 22.VI.85, B.F. and J.L. Carr, 1M, 1F (JLCC); Hudson Bay, 9.VII.54, Brooks-Wallis, 1M, 1F (CNCI).

UNITED STATES OF AMERICA: Colorado: Douglas County, 19.IV.15, 1M (CASC). Minnesota: 1F (USNM). Nebraska: Lincoln, 29.VI., Wolcott, 1F (UNSM). South Dakota: Englewood, Haggard, 1M (UNSM).

Derivation of specific epithet.--- From the Latin "im"

meaning not, and "*macula*" meaning spot or mark. This name is used in reference to the lack of distinct elytral markings in most individuals of this species.

**Problems in recognition.**--- This is the only species of North American *Stephanocleonus* in which the elytra are more or less uniformly covered with small white scales and in which most specimens lack a distinct pattern of elytral maculations (Fig. 21). Most specimens also lack the small triangular shiny glabrous area at the confluence of intervals 4 to 6 which is found in all other North American *Stephanocleonus* (Fig. 21). The shape of the male aedeagus and form of the internal sac are also distinctive (Figs. 33, 45).

**Description.**--- Data about variation in LR, WF, WRA, LP, WPB, WE1H, WE1M, LE1, LEY, WEY, WEY/LEY, WPB/LP, WRA/LR, WRA/WF, WE1M/LE1, and LP/LE1 are presented in Tables 5 and 6. **Size.** Length, male, 13.3-15.6 mm; female, 14.8-16.8 mm. Width, male, 5.9-6.8 mm; female, 6.8-7.5 mm. **Head.** Frons with punctures large, moderately dense, moderately deep, many punctures longitudinally confluent giving a longitudinally rugose appearance; also with small broad appressed white scales, sparse laterally, absent to very sparse medially. Most specimens with a slightly raised carina on vertex. **Rostrum.** Moderately robust, slightly more so in females (width at apex 0.800-0.950 times length in

male; 0.792-0.955 in female) (Fig. 27). With well-developed, sharp, steeply declivous, high, median carina from above point of antennal insertion to base of rostrum; both basally and apically terminated at small, moderately deep fovea (Fig. 27). Rostrum flat from immediately laterad of median carina to lateral margins, lateral margins rounded to sharp from above point of antennal insertion to anterior margin of eye. Dorsal punctures large, moderately dense, moderately deep, many punctures confluent. With scales small uniformly dense, elongate-narrow appressed, white. Pronotum. Dorsal (apical margin not or shallowly emarginate at middle. With low broad distinct median carina in anterior one-half. Subapical constriction slightly to well defined laterally, not so dorsally; with shallow impressions laterad of median carina, but discontinuous with lateral portion of subapical constriction. Median basal area narrowly, shallowly impressed. Dorsal punctures moderately large, dense and moderately deep; medially, the distance between punctures less than the diameter of an individual puncture, some punctures confluent or nearly so and irregularly impressed; laterally, the distance between punctures subequal to diameter of an individual puncture. Prosterhum. With only very shallow impression anterior to each procoxal cavity. Elytra. Moderately robust in general form (width at midlength 0.584-0.684 times length in male; 0.561-0.670 in female) (Fig. 21). In dorsal view with lateral margins straight, slightly convergent from slightly past midlength

to base, humerus acute, very distinct (Fig. 21). All elytral intervals more or less flat. Scales white, small, dense, no distinct maculations present (Fig. 21) Some specimens with a slightly elevated, small, triangular, glabrous, shiny area at apical one-quarter at confluence of intervals 4 to 6. Suberect or erect vestiture lacking. Wings. Long (greater than length of elytra). Legs. Foretibia of female with inner margin with small denticles in apical one-half; subapical tooth moderately large, distinct from apical denticles. Foretibial and mesotibial unci of both sexes moderately large, metatibial uncus small. Tarsal claws connate in basal one-quarter to one-third, divergent, each with basal internal flange well-developed. Ventral tarsal pilose vestiture of foretarsus of male present as elongate-narrow pads covering apical one-quarter to one-half on article 1, as large elongate-oval pads covering apical three-quarters to more or less entire ventral surface on articles 2 and 3; of female, pilose vestiture lacking from article 1, as apical tufts to elongate-narrow pads covering apical one-half on article 2, as elongate-oval pads covering apical three-quarters on article 3; of mesotarsus of male, lacking from, to as an elongate-narrow pad covering apical one-third on outer lobe on article 1, as elongate-narrow pads covering apical three-quarters on article 2, as large elongate-oval pads covering apical three-quarters to more or less entire ventral surface on article 3; of mesotarsus of female, lacking from article 1, lacking from, to as an

elongate-narrow pad covering apical one-half on outer lobe on article 2, as large elongate-oval pads covering apical two-thirds on article 3; of metatarsus of male, lacking from articles 1 and 2, as elongate-narrow pads covering apical two-thirds on article 3; of metatarsus of female, lacking from articles 1 and 2, as apical tufts to small elongate-oval pads covering apical one-half on article 3.

**Abdomen.** Abdominal sternum VII in female evenly rounded, not tumescent; in male with apical margin slightly emarginate medially. **Genitalia.** Female (five examined). Abdominal sternum VIII with basal arm expanded at apex; lateral arms broad, inwardly arcuate at midlength (Fig. 39). Male (six examined). Abdominal sternum VIII with paired sclerite with inner apices rounded (Fig. 33c). Aedeagus elongate-narrow; in lateral view slightly and evenly arcuate throughout length, narrowed from approximately apical one-third to apex, apex slightly spatulate; in ventral view with apex symmetrical, medially produced into truncate apical projection (Figs. 33a,b). Base of aedeagus with at most a small ventral tubercle (Fig. 33a). Internal sac elongate with small laterally directed paired lobe near ventral margin at or slightly beyond midlength; median dorsal pocket high, with large dorsolaterally directed paired lobe at basal one-third near ventral margin, separated from base of median pocket by distinct trough (Fig. 45). In lateral view with basal paired sclerite and apical sclerite complex widely separated (Fig. 45a).

**Geographic variation.**--- None noted.

**Geographic distribution.**--- This species is known from widely scattered localities in the southwestern Northwest Territories, northern Manitoba, and central Ontario, south in the west to Colorado and Nebraska (Fig. 202).

**Natural history.**--- A single adult has been collected on *Rosa* sp. (Rosaceae).

**Chorological relationships.**--- This species is broadly sympatric throughout the northern part of its range with *S. plumbeus* and *S. parshus*, and is narrowly sympatric in the western part of its range with *S. confusus*.



based on eight males from throughout the species range

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	2.00-2.30	2.138	0.195	0.092
WF	1.80-2.00	1.950	0.113	0.053
WRA	1.60-2.00	1.850	0.196	0.092
LP	2.90-3.40	3.163	0.310	0.146
WPB	3.80-4.40	4.138	0.288	0.136
WE1H	4.80-5.80	5.400	0.488	0.230
WE1M	5.50-6.60	6.238	0.555	0.262
LE1	9.00-10.90	10.112	0.850	0.400
LEy	1.40-1.80	1.588	0.169	0.080
WEy	0.70-0.90	0.800	0.080	0.038
B. Proportions.				
WEy/LEy	0.50-0.53	0.504	0.018	0.008
WPB/LP	1.24-1.40	0.310	0.078	0.037
WRA/LR	0.80-0.95	0.866	0.069	0.033
WRA/WF	0.89-1.00	0.948	0.072	0.034
WE1M/LE1	0.58-0.64	0.617	0.030	0.014
LP/LE1	0.30-0.34	0.313	0.019	0.009

based on eight females from throughout the species range

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	2.20-2.40	2.275	0.133	0.063
WF	2.00-2.40	2.150	0.196	0.092
WRA	1.90-2.10	2.000	0.139	0.065
LP	3.20-3.60	3.438	0.226	0.106
WPB	4.20-4.70	4.425	0.328	0.155
WE1H	5.50-6.40	6.000	0.488	0.230
WE1M	6.30-7.30	6.788	0.522	0.246
LE1	9.40-12.00	11.075	1.254	0.591
LEy	1.50-1.80	1.675	0.155	0.073
WEy	0.80-1.00	0.900	0.113	0.053
B. Proportions.				
WEy/LEy	0.47-0.60	0.538	0.069	0.032
WPB/LP	1.19-1.44	1.289	0.121	0.057
WRA/LR	0.79-0.96	0.880	0.072	0.034
WRA/WF	0.88-1.05	0.932	0.081	0.038
WE1M/LE1	0.56-0.67	0.614	0.047	0.022
LP/LE1	0.30-0.35	0.311	0.027	0.012

4.3.2.4 *Stephanocleonus parshus* Anderson, new species

(Figs. 28, 34, 40, 46, 203)

*Cleonus* (*Stephanocleonus*) *plumbeus*; Casey 1891 (key). Csiki 1934 (in part; catal., misident.). Kissinger 1964 (biol., misident.).

*Stephanocleonus plumbeus*; Blatchley and Leng 1916 (in part; key, redesc., misident.). Leng 1919 (in part; check., poss. misident.). Leng 1920 (in part; catal., misident.). Marcovitch 1923 (biol., redesc., larvae, misident.). Danks 1981 (in part; check., poss. misident.).

*Cleonus plumbeus*; Leonard 1926 (in part; check., poss. misident.). Bleasdel 1937 (check., prob. misident.). Krombein 1979 (in part; prey, poss. misident.).

*Cleonis plumbeus*; O'Brien and Wibmer 1982 (in part; catal., distn., misident.).

Type material.--- Holotype, male, with a red label

"HOLOTYPE", "Chalk River, Ont./ May 16, 1938/ N.R. Brown",

"HOLOTYPE/ *Stephanocleonus/ parshus/ Anderson*" and with abdomen on card and genitalia in microvial attached to pin (CNCI). Allotype, female, with a red label "ALLOTYPE", "Tp.

36 Rge. 15/ W.5 Mer Alberta/ 16.VII.1973/ B&J Carr Lot 3",

"ALLOTYPE/ *Stephanocleonus/ parshus/ Anderson*" and with abdomen on card and genitalia in microvial attached to pin (CNCI). Type locality, Chalk River, Ontario.

Paratypes. 39 males, 43 females. CANADA: Alberta:  
 Township 24, Range 8, West 5 Meridian, 8.VII.82, B.F. and  
 J.L. Carr, 1M (JLCC); Bragg Creek, 31.V.75, F.A.H. Sperling,  
 1M, 1F (RSAN); Pincher Creek, 20.VI.61, H.E. Milliron, 1M  
 (CNCI); McMurray, 4.VI.53, W.J. Brown, 1M (CNCI), 6.VI.53,  
 W.J. Brown, 1F (CNCI); Township 25, Range 3, West 5  
 Meridian, 7.VI.73, B.F. and J.L. Carr, 2M (JLCC); Township  
 29, Range 5, West 5 Meridian, 28.VI.64, B.F. and J.L. Carr,  
 1M (JLCC); Calgary, 28.VI.64, B.F. and J.L. Carr, 1M (JLCC);  
 Exshaw, 27.VI.54, B.F. and J.L. Carr, 1M (JLCC), 19.V.60,  
 J.L. Carr, 1M (JLCC); Willow Creek, 9.VIII.28, H. Richmond,  
 1M (CASC); Ghost Dam, B.F. and J.L. Carr, 1F (JLCC);  
 Edmonton, 26.V.19, F.S. Carr, 1M, 1F (UASM), 14.V.20, F.S.  
 Carr, 1F (UASM), 13.V.18, F.S. Carr, 1M (UASM); Medicine  
 Hat, 13.VI.32, F.S. Carr, 2M, 1F (UASM); Crow's Nest Pass,  
 9.VI.30, J.H. Pepper, 1M (CNCI); Beaverlodge, 1F (UASM);  
 near Josephburg, 53°41'N 113°55'W, 23.V.79, K. Shaw and J.  
 Sutcliffe, 1F (UASM). British Columbia: Rolla, 21.VII.27,  
 Vroom, 1F (CASC); Pouce Coupe, 1.VII.27, Vroom, 1F (CASC);  
 Baynes Lake, 9.V.76, B.F. and J.L. Carr, 1F (JLCC); Canal  
 Flats, 13.V.82, B.F. and J.L. Carr, 1M, 1F (JLCC);  
 Skookumchuck, 30.V.84, B.F. and J.L. Carr, 1M (JLCC).  
 Manitoba: Aweme, 12.VIII.29, R.H. Handford, 1F (CNCI),  
 2.VII.29, N. Criddle, 1M (CNCI), 20.IV.06, N. Criddle, 1F  
 (CNCI), 31.V.10, N. Criddle, 1F (USNM), 9.VI.14, N. Criddle,  
 1M, 1F (CNCI), 24.IX.28, N. Criddle, 1M (CNCI), VI, N.  
 Criddle, 1M (USNM), 1.VII.05, N. Criddle, 1F (USNM), V.12,

N. Criddle, 1M (USNM), 3.VIII., N. Criddle, 1F (USNM);  
 Township 14, Range 105, 12.V.24, J.B. Wallis, 1F (CNCI);  
 Riding Mountain National Park, 12.VI.38, W.J. Brown, 1F  
 (CNCI); Melita, 8.VI.20, N. Criddle, 1F (CNCI); Shell River,  
 VII.27, E. Criddle, 1F (CNCI); Winnipeg, 1F (USNM);  
 Treesbank, 24.VI.48, Criddle, 1F (CNCI); Sandilands,  
 22.VI.30, W.J. Brodie, 1F (CNCI); 24km. w. Haddashville,  
 27.VI.84, I. Askevold, 1M (CWOB). Northwest Territories:  
 Fort Wrigley, 27.IX.29, O. Bryant, 1M (CASC); Fort Simpson,  
 Manners Creek, 11.VI.72, Smetana, 1F (CNCI). Ontario:  
 Blackburn, 6.VI.32, W.J. Brown, 1F (CNCI); Ridgeville,  
 VI.35, S.D. Hicks, 1F (CNCI); Toronto, 1M (USNM); Dunrobin,  
 18.V.77, J.E. O'Hara, 1M (UASM). Quebec: Kazubazua, 25.V.33,  
 W.J. Brown, 1M (CNCI); Fort Coulonge, 25.VI.19, J. Beaulne,  
 1M (CNCI). Saskatchewan: 8mi. e. Saskatoon, 17.VI.73, C.K.  
 Starr, 1F (CWOB); Township 35, Range 6, West 3 Meridian,  
 20.VII.85, B.F. and J.L. Carr, 1M (JLCC); Pike Lake, 5.I.42,  
 King & Glen, 1F (CNCI); Prince Albert, 4.VII.54,  
 Brooks-Wallis, 1F (CNCI).

UNITED STATES OF AMERICA: Connecticut: New Hartford,  
 26.IV.20, Zappe, 1F (CWOB). Iowa: Ames, 1F (USNM). Maine: 1M  
 (USNM). Massachusetts: Woodshole, 1F (CASC). Michigan: Emmet  
 County, Mackinaw City, 28.VI.20, M.H. Hatch, 1M (USNM).  
 Minnesota: 1F (CASC); Crow Wing County, Pelican Lake,  
 25.VIII.10, Wolcott, 1F (UNSM), 28.VIII.08, Wolcott, 1F  
 (UNSM). Montana: Missoula, 16.V.04, 1F, 11.VII.04, 1M  
 (USNM); Helena, Hubbard and Schwartz, 1F (USNM); Flathead

County, Echo Lake, 31.V.35, Eichmann, 1F (MSU); Hamilton, 16.VIII.26, 1M (MSU); Lake County, 6.VII.36, 1M (MSU).  
 Nebraska: Lincoln, VI., 1F (UNSM); Bellevue, 20.V.02, Bruner, 1M (UNSM). New Mexico: 1F (MCZC). South Dakota: Englewood, Haggard, 1F (UNSM). Tennessee: Knoxville, 12.IV.55, H. and A. Howden, 1F (HAHC), 17.V.57, H. and A. Howden, 1M (HAHC). Vermont: Bennington County, G.H. Horn, 1M (USNM); Lyndon, 22.VIII.00, Melander, 1M, 1F (OCUC).  
 Country unknown: Bridge, 31.V.14, 1F (CNCI).

**Derivation of specific epithet.**--- An arbitrary combination of letters.

**Problems in recognition.**--- Among macropterous North American *Stephanocleonus* species with elytral maculations, individuals of *S. parshus* can be recognized by the large deep and dense pronotal punctures, the distance between punctures distinctly less than the diameter of a puncture (Fig. 28a). As in individuals of *S. plumbeus*, the pronotum has the lateral subapical constriction not defined dorsally and has the apical margin entire and not emarginate. In addition to other features, females of *S. parshus* can further be distinguished from those of other North American macropterous species by the metatarsus with article 3 with elongate pilose pads; males can be distinguished from males of other species by the distinctive shape of the aedeagus

and form of the internal sac (Figs. 34, 46).

**Description.**--- Data about variation in LR, WF, WRA, LP, WPB, WE1H, WE1M, LE1, LEy, WEy, WEy/LEy, WPB/LP, WRA/LR, WRA/WF, WE1M/LE1, and LP/LE1 are presented in Tables 7 and 8. **Size.** Length, male, 10.4-12.0 mm; female, 10.9-13.6 mm. Width, male, 4.8-5.6 mm; female, 5.0-5.9 mm. **Head.** Frons with punctures large, dense, moderately deep, many punctures confluent giving a longitudinally rugose appearance; also with small elongate-narrow appressed white, dense scales. Most specimens with a slightly raised carina on vertex. **Rostrum.** Moderately elongate-narrow, slightly more so in males (width at apex 0.762-0.842 times length in male; 0.773-0.842 in female) (Fig. 28). With well-developed, sharp, gradually declivous, high, median carina from above point of antennal insertion to base of rostrum; both basally and apically terminated at small, shallow fovea (Fig. 28). Rostrum rounded to flat from immediately laterad of median carina to lateral margins, lateral margins rounded to sharp from above point of antennal insertion to anterior margin of eye, indicated in very few specimens by slightly raised rounded carina. Dorsal punctures large, moderately dense, moderately deep, many confluent. With scales small, elongate-narrow, appressed, white, dense. **Pronotum.** Dorsal apical margin not or shallowly emarginate at middle. With low, broad, distinct median carina in anterior one-half. Subapical constriction slightly to well defined laterally,

not so dorsally; with shallow impressions laterad of median carina, but discontinuous with lateral portion of subapical constriction. Median basal area narrowly, shallowly impressed. Dorsal punctures large, dense and deep; medially, most confluent and irregularly impressed; laterally, distance between punctures much less than diameter of a puncture (Fig. 28a). Prosternum. With shallow impression anterior to each procoxal cavity, and with or without slightly developed transverse swelling anterior to each impression. Elytra. Moderately robust in general form (width at midlength 0.610-0.682 times length in male; 0.591-0.646 in female) (Fig. 22). In dorsal view with lateral margins straight and convergent from midlength to base, humerus obtuse, distinct. All elytral intervals more or less uniformly flat. Scales white small moderately dense; smaller and less dense in two posteromedially directed oblique patches, one at basal one-third from intervals 2 to 6, the other at apical one-third from intervals 2 to 4. With slightly elevated triangular glabrous shiny area at apical one-quarter at confluence of intervals 4 to 6. Suberect or erect vestiture lacking. Wings. Long (equal to or greater than length of elytra). Legs. Foretibia of female with inner margin with small denticles in apical one-half; subapical tooth small to moderately large, indistinct to distinct from apical denticles. Foretibial and mesotibial unci of both sexes moderately large, metatibial uncus small. Tarsal claws connate in basal one-quarter to one-third, divergent, each



with basal internal flange well-developed. Ventral tarsal pilose vestiture of foretarsus of male present as elongate-oval pads covering apical one-half on article 1, as large elongate-oval pads covering apical three-quarters on article 2, and as large elongate-oval pads covering more or less the entire ventral surface on article 3; of female, pilose vestiture lacking from article 1, present as elongate-oval pads covering apical one-half on article 2, as elongate-oval pads covering more or less entire ventral surface on article 3; of mesotarsus of male, lacking from to present as an apical tuft on outer lobe on article 1, as elongate-narrow pads covering apical three-quarters on article 2, as large elongate-oval pads covering more or less entire ventral surface on article 3; of mesotarsus of female, lacking from article 1, lacking from to present as an elongate-narrow pad covering apical one-half on outer lobe on article 2, as large elongate-oval pads covering more or less entire ventral surface on article 3; of metatarsus of male, lacking from articles 1 and 2, as elongate-oval pads covering apical one-third to one-half on article 3; of metatarsus of female, lacking from articles 1 and 2, as an elongate-oval pad covering apical one-third to one-half on outer lobe and apical one-half to two-thirds on inner lobe on article 3. **Abdomen.** Abdominal sternum VII in female evenly rounded, not tumescent; in male with apical margin slightly emarginate medially. **Genitalia.** Female (three examined). Abdominal sternum VIII with basal arm expanded at

apex; lateral arms narrow, inwardly arcuate at midlength (Fig. 40). Male (12 examined). Abdominal sternum VIII with paired sclerite with inner apices rounded (Fig. 34c). Aedeagus elongate-narrow; in lateral view slightly and evenly arcuate throughout length, narrowed from approximately apical one-third to apex, apex slightly spatulate; in ventral view with apex symmetrical, medially produced into rounded apical projection (Figs. 34a,b). Base of aedeagus with at most a small ventral tubercle. Internal sac elongate with large dorsally directed paired lobe on dorsal surface immediately beyond midlength; median dorsal pocket lacking, with large dorsolaterally directed paired lobe at basal one-third near dorsal margin (Fig. 46). In lateral view with basal paired sclerite and apical sclerite complex widely separated (Fig. 46a).

**Geographic variation.**--- None noted.

**Geographic distribution.**--- This species is widely distributed in North America from Maine and Massachusetts south to Tennessee, west to Nebraska and Iowa in the south, British Columbia and Alberta in the north (Fig. 203). A single state record for New Mexico is in the Horn collection at the Museum of Comparative Zoology, Harvard University.

**Natural history.**--- At Knoxville, Tennessee, Marcovitch (1923) records larvae of this species feeding on roots of

strawberry in June, July, December and March. Eggs were found on July 13 and a single adult emerged on July 25. Although identified as *S. plumbeus*, the description provided notes that the thorax is coarsely punctured. This, and the verified occurrence of only *S. parshus* at Knoxville, indicates that this is the species to which he referred.

**Chorological relationships.**--- This, the most widespread species of North American *Stephanocleonus*, is sympatric in the northern part of its range with *S. plumbeus*; in the northeastern part of its range with *S. immaculatus*; and, in the western part of its range with *S. cristaticollis* and *S. confusus*.

eight males from southern Manitoba and Ontario

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.70-2.10	1.850	0.196	0.013
WF	1.50-1.90	1.675	0.192	0.091
WRA	1.30-1.60	1.488	0.187	0.088
LP	2.50-3.00	2.725	0.275	0.130
WPB	3.00-3.80	3.363	0.453	0.214
WE1H	3.80-5.30	4.450	0.717	0.338
WE1M	4.50-5.80	5.025	0.825	0.388
LE1	7.20-8.60	7.913	0.909	0.428
LEy	1.20-1.50	1.350	0.179	0.084
WEy	0.60-0.80	0.688	0.125	0.059
B. Proportions.				
WEy/LEy	0.46-0.58	0.509	0.060	0.028
WPB/LP	1.15-1.31	1.233	0.079	0.037
WRA/LR	0.76-0.84	0.804	0.055	0.026
WRA/WF	0.81-0.94	0.889	0.071	0.034
WE1M/LE1	0.61-0.68	0.634	0.042	0.020
LP/LE1	0.33-0.37	0.345	0.016	0.007

eight females from southern Manitoba and Ontario

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.70-2.20	1.913	0.233	0.110
WF	1.60-1.90	1.738	0.178	0.084
WRA	1.40-1.70	1.550	0.160	0.076
LP	2.60-3.30	2.875	0.399	0.188
WPB	3.20-4.00	3.625	0.466	0.220
WE1M	4.20-5.40	4.838	0.708	0.334
WE1M	4.70-6.20	5.400	0.765	0.360
LE1	7.50-9.60	8.613	1.132	0.534
LEy	1.30-1.60	1.425	0.175	0.082
WEy	0.60-0.80	0.738	0.112	0.053
B. Proportions.				
WEy/LEy	0.46-0.54	0.517	0.041	0.019
WPB/LP	1.18-1.36	1.262	0.076	0.036
WRA/LR	0.77-0.84	0.812	0.041	0.020
WRA/WF	0.88-0.94	0.892	0.030	0.014
WE1M/LE1	0.59-0.65	0.627	0.028	0.013
LP/LE1	0.32-0.35	0.334	0.018	0.008

*Cleonus obliquus*; LeConte 1850 (check., misident.).

*Stephanocleonus plumbeus* LeConte 1876a:146. Lectotype (here designated), male, one of an undetermined number of syntypes, labelled with a pale blue circle (=Lake Superior), "Type/ 5259", and with card with abdomen attached, microvial containing genitalia, and my designation label "*Stephanocleonus plumbeus* LeC./ LECTOTYPE/ desig. Anderson" (MCZC). Type locality, north shore of Lake Superior. LeConte 1876a (in part; desc., misident. *Stephanocleonus confusus*, mixed type series). Henshaw 1881-1882 (in part; check., misident. *Stephanocleonus confusus*). Henshaw 1885 (in part; check., misident. *Stephanocleonus confusus*). Wickham 1902 (check., misident. *Stephanocleonus confusus*). Wickham 1909 (check., misident. *Stephanocleonus confusus*). Blatchley and Leng 1916 (in part; key, redesc., misident. *Stephanocleonus immaculatus*, *Stephanocleonus parshus*). Leng 1919 (in part; check., poss. misident. *Stephanocleonus immaculatus*, *Stephanocleonus parshus*, *Stephanocleonus confusus*, *Stephanocleonus stenothorax*). Leng 1920 (in part; catal., misident. *Stephanocleonus immaculatus*, *Stephanocleonus parshus*, *Stephanocleonus confusus*). Marcovitch 1923 (biol., redesc., larvae, misident. *Stephanocleonus parshus*). Danks 1981 (in part; check.,

*Stephanocleonus stenothorax*).

*Cleonus* (*Stephanocleonus*) *plumbeus*; Casey 1891 (key). Fall and Cockerell 1907 (check., misident. *Stephanocleonus immaculatus*, *Stephanocleonus confusus*). Csiki 1934 (in part; catal., misident. *Stephanocleonus immaculatus*, *Stephanocleonus parshus*, *Stephanocleonus confusus*).

*Cleonus plumbeus*; Leonard 1926 (in part; check., poss. misident. *Stephanocleonus immaculatus*, *Stephanocleonus parshus*). Bleasdel 1937 (check., poss. misident. *Stephanocleonus parshus*). Kissinger 1964 (biol., misident. *Stephanocleonus parshus*). Krombein 1979 (in part; as prey, poss. misident. *Stephanocleonus immaculatus*, *Stephanocleonus parshus*, *Stephanocleonus confusus*).

*Cleonis plumbeus*; O'Brien and Wibmer 1982 (in part; catal., distn., misident. *Stephanocleonus immaculatus*, *Stephanocleonus parshus*, *Stephanocleonus confusus*).

**Problems in recognition.**--- Among North American *Stephanocleonus* species that are macropterous and possess maculated elytra, this species can be recognized by the following combination of characters: pronotum with the lateral subapical constriction not defined dorsally; the apical margin of the pronotum entire, not emarginate at

an individual puncture" (Fig. 29a). Females of this species have the metatarsus with article 3 lacking ventral pilose pads.

Individuals of this species are most likely to be confused with those of *S. parshus*, but the two can be readily separated by characters noted in the key and above.

Notes about synonymy.--- I have been unable to ascertain the exact constitution of the type series of *S. plumbeus* LeConte. The species was described from an unspecified number of specimens from the north shore of Lake Superior and from New Mexico. In the LeConte collection at the Museum of Comparative Zoology, there are three specimens labelled as types; two specimens with pale blue circles attached (= Lake Superior), and a single specimen labelled "N.M." (= New Mexico). I have selected one of the two Lake Superior specimens as lectotype and believe that these three specimens likely initially constituted the type series of *S. plumbeus*. The other specimen from Lake Superior is conspecific with the lectotype, but the specimen labelled "N.M." is *S. confusus*.

Other specimens in the LeConte and Horn collections, but not labelled as types and therefore questionably part of the type series, includes a specimen of *S. immaculatus* with



no locality data, two specimens of *S. plumbeus* with no locality data, and a specimen of *S. parshus* from New Mexico.

**Description.**--- Specimens examined. 13 males, 11 females. Data about variation in LR, WF, WRA; LP, WPB, WELH, WELM, LE1, LEy, WEy, WEy/LEy, WPB/LP, WRA/LR, WRA/WF, WELM/LE1, and LP/LE1 are presented in Tables 9 and 10. Size. Length, male, 10.9-13.0 mm; female, 11.2-15.6 mm. Width, male, 4.9-6.0 mm; female, 5.2-7.2 mm. **Head.** Frons lacking distinct large punctures; uniformly covered with small elongate-narrow appressed white scales. Some specimens with slightly raised carina on vertex. **Rostrum.** Moderately elongate-narrow to moderately robust; (width at apex 0.824-0.895 times length in male; 0.818-0.944 in female) (Fig. 29). With well-developed, sharp, gradually declivous, high, median carina from above point of antennal insertion to base of rostrum; basally terminated at small, shallow fovea (Fig. 29a). Rostrum rounded from immediately laterad of median carina to lateral margins, lateral margins rounded from above point of antennal insertion to anterior margin of eye. Dorsally lacking distinct large punctures. With small elongate-narrow appressed white, dense scales. **Pronotum.** Dorsal apical margin not or shallowly emarginate at middle. With sharp distinct median carina in anterior one-half. Subapical constriction slightly defined laterally, not so dorsally; with shallow impressions laterad of median carina, but discontinuous with lateral portion of subapical

constriction. Median basal area broadly, shallowly to moderately deeply impressed. Dorsal punctures uniformly small, sparse and shallow, distance between punctures greater than diameter of one puncture (Fig. 29a).

**Prosternum.** With shallow impression anterior to each procoxal cavity, and slightly developed transverse swelling anterior to each impression. **Elytra.** Moderately robust in general form (width at midlength 0.600-0.635 times length in male; 0.566-0.615 in female) (Fig. 22). In dorsal view with lateral margins straight and convergent from midlength to base, humerus obtuse, distinct (Fig. 22). All elytral intervals more or less uniformly flat. Scales white small moderately dense; smaller and less dense in two posteromedially directed oblique patches, one at basal one-third from intervals 2 to 6, the other at apical one-third from intervals 2 to 4 (Fig. 22). With a slightly elevated triangular glabrous shiny area at apical one-quarter at confluence of intervals 4 to 6. Suberect or erect vestiture lacking. **Wings.** Long (equal to or greater than length of elytra). **Legs.** Foretibia of female with inner margin with small denticles in apical one-half; subapical tooth small to moderately large, indistinct to distinct from apical denticles. Foretibial and mesotibial unci of both sexes moderately large, metatibial uncus small. Tarsal claws connate in basal one-quarter to one-third, divergent, each with basal internal flange well-developed. Ventral tarsal pilose vestiture of foretarsus of male present as

elongate-oval pads covering apical one-third on article 1, as large elongate-oval pads covering apical three-quarters on article 2, and as large elongate-oval pads covering more or less the entire ventral surface on article 3; of female, pilose vestiture lacking from articles 1 and 2, present as elongate-oval pads covering apical three-quarters on article 3; of mesotarsus of male, lacking from to as an apical tuft on outer lobe on article 1, as elongate-narrow pads covering apical three-quarters on article 2, as large elongate-oval pads covering apical three-quarters on article 3; of mesotarsus of female, lacking from articles 1 and 2, as small elongate-narrow pads covering apical one-third on article 3; of metatarsus of male, lacking from articles 1 and 2, as elongate-oval pads covering apical three-quarters on article 3; of metatarsus of female, lacking from articles 1 to 3. **Abdomen.** Abdominal sternum VII in female evenly rounded, not tumescent; in male with apical margin slightly emarginate medially. **Genitalia.** Female (three examined). Abdominal sternum VIII with basal arm expanded at apex; lateral arms narrow, inwardly arcuate at midlength (Fig. 41). Male (seven examined). Abdominal sternum VIII with paired sclerite with inner apices truncate (Fig. 35c). Aedeagus elongate-narrow; in lateral view markedly arcuate, more so in basal one-half, narrowed from approximately apical one-third to apex, apex slightly spatulate; in ventral view with apex asymmetrical, produced into rounded apical projection (Figs. 35a,b). Base of aedeagus with at

most small ventral tubercle. Internal sac elongate with small dorsolaterally directed paired lobe on dorsal surface immediately beyond midlength; median dorsal pocket high, with large dorsolaterally directed paired lobe at basal one-third near ventral margin, separated from base of median pocket by trough (Fig. 47). In lateral view with basal paired sclerite and apical/sclerite complex widely separated (Fig. 47a).

Geographic variation.--- None noted.

Geographic distribution.--- This species is known from widely scattered localities from Newfoundland west to Alberta (Fig. 205).

Natural history.--- No information is available about the natural history of this species.

Chorological relationships.--- This species is sympatric throughout the central and western part of its range with *S. parshus* and *S. immaculatus*. Individuals of these three species have been collected within a few days of each other at Fort McMurray, Alberta.

Table 9. Descriptive statistics for *S. plumbeus* based on eight males from Rivière-au-Tonnerre, Québec

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.70-1.90	1.844	0.109	0.048
WF	1.60-1.80	1.722	0.125 <sup>u</sup>	0.055
WRA	1.40-1.70	1.567	0.130	0.058
LP	2.50-3.00	2.700	0.281	0.125
WPB	3.00-3.60	3.356	0.368	0.164
WE1H	4.10-5.20	4.600	0.618	0.275
WE1M	4.60-5.60	5.067	0.586	0.260
LE1	7.40-9.00	8.233	0.891	0.396
LEy	1.20-1.50	1.356	0.170	0.075
WEy	0.70-0.90	0.767	0.106	0.047
B. Proportions.				
WEy/LEy	0.50-0.64	0.567	0.068	0.030
WPB/LP	1.20-1.32	1.243	0.062	0.028
WRA/LR	0.82-0.90	0.849	0.038	0.017
WRA/WF	0.88-0.94	0.910	0.045	0.020
WE1M/LE1	0.60-0.64	0.615	0.015	0.007
LP/LE1	0.31-0.34	0.328	0.012	0.005

Table 10. Descriptive statistics for *S. plumbeus* based on eight females from Rivière-au-Tonnerre, Québec

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.80-2.40	2.100	0.287	0.145
WF	1.90-2.00	1.971	0.073	0.037
WRA	1.70-2.00	1.786	0.160	0.081
LP	2.80-3.30	3.057	0.272	0.137
WPB	3.60-4.20	3.886	0.351	0.177
WE1H	4.90-6.00	5.386	0.551	0.277
WE1M	5.40-6.60	5.857	0.567	0.286
LE1	9.10-11.00	9.957	1.017	0.512
LEy	1.30-1.60	1.486	0.182	0.092
WEy	0.70-0.80	0.757	0.080	0.040
B. Proportions.				
WEy/LEy	0.44-0.57	0.511	0.063	0.032
WPB/LP	1.22-1.32	1.272	0.054	0.027
WRA/LR	0.82-0.94	0.853	0.065	0.033
WRA/WF	0.85-1.00	0.906	0.068	0.034
WE1M/LE1	0.57-0.62	0.589	0.030	0.015
LP/LE1	0.29-0.33	0.307	0.021	0.010

4.3.2.6 *Stephanocleonus stenothorax* Anderson, new species

(Figs. 23, 30, 36, 42, 48, 204)

*Stephanocleonus plumbeus*; Leng 1919 (in part; check., poss. misident.). Danks 1981 (in part; check., poss. misident.).

Type material.--- Holotype, male, with a red label "HOLOTYPE", "YT. Bluefish caves/ 67°08'N 140°48'W/ 2000' 4.vii.1983/R.J. Cannings", "HOLOTYPE/ *Stephanocleonus/ stenothorax/* Anderson" and with abdomen on card and genitalia in microvial attached to pin (CNCI). Allotype, female, with a red label "ALLOTYPE", "YUKON: Dog. Ck./ 30.VII.77/ R.E. Roughley", "ALLOTYPE/ *Stephanocleonus/ stenothorax/* Anderson" and with abdomen on card and genitalia in microvial attached to pin (CNCI). Type locality, Bluefish Caves archaeological site, Yukon Territory.

Paratypes. 12 males, 11 females. Canada: Yukon Territory, Keele Range, Bluefish Caves archaeological site, 67°08'N 140°47'W, 1983, J. Cinq-Mars, 12M, 11F. (CNC, CWOB, JVM, RSAN, USNM, ZIL.)

Derivation of specific epithet.--- From the Greek "stenos" meaning narrowed, and "thorax" meaning breastplate. This name is used in reference to the apically narrowed form of

the pronotum.

Problems in recognition.--- Among North American species of *Stephanocleonus* with maculate elytra, individuals of this species are recognized by the pronotum with the lateral subapical constriction continued and well-defined dorsally, and with the apical margin shallowly emarginate at the middle. The glabrous shiny tubercle at the confluence of elytral intervals 4 to 6 is markedly elevated and lateral arms of abdominal sternum VIII of the female are very broad and medially approximate to contiguous near their apices (Fig. 42). The shape of the aedeagus of the male is also distinctive (Figs. 36a,b).

Description.--- Data about variation in LR, WF, WRA, LP, WPB, WE1H, WE1M, LE1, LEy, WEy, WEy/LEy, WPB/LP, WRA/LR, WRA/WF, WE1M/LE1, and LP/LE1 are presented in Tables 11 and 12. Size. Length, male, 12.5-13.1 mm; female, 12.8-14.9 mm. Width, male, 5.3-6.2 mm; female, 5.6-6.2 mm. Head. Frons with punctures large, moderately dense and deep, many punctures confluent in some specimens, surface thus longitudinally rugose; uniformly covered with small broad appressed white scales, dense laterally, sparse medially. Most specimens with slightly raised carina on vertex. Rostrum. Robust, (width at apex 0.900-0.950 times length in male; 0.900-1.000 in female) (Fig. 30). With well-developed, sharp, steeply declivous, low, median carina from above



apically terminated at small, shallow fovea (Fig. 30a).  
Rostrum flat from immediately laterad of median carina to lateral margins, lateral margins rounded to sharp from above point of antennal insertion to anterior margin of eye. Dorsally with punctures large moderately dense and deep, many confluent in some specimens, surface thus longitudinally rugose. With small elongate-narrow appressed white, scales; dense laterally, sparse medially. Pronotum. Dorsal apical margin broadly and moderately deeply emarginate at middle. With sharp distinct median carina in anterior one-half. Subapical constriction well defined laterally and dorsally; with distinct moderately deep impressions laterad of median carina continuous with lateral portion of subapical constriction. Median basal area broadly and moderately deeply impressed. Dorsal punctures small, sparse and shallow laterally and medially (except basally at middle where punctures are dense, deep, approximate and irregularly impressed), distance between punctures subequal to or greater than diameter of one puncture. Prosternum. With shallow impression anterior to each procoxal cavity, and with or without slightly to moderately developed transverse swelling anterior to each impression. Elytra. Moderately elongate-narrow in general form (width at midlength 0.566-0.600 times length in male; 0.554-0.608 in female) (Fig. 23). In dorsal view with lateral margins straight and convergent from midlength to base, humerus

less uniformly flat. Scales white small moderately dense; smaller and less dense in two posteromedially directed oblique patches, one at basal one-third from intervals 2 to 6, other at apical one-third from intervals 2 to 4 (Fig. 23). With markedly elevated triangular glabrous shiny area at apical one-quarter at confluence of intervals 4 to 6. Suberect or erect vestiture lacking. Wings. Long (equal to or greater than length of elytra). Legs. Foretibia of female with inner margin with small denticles in apical one-half; subapical tooth moderately large, distinct from apical denticles. Foretibial and mesotibial unci of both sexes moderately large, metatibial uncus small. Tarsal claws connate in basal one-quarter to one-third, divergent, each with basal internal flange well-developed. Ventral tarsal pilose vestiture of foretarsus of male present as elongate-oval pads covering apical one-half on article 1, as large elongate-oval pads covering apical two-thirds on article 2, and as large elongate-oval pads covering more or less the entire ventral surface on article 3; of female, pilose vestiture lacking from articles 1 and 2, present as elongate-oval pads covering apical one-half on article 3; of mesotarsus of male, lacking from to as an apical tuft on outer lobe on article 1, as elongate-narrow pads covering apical two-thirds on outer lobe and apical one-half on inner lobe on article 2, as large elongate-oval pads covering apical three-quarters on article 3; of mesotarsus of female,

lobe on article 3; of metatarsus of male, lacking from articles 1 and 2, as elongate-oval pads covering apical two-thirds on article 3; of metatarsus of female, lacking from articles 1 to 3. **Abdomen.** Abdominal sternum VII in female evenly rounded, not tumescent; in male with apical margin slightly emarginate medially. **Genitalia.** Female (two examined). Abdominal sternum VIII with basal arm expanded at apex; lateral arms very broad, medially approximate or contiguous near apices, inwardly arcuate at midlength (Fig. 42). Male (four examined). Abdominal sternum VIII with paired sclerite with inner apices truncate (Fig. 36c). Aedeagus elongate-narrow; in lateral view slightly and evenly arcuate throughout length, narrowed from approximately apical one-third to apex, apex not spatulate; in ventral view with apex symmetrical, medially rounded but not produced into distinct apical projection (Figs. 36a,b). Base of aedeagus with at most small ventral tubercle. Internal sac elongate with small dorsolaterally directed paired lobe on dorsal surface at midlength; median dorsal pocket moderately high, with moderately large dorsolaterally directed paired lobe at basal one-third near dorsal margin (Fig. 48). In lateral view with basal paired sclerite and apical sclerite complex widely separated (Fig. 48a).

**Geographic variation.**--- None noted.

Natural history.--- Most specimens of the type series were collected in unbaited pitfall traps set in a partly forested limestone upland and fell-field. They were collected in association with *Lepidophorus lineaticollis* Kirby and *Vitavitus thullis* Kissinger and several species of dry-tundra Carabidae.

Chorological relationships.--- This species is allopatric with all other *Stephanocleonus* species.

Site, Yukon Territory

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.80-2.10	1.975	0.133	0.063
WF	1.80-2.10	2.000	0.139	0.065
WRA	1.70-1.90	1.813	0.096	0.045
LP	2.60-2.90	2.825	0.155	0.073
WPB	3.50-3.90	3.700	0.196	0.093
WE1H	4.60-5.40	5.025	0.365	0.172
WE1M	5.30-6.10	5.688	0.388	0.183
LE1	9.00-10.40	9.713	0.745	0.351
LEy	1.60-1.80	1.713	0.125	0.059
WEy	0.80-0.90	0.875	0.069	0.033
B. Proportions.				
WEy/LEy	0.47-0.56	0.512	0.042	0.020
WPB/LP	1.24-1.35	1.310	0.051	0.024
WRA/LR	0.90-0.95	0.918	0.036	0.017
WRA/WF	0.86-0.95	0.907	0.044	0.020
WE1M/LE1	0.57-0.60	0.586	0.017	0.008
LP/LE1	0.28-0.32	0.291	0.020	0.009

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.80-2.10	1.988	0.169	0.080
WF	1.90-2.20	2.088	0.169	0.080
WRA	1.70-2.00	1.888	0.169	0.080
LP	2.70-3.20	2.988	0.233	0.110
WPB	3.50-4.20	3.925	0.356	0.168
WE1H	4.70-5.80	5.388	0.575	0.271
WE1M	5.40-6.40	6.088	0.450	0.212
LE1	9.60-11.50	10.512	0.930	0.438
LEy	1.50-1.90	1.750	0.212	0.100
WEy	0.80-0.90	0.863	0.078	0.037

B. Proportions.

WEy/LEy	0.47-0.53	0.494	0.031	0.015
WPB/LP	1.27-1.36	1.314	0.042	0.020
WRA/LR	0.90-1.00	0.950	0.040	0.019
WRA/WF	0.86-0.95	0.904	0.038	0.018
WE1M/LE1	0.55-0.61	0.580	0.030	0.014
LP/LE1	0.27-0.30	0.284	0.012	0.006

#### 4.4 Genus *Apleurus* Chevrolat

*Apleurus* Chevrolat 1873:78. Gender, masculine. Type species *Apleurus fossus* Chevrolat (= *Cleonus lutulentus* LeConte) by subsequent designation (Casey 1891:186). Chevrolat 1873 (in part; catal., sp. desc.). Faust 1904 (key).

Notes about synonymy.--- See "Notes about synonymy" section under subgenus *Apleurus*.

Diagnosis.--- Adult Cleoninae with moderately elongate-narrow to very robust body form (Figs. 24, 68-80). Eye elongate teardrop shaped, slightly to moderately prominent and convex in dorsal view (Figs. 81-88). Rostrum with median carina lacking to variously developed (Figs. 84-86). Pronotum punctate, anterolateral margin of pronotum straight to with variously developed acute postocular projection; postocular vibrissae long, of unequal length and longest behind base of eye (Figs. 81-88). Disk with elongate white moderately large moderately dense scales present in lateral stripe of variable width, small and fine in moderately broad to very broad apically narrowed median stripe, and variously small and fine to moderately large and elongate along lateral margins; median area largely black in color, underlying dark cuticle not obscured by overlying scales (Figs. 24, 68-80). Antenna with funiculus with

article 1 longer than, to more or less equal in length to, article 2; article 2 more or less as long as wide, to slightly longer than wide (Figs. 81-88). Ventral tarsal pilosity various but reduced to some extent (not covering entire ventral surface of a tarsal article) to lacking entirely from at least more basal tarsal articles (especially of metatarsus). Tibia with corbel ridge sharp (Fig. 6). Prosternum with swellings absent or present, if present, located immediately anterior to each prosternal impression (Fig. 7). Female with abdominal sternum VIII lacking basal arm (Figs. 89-96).

**Description.**--- **Size.** Small to large, moderately elongate-narrow to very robust in body form. **Mouthparts.** Prementum flat to slightly swollen, lacking or with up to five large setae on each side (Figs. 61-63). Maxillary palpus with palpifer with large seta, stipes with or without large seta (Figs. 64-65). Labial palpi separated by a distance variable from subequal to, to twice width of basal article of a labial palpus (Figs. 61-63). **Rostrum.** Moderately to markedly robust, not medially tumescent (Figs. 81-88). Median carina lacking to low, moderately rounded to sharp (Figs. 81-88). Epistoma not swollen, produced anteriorly, with apical margin emarginate (Fig. 81b) or rounded medially (Figs. 82b-88b). Antenna with funiculus with article 1 subequal in length to, to slightly longer than article 2 (Figs. 81a-88a); apical three articles of



club with placoidal sensillae. Head. Eye elongate teardrop shaped, slightly to moderately prominent and convex in dorsal view (Figs. 81-88). Area behind eye with moderately deep irregular punctures. Upper margin of eye rounded to sharp, frons convex to more or less flat. Vestiture. Dorsum with suberect or erect vestiture lacking to very long and dense; with simple elongate-narrow appressed white scales of variable size and density. Prothorax. Dorsal surface of pronotum punctate. Pronotum with median basal area shallowly to deeply impressed; anterolateral margin, behind eyes, straight to with variously developed acute postocular projection; postocular vibrissae long, of unequal length and longest behind base of eye (maximum length greater than or equal to one-half width of eye) (Figs. 81a-88a); disk with elongate white moderately large moderately dense scales present in lateral stripe of various width, small and fine in moderately broad to very broad apically narrowed median stripe, and variously small and fine to moderately large and elongate along lateral margins; median area largely black in color, underlying dark cuticle not obscured by overlying scales (Figs. 24, 68-80). Prosternum with slight to moderately deep impression anterolaterad of each procoxal cavity; with or without slight to markedly developed swelling immediately anterior to each prosternal impression (Fig. 7). Legs. Foretarsus moderately broad, articles 2 and 3 more or less subequal in length, at most only slightly longer than broad; article 1 only slightly longer than

articles 2 or 3; article 3 moderately deeply bilobed. Meso- and especially metatarsus slightly more elongate-narrow, article 2 slightly to distinctly longer than article 3; article 1 distinctly longer than articles 2, or 3; article 3 moderately deeply bilobed. Ventral tarsal pilosity various but reduced to some extent (not covering entire ventral surface of a tarsal article) to lacking entirely from at least more basal tarsal articles (especially of metatarsus). Claws variously connate from near base to through basal one-third, slightly to markedly divergent. Foretibia with inner margin with at most only small denticles in apical one-half; near apex with second spur very slightly to moderately well-developed. Metatibia of male with ventral margin of uncus evenly rounded. Tibia with corbel ridge sharp (Fig. 6). Wings. Absent or present (various in length). Elytra. Intervals, except humerus and bases of intervals 3 and 5 flat, to with sutural interval and intervals 3, 5, 7, and 9 variously slightly to markedly swollen and convex. Striae distinctly punctate. Humeri acute to rounded. Scale pattern various (Figs. 24, 68-80).

**Abdomen.** Ventral surface with small shiny glabrous patches, each with single small appressed scale to uniformly covered with fine dense hair-like scales. Abdominal sternum VII in males with apical margin at middle with or without small dorsally directed median tooth. Abdominal terga VII and VIII of female not, or slightly to markedly longitudinally carinate. **Genitalia.** Female. Abdominal sternum VIII lacking

basal arm (Figs. 89a-96a). Gonopocoxite II elongate triangular in form, apex not prolonged into marked lobe; stylus absent or present, small to moderately large in size; apical to slightly anteapical in position (Figs. 89b-96b).

Spermathecal gland round (Fig. 60). Male. Abdominal sternum VIII with interior angle of each sclerite lacking basal projection to with basal projection slightly to markedly developed (Figs. 97f, 101e, 103e). Aedeagus moderately robust, in lateral view more or less evenly arcuate throughout length; apex not spatulate (Figs. 97d, 98c-104c). Internal sac various; with dorsal median pocket low to high; apical and dorsal median pockets individually distinct; various lobes present, lobe A absent (Figs. 97a,b-104a,b). Apical sclerite complex with individual sclerites scythe-like (Figs. 97c, 98d-104d).

**Comparisons.**--- Species of *Apleurus* are most likely to be confused with the Palearctic *Chromoderus* and the South African *Cneomdontus*. Members of these three genera either lack pronotal postocular lobes entirely or possess variously developed acute postocular projections. *Apleurus* species are separated from species of these other two genera by the ventral tarsal pilosity reduced in extent and also by characters in the male genitalia (see "Phylogenetic Analysis" section). Species of *Pleurocleonus* also lack a pronotal postocular lobe but are readily separated from *Apleurus* by the possession of a low broad longitudinally

sulcate median rostral carina, flat eyes, and characters of male and female genitalia.

In the New World, some *Apleurus* species may be confused with large *Cleonidius* species which lack a pronotal postocular lobe. Consultation of diagnoses and key characters will readily separate the two.

**Checklist of included species.**--- Eight species, assigned to two subgenera, are included in this endemic New World genus. These species are:

Subgenus *Gibbostethus* Anderson, new subgenus

1. *A.(G.) hystrix* (Fall)

Subgenus *Apleurus* Chevrolat

2. *A.(A.) aztecus* (Champion)
3. *A.(A.) lutulentus* (LeConte)
4. *A.(A.) porosus* (LeConte)
5. *A.(A.) angularis* (LeConte)
6. *A.(A.) jacobinus* (Casey)
7. *A.(A.) albovestitus* (Casey)
8. *A.(A.) saginatus* (Casey)

**Phylogenetic relationships.**--- See "Phylogenetic analysis" section.

#### 4.4.1 Subgenus *Gibbostethus* Anderson, new subgenus

*Dinocleus*; Fall 1913 (in part; sp. desc.). Leng 1920 (in part; catal.).

*Cleonus* (*Dinocleus*); Csiki 1934 (in part; catal.).

*Cleonis*; O'Brien and Wibmer 1982 (in part; catal., distn.).

Derivation of name.--- Derived from the Greek "*gibbos*" meaning protuberant or swollen, and "*stethos*" meaning chest or breast. This name is used in reference to the tumescent mesosternal process of members of the single species placed in this subgenus.

Type species.--- *Dinocleus hystrix* Fall by monotypy.

Diagnosis.--- *Apleurus* with moderately elongate-narrow body form (Fig. 24). Dorsal erect vestiture dense and long. Prementum with bases of labial palpi separated by distance more or less equal to twice width of basal article of labial palpus (Fig. 61). Maxillary palpus with palpifer with large seta (Fig. 65). Rostrum with epistoma with apical margin emarginate medially (Fig. 81b). Mesosternum with mesosternal process markedly tumescent and convex (Fig. 58). Female with abdominal sternum VIII with lateral arms arcuate (Fig. 89a); gonocoxite II with stylus moderately large, apical in position (Fig. 89b). Male with abdominal sternum VIII with each sclerite with rounded interior angles, basal

projections, lacking (Fig. 97f); aedeagus in dorsal view markedly laterally expanded from midlength to apical one-third (Fig. 97e).

**Included species.**--- *Apleurus (Gibbostethus) hystrix* Fall from southern California is the only species in the subgenus.

**Phylogenetic relationships.**--- This subgenus is sister to the remainder of *Apleurus*. Recognition as a distinct subgenus is warranted by the primitive phylogenetic position of the species and the sister-group relationship with the remainder of *Apleurus*, by a restricted and likely relictual distribution (Fig. 212), and by various autapotypic character states and character states exhibited by *Gibbostethus* which although interpreted as evolved independently, are homoplasious in derived *Apleurus* species (see "Phylogeny" section).

#### 4.4.1.1 Species accounts

##### 4.4.1.1.1 *Apleurus (Gibbostethus) hystrix* (Fall), new combination

(Figs. 24, 58, 61, 65, 81, 89, 97, 212)

*Dinocleus hystrix* Fall 1913:41. Holotype (examined), male, labelled "Pasadena/ Cal.", "Feb.", "Not in/ Lec.Coll.",

"TYPE", "M.C.Z./ Type/ 25191", "H.C.FALL/ COLLECTION",  
"Dinocleus/ hystrix/ Fall" (MCZC). Type locality,  
Pasadena, California. Leng 1920 (catal.).

*Cleonus* (*Dinocleus*) *hystrix*; Csiki 1934 (catal.).

*Cleonis hystrix*; O'Brien and Wibmer 1982 (catal., distn.).

**Problems in recognition.**--- A distinctive and easily recognized species, *A. hystrix* is characterized by the markedly tumescent mesosternal process (Fig. 58) and the rostrum with the apical margin of the epistoma emarginate medially (Fig. 81b). The markedly laterally expanded aedeagus of males is also characteristic of this species (Fig. 97e).

Although it is unlikely that *A. hystrix* may be confused with other species of *Apleurus*, individuals may be confused with species of *Cleonidius* because of their small size and more elongate general body form (Fig. 24). Among other characters, *A. hystrix* individuals can be separated from *Cleonidius* individuals by the presence in the former of low rounded swellings on the prosternum immediately anterior to the prosternal impressions (Fig. 7). Swellings may or may not be present in *Cleonidius* species, but if present are situated immediately anterior to the procoxal cavities, adjacent to the prosternal impression (Fig. 8).

**Description.**--- Specimens examined. 18 males, 11 females.

Data about variation in LR, WF, WRA, LP, WPB, WPT, WELM, LE1, WPB/LP, WPT/WPB, WRA/LR, WRA/WF, WELM/LE1, and LP/LE1 are presented in Tables 13 and 14. Size. Length, male, 7.0-9.5 mm; female, 8.0-10.6 mm. Width, male, 2.8-3.5 mm; female, 3.0-4.2 mm. **Head.** Eye very prominent and convex in dorsal view: Frons and vertex with dense, large, irregularly impressed, deep punctures. Frons also with dense, very long erect white hair-like scales. Area immediately behind posterior margin of eye with very large, deep, irregularly impressed punctures. Area above eyes slightly to moderately elevated above rest of frons (eyes appearing browed in anterior view). Width of frons greater than width at apex of rostrum. **Rostrum.** Moderately robust (width at apex 0.692-0.852 times length in male; 0.703-0.816 in female) (Fig. 81). Median carina absent. Dorsal and lateral punctation dense, large, deep and irregularly impressed, some punctures longitudinally confluent medially. Dorsally, excluding epistoma, with dense, very long, erect white hair-like scales and dense very elongate-narrow recumbent white scales. In lateral view only slightly declivous from point of antennal insertion to apex (Fig. 81a). Epistoma with apical margin shallowly emarginate at middle (Fig. 81b). **Mouthparts.** Maxillary palpus with stipes lacking large seta on outer margin (Fig. 65). Labial palpi separated by more or less twice width of base of a labial palpus (Fig. 61). Prementum with one pair of large setae (Fig. 61). **Pronotum.** In dorsal view with lateral margins subparallel to



slightly divergent from base to apical one-quarter; constricted at oblique angle anterior to apical one-quarter; apical one-quarter and base subequal in width to slightly wider at apical one-quarter; distinct lateral tubercles not evident (Fig. 24). Dorsal punctation large to very large, dense and deep; punctures sparser smaller and shallower laterally and on flanks. Scales white, elongate-narrow, recumbent, sparse and small or lacking medially from disk (except for midline and along apical margin), uniformly dense laterally, sparser at lateral margins and on flanks (Fig. 24). Median carina lacking. Dorsally covered throughout with dense very long erect white hair-like scales each situated within a large puncture. Anterolateral margin with postocular projection absent or at most very slightly developed (Fig. 81a). Prosternum, With very shallow impression anterior to each procoxal cavity and with moderately developed rounded swelling anterior to each prosternal impression (Fig. 81a). Elytra. Moderately elongate-narrow in general form (width at midlength 0.526-0.655 times length in males; 0.543-0.641 in females) (Fig. 24). In dorsal view with lateral margins straight to very slightly arcuate from apical one-third to humerus; moderately and evenly arcuate from apical one-third to apex (Fig. 24). Humerus rounded, indistinct. Dorsally with dense, very long erect hair-like scales. Sutural interval, and intervals 3 and 5 very slightly elevated and convex, especially basally. Scales elongate-narrow, recumbent,

white, uniformly moderately dense except absent to sparse small and fine on interval 4 at basal one-third and apical one-third (Fig. 24) Wings. Absent. Legs. Foretibia of female with inner margin with small denticles in apical one-third; subapical tooth small, indistinct from tibial denticles. Foretibial, mesotibial and metatibial unci of both sexes large. Tarsal claws not to very slightly divergent, basal internal flange distinct. Ventral tarsal pilose vestiture of foretarsus of male and female present as small rounded apical pads on article 1, as small rounded pads covering apical one-third of article 2 and apical one-half on article 3; of mesotarsus of male present as apical tufts to small rounded apical pads on article 1, as small rounded apical pads to small rounded pads covering apical one-third of article 2 and apical one-third to one-half on article 3; of mesotarsus of female present as apical tufts on article 1, as small rounded apical pads on article 2, and as rounded pads covering apical one-third on article 3; of metatarsus of male and female present as apical tufts on article 1, as small rounded apical pads on article 2, and as rounded pads covering apical one-third on article 3. **Mesosternum.** Mesosternal process markedly swollen and tumescent (Fig. 58). **Abdomen.** Ventral surface with moderately dense, elongate-fine recumbent white scales and short suberect white hair-like scales. Abdominal sterna lacking distinct glabrous patches. Apex of abdominal sternum VII of male lacking dorsally directed tooth. Abdominal tergum VIII

uniformly convex, apical margin slightly reflexed. Genitalia. Female (two examined). Abdominal sternum VIII with lateral arms narrow, straight and divergent in basal one-half, moderately and evenly inwardly arcuate from midlength to apex (Fig. 89a). Gonocoxite II with stylus moderately large, slightly anteapical in position (Fig. 89b). Male (two examined). Abdominal sternum VIII with paired sclerite with inner apices lacking ventral projections (Fig. 97f). Aedeagus elongate-narrow, in lateral view thickest at midlength; in ventral view markedly laterally expanded at approximately midlength (Figs. 97d,e). Internal sac short and high, markedly apically deflexed; median dorsal pocket high, with moderately large dorsolaterally directed paramedial lobe at midheight and midlength and small laterally directed lobes near basal margin at basal one-third (Figs. 97a,b). Eversible apical sclerite complex with paired scythe-like sclerites only slightly sclerotized, each with small median projection (Fig. 97c); adjacent ventral surface and basal portion of sides of apex of internal sac unsclerotized (Fig. 97a).

**Geographic variation.**--- None noted.

**Geographic distribution.**--- This species has the most restricted distribution of any species of *Apleurus*. It has been found only in southern California, primarily in Los Angeles County at the El Segundo sand dunes but also at

County (Fig. 212).

**Natural history.**--- This species appears to be restricted to sand dune habitats in the Pacific semi-desert region of southern California. Adults have been collected on *Chaenactis glabriuscula* DC., *Aplopappus ericoides* (Less.) H. & A. (Compositae); *Croton californicus* Muell.-Arg. (Euphorbiaceae); and, *Lupinus albifrons* Benth. (Leguminosae). Definite host plants have yet to be determined. Adults have been collected from February to July at elevations of from 33-313 m ( $\bar{X}=166$  m,  $N=3$ ) (Fig. 232).

**Chorological relationships.**--- This species is sympatric with extreme western *A. angularis* and *A. albovestitus*, and southern *A. jacobinus*.

**Phylogenetic relationships.**--- This species is the sister-group to the rest of *Apleurus*.

Table 13. Descriptive statistics for *A. hystrix* based on eight males from the El Segundo Sand Dunes, Los Angeles County, California

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.08-1.56	1.345	0.283	0.133
WF	0.88-1.24	1.105	0.181	0.085
WRA	0.88-1.12	1.040	0.132	0.062
LP	2.08-2.72	2.530	0.384	0.181
WPB	2.00-2.80	2.530	0.444	0.209
WPT	2.00-2.88	2.560	0.411	0.194
WE1M	2.40-3.68	3.280	0.615	0.290
LE1	4.56-6.24	5.540	0.918	0.433
B. Proportions.				
WPB/LP	0.96-1.06	0.999	0.050	0.024
WPT/WPB	1.00-1.04	1.013	0.028	0.013
WRA/LR	0.69-0.85	0.780	0.088	0.041
WRA/WF	0.87-1.05	0.945	0.090	0.043
WE1M/LE1	0.53-0.66	0.592	0.062	0.029
LP/LE1	0.42-0.47	0.457	0.024	0.011

Table 14. Descriptive statistics for *A. hystrix* based on eight females from the El Segundo Sand Dunes, Los Angeles County, California

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.36-1.64	1.500	0.124	0.059
WF	1.04-1.40	1.230	0.162	0.077
WRA	1.04-1.24	1.125	0.113	0.053
LP	2.48-3.28	2.800	0.345	0.163
WPB	2.48-3.20	2.840	0.333	0.157
WPT	2.56-3.28	2.860	0.325	0.153
WE1M	3.12-4.24	3.720	0.525	0.247
LE1	5.60-7.28	6.280	0.775	0.365
B. Proportions.				
WPB/LP	0.97-1.06	1.015	0.060	0.028
WPT/WPB	0.97-1.03 <sub>0</sub>	1.008	0.038	0.018
WRA/LR	0.70-0.82	0.751	0.058	0.028
WRA/WF	0.84-1.00	0.918	0.086	0.040
WE1M/LE1	0.54-0.64	0.592	0.047	0.022
LP/LE1	0.43-0.47	0.446	0.017	0.008

4.4.2 Subgenus *Apleurus* Chevrolat

*Cleonus*; LeConte 1858 (sp. desc.). LeConte 1859 (sp. desc.). Gemminger and von Harold 1871 (in part; catal.). Leng 1920 (in part; catal.). Bradley 1930 (in part; key). Blackwelder 1947 (in part; check.). Arnett 1960-1962 (in part; key, catal.). Tanner 1966 (in part; sp. redesc., biol.). Krombein 1979 (in part).

*Apleurus* Chevrolat 1873:78. Gender, masculine. Type species *Apleurus fossus* Chevrolat (= *Cleonus lutulentus* LeConte) by subsequent designation (Casey 1891:186). Chevrolat 1873 (in part; catal., sp. desc.). Faust 1904 (key).

*Centrocleonus* LeConte 1876a:145 [not Chevrolat 1873:62]. New synonymy. Type species *Cleonus angularis* LeConte, according to article 67 (h) of the International Code of Zoological Nomenclature. Gender, masculine. Henshaw 1881-1882 (check.). LeConte and Horn 1883 (key). Henshaw 1885 (check.). Wickham 1889. Horn 1894 (check.).

*Cleonopsis* LeConte 1876a:147. New synonymy. Gender, feminine. Type species *Cleonus pulvereus* LeConte (= *Cleonus lutulentus* LeConte) by monotypy. LeConte and Horn 1883 (key). Henshaw 1885 (check.). Wickham 1889. Wickham 1896 (check.).

*Cleonaspis* LeConte 1876a:153. New synonymy. Gender, feminine. Type species *Cleonus lutulentus* LeConte by monotypy. Henshaw 1881-1882 (check.). LeConte and Horn

1883 (key). Henshaw 1885 (check.). Wickham 1902 (check.).

*Stephanocleonus*; Henshaw 1881-1882 (in part; check.).

*Dinocleus* Casey 1891:176. New synonymy. Gender, masculine.

New name for *Centrocleonus* LeConte. Type species *Cleonus angularis* LeConte by subsequent designation (Faust 1904:190, error as "*Typus angularatus* Lac", subsequently listed, p. 274, as "*angularis* Lec").

Wickham 1896 (check.). Fall 1901 (check.). Wickham 1902 (check.). Champion 1902-1906 (sp. desc., notes). Casey 1904 (sp. desc.). Faust 1904 (key). Fall and Cockerell 1907 (check.). Leng 1920 (catal.). Bradley 1930 (key). Tanner 1934 (check.). Ting 1936 (morphol.). Bruhn 1947 (morphol.). Van Dyke 1953 (sp. desc.). Essig 1958 (biol.). Sanders 1960 (morphol.). Arnett 1960-1962 (key, catal.). Hatch 1971 (key).

*Cleonus (Apleurus)*; Casey 1891 (key). Champion 1902-1906 (sp. desc., redescs.). Fall and Cockerell 1907 (check.). Csiki 1934 (catal.).

*Cleonurus* Faust 1904:274. *Nomem nudum*, incorrectly attributed to LeConte 1876a:152.

*Cleonus (Dinocleus)*; Csiki 1934 (catal.). Blackwelder 1939 (check.). Kissinger 1964 (check.).

*Cleonus (Cleonopsis)*; Kissinger 1964 (check.).

*Cleonis*; O'Brien and Wibmer 1982 (in part; catal., distn.).

O'Brien and Wibmer 1984 (in part).



Notes about synonymy.--- The genus *Apleurus* was initially established by Chevrolat (1873) to include three new Mexican and southwestern United States species, and one previously described southwestern United States species. Three of these species, *A. boucardi*, *A. trivittatus* and *A. quadrilineatus*, were regarded as "espèces lyxiformes" by Chevrolat (1873:109) thereby indicating their similarity to members of the genus *Lixus* and focusing attention on their lack of similarity to *A. fossus*, the fourth member of the genus.

No type species was designated by Chevrolat although he (1873:80) stated in his treatment of *A. boucardi* that "le corps de cet insecte est plus ovalaire que chez le type". This suggests that he considered one of the other two species (*Cleonidius trivittatus* was not treated in detail but merely placed in *Apleurus* by Chevrolat), likely *A. fossus* (because the description of it immediately follows the generic description and it is the only species excluded from his "espèces lyxiformes" grouping) as the type species. *A. fossus* was subsequently clearly designated as the type species by Casey (1891:76).

LeConte (1876a) undertook the first comprehensive revision of North American Cleoninae, wherein he proposed three new genera; *Centrocleonus* (a junior homonym of *Centrocleonus* Chevrolat), *Cleonopsis* and *Cleonaspis*. Species of *Centrocleonus* LeConte were characterized by the supposed unique presence of prosternal spines in front of the procoxae. Both *Cleonopsis* and *Cleonaspis* were monotypic

genera distinguished from other Cleoninae, most notably *Cleonus*, primarily on the basis of features of the tarsi and antennae. I find LeConte's proposal of *Cleonopsis* and *Cleonaspis* puzzling for they are based on forms which I consider to be conspecific, differing only slightly in the extent of ventral tarsal pilosity, but otherwise not to the extent discussed by LeConte (1876a:144-145, in key) as characteristic (see also "Notes on synonymy" of *A. lutulentus*). Apparently LeConte (1876a) was not concerned that these two genera might be confused, for in his descriptions of each, only characters permitting separation from *Cleonus* are discussed.

LeConte (1876a) assigned the "espèces lyxiformes" of Chevrolat (1873) to *Cleonus* along with a number of newly described North American species which Casey (1891) subsequently placed in his new subgenus *Cleonidius*. No mention is made of *A. fossus* by LeConte (1876a) and it is likely that he did not see this species for it did not occur within the geographic area of immediate concern to him.

In the next major work on North American species, Casey (1891) recognized that the distribution of the character states considered diagnostic of *Apteurus* also necessitated inclusion of the species of *Cleonopsis* and *Cleonaspis*, and that these latter two taxa did not warrant separate generic group status. He did not however consider *Apteurus* as warranting any more than subgeneric status under *Cleonus*. *Dinocleus*, proposed as a new name for *Centrocleonus* LeConte,

and considered as of generic rank, included all of LeConte's *Centrocleonus* as well as many new species.

My use of *Apleurus* is decidedly broader than that of any of these authors, including not only those species regarded as *Apleurus* by Casey (1891) but also all those placed in *Dinocleus* by Casey (1891), all subsequently described *Dinocleus* species and lastly, *Cleonus aztecus* Champion. Support for this broader definition is seen in the reconstructed phylogeny of these species wherein species of *Dinocleus* are placed as no more than derived species of *Apleurus*. Recognition of *Dinocleus* as a distinct genus would make *Apleurus* paraphyletic, an undesirable result because *Apleurus* are insufficiently structurally or biologically distinct from *Dinocleus* to warrant recognition as a formal paraphyletic taxon. The broader definition of *Apleurus* employed herein was also indirectly suggested by Van Dyke (1953) in his description of *Dinocleus bryanti*. Because *D. bryanti* is clearly conspecific with forms then regarded by Casey (1891), and I suspect also by Van Dyke, as *Apleurus*, Van Dyke's placement of his new species suggests a broader definition of *Dinocleus* and implies recognition of the similarities of members of that genus to those previously assigned to *Apleurus* even though it is doubtful he examined representatives of the latter.

**Diagnosis.**--- Adult *Apleurus* with moderately robust to very

robust body form (Figs. 68-80). Dorsal suberect or erect vestiture absent to very long and dense. Prementum with labial palpi separated by distance subequal to width of basal article of labial palpus (Figs. 62-63). Maxillary palpus with palpifer and stipes each with large seta, or with only palpifer with large seta (Fig. 64). Rostrum with epistoma with apical margin rounded medially (Figs. 82b-88b). Mesosternum with mesosternal process flat to only slightly convex (Fig. 59). Female with abdominal sternum VIII with lateral arms straight or arcuate (Figs. 90a-96); gonocoxite II with stylus absent or present, small to moderately large, apical to antepical in position (Figs. 90b-96b). Male with abdominal sternum VIII with each sclerite with basal projection at interior angle slightly to markedly developed (Figs. 101e, 103e). Aedeagus in dorsal view more or less uniform in width throughout median portion of length.

**Included species.**--- Seven species are placed in the subgenus *Apleurus*. They are found in the southwestern United States of America south to southern Mexico.

**Phylogenetic relationships.**--- This subgenus is the monophyletic sister-group of the subgenus *Gibbostethus* (see "Phylogeny" section).

1. Elytra with sutural interval, intervals 3, 5, 7, and 9 markedly elevated and convex throughout the greater part of their lengths. Metathoracic wings absent.....2
- 1'. Elytra with all intervals flat or with sutural interval, intervals 3, 5, and 7 only very slightly elevated and convex. Metathoracic wings present, variable in length from short (approximately one-half length of elytra) to long (equal to or greater than elytra in length).....6
- 2(1). Prosternum lacking swelling or with at most a low rounded swelling anterior to each prosternal impression (Fig. 82a). Rostrum lacking median carina. Elytra very robust (width at midlength greater than 0.70 times length) (Fig. 68).....  
.....*A. aztecus* (Champion)
- 2'. Prosternum with prominent swelling anterior to each prosternal impression (Figs. 84a-88a). Rostrum with median carina (obscured in some specimens by large deep punctures) (Figs. 84-86). Elytra less robust (width at midlength less than 0.75 times length) (Figs. 70-73).....3
- 3(2'). Tarsal claws slender, widely divergent, each lacking basal internal flange, tapered evenly from base to apex (Fig. 66). Dorsal erect vestiture of pronotum and elytra short.....4

- with basal internal flange, tapered abruptly from midlength to apex (Fig. 67). Dorsal erect vestiture of pronotum and elytra short or long.....5
- 4(3). Abdomen ventrally with scales broad, moderately dense; at least sterna III and IV with distinct glabrous shiny patches, each with a single long and erect hair .....*A. angularis* (LeConte) (in part)
- 4'. Abdomen ventrally with scales fine and elongate, extremely dense; no glabrous patches present .....  
.....*A. porosus* (LeConte) (in part)
- 5(3'). Elytra with sutural interval, intervals 3, 5, 7, and 9 (especially 3 and 5 near base) markedly elevated and convex. Most specimens with dorsal vestiture of head, pronotum and elytra very long and erect and with rostrum with carina moderately well-developed (Fig. 86). Abdomen ventrally with scales fine and elongate, moderately dense; sterna III and IV with distinct glabrous shiny patches, each with single long and erect hair.....  
.....*A. jacobinus* (Casey)
- 5'. Elytra with sutural interval, intervals 3, 5, 7, and 9 only slightly elevated and convex. Dorsal vestiture of head, pronotum and elytra short and suberect. Rostrum with carina only slightly developed, almost entirely obscured in some specimens by large deep punctures (Fig. 84).

- elongate, extremely dense; no glabrous patches present.....*A. porosus* (LeConte) (in part)
- 6(1'). Pronotum laterally expanded immediately posterior to subapical constriction (giving a distinctly tuberculate appearance); with base and apical one-quarter approximately equal in width to distinctly widest at apical one-quarter (Figs. 72-73). Tarsal claws slender, widely divergent, each lacking basal internal flange and tapered evenly from base to apex (Fig. 66).....
- .....*A. angularis* (LeConte) (in part)
- 6'. Pronotum not laterally expanded immediately posterior to subapical constriction (not appearing distinctly tuberculate); with base and apical one-quarter approximately equal in width to distinctly widest at base (Figs. 69, 74-80). Tarsal claws robust, not widely divergent; each with basal internal flange and tapered abruptly from midlength to apex (as in Fig. 67).....7
- 7(6'). Tarsus of hind leg with venter of article 3 with large pubescent pad. Prosternum lacking swelling or with swelling anterior to each prosternal impression only slightly developed (Fig. 83a). Female with tergum VII not longitudinally carinate; gonocoxite II with stylus large (Fig. 91b) .....
- .....*A. lutulentus* (LeConte)

- of pubescence. Prosternum with prominent swelling anterior to each prosternal impression (Figs. 87-88). Female with tergum VII longitudinally carinate; gonocoxite II lacking stylus (Figs. 95-96) .....8
- 8(7'). Tarsus of fore- and middle leg with venter of article 3 with large pubescent pad.....
- .....*A. saginatus* (Casey) (in part; males)
- 8'. Tarsus of fore- and middle leg with venter of article 3 lacking large pubescent pad, with at most small apical tuft of pubescence.....9
- 9(8'). Pronotum with lateral margins with outer margins of large punctures swollen, shiny and glabrous (appearing as small distinct tubercles). Abdomen ventrally (especially sterna III and IV) with distinct large glabrous patches, each with single short and suberect scale situated in the center....
- .....*A. saginatus* (Casey) (in part; females)
- 9'. Pronotum with lateral margins with outer margins of large punctures not swollen, glabrous or shiny. Abdomen ventrally either lacking glabrous patches and with scales extremely dense; with small glabrous patches; or, with most scales abraded....
- .....*A. albvestitus* (Casey)



combination

(Figs. 68, 82, 90, 98, 210)

*Cleonus aztecus* Champion 1902-1906:99. Lectotype (here designated), male, one of two syntypes, labelled "Type", inverted "Sp. figured", "Refugio/ Durango/ Hoge.", "B.C.A. Col. IV.4./ *Cleonus/ aztecus/* Champ." and with my designation label "LECTOTYPE/ *Cleonus aztecus/* Champ. desig./ Anderson" (BMNH). Type locality, Refugio, Durango, Mexico.

*Cleonis aztecus*; O'Brien and Wibmer 1982 (catal., distn.).

Notes about synonymy.--- None.

Problems in recognition.--- Members of this species are most likely to be confused with brachypterous specimens of *A. lutulentus*. In these latter specimens, the alternate elytral intervals are slightly elevated and convex and the elytra are moderately robust (more so than in macropterous *A. lutulentus*). These individuals are sympatric with *A. aztecus* and although superficially similar, *A. aztecus* individuals can be distinguished by the tarsal claws, each lacking a basal internal flange (as in Fig. 66), lack of metathoracic wings, and lack of a dorsally directed tooth on the apical

other species of *Apieurus* in which alternate elytral intervals are elevated and convex. Key characters should readily serve to separate these species.

Description.--- Specimens examined. 16 males, 7 females. Data about variation in LR, WF, WRA, LP, WPB, WPT, WE1M, LE1, WPB/LP, WPT/WPB, WRA/LR, WRA/WF, WE1M/LE1, and LP/LE1 are presented in Tables 15 and 16. Size. Length, male, 5.6-9.4 mm; female, 5.2-9.4 mm. Width, male, 2.9-4.6 mm; female, 2.5-4.8 mm. Head. Eye prominent and convex in dorsal view. Frons and vertex with moderately dense, small to moderately large, deep punctures. Frons also with sparse, short suberect white hair-like scales immediately above eyes. Area immediately behind posterior margin of eye with large, deep, irregularly impressed punctures. Area above eyes slightly elevated above rest of frons (eyes appearing browed in anterior view). Width of frons greater than width at apex of rostrum. Rostrum. Robust (width at apex 0.789-1.000 times length in male; 0.778-0.909 in female) (Fig. 82). Median carina absent. Dorsal and lateral punctation moderately dense, small to large, deep. Dorsally, excluding epistoma, with moderately dense, short, suberect white hair-like scales and moderately dense elongate-narrow appressed white scales. In lateral view with apical portion flat to only very slightly declivous from point of antennal

insertion to apex (Fig. 82a). Epistoma with apical margin rounded at middle (Fig. 82b). Mouthparts. Maxillary palpus with stipes lacking large seta on outer margin (as in Fig. 65). Labial palpi separated by more or less width of basal article of labial palpus (as in Fig. 62-63). Prementum with one pair of large setae (as in Fig. 62). Pronotum. In dorsal view with lateral margins subparallel to slightly divergent from base to apical one-quarter; constricted at an oblique angle anterior to apical one-quarter; apical one-quarter and base subequal in width to slightly wider at apical one-quarter; no distinct lateral tubercles evident (Fig. 68). Dorsal and lateral punctation small, sparse to moderately dense and shallow; punctures sparser and smaller on flanks. Scales white, elongate-narrow, appressed, sparse and small or lacking medially from disk and dorsally from flanks (except for midline and area anterior to low transverse carina if present), dense laterally to lateral margins in anterior two-thirds and ventrally on flanks. Median carina lacking; some specimens with low median transverse carina or impunctate area at midlength. Dorsally with dense very short fine suberect white hair-like scales each situated within large puncture. Anterolateral margin with postocular projection absent or at most very slightly developed (Fig. 82a). Prosternum. With very shallow impression anterior to each procoxal cavity and with at most a low slightly developed rounded swelling anterior to each prosternal impression (Fig. 82a). Elytra. Very robust in

general form (width at midlength 0.701-0.896 times length in males; 0.718-0.800 in females) (Fig. 68). In dorsal view with lateral margins moderately arcuate throughout length (Fig. 68). Humerus rounded, indistinct. Dorsally with sutural interval and intervals 3, 5, 7, and 9 with multiple rows of short fine suberect hair-like scales; intervals 2, 4, 6, and 8 with only single row of similar hair-like scales. Sutural interval, and intervals 3, 5, 7, and 9 markedly elevated and convex throughout their lengths. Scales various in density and size; elongate-narrow, pattern various but with at least small glabrous or nearly so area immediately apicad of midlength on interval 4 (Fig. 68).

**Wings.** Absent. **Legs.** Foretibia of female with inner margin with small denticles in apical one-third; subapical tooth large, distinct from tibial denticles. Foretibial and mesotibial unci of both sexes large, metatibial uncus of both sexes small. Tarsal claws widely divergent, no basal internal flange present (as in Fig. 66). Ventral tarsal pilose vestiture of foretarsus of male present as small rounded apical pads on article 1, as elongate-oval pads covering apical one-half of article 2, and as large elongate-oval pads covering apical two-thirds of article 3; of foretarsus of female present as small rounded apical pads on articles 1 and 2, as rounded pads covering apical one-third to one-half of article 3; of mesotarsus of male present as minute apical tufts on articles 1 and 2, and as small rounded apical pads to small rounded pads covering

apical one-half on article 3; of mesotarsus of female absent to present as minute apical tufts on articles 1 and 2, and as small rounded apical pads on article 3; of metatarsus of male present as minute apical tufts on articles 1 and 2, and as minute to small apical tufts on article 3; of metatarsus of female absent on articles 1 and 2, and present as minute apical tuft on article 3. **Mesosternum.** Mesosternal process moderately convex but not at all tumescent (as in Fig. 59). **Abdomen.** Ventral surface with moderately dense, elongate-fine appressed white scales. Abdominal sterna III to VI (especially III and IV) with moderately dense, small, individually indistinct to distinct rounded glabrous shiny patches, each with large puncture and single long erect hair-like scale situated in or near the center. Apex of abdominal sternum VII of male lacking dorsally directed tooth. Abdominal tergum VIII uniformly convex, apical margin slightly elevated and reflexed. **Genitalia.** Female (two examined). Abdominal sternum VIII with lateral arms narrow, very slightly sinuate and markedly inwardly arcuate near apex (Fig. 90a). Gonocoxite II with stylus moderately large, slightly anteapical in position (Fig. 90b). Male (three examined). Abdominal sternum VIII with paired sclerite with inner apices lacking or with only slightly developed ventral projections. Aedeagus elongate-narrow, in lateral view thickest at midlength; in ventral view more or less parallel sided throughout length. Internal sac elongate and low, only slightly apically deflexed; median dorsal pocket low, with

moderately large dorsolaterally directed paramedial lobe at midheight at basal one-third, small median lobe on dorsal surface immediately anterior to crest of median dorsal pocket, and moderately large dorsolaterally directed paramedial lobes at midheight at apical one-third (Figs. 98a,b). Eversible apical sclerite complex with paired narrow scythe-like sclerites well sclerotized, each lacking median projection (Fig. 98d); adjacent ventral surface and basal portion of sides of apex of internal sac with pair of well-developed transverse sclerites (Fig. 98a).

**Geographic variation.**--- None noted.

**Geographic distribution.**--- This species is found throughout the highlands of central Mexico (Fig. 210).

**Natural history.**--- This species appears to be restricted to the xeric high-elevation mesquite-grasslands of central Mexico (pastizal and perhaps pastizal-matorral xerofilo transition of Rzedowski [1978]). Adults have been collected on *Solidago* sp. (Compositae). Other adults have been collected from under prickly-pear pads, stones and dry cow dung. Definite host plants are unknown. Adults have been collected from June to August at elevations from 2250-3385 m ( $\bar{X}=2675$  m,  $N=8$ ) (Fig. 232).

**Chorological relationships.**--- This species is sympatric

throughout its range with *A. lutulentus*. Adult specimens of the two species have been collected together at 1.5 mi. S. Fresnillo and 20 mi. NE. Dolores Hidalgo.

**Phylogenetic relationships.**--- This species is the sister-species of the rest of the species in the subgenus *Apleurus* (Figs. 234-235).

Table 15. Descriptive statistics for *A. aztecus* based on 13 males from throughout the species range

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	0.96-1.52	1.182	0.251	0.093
WF	0.92-1.36	1.172	0.224	0.083
WRA	0.80-1.20	1.015	0.198	0.073
LP	1.76-2.88	2.209	0.607	0.225
WPB	1.84-3.20	2.566	0.658	0.244
WPT	1.84-3.28	2.622	0.674	0.249
WE1M	2.88-4.56	3.699	0.856	0.317
LE1	3.60-6.40	4.701	1.300	0.481
B. Proportions.				
WPB/LP	1.05-1.27	1.165	0.094	0.035
WPT/WPB	1.00-1.04	1.021	0.023	0.008
WRA/LR	0.79-1.00	0.862	0.081	0.030
WRA/WF	0.82-0.92	0.866	0.036	0.013
WE1M/LE1	0.70-0.90	0.792	0.074	0.028
LP/LE1	0.37-0.54	0.473	0.073	0.027



Table 16. Descriptive statistics for *A. aztecus* based on six females from throughout the species range

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	0.88-1.60	1.387	0.384	0.209
WF	0.96-1.52	1.333	0.310	0.169
WRA	0.80-1.28	1.147	0.270	0.147
LP	1.52-2.88	2.520	0.754	0.411
WPB	1.76-3.44	2.947	0.899	0.490
WPT	1.84-3.60	3.053	0.928	0.505
WELM	2.56-4.80	4.173	1.211	0.659
LE1	3.20-6.24	5.413	1.678	0.914
B. Proportions.				
WPB/LP	1.12-1.21	1.169	0.053	0.029
WPT/WPB	0.98-1.05	1.037	0.046	0.025
WRA/LR	0.78-0.91	0.833	0.082	0.045
WRA/WF	0.83-0.88	0.859	0.034	0.019
WELM/LE1	0.72-0.80	0.774	0.049	0.026
LP/LE1	0.45-0.49	0.467	0.022	0.012

4.4.2.2.2 *Apleurus (Apleurus) lutulentus* (LeConte), new combination

(Figs. 6, 59, 62, 69, 83, 91, 99, 207)

*Cleonus lutulentus* LeConte 1859a:18. Lectotype (here

designated), female, one of two syntypes (only one examined), labelled with a dark green circle (=New Mexico), "Type/ 5186", "*Cleonaspis/ lutulentus/ (Lec)*" and with my designation label "*Cleonus/ lutulentus/ Lec. LECTOTYPE/ desig. Anderson*" (MCZC). Type locality, Santa Fe, New Mexico. Gemminger and von Harold 1871 (catal.). Leng 1920 (catal.).

*Cleonus pulvereus* LeConte 1859a:18. New synonymy. Holotype (examined), female, labelled "Dallas/ Tex/ Boll", "Type/ 5185", "*Cleonopsis/ pulvereus/ (Lec)*" (MCZC). Type locality, Dallas, Texas. Gemminger and von Harold 1871 (catal.). Leng 1920 (catal., error as *Cleonus pluvareus*). Blackwelder 1947 (check.). Krombein 1979 (as prey).

*Apleurus fossus* Chevrolat 1873:78. New synonymy. Lectotype (here designated), female, one of two syntypes, labelled "12....", "Mexico/ Au. Salle", "TYPUS", "40", "473/ 85", "Riksmuseum/ Stockholm" and with my designation label "*Apleurus fossus/ Chevrolat/ LECTOTYPE/ desig. Anderson*" (Riksmuseum, Stockholm; other syntype in BMNH). Type locality, Mexico. Faust 1904.

*Cleonaspis lutulentus*; LeConte 1876a (redesc.). Henshaw 1881-1882 (check.). Henshaw 1885 (check.). Wickham 1902 (check.).

*Cleonopsis pulvereus*; LeConte 1876a (redesc.). Henshaw 1885 (check.). Wickham 1889. Wickham 1896 (check.).

- Stephanocleonus pulvereus*; Henshaw 1881-1882 (check.).
- Cleonus (Apleurus) fossus*; Casey 1891 (designation as type species of *Apleurus* Chevrolat). Champion 1902-1906 (syn., distn.). Csiki 1934 (catal.).
- Cleonus (Apleurus) lutulentus*; Casey 1891 (key). Fall and Cockerell 1907 (check.). Csiki 1934 (catal.).
- Cleonus (Apleurus) pulvereus*; Casey 1891 (key). Champion 1902-1906 (syn., distn.). Fall and Cockerell 1907 (check.). Csiki 1934 (catal.).
- Cleonus fossus*; Blackwelder 1947 (check.).
- Dinocleus bryanti* Van Dyke 1953:101. **New synonymy.** Holotype (examined), male, labelled "Seligman, / Ariz. VIII 3 36/ Bryant.112.", "From the/ O. Bryant/ Collection", "Holotype" (CASC). Type locality, Seligman, Arizona.
- Cleonis bryanti*; O'Brien and Wibmer 1982 (catal., distn.).
- Cleonis fossus*; O'Brien and Wibmer 1982 (catal., distn.).
- Cleonis lutulentus*; O'Brien and Wibmer 1982 (catal., distn.).
- Cleonis pulvereus*; O'Brien and Wibmer 1982 (catal., distn.).
- lutulentus, Incertae sedis*; Chevrolat 1873 (error, misspelling).
- pulverosus, Incertae sedis*; Chevrolat 1873 (error, misspelling).

**Notes about synonymy.**--- I can find no consistent differences that warrant separation of *Cleonus pulvereus*

LeConte, *Cleonus lutulentus* LeConte and *Dinocleus bryanti* Van Dyke as distinct species. LeConte (1859a) emphasized characters (extent of ventral tarsal vestiture and degree of development of rostral carina) which are intraspecifically variable in describing *C. pulvereus* and *C. lutulentus* and subsequently placing them in separate genera (LeConte 1876a) (see "Notes about synonymy" section for *Apleurus*). I have not seen any specimens of *A. lutulentus* in which the ventral tarsal pilose vestiture is entirely lacking and in which the tarsal articles are not bilobed, as was given by LeConte (1876a) as, in part characteristic of *C. lutulentus*, and I do not find that the characters subsequently given by Casey (1891) to separate the two species represent anything more than intraspecific variation.

*Dinocleus bryanti* Van Dyke is clearly *A. lutulentus*. Van Dyke (1953) considered *D. bryanti* a member of the genus *Dinocleus*, however, since he did not mention either *Cleonus pulvereus* or *Cleonus lutulentus*, and since both were not then considered to be *Dinocleus*, I suspect that Van Dyke did not look at representatives of these forms in describing *D. bryanti*, but only at those forms already placed in *Dinocleus*. As a result, he found his new species to be very distinct from those then included in *Dinocleus*. As discussed elsewhere ("Notes about synonymy" section for *Apleurus*) this placement of *D. bryanti* in *Dinocleus* concurs with my classification of species of *Dinocleus* and *Apleurus* (including *Cleonaspis* and *Cleonopsis*) herein as congeneric.

I also consider the Mexican brachypterous *A. fossus* as conspecific with the other forms discussed previously in this section. No specific type locality was given for this species but because brachypterous individuals are otherwise only known from central and southern Mexico I suspect this is the area of collection of the type series. These brachypterous forms are otherwise distinguished from typical *C. lutulentus* by a more robust and globose elytral form, possession of elytral intervals 3, 5 and 7 slightly elevated and convex, and possession of deep and large punctures on the head, pronotum and elytra. Although only this brachypterous form occurs in the southern Mexican highlands of Oaxaca and Puebla, some of these similarly brachypterous individuals are also known to be sympatric with typically macropterous individuals in central Mexico. Because wing length polymorphism and associated variation in elytral form is found in sympatric individuals of other species of *Apleurus* (*A. angularis* and *A. albovestitus*), I concur with Champion (1902-1906:98) that *A. fossus* is "probably nothing more than a southern form of *C. pulvereus* LeC."

**Problems in recognition.**--- Brachypterous specimens of this species may be confused with individuals of *A. aztecus* as noted in the "Problems in recognition" section for that species. They may also be confused with small *A. saginatus* or *A. albovestitus*, or macropterous *A. angularis*, but the presence of distinct prosternal swellings anterior to the

prosternal impressions in members of the latter three species (Figs. 85, 87-88), among other key characters, will readily separate them from *A. lutulentus*.

**Description.**--- Specimens examined. 334 males, 369 females. Data about variation in LR, WF, WRA, LP, WPB, WPT, WELM, LE1, WPB/LP, WPT/WPB, WRA/LR, WRA/WF, WELM/LE1, and LP/LE1 are presented in Tables 17 and 18. Size. Length, male, 5.3-10.3 mm; female, 6.4-11.4 mm. Width, male, 2.4-4.8 mm; female, 2.9-5.5 mm. Head. Eye prominent and convex in dorsal view. Frons and vertex with moderately dense, small to large, deep punctures. Frons lacking suberect or erect vestiture; with only sparse elongate-narrow appressed white scales. Area immediately behind posterior margin of eye with large, deep, irregularly impressed punctures. Area above eyes flat, not distinct from rest of frons (eyes not appearing distinctly browed in anterior view). Width of frons greater than to subequal to width at apex of rostrum. Rostrum. Moderately robust (width at apex 0.647-0.909 times length in male; 0.667-0.842 in female) (Fig. 83). Median carina absent or at most only slightly developed and indistinct. Dorsal and lateral punctation moderately dense, small to large, deep; sparser apically and irregularly impressed medially in some specimens). Dorsally, excluding epistoma, with scattered very short, fine, suberect hair-like scales towards apex, and sparse elongate-narrow appressed white scales. In lateral view with apical portion

flat to only very slightly declivous from point of antennal insertion to apex (Fig. 83a). Epistoma with apical margin rounded at middle (Fig. 83b). Mouthparts. Maxillary palpus with stipes with large seta on outer margin (as in Fig. 64). Labial palpi separated by more or less width of basal article of a labial palpus (Fig. 62). Prementum with one pair of large setae (Fig. 62). Pronotum. In dorsal view with lateral margins slightly arcuate from base to apical one-quarter; constricted at oblique angle anterior to apical one-quarter; pronotum widest at base to subequal in width at base and at apical one-quarter; distinct lateral tubercles not evident (Fig. 69). Dorsal and lateral punctation small to large, moderately dense to dense, and deep; punctures sparser and smaller on flanks. Scales white, elongate-narrow, appressed, sparse and small or lacking medially from disk in a broad, apically narrowed patch (except for midline in some specimens), and at lateral margins near base; uniformly dense laterally to lateral margins in anterior one-half and ventrally on flanks. Median carina lacking. Dorsally lacking suberect or erect vestiture. Anterolateral margin with postocular projection absent or at most very slightly developed (Fig. 83a). Prosternum. With very shallow impression anterior to each procoxal cavity and with at most low slightly developed rounded swelling anterior to each prosternal impression (Fig. 83a). Elytra. Moderately robust in general form (width at midlength 0.568-0.724 times length in males; 0.521-0.761

in females) (Fig. 69). In dorsal view with lateral margins slightly arcuate and convergent to subparallel from apical one-third to humerus; markedly arcuate from apical one-third to apex; humerus obtuse to acute, moderately distinct to distinct (Fig. 69). Dorsal suberect or erect vestiture absent or at most with sutural interval and intervals 2 and 3 with very short suberect hair-like scales. Dorsally with all elytral intervals flat to slightly convex, to with sutural interval and intervals 3, 5, and 7 slightly swollen and convex in brachypterous individuals. Scales various in density and size; elongate-narrow, pattern various with numerous irregularly distributed small glabrous or nearly so areas (Fig. 69). **Wings.** Short (very slightly shorter than elytra in length [2.1%, N=15]) to long (greater than elytra in length [97.9%, N=688]). Branches of 2A not joined at base. **Legs.** Foretibia of female with inner margin with small to moderately large denticles throughout most of length; subapical tooth moderately large, distinct from tibial denticles. Foretibial and mesotibial unci of both sexes large, metatibial uncus of both sexes moderately large. Tarsal claws not divergent, with well-developed basal internal flange present. Ventral tarsal pilose vestiture of foretarsus of male present as small elongate apical pads on article 1, as elongate-oval pads covering apical one-half of article 2, and as large elongate-oval pads covering apical two-thirds of article 3; of foretarsus of female present as small elongate apical pads on article 1, as moderately large



elongate-oval pads covering apical one-half on article 2, and as large elongate-oval pads covering apical one-half to two-thirds of article 3; of mesotarsus of male present as elongate-narrow pads covering apical one-third on article 1 and apical one-half on article 2, and as large elongate-oval pads covering apical one-half to two-thirds on article 3; of mesotarsus of female present as elongate-narrow pads covering apical one-third on article 1, as elongate-oval pads covering apical one-half on article 2, and as large rounded pads covering apical one-half on article 3; of metatarsus of male absent on article 1, present as elongate very narrow pads on article 2, and as elongate-oval pads covering apical one-half on article 3; of metatarsus of female absent to as small elongate-narrow pads covering apical one-quarter on article 1, as elongate very narrow pads covering apical one-third on article 2, and as large rounded pads covering apical one-half to two-thirds on article 3. **Mesosternum.** Mesosternal process flat to very slightly convex but not at all tumescent (Fig 59). **Abdomen.** Ventral surface with dense, elongate-fine appressed white scales. Abdominal sterna III to VI (especially III and IV) with sparse to moderately dense, small, individually indistinct to distinct rounded glabrous shiny patches, each with large puncture and single short erect hair-like scale situated in or near the center. Apex of abdominal sternum VII of male with large dorsally directed median truncate tooth. Abdominal tergum VIII uniformly convex, apical margin

slightly elevated and reflexed. **Genitalia.** Female (three examined). Abdominal sternum VIII with lateral arms narrow, very slightly inwardly arcuate in basal three-quarters, then markedly inwardly arcuate to apex (Fig. 91a). Gonocoxite II with stylus moderately large, antepical in position by more or less length of stylus (Fig. 91b). Male (12 examined). Abdominal sternum VIII with paired sclerite with inner apices with slightly to moderately developed ventral projections. Aedeagus elongate-narrow, in lateral view subequal in thickness from base to midlength (Fig. 99c); in ventral view more or less parallel sided throughout length. Internal sac elongate and low, only slightly apically deflexed; median dorsal pocket low, with moderately large dorsolaterally directed paramedial lobe at midheight at basal one-third and small median lobe on dorsal surface immediately anterior to crest of median dorsal pocket (Figs. 99a,b). Eversible apical sclerite complex with paired narrow scythe-like sclerites well sclerotized, each with short median projection (Fig. 99d); adjacent ventral surface and basal portion of sides of apex of internal sac with pair of well-developed transverse sclerites (Fig. 99a).

**Geographic variation.**--- Some specimens from central Mexico have larger and deeper punctures of the head, pronotum and elytra; more robust and globose elytra; elytral intervals 3, 5, and 7 slightly elevated and convex; and short wings. Only specimens with extreme states in all of these characters are

found in the area of the southern border of the species range in the Mexican states of Oaxaca and Puebla.

**Geographic distribution.**--- This species is found from Kansas, Oklahoma and eastern Texas, west to Arizona, disjunct to central and southern Mexico (Fig. 207). The species does not appear to be found in the Chihuahuan desert region in Mexico. There is a single questionable record from northeastern California.

**Natural history.**--- This is the most widely distributed species of *Apleurus*. It is found in mesquite-grassland, grassland, and desert-grassland transitional habitats in Mexico and the United States, and in the grassland-deciduous forest transitional zone. Adults have been collected in dry upland desert washes and grasslands on a wide variety of plants, but mostly Compositae, as follows: *Gossyplum hirsutum* L. (cotton, Malvaceae); *Chenopodium* sp. (Chenopodiaceae); *Asclepias* sp., *A. subverticillata* (Gray) Vail (Asclepiadaceae); *Koeberlinia spinosa* Zucc. (Capparaceae); *Acacia* sp. (Leguminosae); *Baccharis* sp., *B. glutinosa* Pers., *Balleya pleniradiata* Harv. and Gray, *Chrysothamnus nauseosus* (Pall.) Britton, *Flourensia cernua* DC., *Gutierrezia* sp., *Solidago* sp., *Verbesina encelloides* (Cav.) Benth. and Hook., *V. oreophila* Woot. and Standl., *Xanthium* sp. (all Compositae). Definite host plants are not known. Adults have been collected from March to October at

elevations from 9-3058 m (X=1726 m, N=77) (Fig. 232).

**Chorological relationships.**--- This species is sympatric with *A. saginatus* in southern Arizona and New Mexico; with *A. aztecus* in central Mexico; with *A. angularis* in New Mexico, western Texas, Colorado, western Oklahoma, and Arizona; and with *A. albovestitus* in Arizona, New Mexico, and western Texas.

Adults of *A. tutulentus* and *A. albovestitus* have been caught together near Portal, Arizona, on *Chrysothamnus nauseosus*. Adult specimens of *A. tutulentus* and *A. aztecus* have been collected together 1.5 mi. S. Fresnillo, on *Solidago* sp., and 20 mi. NE. Dolores Hidalgo, Mexico.

**Phylogenetic relationships.**--- This species is the sister-species of the *A. porosus*-*A. angularis*-*A. jacobinus*-*A. saginatus*-*A. albovestitus* lineage of *Apleurus* (Figs. 234-235).

Table 17. Descriptive statistics for *A. lutulentus*  
based on eight males from 2 mi. NE. Portal, Arizona

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.44-1.76	1.660	0.167	0.079
WF	1.20-1.44	1.300	0.140	0.066
WRA	1.04-1.28	1.180	0.140	0.066
LP	2.56-3.36	3.050	0.478	0.202
WPB	2.72-3.52	3.160	0.406	0.191
WPT	2.56-3.20	2.930	0.327	0.154
WE1M	3.36-4.80	4.340	0.693	0.327
LE1	5.92-7.04	6.560	0.588	0.277
B. Proportions.				
WPB/LP	1.00-1.08	1.037	0.038	0.018
WPT/WPB	0.90-0.95	0.928	0.024	0.011
WRA/LR	0.67-0.76	0.711	0.045	0.021
WRA/WF	0.87-0.94	0.908	0.048	0.023
WE1M/LE1	0.57-0.72	0.661	0.072	0.034
LP/LE1	0.43-0.50	0.464	0.039	0.018

Table 18. Descriptive statistics for *A. lutulentus*  
based on eight females from 2 mi. NE. Portal, Arizona

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.60-1.84	1.750	0.119	0.056
WF	1.20-1.44	1.350	0.150	0.070
WRA	1.12-1.36	1.250	0.127	0.060
LP	2.80-3.44	3.160	0.368	0.174
WPB	2.88-3.92	3.350	0.478	0.225
WPT	2.72-3.52	3.180	0.419	0.198
WE1M	3.92-5.20	4.540	0.562	0.265
LE1	6.48-7.92	7.110	0.763	0.360
B. Proportions.				
WPB/LP	1.03-1.14	1.060	0.056	0.026
WPT/WPB	0.90-1.00	0.950	0.052	0.024
WRA/LR	0.67-0.74	0.714	0.036	0.017
WRA/WF	0.89-0.94	0.926	0.035	0.017
WE1M/LE1	0.61-0.66	0.639	0.028	0.013
LP/LE1	0.42-0.47	0.445	0.028	0.013

4.4.2.2.3 *Apleurus (Apleurus) porosus* (LeConte), new  
combination

(Figs. 70, 84, 92, 102, 210)

*Centrocleonus porosus* LeConte 1876a:146. Holotype

(examined), female, labelled with a silver circle with portion cut away (=Baja California) "Type/ 5239", "*C. porosus*/ Lec" (MCZC). Type locality, Cape San Lucas, Baja California Sur, Mexico. Henshaw 1881-1882 (check.). Henshaw 1885 (check.). Horn 1894 (check.).

*Dinocleus farctus* Casey 1891:181. New synonymy. Holotype (examined), male, labelled "Cal", "CASEY/ bequest/ 1925", "TYPE USNM/ 37271", "*D./ farctus*/ Cas." (USNM). Type locality, California, near the southern border. Fall 1901 (check.). Leng 1920 (catal.).

*Dinocleus porosus*; Casey 1891 (key, redesc.). Leng 1920 (catal.).

*Cleonus (Dinocleus) farctus*; Csiki 1934 (catal.).

*Cleonus (Dinocleus) porosus*; Csiki 1934 (catal.).

*Cleonus porosus*; Blackwelder 1947 (check.).

*Cleonus farctus*; Hardy and Andrews 1976 (distn., prob. misident. *A. angularis*).

*Cleonis farctus*; O'Brien and Wibmer 1982 (catal., distn.).

*Cleonis porosus*; O'Brien and Wibmer 1982 (catal., distn.).

Notes about synonymy.--- Both LeConte (1876a) and Casey (1891) based their species descriptions of *Centrocleonus porosus* and *Dinocleus farctus* on unique specimens. Having examined larger numbers of individuals from various localities, I conclude that the two are conspecific and the differences noted by Casey (1891) as species specific are

due to intraspecific variation or secondary sexual characteristics (the holotype of *C. porosus* is a female; that of *D. farctus*, a male).

The type locality of *D. farctus* is given by Casey (1891) as California, near the southern border. I have not seen any *A. porosus* from the state of California.

**Problems in recognition.**--- Members of this species are likely to be confused with *A. angularis* but can be distinguished from individuals of that species by the robust form of the tarsal claws, each possessing a basal internal flange in most specimens (as in Fig. 67); by the lack of distinct glabrous patches on the abdominal sterna; and by presence of only very short suberect dorsal vestiture in the former. Some specimens of *A. porosus* from various localities however, possess tarsal claws, which although moderately robust, lack a distinct internal flange.

*Apleurus porosus* specimens are also likely to be confused with *A. jacobinus* individuals particularly in Baja California Norte and in southern California where the species appear to be narrowly sympatric. *Apleurus jacobinus* are distinguished from *A. porosus* specimens by the long dorsal erect vestiture; alternate elytral intervals markedly elevated and convex; and abdominal sterna with distinct glabrous patches in the former. However, in the southern part of the range of *A. jacobinus* in Baja California Norte, individuals I examined have shorter dorsal erect vestiture.



and less markedly elevated elytral intervals than elsewhere throughout the species range, thus, approaching the states in *A. porosus* in these characters. Although the differences in degree of elevation of the elytral intervals and length of dorsal vestiture in these individuals should still allow for separation of *A. jacobinus* and *A. porosus*, the presence of distinct glabrous patches on the abdominal sterna of the former will allow for unequivocal separation of the two species. *Aptaurus porosus* specimens also have the pronotum with scales uniformly very dense laterally and on the flanks, more or less obscuring the individual large punctures.

Many *A. porosus* can also be separated from *A. jacobinus* and *A. angularis* by their larger size, more elongate-narrow rostrum, and by the pronotum being widest at the base (see Figs. 105-106, 109).

**Description.**--- Specimens examined. 23 males, 32 females. Data about variation in LR, WF, WRA, LP, WPB, WPT, WELM, LE1, WPB/LP, WPT/WPB, WRA/LR, WRA/WF, WELM/LE1, and LP/LE1 are presented in Tables 19 and 20. Size. Length, male, 5.8-13.6 mm; female, 6.9-15.2 mm. Width, male, 2.6-6.1 mm; female, 3.0-7.2 mm. Head. Eye very prominent and convex in dorsal view. Frons and vertex with dense, large to very large, deep punctures. Frons and area immediately above eyes with sparse to moderately dense short suberect hair-like scales. Area immediately behind posterior margin of eye with

large, deep, irregularly impressed punctures. Area above eyes slightly elevated above rest of frons (eyes appearing distinctly browed in anterior view). Width of frons greater than width at apex of rostrum. Rostrum. Moderately elongate-narrow (width at apex 0.612-0.684 times length in male; 0.616-0.693 in female) (Fig. 84). Median carina absent to slightly developed but distinct, almost entirely obscured in some specimens by large deep punctures. Dorsal and lateral punctation moderately dense, large, deep. Dorsally, excluding epistoma, with scattered short, fine, suberect hair-like scales towards apex, and dense elongate-narrow appressed white scales. In lateral view with apical portion steeply declivous from point of antennal insertion to apex (Fig. 84a). Epistoma with apical margin rounded at middle (Fig. 84b). Mouthparts. Maxillary palpus with stipes lacking large seta on outer margin (as in Fig. 65). Labial palpi separated by more or less width of basal article of a labial palpus (as in Figs. 62-63). Prementum with two or three pairs of large setae (as in Fig. 63). Pronotum. In dorsal view with lateral margins subparallel to slightly convergent from base to apical one-quarter; constricted at an acute angle anterior to apical one-quarter; widest at base to subequal in width at base and apical one-quarter; distinct lateral tubercles evident (Fig. 70). Dorsal and lateral punctation large, moderately deep to deep, dense; punctures sparser and smaller on flanks. Scales white, elongate-narrow, appressed, sparse and small or lacking

medially from disk in broad, apically narrowed patch (except for midline in some specimens); uniformly dense laterally, at lateral margins and ventrally on flanks, more or less obscuring large punctures. Median carina lacking. Dorsally covered throughout with sparse, fine, short, erect hair-like scales. Anterolateral margin with postocular projection absent or at most very slightly developed (Fig. 84a).

**Prosternum.** With shallow impression anterior to each procoxal cavity and with well-developed rounded swelling anterior to each prosternal impression (Fig. 84a). **Elytra.** Moderately robust in general form (width at midlength 0.651-0.700 times length in males; 0.645-0.732 in females) (Fig. 70). In dorsal view with lateral margins moderately arcuate throughout length; humerus rounded, not distinct (Fig. 70). Sutural interval and intervals 3, 5, 7, and 9 with sparse, very short, suberect hair-like scales. Dorsally with sutural interval and intervals 3, 5, 7, and 9 slightly swollen and convex (more so basally in some specimens).

Scales elongate-narrow, various in density and size; pattern various but in most specimens with 2 extensive glabrous or nearly so areas, one from area behind humerus, posteromedially directed to sutural interval at midlength, other continuous with the former at midlength and directed laterally from sutural interval to interval 9; some specimens also with glabrous or nearly so area of variable extent at confluence of intervals 4 to 7 (Fig. 70). **Wings.**

**Absent.** **Legs.** Foretibia of female with inner margin with

small denticles in apical one-third; subapical tooth small, not distinct from tibial denticles. Foretibial and mesotibial unci of both sexes large, metatibial unci of both sexes moderately large. Tarsal claws not divergent, basal internal flange absent to present, well-developed. Ventral tarsal pilose vestiture of foretarsus of male absent on article 1, present as minute apical tufts on article 2, as moderately large rounded pads covering apical one-third to one-half of article 3; of foretarsus of female, mesotarsus of male and female, and metatarsus of male, lacking from articles 1 and 2, present as small rounded pads at apex on article 3; of metatarsus of female lacking from articles 1 and 2, present as minute pads at apex of article 3. **Mesosternum.** Mesosternal process flat to very slightly convex but not at all tumescent (as in Fig. 59). **Abdomen.** Ventral surface with very dense, elongate-fine appressed white scales and scattered suberect hair-like scales. Abdominal sterna lacking distinct glabrous patches, with at most only lateral margins of abdominal sterna III and IV with very small indistinct glabrous patches. Apex of abdominal sternum VII of male with small dorsally directed median truncate tooth. Abdominal tergum VIII uniformly convex, apical margin slightly elevated and reflexed. **Genitalia.** Female (four examined). Abdominal sternum VIII with lateral arms narrow, slightly outwardly arcuate in basal one-half, then parallel to near apex, markedly inwardly arcuate immediately before apex (Fig. 92a).

Gonocoxite II with stylus very small, antepical in position by more or less one-half length of stylus (Fig. 92b). Male (five examined). Abdominal sternum VIII with paired sclerite with inner apices with slightly to moderately developed ventral projections (as in Fig. 101e). Aedeagus elongate-narrow, in lateral view thickest at midlength; in ventral view more or less parallel sided throughout length. Internal sac short and high, markedly apically deflexed; median dorsal pocket high, with moderately large dorsolaterally directed paramedial lobe at midheight and midlength; small median lobe on dorsal surface immediately anterior to crest of median dorsal pocket (Figs. 102a,b). Eversible apical sclerite complex with paired narrow scythe-like sclerites lightly sclerotized, each lacking median projection (as in Figs. 100d-101d); adjacent ventral surface and basal portion of sides of apex of internal sac with pair of indistinct lightly sclerotized transverse sclerites (Fig. 102a).

**Geographic variation.**--- None noted, but this may be due to the small number of specimens examined, especially from Baja California Norte where this species is at most narrowly sympatric with *A. angularis* and *A. jacobinus*.

**Geographic distribution.**--- This species is found throughout Baja California Sud and Baja California Norte, Mexico (Fig. 210). It may occur in the extreme south of California.

**Natural history.**--- Label data indicates that most adults have been caught in sand dune areas in the "matorral xerofilo" of Baja California (Sonoran desert) and Pacific semi-desert region at elevations of from 6-144 m ( $X=75$  m,  $N=5$ ) (Fig. 232). Sleeper (pers. comm.) has collected numerous specimens from a woody *Encelia* species and a single specimen from *Encelia yacintata* Vasey and Rose (Compositae). Definite host plants are not known.

**Geographical relationships.**--- This species has one of the more restricted distributions of *Apleurus* species. It is sympatric with extreme southern *A. albovestitus*, and at most narrowly sympatric to parapatric with its close relatives *A. angularis* and *A. jacobinus*.

**Phylogenetic relationships.**--- Primary characters allow only placement of this species as a member of an unresolved trichotomy also including *A. angularis* and *A. jacobinus*. This lineage is the sister-group to the *A. albovestitus*-*A. saginatus* lineage; these two lineages sister to *A. luteiventris* (Fig. 234).

Distribution of the apotypic states of secondary characters of length of erect vestiture and development of lateral pronotal tubercles allows for recognition of *A. porosus* as the sister-species of the *A. angularis*-*A. jacobinus* sister-species pair (Fig. 235). Presence of the apotypic state of tarsal claws each lacking a basal internal

flange in some *A. porosus* and in *A. angularis* (Fig. 66) is either a result of hybridization or of homoplasy, more likely the former.

Table 19. Descriptive statistics for *A. porosus* based on seven males from 68.0 km. W. Ejido Viscaino, Baja California Sur, Mexico

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	2.37-3.20	2.904	0.473	0.238
WF	1.86-2.32	2.124	0.261	0.132
WRA	1.55-2.12	1.887	0.293	0.148
LP	3.82-5.57	4.747	0.890	0.448
WPB	4.13-5.88	5.189	0.934	0.471
WPT	4.02-5.68	4.984	0.864	0.435
WE1M	5.57-8.05	7.106	1.266	0.638
LE1	8.46-11.76	10.467	1.820	0.917
B. Proportions.				
WPB/LP	1.02-1.17	1.095	0.088	0.044
WPT/WPB	0.93-1.00	0.961	0.041	0.021
WRA/LR	0.61-0.68	0.650	0.033	0.017
WRA/WF	0.83-0.96	0.887	0.061	0.031
WE1M/LE1	0.65-0.70	0.679	0.026	0.013
LP/LE1	0.41-0.48	0.454	0.035	0.018



Table 20. Descriptive statistics for *A. porosus* based on eight females from 68.0 km. W. Ejido Viscaïno, Baja California Sur, Mexico

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	2.32-3.51	3.095	0.580	0.274
WF	1.86-2.63	2.329	0.389	0.183
WRA	1.60-2.32	2.006	0.356	0.168
LP	4.02-5.68	5.096	0.866	0.408
WPB	4.23-6.40	5.549	1.155	0.544
WPT	4.23-6.09	5.366	0.960	0.452
WE1M	5.99-8.88	7.858	1.508	0.711
LE1	8.46-13.00	11.430	2.411	1.136
B. Proportions.				
WPB/LP	1.05-1.13	1.086	0.054	0.025
WPT/WPB	0.90-1.02	0.970	0.058	0.027
WRA/LR	0.62-0.69	0.650	0.043	0.020
WRA/WF	0.83-0.91	0.861	0.047	0.022
WE1M/LE1	0.65-0.73	0.689	0.041	0.019
LP/LE1	0.42-0.48	0.448	0.028	0.013

4.4.2.2.4 *Apleurus (Apleurus) angularis* (LeConte), new  
combination

(Figs. 66, 72-74, 85, 93, 100, 211)

*Cleonus angularis* LeConte 1859a:18. Lectotype (here designated), male, one of two syntypes (only one examined), labelled with a pale green circle (=Wyoming Territory) "*C. angularis*/ Beckwith Lec.", "Type/ 5182" and with my designation label "*Cleonus/ angularis*/ LeC. LECTOTYPE/ desig. Anderson" (MCZC). Type locality, Kansas. LeConte 1858 (*nomen nudum*). Gemminger and von Harold 1871 (catal.). Hatch 1971.

*Centrocleonus angularis*; LeConte 1876a (key, diag.). Henshaw 1881-1882 (check.). Henshaw 1885 (check.). Wickham 1889.

*Dinocleus denticollis* Casey 1891:180. New synonymy.

Lectotype (here designated), female, one of five syntypes, labelled "Ari", "CASEY/ bequest/ 1925", "TYPE USNM/ 37269", "*D./ denticollis*/ Cas." and with my designation label "*Dinocleus/ denticollis*/ Csy. LECTOTYPE/ desig. Anderson" (USNM). Type locality, Peach Springs, Arizona. Wickham 1896 (check.). Casey 1904. Van Dyke 1953. Hatch 1971 (redesc.).

*Dinocleus angularis*; Casey 1891 (key, redescr.). Wickham 1902 (check.). Faust 1904 (desig. as type species of *Apleurus* Chevrolat; error as "*angulatus* Lac").

*Dinocleus porcatus* Casey 1904:321. New synonymy. Lectotype (here designated), female, one of two syntypes,

labelled "Ogden/ Ut. Solt", "CASEY/ BEQUEST/ 1925",  
 "TYPE USNM/ 37270", "*porcatus*" and with my designation  
 label "*Dinocleus/ porcatus/ Csy. LECTOTYPE/ desig.*  
 Anderson" (USNM). Type locality, Ogden, Utah. Fall and  
 Cockerell 1907 (check.). Leng 1920 (catal.).

*Cleonus denticollis*; Leng 1920 (catal.). Tanner 1966  
 (biol.).

*Cleonus (Dinocleus) angularis*; Csiki 1934 (catal.).

*Cleonus (Dinocleus) denticollis*; Csiki 1934 (catal.).

*Cleonus (Dinocleus) porcatus*; Csiki 1934 (catal.).

*Cleonus farctus*; Hardy and Andrews 1976 (distn., prob.  
 misident.).

*Cleonis angularis*; O'Brien and Wibmer 1982 (catal., distn.).

*Cleonis denticollis*; O'Brien and Wibmer 1982 (catal.,  
 distn.).

*Cleonis porcatus*; O'Brien and Wibmer 1982 (catal., distn.).

*angularis, incertae sedis*; Chevrolat 1873.

**Notes about synonymy.**--- Geographic patterns of variation in  
 states of structural features considered characteristic of  
*Cleonus angularis* LeConte, *Dinocleus denticollis* Casey, and  
*Dinocleus porcatus* Casey, suggest that only a single species  
 warrants recognition and that these three names be placed in  
 synonymy (see "Geographic variation" section).

I have only been able to locate and examine one of the  
 two syntypes of *Cleonus angularis* LeConte.

Problems in recognition.--- *Apleurus angularis* and *A. aztecus* are the only two species of *Apleurus* in which all individuals have the tarsal claws widely divergent and each lacking a basal internal flange (Fig. 66). This character state should serve to separate nearly all *A. angularis* from *A. jacobinus* and *A. porosus*. Most *A. angularis* can further be separated from most *A. jacobinus* by the presence of long dorsal erect vestiture and markedly elevated alternate elytral intervals in the latter. A very few specimens of *A. jacobinus* from Baja California Norte have less markedly elevated alternate elytral intervals and have shorter dorsal erect vestiture than throughout the rest of the species range and are thus difficult to separate from *A. angularis*. Although these character states are such that they still allow for recognition of these individuals as *A. jacobinus*, they can also be more easily recognized by the robust and less markedly divergent tarsal claws. (See also "Problems in recognition" section for *A. porosus*).

Description.--- Specimens examined. 218 males, 246 females. Data about variation in LR, WF, WRA, LP, WPB, WPT, WE1M, LE1, WPB/LP, WPT/WPB, WRA/LR, WRA/WF, WE1M/LE1, and LP/LE1 are presented in Tables 21 and 22. Size. Length, male, 5.7-10.6 mm; female, 6.9-12.2 mm. Width, male, 2.6-5.2 mm; female, 3.0-5.6 mm. Head. Eye very prominent and convex in dorsal view. Frons and vertex with moderately dense, small to moderately large, deep punctures. Frons and area

immediately above eyes with sparse short erect hair-like scales. Area immediately behind posterior margin of eye with large, shallow to deep, irregularly impressed punctures. Area above eyes markedly elevated above rest of frons (eyes appearing distinctly browed in anterior view). Width of frons greater than to subequal to width at apex of rostrum. **Rostrum.** Moderately robust (width at apex 0.664-0.908 times length in male; 0.619-0.919 in female) (Fig. 85). Median carina slightly to well-developed, low and rounded but distinct, to high and sharp (Fig. 85). Dorsal and lateral punctation small, moderately deep, sparse to moderately dense. Dorsally, excluding epistoma, with dense short, suberect hair-like scales, and dense elongate-narrow to broad appressed white scales. In lateral view with apical portion steeply declivous from point of antennal insertion to apex (Fig. 85a). Epistoma with apical margin rounded at middle (Fig. 85b). **Mouthparts.** Maxillary palpus with stipes lacking large seta on outer margin (as in Fig. 65). Labial palpi separated by more or less width of basal article of a labial palpus (as in Figs. 62-63). Prementum with two or three pairs of large setae (as in Fig. 63). **Pronotum.** In dorsal view with lateral margins subparallel from base to almost apical one-quarter, then divergent to apical one-quarter (giving a distinctly laterally tuberculate appearance); markedly constricted at an acute angle anterior to apical one-quarter, then straight or slightly convergent to apex; widest at tubercles to subequal in width at base

and apical one-quarter (Figs. 72-74). Dorsal and lateral punctation moderately large to large, deep, sparse to moderately dense, in some specimens the areas between punctures irregularly elevated thus appearing sculptured; punctures sparser and smaller on flanks. Scales white, elongate-narrow to broad, appressed, sparse and small or lacking medially from disk in broad, apically narrowed patch (except for midline in some specimens); dense laterally in pair of posterolaterally directed arcuate stripes; scales sparser at lateral margins and ventrally on flanks, to uniformly dense laterally, at lateral margins and on flanks, not obscuring individual large punctures. Median carina lacking. Dorsally covered throughout with moderately dense, fine, short, erect hair-like scales. Anterolateral margin with postocular projection absent or at most very slightly developed (Fig. 85a). Prosternum. With shallow impression anterior to each procoxal cavity and with moderately to well-developed rounded swelling anterior to each prosternal impression (Fig. 85a). Elytra. Moderately robust in general form (width at midlength 0.567-0.761 times length in males; 0.585-0.753 in females) (Figs. 72-74). In dorsal view with lateral margins slightly arcuate throughout length or with margins slightly convergent to subparallel from apical one-third to humerus; humerus rounded to obtuse, not distinct to distinct (Figs. 72-74). Sutural interval and intervals 3, 5, 7, and 9 with moderately dense, short, erect hair-like scales; sparser on other intervals. Dorsally with

sutural interval and intervals 3, 5, 7, and 9 slightly to moderately swollen and convex (more so basally in some specimens). Scales elongate-narrow, various in density and size. Pattern extremely various; many specimens with two variously sized posteromedially directed glabrous or nearly so areas, one at basal one-third from intervals 4 to 6, other at apical one-third from intervals 2 to 7, and with various number of scattered small glabrous or nearly so patches (Fig. 72); other specimens with glabrous areas of various extent on interval 2 at basal one-quarter and from midlength to apical one-quarter, interval 4 at basal one-third, interval 6 at basal one-quarter and apical one-quarter, and interval 8 at midlength; scales confined to median portion of intervals, striae and immediately adjacent portion of intervals lacking scales (Fig. 73). Wings. Absent (95.5%, N=443) or present, long (greater than elytra in length [4.5%, N=21]). Branches of 2A not complete, not joined at base. Legs. Foretibia of female with inner margin with small denticles in apical one-third; subapical tooth small, not distinct from tibial denticles. Foretibial and mesotibial unci of both sexes large, metatibial unci of both sexes small. Tarsal claws widely divergent, basal internal flange absent (Fig. 66). Ventral tarsal pilose vestiture of foretarsus of male absent on article 1, present as minute apical tufts on article 2, as small apical pads on article 3; of foretarsus of female, mesotarsus of male and female, and metatarsus of female, lacking from articles 1

and 2, present as minute to small pads at apex on article 3; of metatarsus of male lacking from articles 1 and 2, lacking from to present as minute pads at apex of article 3.

**Mesosternum.** Mesosternal process flat to very slightly convex but not at all tumescent (as in Fig. 59). **Abdomen.** Ventral surface with moderately dense, elongate-fine appressed white scales. Abdominal sterna III to VI (especially III and IV) with moderately dense, individually distinct circular glabrous patches, each with large puncture and single long erect hair-like scale situated in or near the center. Apex of abdominal sternum VII of male with small dorsally directed median truncate tooth. Abdominal tergum VIII uniformly convex, apical margin slightly elevated and reflexed. **Genitalia.** Female (nine examined). Abdominal sternum VIII with lateral arms narrow, slightly outwardly arcuate in basal one-half, then parallel to near apex, markedly inwardly arcuate immediately before apex (Fig. 93a). Gonocoxite II with stylus very small, antepical in position by more or less one-half length of stylus (Fig. 93b). Male (10 examined). Abdominal sternum VIII with paired sclerite with inner apices with slightly to moderately developed ventral projections (as in Fig. 101e). Aedeagus elongate-narrow, in lateral view thickest at midlength; in ventral view more or less parallel-sided throughout length. Internal sac short and high, markedly apically deflexed; median dorsal pocket high, with moderately large dorsolaterally directed paramedial lobe at midheight and



midlength; small median lobe on dorsal surface immediately anterior to crest of median dorsal pocket (Figs. 100a,b). Eversible apical sclerite complex with paired narrow scythe-like sclerites lightly sclerotized, each lacking median projection (Fig. 100d); adjacent ventral surface and basal portion of sides of apex of internal sac with pair of indistinct lightly sclerotized transverse sclerites (Fig. 100a).

**Geographic variation.**--- Geographic variation is extensive in a number of characters in members of this species. As a result there has been a lot of confusion regarding the specific limits of the previously recognized species (the names of which are here placed in synonymy) and identity of various specimens. Specimens I examined bear label notes either indicating a questionable identification or indicating that the specimen represents a probable new species. One manuscript name is also represented on these labels.

Both macropterous and apterous individuals are known. Macropterous individuals have only slightly elevated and convex alternate elytral intervals and have more or less parallel-sided elytra with distinct humeri. Apterous individuals on the other hand have alternate elytral intervals that are more markedly elevated and convex, and have elytra with the lateral margins more arcuate with indistinct humeri. There appears to be no geographic

component to this variation because macropterous individuals, although few in number, are found throughout most of the species range, with the notable exception of Colorado. This has no doubt contributed to confusion, but the main cause of the confusion is due to the patterns of geographic variation in other structural features. Specimens from eastern and central Colorado, western Oklahoma, and western Texas (called hereafter the "eastern morph" and including typical *Cleonus angularis* LeConté), are different from those to the west in California, Nevada, Utah, and northern Arizona (called hereafter the "western morph" and including typical *Dinocleus denticollis* Casey and *Dinocleus porcatus* Casey).

Eastern morphs (Fig. 73) differ from western morphs (Fig. 72) in states of the following characters: shorter and more robust rostrum; rostral carina higher and sharper; eyes slightly less prominent and convex in dorsal view; pronotal punctures denser, larger, deeper and more irregularly impressed; slightly more markedly elevated alternate elytral intervals; generally a more globose and robust elytral form; elytra with glabrous patches primarily transversely oriented (Fig. 72) not longitudinally oriented (Fig. 73). Specimens from southern Arizona and southern New Mexico however, are not reliably assignable to either form. They are intermediate in many structural features which characterize the eastern and western morphs and it is in this area that apparent intergradation takes place. Elytral scale patterns

are not clearly of one or the other form (Fig. 74); rostra are variable in form, and carinae both in form and in degree of elevation; pronotal punctation is variable in depth, density and regularity of impression; and elytral characters (other than scale pattern) are also variable. Based upon this intergradation, I conclude that the eastern and western morphs and the individuals from southern New Mexico and southern Arizona are all conspecific.

It appears that members of certain species of *Apleurus* (and also *Cleonidius*) are largely restricted to sandy habitats such as dune fields, sand hills, dry washes, stream beds, and riparian habitats. These are either localized and discontinuous in the case of dune fields, or although widespread, restricted in their degree of continuity in the case of washes and riparian communities. Dispersal possibilities are therefore, I believe, markedly influenced by the continuity of drainage patterns or other suitable habitats, and also as is true for all insects, by the presence or absence of wings. Population distinctiveness and apparent lack of intergradation in the north in Colorado and adjacent Utah and northern Arizona, is perhaps due to a combination of a higher continental divide in this area with a more marked discontinuity between eastern and western drainage patterns, consequently more localized and disjunct habitats, and general overall low number of macropterous individuals including presence of only apterous individuals in Colorado. These features may all combine to reduce the

probability of dispersal between eastern and western drainages resulting in local selection for differences between the two morphs. On the other hand, this is not so in southern Arizona and New Mexico where the continental divide is much lower, drainage patterns are not as markedly discontinuous and localized and where macropterous individuals although still few in number, are present in both drainages. Dispersal and interbreeding of individuals is therefore possible in this area with the result that there is no local selection for differences between the two drainages.

This pattern is similar to that found in *C. canescens* which also differs between Colorado and adjacent Utah in the north but not so in the south in New Mexico and Arizona, and all individuals of which are brachypterous and incapable of flight. Species which are found in these same areas but which are macropterous do not show distinctive populations on the eastern and western sides of the continental divide.

**Geographic distribution.**--- This species is widespread in the southwestern United States of America from Colorado, western Oklahoma and western Texas, west to Nevada in the north, extreme southern coastal California and Baja California Norte, Mexico in the south (Fig. 211).

**Natural history.**--- This species is associated with grassland, desert-grassland transitional habitats, Pacific

semi-desert, and Great Basin, desert. Specimens have been collected in sand dune habitats in California, Arizona, Utah, and New Mexico, and otherwise in dry washes and streambeds throughout the species range. Adults have been collected largely on Compositae as follows: *Chaenactis stevioides* Hook. and Arn., *Chrysothamnus nauseosus* (Pall.) Britton, *C. viscidiflorus* (Hook.) Nutt. var. *pumilus*, *Gutierrezia luclida* Greene, *G. microcephala* (DC.) Gray, *Aplopappus arcadenius* (Greene) Blake, *A. linearifolius* DC (all Compositae); *Atriplex lentiformis* (Torr.) Wats. (Chenopodiaceae). Adults have also been collected from roots of *Aplopappus tenuisectus* (Greene) Blake and *Psilostrophe cooperi* (Gray) Greene (Compositae). It is likely that many shrubby Compositae, especially those on which adults are commonly found, act as hosts for this species. Adults have been collected throughout the year at elevations of from 15-2876 m (X=1591 m, N=63) (Fig. 232).

Adults have been collected in the stomach contents of the roadrunner, *Geococcyx californicus* (Aves) at Carlsbad, New Mexico.

**Chorological relationships.**--- This species is sympatric throughout nearly its entire range with *A. albovestitus*; throughout Arizona, New Mexico, and western Texas with *A. lutulentus*; and throughout southeastern Arizona with *A. saginatus*. It is at most narrowly sympatric with its close relatives *A. jacobinus* in southeastern California, and *A.*

*porosus* in Baja California Norte, Mexico.

**Phylogenetic relationships.**--- Primary characters allow only for placement of this species as a member of an unresolved trichotomy also including *A. angularis* and *A. jacobinus*. This lineage is the sister-group to the *A. albovestitus*-*A. saginatus* lineage, these two lineages sister to *A. lutulentus* (Fig. 234).

Distribution of the apotypic states of secondary characters of length of erect vestiture and development of lateral pronotal tubercles allows for recognition of *A. angularis* as the sister-species of *A. jacobinus* (Fig. 235). Presence of the apotypic state of tarsal claws each lacking a basal internal flange in some *A. porosus* and in *A. angularis* is either a result of hybridization or of homoplasy, more likely the former.

Table 21. Descriptive statistics for *A. angularis* based on 15 males from Mercury, Nevada

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.55-2.23	1.861	0.283	0.097
WF	1.44-1.75	1.615	0.190	0.065
WRA	1.14-1.65	1.430	0.239	0.082
LP	2.48-3.82	3.111	0.573	0.197
WPB	2.99-4.33	3.563	0.606	0.209
WPT	3.20-4.54	3.818	0.586	0.201
WE1M	3.72-5.78	4.918	0.884	0.304
LE1	5.68-8.57	7.376	1.183	0.407
B. Proportions.				
WPB/LP	1.07-1.29	1.148	0.081	0.028
WPT/WPB	1.03-1.13	1.073	0.045	0.016
WRA/LR	0.66-0.84	0.769	0.075	0.026
WRA/WF	0.79-0.94	0.884	0.068	0.023
WE1M/LE1	0.57-0.72	0.670	0.062	0.021
LP/LE1	0.36-0.46	0.422	0.034	0.012

Table 22. Descriptive statistics for *A. angularis* based on 10 females from Mercury, Nevada

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.55-2.33	1.892	0.356	0.150
WF	1.44-1.86	1.651	0.208	0.088
WRA	1.24-1.65	1.464	0.204	0.086
LP	2.79-3.61	3.168	0.399	0.168
WPB	3.10-4.13	3.562	0.511	0.216
WPT	3.10-4.33	3.818	0.583	0.246
WE1M	3.92-5.78	4.984	0.894	0.377
LE1	6.09-9.49	7.565	1.503	0.634
B. Proportions.				
WPB/LP	1.03-1.17	1.123	0.055	0.023
WPT/WPB	1.00-1.13	1.066	0.062	0.026
WRA/LR	0.71-0.84	0.777	0.060	0.025
WRA/WF	0.82-0.94	0.887	0.057	0.024
WE1M/LE1	0.61-0.71	0.660	0.047	0.020
LP/LE1	0.38-0.46	0.421	0.036	0.015



4.4.2.2.5 *Apleurus (Apleurus) jacobinus* (Casey), new  
combination

(Figs. 67, 71, 86, 94, 101, 208)

*Centrocleonus pilosus* LeConte 1876a:145 [not Chevrolat 1873:42]. New synonymy. Holotype (examined), male, labelled "Calif.", "type/ 5240", "*Centrocleonus pilosus* Lec." (MCZC). Type locality, California. Henshaw 1881-1882 (check.). Henshaw 1885 (check.).

*Dinocleus jacobinus* Casey 1891:179. Lectotype (here designated), female, one of eight syntypes, labelled "Cal.", "Casey/ bequest/ 1925", "TYPE USNM/ 37268", "*D. jacobinus/ Cas.*" and with my designation label "*Dinocleus jacobinus/ Casey LECTOTYPE/ desig. Anderson*" (USNM). Type locality, San Diego, California. Fall 1901 (check.). Fall 1913. Leng 1920 (catal.).

*Dinocleus pilosus*; Casey 1891 (key, redesc.). Fall 1901 (check.). Fall 1913. Leng 1920 (catal.). Essig 1958 (biol., misident. of *Apleurus albovestitus*).

*Cleonus (Dinocleus) capillosus* Csiki 1934:66. New synonymy. New name for *Centrocleonus pilosus* LeConte. Blackwelder 1939 (check.).

*Cleonus (Dinocleus) jacobinus*; Csiki 1934 (catal.).

*Cleonus (Dinocleus) pilosus*; Csiki 1934 (catal., as jr. homonym).

*Cleonis capillosus*; O'Brien and Wibmer 1982 (catal., distn.).

*Cleonis jacobinus*; O'Brien and Wibmer 1982 (catal., distn.).

*Cleonis pillosus*; O'Brien and Wibmer 1982 (catal., distn., as jr. homonym).

**Notes about synonymy.**--- Based upon examination of very few specimens, Casey (1891) distinguished *Dinocleus jacobinus* Casey from *Dinocleus pillosus* (LeConte) on the basis of the smaller size, coarser darker and sparser elytral erect vestiture, more parallel-sided elytra with exposed humeri, and less markedly elevated alternate elytral intervals of the former. Having now examined large numbers of individuals I find that unequivocal assignment to either of the forms is not possible because of variation in all but one of the above characters, and conclude that the two forms are conspecific. I can see no differences in degree of coarseness or color of the erect elytral vestiture as noted by Casey (1891).

**Problems in recognition.**--- Specimens of *A. jacobinus* are often confused with individuals of *A. angularis* and *A. porosus*. Characters and their states allowing for the separation of these species from *A. jacobinus* are discussed in the respective "Problems in recognition" sections for those species.

**Description.**--- **Specimens examined.** 123 males, 255 females.

Data about variation in LR, WF, WRA, LP, WPB, WPT, WE1M, LE1, WPB/LP, WPT/WPB, WRA/LR, WRA/WF, WE1M/LE1, and LP/LE1 are presented in Tables 23 and 24. Size. Length, male, 7.8-11.8 mm; female, 5.8-14.3 mm. Width, male, 3.7-5.5 mm; female, 2.8-6.5 mm. Head. Eye very prominent and convex in dorsal view. Frons and vertex with sparse to dense, large to very large, deep punctures. Frons and area immediately above eyes with dense very long erect hair-like scales (short or lacking in few specimens). Area immediately behind posterior margin of eye with large, deep, irregularly impressed punctures. Area above eyes markedly elevated above rest of frons (eyes appearing distinctly browed in anterior view). Width of frons greater than width at apex of rostrum. Rostrum. Moderately robust (width at apex 0.667-0.792 times length in male; 0.649-0.800 in female) (Fig. 86). Median carina slightly to well-developed, low to high, rounded, in part obscured in many specimens by large deep punctures (Fig. 86). Dorsal and lateral punctation large to very large, deep, moderately dense to dense. Dorsally, excluding epistoma, with dense very long, erect hair-like scales (short or lacking from few specimens), and dense elongate-narrow appressed white scales. In lateral view with apical portion steeply declivous from point of antennal insertion to apex (Fig. 86a). Epistoma with apical margin rounded at middle (Fig. 86b). Mouthparts. Maxillary palpus with stipes lacking large seta on outer margin (as in Fig. 65). Labial palpi separated by more or less width of basal

article of labial palpus (as in Figs. 62-63). Prementum with two or three pairs of large setae (as in Fig. 63). Pronotum. In dorsal view with lateral margins subparallel from base to almost apical one-quarter, then divergent to apical one-quarter (thus distinctly laterally tuberculate in appearance); markedly constricted at acute angle anterior to apical one-quarter, then straight or slightly convergent to apex; widest at tubercles to subequal in width at base and apical one-quarter (Fig. 71). Dorsal and lateral punctation moderately large to large, deep, sparse to moderately dense, in some specimens the areas between punctures irregularly elevated thus appearing sculptured; punctures sparser and smaller on flanks. Scales white, elongate-fine to, elongate-narrow, appressed, sparse and small or lacking medially from disk in broad, apically narrowed patch (except for midline in most specimens); dense laterally in pair of posterolaterally directed arcuate stripes; scales sparser at lateral margins and ventrally on flanks, to uniformly dense laterally, at lateral margins and on flanks, not obscuring individual large punctures. Median carina absent to present and distinct in some specimens. Dorsally covered throughout with sparse to moderately dense, fine, very long, erect hair-like scales, each situated in large puncture. Anterolateral margin with postocular projection absent or at most very slightly developed (Fig. 86a). Prosternum. With shallow impression anterior to each procoxal cavity and with well-developed rounded swelling anterior to each prosternal

impression (Fig. 86a). Elytra. Moderately robust in general form (width at midlength 0.650-0.750 times length in males; 0.609-0.764 in females) (Fig. 71). In dorsal view with lateral margins slightly arcuate throughout length; humerus rounded, not distinct (Fig. 71). Sutural interval and intervals 3, 5, 7, and 9 with dense, very long, erect hair-like scales; sparser and slightly shorter on other intervals (short or lacking in some specimens). Dorsally with sutural interval and intervals 3, 5, 7, and 9 moderately to markedly swollen and convex (interval 5 at basal one-third and intervals 7 and 9 near base only slightly swollen in some specimens). Scales elongate-narrow, various in density and size; pattern slightly various, but generally with 2 variously sized posteromedially directed glabrous or nearly so areas, one at basal one-third from intervals 4 to 6, other at apical one-third from intervals 2 to 7, and with various number of scattered small glabrous or nearly so patches (Fig. 71). Wings. Absent. Legs. Foretibia of female with inner margin with small denticles in apical one-third; subapical tooth small, not distinct from tibial denticles. Foretibial and mesotibial unci of both sexes moderately large to large, metatibial unci of both sexes small. Tarsal claws slightly divergent, basal internal flange present, slightly to well-developed (Fig. 67). Ventral tarsal pilose vestiture of foretarsus of male absent on article 1, present as minute apical tufts on article 2, as small apical pads to moderately large pads covering

apical one-third on article 3; of foretarsus of female, lacking from article 1, present as minute to small pads at apex of articles 2 and 3; of mesotarsus and metatarsus of male, lacking from all three articles to as minute apical pads on article 2, and as small apical pads to moderately large pads covering apical one-third on article 3; of mesotarsus and metatarsus of female, lacking from articles 1 and 2, present as minute apical tufts on article 3.

**Mesosternum.** Mesosternal process flat to very slightly convex but not at all tumescent (as in Fig. 59). **Abdomen.** Ventral surface with moderately dense, elongate-fine appressed white scales. Abdominal sterna III to VI (especially III and IV) with moderately dense, individually distinct circular glabrous patches, each with large puncture and single long erect hair-like scale situated in or near the center. Apex of abdominal sternum VII of male with small dorsally directed median truncate tooth. Abdominal tergum VIII uniformly convex, apical margin slightly elevated and reflexed. **Genitalia.** Female (six examined). Abdominal sternum VIII with lateral arms narrow, slightly outwardly arcuate in basal one-half, then parallel to near apex, markedly inwardly arcuate immediately before apex (Fig. 94a). Gonocoxite II with stylus very small, antepical in position by more or less one-half length of stylus (Fig. 94b). Male (seven examined). Abdominal sternum VIII with paired sclerite with inner apices with slightly to moderately developed ventral projections (Fig. 101e).

Aedeagus elongate-narrow, in lateral view thickest at midlength; in ventral view more or less parallel-sided throughout length. Internal sac short and high, markedly apically deflexed; median dorsal pocket high, with moderately large dorsolaterally directed paramedial lobe at midheight and midlength; small median lobe on dorsal surface immediately anterior to crest of median dorsal pocket (Figs. 101a,b). Eversible apical sclerite complex with paired narrow scythe-like sclerites lightly sclerotized, each lacking median projection (Fig. 101d); adjacent ventral surface and basal portion of sides of apex of internal sac with pair of indistinct lightly sclerotized transverse sclerites (Fig. 101a).

**Geographic variation.**--- Specimens from the southern part of the species range in extreme southern California and Baja California Norte, Mexico have shorter and sparser dorsal erect vestiture than do individuals from the northern inland portions of the range. These same specimens tend to have the elytral intervals less markedly elevated throughout (Baja California Norte) or have only intervals 5 and 7 variously swollen basally (extreme southern California). Tarsal claws of the Baja California Norte individuals, although robust and not widely divergent, each have only a slightly developed basal internal flange. All of these southern individuals possess distinct glabrous patches on the abdominal sterna.

Two specimens from Point Reyes National seashore and two specimens from Redwoods Regional Park near Oakland lack or else have extremely short dorsal erect vestiture (apparently not due to abrasion), but otherwise are typical *A. jacobinus*.

**Geographic distribution.**--- This species is found in California from the San Francisco Bay area south through coastal and central California to northern coastal Baja California Norte, Mexico (Fig. 208).

**Natural history.**--- This species is associated exclusively with Pacific semi-desert habitat. Adults of have been collected on *Aster* sp., *Hemizonia* sp., *H. pungens* (H. and A.) T. and G. (Compositae); beans (Leguminosae); *Gossypium hirsutum* L. (cotton; Malvaceae); and carrots (Umbelliferae). They have also been found in various shipments of beans and raisins. Definite hosts are not known. Adults have been collected throughout the year at elevations of from 5-424 m ( $\bar{X}=87$  m,  $N=15$ ) (Fig. 232).

**Chorological relationships.**--- This species is sympatric throughout its range with western *A. albovestitus*, and is at most narrowly sympatric with southern Californian *A. angularis* and *A. porosus* from southern California or northern Baja California Norte, Mexico.



**Phylogenetic relationships.**--- Primary characters allow only for placement of this species as a member of an unresolved trichotomy also including *A. angularis* and *A. jacobinus*. This lineage is the sister-group to the *A. albovestitus*-*A. saginatus* lineage, these two lineages sister to *A. lutulentus* (Fig. 234).

Distribution of the apotypic states of secondary characters of length of erect vestiture and development of lateral pronotal tubercles allows for recognition of *A. jacobinus* as the sister-species of *A. angularis* (Fig. 235).

Table 2. Descriptive statistics for *A. jacobinus* based on eight males from Turlock, California

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	2.16-2.88	2.530	0.314	0.148
WF	1.76-2.24	2.070	0.226	0.107
WRA	1.52-1.92	1.770	0.175	0.082
LP	3.36-4.32	3.920	0.425	0.201
WPB	3.68-4.64	4.320	0.497	0.234
WPT	3.84-4.96	4.440	0.525	0.247
WE1M	5.12-6.88	6.140	0.769	0.362
LE1	7.20-10.24	9.000	1.325	0.625
B. Proportions.				
WPB/LP	1.07-1.16	1.102	0.044	0.021
WPT/WPB	0.97-1.07	1.028	0.047	0.022
WRA/LR	0.67-0.73	0.701	0.035	0.017
WRA/WF	0.79-0.89	0.856	0.056	0.026
WE1M/LE1	0.65-0.71	0.684	0.032	0.015
LP/LE1	0.40-0.47	0.437	0.033	0.016

Table 24. Descriptive statistics for *A. jacobinus* based on eight females from Turlock, California

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	2.08-3.04	2.670	0.471	0.222
WF	1.84-2.48	2.150	0.336	0.158
WRA	1.60-2.08	1.840	0.213	0.100
LP	3.36-4.80	4.120	0.723	0.341
WPB	3.68-4.96	4.420	0.666	0.314
WPT	3.84-5.28	4.580	0.736	0.347
WE1M	5.12-7.52	6.480	1.183	0.557
LE1	7.52-10.88	9.600	1.702	0.802

B. Proportions.

WPB/LP	1.03-1.13	1.075	0.055	0.026
WPT/WPB	1.00-1.07	1.036	0.039	0.018
WRA/LR	0.65-0.77	0.693	0.056	0.026
WRA/WF	0.79-0.92	0.859	0.058	0.027
WE1M/LE1	0.66-0.69	0.675	0.014	0.006
LP/LE1	0.42-0.45	0.429	0.018	0.008

4.4.2.2.6 *Apleurus (Apleurus) albovestitus* (Casey), new combination

(Figs. 75-79, 88, 95, 103, 209)

*Cleonus molitor* LeConte 1858:78 [not Gyllenhal 1834:174].

New synonymy. Holotype (examined), female, labelled with a gold circle (=California), "*Cleonus/ molitor/ Lec.*", "Type/ 5183", "Anobium!/ parasitic!" (MCZC). Type locality, California. Gemminger and von Harold 1871 (catal.).

*Centrocleonus molitor*; LeConte 1876a (key, diag.). Henshaw 1881-1882 (check.). Henshaw 1885 (check.). Wickham 1889.

*Dinocleus albovestitus* Casey 1891:183. Lectotype (here designated), male, one of an undetermined number of syntypes, labelled "Los Angeles/ Co. CAL", "TYPE USNM/ 37273", "CASEY/ bequest/ 1925", "D./ albovestitus/ Cas." and with my designation label "*Dinocleus albovestitus/ Csy. LECTOTYPE/ desig. Anderson*" and genitalia in microvial under labels (USNM). Type locality, Los Angeles, California. Fall 1901 (check.). Casey 1904. Leng 1920 (catal.). Ting 1936 (morphol.). Bruhn 1947 (morphol.). Sanders 1960 (morphol.).

*Dinocleus densus* Casey 1891:185. New synonymy. Lectotype (here designated), male, one of four syntypes, labelled "Winslow", "CASEY/ bequest/ 1925", "TYPE USNM/ 37277", "D./ densus/ Cas." and with my designation label "*Dinocleus/ densus Csy./ LECTOTYPE/ desig. Anderson*" (USNM). Type locality, Winslow, Arizona. Wickham 1896 (check.). Leng 1920 (catal.).

*Dinocleus molitor*; Casey 1891 (key, redesc.). Fall 1901 (check.). Champion 1902-1906 (syn., distn.). Casey

1904. Leng 1920 (catal.).

*Dinocleus wickhami* Casey 1891:184. **New synonymy.** Holotype (examined), male, labelled "Indio/ California/ Wickham", "CASEY/ bequest/ 1925", "TYPE USNM/ 37276", "D./ wickhami/ Cas." (USNM). Type locality, Indio, California. Fall 1901 (check.). Leng 1920 (catal.). Tanner 1934 (check.).

*Dinocleus interruptus* Casey 1904:322 [not Zoubkoff 1829:162]. **New synonymy.** Lectotype (here designated), male, one of three syntypes, labelled "Ut", "CASEY/ bequest/ 1925", "TYPE USNM/ 37274", "interruptus" and with my designation label "*Dinocleus interruptus* Csy./ LECTOTYPE/ desig. Anderson" (USNM). Type locality, Utah. Leng 1920 (catal.).

*Dinocleus mexicanus* Casey 1904:322. **New synonymy.** Lectotype (here designated), male, one of two syntypes, labelled "Guer.", "CASEY/ bequest/ 1925", "TYPE USNM/ 37275", "mexicanus" and with my designation label "*Dinocleus mexicanus* Csy./ LECTOTYPE/ desig. Anderson" (USNM). Type locality, Guerrero, Mexico. Champion 1902-1906 (distn.).

*Cleonus (Dinocleus) albovestitus*; Csiki 1934 (catal.).

*Cleonus (Dinocleus) densus*; Csiki 1934 (catal.).

*Cleonus (Dinocleus) interruptus*; Csiki 1934 (catal.).

*Cleonus (Dinocleus) mexicanus*; Csiki 1934 (catal.).

*Cleonus (Dinocleus) structor* Csiki 1934:67. **New synonymy.**

New name for *Cleonus molitor* LeConte. Blackwelder 1939

(check.).

*Cleonus (Dinocleus) wickhami*; Csiki 1934 (catal.).

*Cleonus mexicanus*; Blackwelder 1947 (check.).

*Dinocleus pilosus*; Essig 1958 (biol., misident.).

*Cleonus albovestitus*; Hardy and Andrews 1976 (distn.,  
biol.). Andrews et al. 1979 (distn., biol.).

*Cleonis albovestitus*; O'Brien and Wibmer 1982 (catal.,  
distn.).

*Cleonis densus*; O'Brien and Wibmer 1982 (catal., distn.).

*Cleonis Interruptus*; O'Brien and Wibmer 1982 (catal.,  
distn.).

*Cleonis mexicanus*; O'Brien and Wibmer 1982 (catal., distn.).

*Cleonis molitor*; O'Brien and Wibmer 1982 (catal., distn., as  
jr. homonym).

*Cleonis structor*; O'Brien and Wibmer 1982 (catal., distn.).

*Cleonis wickhami*; O'Brien and Wibmer 1982 (catal., distn.).

*molitor, Incertae sedis*; Chevrolat 1873.

**Notes about synonymy.**--- I consider *A. albovestitus* to be composed of a number of largely allopatric morphotypes most of which can be distinguished solely on the basis of patterns of surface vestiture (Figs. 75-79), but which exhibit continuous intergradation in zones of parapatry or narrow sympatry (see "Geographic variation" section).

One of these morphotypes is largely black and glabrous, most of the surface vestiture having been abraded (Figs.

78-79). This is *Centrocleonus molitor* LeConte. *Dinocleus mexicanus* Casey is this same morphotype, but in which the scales have not been extensively abraded. *Dinocleus densus* Casey, is a form in which there has been no abrasion of scales. This morphotype is confined to the Colorado, Virgin, and Gila River drainages.

The second morphotype, that with dense elytral and ventral abdominal scales and lacking abdominal glabrous patches but with small elytral glabrous patches, includes *Dinocleus interruptus* Casey, *D. wickhami* Casey, and *D. albovestitus* Casey (Figs. 76-77).

None of the described species are of the third morphotype, recognized, among other characters, by the larger and more numerous elytral and abdominal glabrous patches (Fig. 75).

Since I have been unable to find other structural features which correlate with the variation in patterns of surface vestiture to allow for reliable separation of the various morphotypes, I conclude that a single species is present, but with allopatric forms having different locally adaptive patterns of surface vestiture.

Champion (1902-1906) incorrectly stated that *D. mexicanus* Casey is a new name for *Centrocleonus molitor* LeConte.

Problems in recognition.--- Individuals of *A. albovestitus* might only be confused with those of *A. saginatus* although

key characters should serve to reliably separate all members of the two species. Special note should be made of the marked secondary sexual dimorphism in extent of ventral tarsal vestiture in *A. saginatus* that is not as marked in *A. albovestitus*. This allows for easy recognition of males of *A. saginatus*.

All *A. albovestitus* that possess glabrous ventral abdominal patches are found only in western California; those *A. albovestitus* found sympatrically with *A. saginatus* have extremely dense scales on the abdominal sterna and lack the glabrous patches present in all *A. saginatus*.

**Description.**--- Specimens examined. 1251 males, 1470 females. Data about variation in LR, WF, WRA, LP, WPB, WPT, WE1M, LE1, WPB/LP, WPT/WPB, WRA/LR, WRA/WF, WE1M/LE1, and LP/LE1 are presented in Tables 25 and 26. Size. Length, male, 11.8-16.8 mm; female, 12.8-21.4 mm. Width, male, 5.1-7.2 mm; female, 5.4-8.8 mm. Head. Eye slightly prominent and convex in dorsal view. Frons and vertex with sparse to moderately dense, small, shallow punctures. Frons largely lacking suberect or erect vestiture, with at most only sparse, small suberect scales immediately above eyes. Area immediately behind posterior margin of eye with small, shallow to moderately deep, irregularly impressed punctures. Area above eyes continuous with and not elevated above rest of frons (eyes not browed in anterior view). Width of frons greater than to subequal in width to apex of rostrum.



Rostrum. Moderately robust (width at apex 0.622-0.775 times length in male; 0.632-0.794 in female) (Fig. 88). Median carina lacking. Dorsal and lateral punctation small to moderately large, moderately deep, sparse to moderately dense. Dorsally, excluding epistoma, either lacking suberect or erect vestiture or with at most scattered short suberect scales; and lacking (abraded) or with dense elongate-narrow appressed white scales. In lateral view with apical portion steeply declivous from point of antennal insertion to apex (Fig. 88a). Epistoma with apical margin rounded at middle (Fig. 88b). Mouthparts. Maxillary palpus with stipes with large seta on outer margin (as in Fig. 64). Labial palpi separated by more or less width of basal article of labial palpus (as in Fig. 62-63). Prementum with two or three pairs of large setae (as in Fig. 63). Pronotum. In dorsal view with lateral margins straight and subparallel to slightly convergent from base to almost apical one-quarter, not to slightly expanded laterally at apical one-quarter (thus indistinctly laterally tuberculate in appearance); moderately constricted at obtuse to subacute angle anterior to apical one-quarter, then straight and slightly convergent to apex; widest at base to subequal in width at base and apical one-quarter (Figs. 75-79). Dorsal and lateral punctation small, shallow, sparse to moderately dense; punctures sparser on flanks; outer margins of larger punctures not swollen or glabrous. Scales white, elongate-narrow to broad, appressed, entirely lacking

(abraded) to sparse and small or lacking medially from disk in broad, apically narrowed patch (except for midline in most specimens); very dense laterally and ventrally on flanks onto prosternum; scales sparser at lateral margins. Median carina lacking to present but irregularly developed but with at least a broad low rounded median swelling in most specimens. Dorsally lacking suberect or erect vestiture or with sparse, short, erect hair-like scales, each situated in large puncture. Anterolateral margin with postocular projection slightly to moderately developed (Fig. 88a). Prosternum. With shallow impression anterior to each procoxal cavity and with well-developed rounded swelling anterior to each prosternal impression (Fig. 88a). Elytra. Moderately robust to moderately elongate-narrow in general form (width at midlength 0.590-0.724 times length in males; 0.559-0.694 in females) (Figs. 75-79). In dorsal view with lateral margins straight, slightly convergent to divergent from apical one-third to humerus; humerus rounded to obtuse, indistinct to distinct, with at most only very few small glabrous shiny tubercles. Dorsal suberect or erect vestiture lacking to sparse, short and fine. Dorsally with all intervals flat. Scales, if present, elongate-narrow to elongate-fine, white to golden. Scale pattern extremely various; with scales entirely or largely lacking (abraded) except near apex on declivity (Figs. 78-79), to present to various extent (not abraded), uniformly dense and large on sutural interval and intervals 3, 5, 7, and 9, uniformly

sparse and small on intervals 2, 4, 6, and 8, distinct glabrous patches absent (Fig. 77); or, to scales not at all abraded, uniformly dense and large on sutural interval and intervals 3, 5, 7, and 9, and sparse and small to various degree on intervals 2, 4, 6, and 8, with at most scattered but few glabrous patches present (Fig. 76); or, to sutural interval with scales sparse and small or large and dense, otherwise with scales of elytral intervals extremely various in size and density, with numerous irregularly distributed small and moderately large glabrous or nearly so patches present (Fig. 75). Wings. Short (slightly shorter than elytra in length [0.2%, N=4]), to long (slightly greater than elytra in length [99.8%, N=1717]). Branches of 2A complete and joined at base. Legs. Foretibia of female with inner margin with small to minute denticles in apical one-third; subapical tooth minute, not distinct from tibial denticles. Foretibial and mesotibial unci of both sexes moderately large to large, metatibial uncus of both sexes small. Tarsal claws slightly divergent, basal internal flange present, well-developed (as in Fig. 67). Ventral tarsal pilose vestiture of foretarsus of male absent to present as minute apical tufts on articles 1 to 3; of mesotarsus of male, lacking from articles 1 and 2, present as minute apical tufts on article 3; lacking from articles 1 to 3 of all tarsi of female and from articles 1 to 3 of metatarsus of male. Mesosternum. Mesosternal process flat to very slightly convex but not at all tumescent (as in Fig.

59). Abdomen. Ventral surface either with scales present, white, appressed, along posterior margins of abdominal sterna, also with scattered, suberect short hair-like scales, each situated in large puncture; or, with white appressed scales uniformly very dense, abdominal sterna III to VI without glabrous patches to with scales moderately dense, individually distinct circular glabrous patches present (especially on abdominal sterna III and IV), each with large puncture and single short suberect hair-like scale situated in or near the center. Apex of abdominal sternum VII of male with large dorsally directed median truncate tooth. Abdominal tergum VII of female slightly medially longitudinally carinate, apical margin moderately elevated and reflexed; abdominal tergum VIII of female markedly medially longitudinally carinate, apical margin moderately elevated and reflexed. Genitalia. Female (12 examined). Abdominal sternum VIII with lateral arms broad, straight and slightly divergent from base to apical one-third, slightly inwardly arcuate from apical one-third to apex (Fig. 95a). Gonocoxite II with stylus absent, with a darkly sclerotized slightly elevated dorsal subapical ridge (Fig. 95b). Male (eight examined). Abdominal sternum VIII with paired sclerite with inner apices with well-developed ventral projections (Fig. 103e). Aedeagus elongate-narrow, in lateral view thickest at midlength; in ventral view more or less parallel-sided throughout length. Internal sac short and high, slightly apically deflexed; median dorsal pocket

high, with moderately large dorsolaterally directed paramedial lobe near dorsal margin at midlength; small median lobe on dorsal surface immediately anterior to crest of median dorsal pocket (Figs. 103a,b). Eversible apical sclerite complex with paired narrow scythe-like sclerites darkly sclerotized, each with long dorsoapically directed median projection (Fig. 103d); adjacent ventral surface and basal portion of sides of apex of internal sac with pair of indistinct lightly sclerotized transverse sclerites (Fig. 103a).

Geographic variation.--- Variation in the pattern of surface vestiture in members of this species is very extensive (Figs. 75-79). There is also slight variation in other structural features but these could not be reliably correlated with the variation in vestiture to allow recognition of more than one species (see also "Notes about synonymy" section).

Surface vestiture varies from largely absent to dense. In individuals in which surface vestiture is variously lacking, this appears to be due to abrasion. Such specimens are primarily black and glabrous (except for the tibiae and tarsi) and possess moderately dense to dense scales only along the lateral margins of the elytra and the posterior margins of the abdominal sterna (especially V and VI), and to a various extent in some specimens, also in the apical one-third of the elytra, onto the elytral declivity (Figs.

78-79). Scattered single scales are also found variably on the thoracic and abdominal sterna, elytra and femora. In individuals in which scales are variably present on the elytra, the scales are uniformly dense and large on all intervals to slightly less dense and smaller on intervals 2, 4, and 6. Scales are very easily abraded on these specimens. Individuals of this morphotype have been collected at elevations of from -67-870 m ( $X=146$  m,  $N=23$ ).

This form is found throughout the southern portion of the Colorado River drainage, the western portion of the Gila River drainage, and throughout Imperial County, California. Individuals from the latter area tend to have the elytral scales denser and not abraded and although lacking distinct glabrous elytral patches, the general scale pattern clearly grades into that pattern to be discussed next in which small elytral glabrous patches are present (Fig. 76). Individuals of this morphotype seem restricted to *Pluchea sericea* (Nutt.) Coville (Compositae) in sand dune habitats along the margins of waterways in these areas.

Another form of *A. albovestitus* is also found in these and other areas, and although the ranges of the two morphotypes narrowly overlap, they do not appear to be found together. Individuals of this form possess large and dense elytral surface vestiture that is not prone to abrasion. All abdominal sterna have uniformly very dense scales; no glabrous patches are present. Elytral scales are either uniformly large and dense to very dense, and are lacking or

very sparse in scattered small to moderately large patches on elytral intervals 2 to 8 (Fig. 76). Scales of the sutural interval of many of these specimens are uniformly small and ~~sparse~~. Punctures of elytral striae are small and indistinct. This form is widespread from Nevada and Utah south to extreme western Texas, New Mexico, Arizona and southeastern California, south into Baja California Sur, Mexico. Individuals of this form in Utah and northern Arizona, along the Colorado and Virgin River drainages differ from those elsewhere by largely lacking glabrous or nearly so patches on the elytra (Fig. 77). Scales of some individuals in these areas are less dense and smaller on intervals 2, 4, and 6, much the same pattern as in specimens of the largely glabrous morphotype in which elytral scales have not been extensively abraded. This similarity in scale pattern suggests the notion that the largely glabrous individuals in the southern portion of the Colorado River drainage may have reached this area as a result of downriver dispersal from parent populations along the Virgin and northern Colorado Rivers. Individuals of this morphotype have been collected at elevations of from 2-1446 m ( $X=231$  m,  $N=49$ ).

A third pattern of vestiture is found in individuals from southern coastal to northern coastal and central California. In this form, scales of the elytra are less dense and smaller than in the previous forms, and the elytral glabrous or nearly so patches are generally larger

and more numerous (Fig. 75). Punctures of the elytral striae are generally slightly larger and more distinct than in the other morphotypes. Abdominal sterna (especially III and IV) of many individuals with this elytral scale pattern possess small to moderately large, indistinct to distinct, glabrous areas, each with a single short suberect hair-like scale situated in the center. Suberect vestiture, lacking from the elytra on the other morphotypes, is short, sparse and fine on the elytra of many of these specimens. Elytral scale patterns of this and of the second morphotype completely intergrade in southern California. Individuals of this third morphotype have been collected at elevations of from 41-3185 m ( $X=998$  m,  $N=61$ ).

The potential adaptive significance of variation in surface vestiture is not known but is suspected to be influenced by thermoregulatory effects, or, more likely, by cryptic effects resulting from similarity in color and pattern to the ground substrate (see "Evolutionary Trends" section).

Brachypterous specimens of *A. albovestitus* are only known from two localities in Baja California Norte, Mexico.

**Geographic distribution.** -- This species is distributed from New Mexico and western Texas west through Utah and Nevada to northern California in the north through Arizona and extreme northwestern Mexico to southern California and Baja California Sur, Mexico in the south (Fig. 209).



**Natural history.**--- This species is associated with a variety of habitats as follows; desert-grassland transitional, Pacific semi-desert, and Great Basin, Chihuahuan, Mojave, and Sonoran deserts. Adults of this species have been collected in sand dune habitats in California, Arizona, Utah, Nevada (label data; Andrews et al. 1979; Hardy and Andrews 1976) and otherwise primarily in dry washes and stream beds throughout the species range. Andrews et al. 1979 tentatively suggest that the species is an obligate sand associate.

Adults have been collected on a wide variety of plants as follows: *Ephedra* sp., *E. californica* Wats., (Ephedraceae); *Ambrosia* sp., *A. psilostachya* DC., *Artemesia tridentata* Nutt., *Baileya multiradiata* Harv. and Gray, *B. pleniradiata* Harv. and Gray, *Baccharis glutinosa* Pers., *Chrysothamnus nauseosus* (Pall.) Britton, *Dicoria* sp., *Franseria* sp., *F. confertiflora* (DC.) Rydb., *F. dumosa* Gray, *Flourensia cernua* DC., *Gutierrezia microcephala* (DC.) Gray, *Aplopappus acradenius* (Greene) Blake, *Hymenoclea* sp., *H. salsola* T. and G., *Hymenothrix wislizeni* Gray, "*Palafoxia arida*", *P. linearis* (Cav.) Lag., *Pluchea sericea* (Nutt.) Coville, *Verbesina enceloides* (Cav.) Benth. and Hook., *Xanthium* sp., (all Compositae); *Larrea tridentata* (DC.) Coville (Zygophyllaceae); *Hordeum* sp., *Sitanion* sp., (Graminae); *Datura* sp., (Solanaceae); *Gossypium* sp., (Malvaceae); *Eriogonum fasciculatum* Benth. var. *polifolium* (Polygonaceae); *Astragalus* sp., *Cercidium floridum* Benth.,

*Medicago sativa* L., *Olneya tesota* Gray, *Prosopis juliflora* (Swartz) DC. (all Leguminosae); *Atriplex* sp., *A. lentifolmis* (Torr.) Wats., *Salsola kali* L., *Sarcobatus vermiculatus* (Hook.) Torr. (all Chenopodiaceae); *Vitus* sp. (Vitaceae); *Prunus persica* Batsch. (peach; Rosaceae); *Raphanus sativus* L. (radish; Cruciferae). No definite hosts are known, but the vast majority of plant association are with Compositae; records of occurrence on members of other families are generally single instances of but one or two individuals and very likely represent chance occurrences. I suspect that a wide variety of Compositae serve as host plants for this species.

Adults have been collected throughout the year at elevations from -67-3185 m (X=567 m, N=133) (Fig. 232).

As discussed in the "Geographic variation" section, individuals of each of the three morphotypes occur at slightly different elevations.

**Chorological relationships.**--- This species is sympatric with *A. luteolentus* in Arizona, New Mexico and western Texas; with *A. angularis* throughout most of its range, with the exception of northern California; with *A. jacobinus* throughout the entire range of that species in California; narrowly sympatric with *A. porosus* in Baja California Norte, Mexico and southern California; and sympatric with its sister-species *A. saginatus* throughout Arizona, southern New Mexico, and extreme northwestern Mexico.

I have caught large numbers of specimens of *A. albovestitus* together with *A. lutulentus* on *Chrysothamnus nauseosus* (Compositae) in the vicinity of Portal, Arizona and Animas, New Mexico.

**Phylogenetic relationships.**--- Distribution of the apotypic states of primary characters of presence or absence of stylus on gonocoxite II in female and degree of development of the ventral projection at the inner angle of each sternite of sternum VIII in male place this species as the sister-species of *A. saginatus*. This lineage is sister to the *A. angularis*-*A. jacobinus*-*A. porosus* lineage (Fig. 234).

Distribution of apotypic states of numerous secondary characters support recognition of *A. albovestitus* and *A. saginatus* as sister-species (Fig. 235).

Table 25. Descriptive statistics for *A. albovestitus* based on eight males from Portal, Arizona.

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	2.24-3.04	2.600	0.449	0.212
WF	1.60-2.00	1.820	0.200	0.094
WRA	1.60-2.00	1.800	0.231	0.109
LP	3.52-4.80	4.280	0.688	0.324
WPB	3.68-5.28	4.520	0.756	0.356
WPT	3.60-4.80	4.290	0.608	0.287
WE1M	5.28-7.36	6.440	1.087	0.512
LE1	8.48-11.68	10.000	1.653	0.779
B. Proportions.				
WPB/LP	1.03-1.10	1.056	0.039	0.018
WPT/WPB	0.91-1.00	0.951	0.045	0.021
WRA/LR	0.63-0.75	0.695	0.061	0.029
WRA/WF	0.91-1.00	0.989	0.048	0.023
WE1M/LE1	0.60-0.72	0.645	0.055	0.026
LP/LE1	0.40-0.45	0.428	0.024	0.011

Table 26. Descriptive statistics for *A. albovestitus*  
based on eight females from Portal, Arizona

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	2.24-3.36	2.770	0.559	0.263
WF	1.60-2.32	2.030	0.400	0.189
WRA	1.52-2.24	1.940	0.414	0.195
LP	3.68-5.12	4.550	0.820	0.387
WPB	3.84-5.60	4.780	1.012	0.477
WPT	3.84-5.44	4.640	0.907	0.428
WE1M	5.60-8.16	6.840	1.450	0.683
LE1	8.48-13.76	10.780	2.623	1.236
B. Proportions.				
WPB/LP	0.98-1.09	1.049	0.061	0.029
WPT/WPB	0.94-1.00	0.972	0.031	0.015
WRA/LR	0.67-0.74	0.700	0.047	0.022
WRA/WF	0.92-1.00	0.955	0.038	0.018
WE1M/LE1	0.95-0.69	0.637	0.054	0.025
LP/LE1	0.37-0.46	0.425	0.039	0.018

4.4.2.2.7 *Apleurus (Apleurus) saginatus* (Casey), new  
combination

(Figs. 63-64, 80, 87, 96, 104, 206)

*Dinocleus saginatus* Casey 1891:182. Holotype (examined), male, labelled "Ariz.", "CASEY/ bequest/ 1925", "TYPE USNM/ 37272", "*D./ saginatus/ Cas.*" (USNM). Type locality, Arizona. Leng 1920 (catal.).

*Dinocleus dentatus* Champion 1902-1906:100. New synonymy. Holotype (examined), female, labelled "Type", inverted "Sp. figured", "Pinos Altos/ Chihuahua,/ Mexico/ Buchan-Hepburn", "B.C.A. Col. IV.4./ *Dinocleus/ dentatus,/ Champ.*", inverted "*dentatus, Ch.*" (BMNH). Type locality, Pinos Altos, Chihuahua, Mexico.

*Cleonus (Dinocleus) dentatus*; Csiki 1934 (catal.).

*Cleonus dentatus*; Blackwelder 1947 (check.).

*Cleonus (Dinocleus) saginatus*; Csiki 1934 (catal.).

*Cleonis dentatus*; O'Brien and Wibmer 1982 (catal., distn.).

*Cleonis saginatus*; O'Brien and Wibmer 1982 (catal., distn.).

**Notes about synonymy.**--- Following examination of types and numerous specimens I regard *Dinocleus saginatus* Casey and *D. dentatus* Casey as clearly conspecific.

**Problems in recognition.**--- Individuals of this species might only be confused with those of *A. albovestitus* (see

"Problems in recognition" section for *A. albovestitus*).

**Description.**--- Specimens examined. 597 males, 657 females. Data about variation in LR, WF, WRA, LP, WPB, WPT, WELM, LEL, WPB/LP, WPT/WPB, WRA/LR, WRA/WF, WELM/LEL, and LP/LEL are presented in Tables 27 and 28. **Size.** Length, male, 14.9-19.5 mm; female, 15.9-21.8 mm. Width, male, 5.8-8.0 mm; female, 6.2-9.0 mm. **Head.** Eye slightly prominent and convex in dorsal view. Frons and vertex with sparse, small, shallow punctures. Frons largely lacking suberect or erect vestiture, with at most only sparse, small suberect scales immediately above eyes; white elongate-narrow appressed scales dense above eyes, sparser medially. Area immediately behind posterior margin of eye with small, shallow to moderately deep, irregularly impressed punctures. Area above eyes continuous with and not elevated above rest of frons (eyes not browed in anterior view). Width of frons greater than to subequal in width to apex of rostrum. **Rostrum.** Moderately robust (width at apex 0.629-0.730 times length in male; 0.632-0.744 in female) (Fig. 87). Median carina lacking to variously developed as low rounded to sharp fine glabrous line. Dorsal and lateral punctation small, shallow, sparse. Dorsally, excluding epistoma, with short scattered suberect scales; and with dense elongate-narrow appressed white scales. In lateral view with apical portion steeply declivous from point of antennal insertion to apex (Fig. 87a). Epistoma with apical margin rounded at middle (Fig.

87b). Mouthparts. Maxillary palpus with stipes with large seta on outer margin (Fig. 64). Labial palpi separated by more or less width of basal article of labial palpus (Fig. 63). Prementum with two or three pairs of large setae (Fig. 63). Pronotum. In dorsal view with lateral margins slightly arcuate to straight and subparallel to slightly convergent from base to almost apical one-quarter, not to slightly expanded laterally at apical one-quarter (thus indistinctly laterally tuberculate in appearance); moderately constricted at obtuse to subacute angle anterior to apical one-quarter, then straight and slightly convergent to apex; widest at base to subequal in width at base and apical one-quarter (Fig. 80). Dorsal and lateral punctation small, shallow, sparse to moderately dense; punctures sparser and shallower on flanks; outer margins of larger punctures swollen, glabrous and shiny (appearing as small tubercles). Scales white, elongate-narrow to broad, appressed sparse and small or lacking medially on disk in broad, apically narrowed patch (except for midline in most specimens); very dense laterally and ventrally on flanks onto prosternum; scales sparser at lateral margins. Median carina lacking to present and irregularly developed in basal one-half as low, broad glabrous line. Dorsally with sparse, short, erect hair-like scales, each situated in large puncture. Anterolateral margin with postocular projection slightly developed (Fig. 87a). Prosternum. With shallow impression anterior to each procóxal cavity and with well-developed rounded swelling



anterior to each prosternal impression (Fig. 87a). Elytra. Moderately robust to moderately elongate-narrow in general form (width at midlength 0.590-0.620 times length in males; 0.563-0.635 in females) (Fig. 80). In dorsal view with lateral margins straight or nearly so, slightly convergent from apical one-third to humerus; humerus obtuse, distinct, with numerous distinct small glabrous shiny tubercles (also present at bases of adjacent elytral intervals) (Fig. 80). Dorsal suberect or erect vestiture lacking. Dorsally with all intervals flat. Scales appressed, white to golden in color, various in size and density. Scale pattern various, with numerous small irregularly distributed glabrous or nearly so patches (Fig. 80). Wings. Long (slightly greater than elytra in length). Branches of 2A complete and joined at base. Legs. Foretibia of female with inner margin with small to minute denticles in apical one-third; subapical tooth minute to small, not distinct from tibial denticles. Foretibial and mesotibial unci of both sexes moderately large to large, metatibial unculus of both sexes small. Tarsal claws slightly divergent, basal internal flange present, well-developed (as in Fig. 67). Ventral tarsal pilose vestiture of foretarsus of male present as small rounded apical pads on article 1, as moderately large rounded pads covering apical one-half on article 2, and as large rounded pads covering apical one-half on article 3; of foretarsus of female, lacking from articles 1 and 2, present as minute apical tufts on article 3; of mesotarsus of male, absent to

present as minute apical tufts on article 1, present as small apical pads on article 2, and present as moderately large round pads covering apical one-third on article 3; of metatarsus of male, absent on article 1, present as minute apical tufts on article 2, and present as small apical pads on article 3; lacking from articles 1 to 3 of mesotarsus and metatarsus of female. Mesosternum. Mesosternal process flat to very slightly convex but not at all tumescent (as in Fig. 959). Abdomen. Ventral surface with very dense elongate-narrow to broad, white, appressed scales; abdominal sterna III to VI (especially III and IV) with moderately dense individually distinct large circular glabrous shiny patches, each with large puncture and single short suberect hair-like scale situated in or near the center. Apex of abdominal sternum VII of male with large dorsally directed median truncate tooth. Abdominal tergum VII of female slightly medially longitudinally carinate, apical margin moderately elevated and reflexed; abdominal tergum VIII of female markedly medially longitudinally carinate, apical margin moderately elevated and reflexed. Genitalia. Female (six examined). Abdominal sternum VIII with lateral arms broad, straight and slightly divergent from base to near apex, slightly inwardly arcuate near apex (Fig. 96a). Gonocoxite II with stylus absent, with a darkly sclerotized slightly elevated dorsal subapical ridge (Fig. 96b). Male (six examined). Abdominal sternum VIII with paired sclerite with inner apices with well-developed ventral projections

(as in Fig. 103e). Aedeagus elongate-narrow, in lateral view thickest at midlength; in ventral view more or less parallel sided throughout length. Internal sac short and high, slightly apically deflexed; median dorsal pocket high, with moderately large dorsolaterally directed paramedial lobe near dorsal margin at midlength (Figs. 104a,b). Eversible apical sclerite complex with paired narrow scythe-like sclerites darkly sclerotized, each with long dorsoapically directed median projection (Fig. 104d); adjacent ventral surface and basal portion of sides of apex of internal sac with pair of indistinct lightly sclerotized transverse sclerites (Fig. 104a).

**Geographic variation.**--- None noted.

**Geographic distribution.**--- This species is found in southeastern Arizona and extreme southwestern New Mexico, south into northwestern Mexico (Fig. 206).

**Natural history.**--- This species appears restricted to dry washes and riparian habitats in the desert-grassland transitional habitat bordering the eastern Sonoran desert region. Adults have been collected on *Baccharis glutinosa* Pers., *Chrysothamnus nauseosus* (Pall.) Britton, *Hymenoclea monogyra* Torr. and Gray (all Compositae). Definite hosts are not known. Adults have been collected throughout the year at elevations of from 283-1820 m ( $\bar{X}$ =1152 m, N=36) (Fig. 232).

**Chorological relationships.**--- This species is sympatric throughout its range with *A. angularis*, *A. lutulentus*, and its sister-species *A. albovestitus*. I know of no records of *A. saginatus* being caught with any of these three species.

**Phylogenetic relationships.**--- Distribution of the apotypic states of primary characters of presence or absence of stylus on gonocoxite II in female and degree of development of the ventral projection at the inner angle of each sternite of sternum VIII in male place this species as the sister-species of *A. albovestitus*. This lineage is sister to the *A. angularis*-*A. jacobinus*-*A. porosus* lineage (Fig. 234).

— Distribution of apotypic states of numerous secondary characters support recognition of *A. saginatus* and *A. albovestitus* as sister-species (Fig. 235).

Table 27. Descriptive statistics for *A. saginatus* based on eight males from 5 mi. SW. Patagonia, Arizona

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	2.72-3.28	2.940	0.285	0.134
WF	1.84-2.16	1.980	0.154	0.072
WRA	1.76-2.16	1.950	0.212	0.100
LP	4.32-5.60	4.840	0.676	0.319
WPB	4.48-5.60	5.080	0.555	0.262
WPT	4.32-5.12	4.720	0.406	0.191
WE1M	6.56-7.84	7.280	0.691	0.326
LE1	11.04-13.12	12.060	1.133	0.534
B. Proportions.				
WPB/LP	1.00-1.11	1.052	0.062	0.029
WPT/WPB	0.88-0.97	0.930	0.044	0.021
WRA/LR	0.63-0.73	0.664	0.056	0.026
WRA/WF	0.92-1.08	0.985	0.079	0.037
WE1M/LE1	0.59-0.62	0.604	0.016	0.007
LP/LE1	0.38-0.43	0.401	0.024	0.011

Table 28. Descriptive statistics for *A. saginatus* based on eight females from 5 mi. SW. Patagonia, Arizona

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	2.96-3.60	3.156	0.288	0.128
WF	2.00-2.32	2.151	0.152	0.068
WRA	1.92-2.40	2.169	0.228	0.101
LP	4.96-5.92	5.351	0.418	0.186
WPB	4.96-6.08	5.511	0.525	0.233
WPT	4.64-5.76	5.156	0.573	0.255
WE1M	7.36-8.96	8.018	0.782	0.348
LE1	12.48-14.72	13.387	1.170	0.520
B. Proportions.				
WPB/LP	1.00-1.06	1.029	0.031	0.014
WPT/WPB	0.91-0.97	0.935	0.030	0.013
WRA/LR	0.63-0.74	0.688	0.056	0.025
WRA/WF	0.96-1.04	1.007	0.046	0.021
WE1M/LE1	0.56-0.64	0.599	0.030	0.013
LP/LE1	0.39-0.43	0.400	0.017	0.008

#### 4.5 Genus *Cleonidius* Casey

*Curculio*; Pallas 1781 (sp. desc.). Herbst 1795.

*Cleonus*; Say 1831 (sp. desc.). Melsheimer 1853 (in part; check.). Gemminger and von Harold 1871 (in part; catal.). LeConte 1876a (key, sp. desc.). Henshaw 1881-1882 (check.). LeConte and Horn 1883 (key). Henshaw 1885 (check.). Wickham 1899. Wickham 1896 (check.). Fall 1897 (sp. desc.). Fall 1901 (check.). Wickham 1902 (check.). Fletcher 1906. Fall and Cockerell 1907 (in part; check.). Pierce 1907 (biol.). Chittenden 1911 (biol.). Ely 1913 (biol.). Gibson 1914. Anderson 1914. Blatchley and Leng 1916 (key, sp. redescs.). Yothers 1916. Leng 1920 (in part; catal.). Leonard 1926 (in part; check.). Böving 1927 (in part; larval key). Bradley 1930 (in part; key). Wilcox et al. 1934. Brimley 1938 (check.). Blackwelder 1947 (in part; check.). Bruhn 1947 (morphol.). Essig 1958 (biol.). Sanders 1960 (morphol.). Tanner 1966 (in part; sp. redesc., biol.). Kingsolver 1972. Kumar et al. 1976 (biol.). Arnett et al. 1980 (key, biol.).

*Rhynchophorus*; Say 1831 (in part; sp. desc.).

*Cleonis*; Kirby 1837 (sp. desc.). O'Brien and Wibmer 1982 (catal., distn.). O'Brien and Wibmer 1984 (in part).

*Lixus*; Zoubkoff 1833 (sp. desc.). Boheman 1836 (sp. desc.). Randall 1838 (in part; sp. desc.). Mannerheim 1843 (sp. desc.). Motschulsky 1845 (sp. desc.). Gemminger and von Harold 1871 (in part; catal.). Capiomont and Leprieur

1874 (sp. redesc.). Sprague and Austin 1875. LeConte 1876a (in part; key, sp. desc.). Henshaw 1881-1882 (in part; check.). LeConte and Horn 1883 (in part; key). Henshaw 1885 (in part; check.). Wickham 1889. Faust 1890. Casey 1891 (in part; key). Horn 1894 (in part; check.). Petri 1905 (key). Petri 1912. Blatchley 1914 (sp. desc.). Blatchley and Leng 1916 (in part; key, sp. redesc.). Leng 1920 (in part; catal.). Lukjanovitsh 1926. Blatchley 1930. Bradley 1930 (in part; key). Arnett 1960-1962 (in part; key). Kissinger 1964 (in part; key).

*Apleurus*; Chevrolat 1873 (in part, espèces lyxiformes; check., key, sp. desc.).

*Cleonidius* Casey 1891:186. As subgenus of *Cleonus*. Gender, masculine. Type species *Cleonis vittatus* Kirby (= *Cleonus (Cleonidius) lobigerinus* Casey) by subsequent designation (Faust 1904:190). Champion 1902-1906 (sp. redesc.). Csiki 1934 (catal.). Blackwelder 1939 (check.). Arnett 1960-1962 (key, catal.). Kissinger 1964 (key). Hatch 1971 (key, sp. redescs.)

*Cleonidius*; Faust 1904 (key, check., type species desig.).

*Lixestus* Reitter 1916:89. As subgenus of *Lixus*. New synonymy. Gender, masculine. Type species *Curculio vibex* Pallas by monotypy. Csiki 1934 (catal.).

*Lixus (Lixesthus)*; Ter-Minasyan 1978 (error, misspelling).



Notes about synonymy.--- The subgenus *Cleonidius* was first proposed by Casey (1891) to accomodate those *Cleonus* with a cylindrical rostrum, more or less vittate elytral scale pattern, and an elongate-narrow body form approaching that of *Lixus*. Indeed species of *Cleonidius* very closely resemble *Lixus* and species of the two genera are often difficult to separate. This is evidenced by the placement of some species, now regarded as *Cleonidius*, originally in *Lixus*, and by the herein proposed new synonymy of *Lixestus* Reitter, a subgenus of *Lixus*, with *Cleonidius*.

Among *Lixus* species examined, only *Lixus* (*Lixestus*) *vibex* Pallas (type species of *Lixestus*), was found to possess a small dorsal median tubercle at the basal margin of variously abdominal sterna V to VII, that is characteristic of *Cleonidius*. However, other *Lixus* species, as yet not examined for this character, may also prove to require inclusion in *Cleonidius* should they possess the apotypic state as noted above. This is not surprising given the similarity of the two genera, their apparent close phylogenetic relationship, and previous lack of consideration of *Cleonidius* species in past studies and resulting classifications of Palearctic *Lixus* species. Thus there is the need for a reevaluation of the validity of at least the subgeneric groupings of *Lixus* in this light. Whether *Cleonidius* will remain as a valid genus when this is done remains to be seen; undoubtedly *Lixus* species will require reclassification.

Prior to the present study, *Cleonidius* had only been given generic status by Faust (1904).

**Diagnosis.**--- Adult Cleoninae with elongate-narrow to moderately robust body form (Figs. 118-136). Eye more or less oval to elongate-oval, flat (Figs. 137-156). Rostrum elongate-narrow to moderately robust, with at most only low median carina (indicated in the majority of species by low glabrous shiny line) (Figs. 137-156). Antenna with funiculus with article 1 longer than article 2; article 2 more or less as wide as long (Figs. 137-156). Pronotal postocular lobes lacking to present and well-developed; postocular vibrissae uniformly short to long but of unequal length, longest immediately behind base of eye (Figs. 137-156). Pronotal disk slightly to distinctly vittate, with white scales largest and/or densest immediately laterad of midline, smaller and/or sparser laterally; largely whitish in color, underlying dark cuticle largely obscured by white scales (Figs. 118-136). Prosternal swellings absent or present, if present, situated immediately anterior to each procoxal cavity (Fig. 8). Ventral tarsal pilosity extensive (covering greater part of ventral surface of each tarsal article) to reduced in extent or lacking from at least the basal tarsal articles of some species. Tibia with corbel ridge rounded (Fig. 5). Abdominal sternum VII (at least) of females dorsally (internally) with variously developed basal median glabrous shiny tubercle, evident externally as shallow

impression. Abdominal sternum VIII of female with basal arm short to long (Figs. 157-176).

**Description.**--- **Size.** Small to moderately large; elongate-narrow to moderately robust in general body form (Figs. 118-136). **Mouthparts.** Prementum flat to slightly swollen ventrally; with up to three large setae on each side. Maxillary palpus with palpifer and stipes each with at least one large seta. Labial palpi separated by a distance subequal to width of basal article of labial palpus. **Rostrum.** Elongate-narrow to robust, not to markedly medially tumescent, with at most only low median carina indicated in majority of species by fine glabrous shiny line (Figs. 137-156). Epistoma not to moderately swollen, with apical margin emarginate medially (Figs. 137b-156b). Antenna with funiculus with article 1 longer than article 2; article 2 more or less as long as wide; apical three articles of club lacking placoidal sensillae. **Head.** Eye oval to elongate-oval; flat (Figs. 137-156). Upper margin of eye rounded, frons flat to variously convex. **Vestiture.** Dorsum with suberect to erect vestiture absent to present and dense, short to very long in length; with simple appressed white scales of various size and density. **Pronotum.** Dorsal surface punctate, median basal area shallowly to deeply impressed; disk with or without various other impressions. Pronotal disk slightly to distinctly vittate, with white scales largest and/or densest immediately laterad of

midline, smaller and/or sparser laterally; largely whitish in color, underlying dark cuticle medially largely obscured by white scales (Figs. 118-136). Lateral margins with white scales large and/or dense. Pronotum widest at base to subequal in width from base to near apex, lateral margins arcuate, more or less parallel or slightly convergent from base to near apex, then variously constricted and more convergent to apex (Figs. 118-136). Pronotal postocular lobes lacking to present and well-developed; postocular vibrissae uniformly short (length less than one-half width of an eye) to long but of unequal length, longest behind base of eye (greatest length greater than one-half width of an eye) (Figs. 137a-156a). Prosternum. With shallow, rounded impression anterolaterad of each procoxal cavity; with or without variously developed rounded swelling immediately anterior to each procoxal cavity (Fig. 8). Legs. Tarsus moderately broad to broad; articles 2 and 3 more or less subequal in length, width and length of each article subequal to slightly wider than long; article 1 only slightly longer than articles 2 or 3; article 3 moderately deeply bilobed. Ventral tarsal pilosity various in extent from dense and covering almost entire ventral surface of each tarsal article, to lacking entirely from at least more basal articles in few species. Claws connate only at very base to from very base through basal one-third to midlength, not to markedly divergent. Tibia with corbel ridge rounded (Fig. 5). Foretibia with inner margin with small to large

denticles in apical one-half to two-thirds; near apex with small, indistinct to moderately large subapical tooth. Metatibia of male with uncus with ventral margin evenly rounded. Wings. Absent or present (various in length). Elytra. All intervals equally flat except for humerus and very base of interval 3 which are variously swollen and convex. Scale pattern various, more or less vittate (Figs. 131, 133-134, 136) or as marginal band of large white scales (Figs. 118-123). Humeri acute to rounded. Abdomen. Ventral surface with small shiny glabrous patches, each with single small appressed to suberect scale-like seta. Female with base of abdominal sternum VII (in some individuals also sterna V and VI) internally (dorsally) with variously developed rounded glabrous shiny median tubercle, evident externally (ventrally) as rounded shallow impression. Genitalia. Female. Abdominal sternum VIII with basal arm short to long; lateral arms various in shape (Figs. 157-176). Gonocoxite II elongate triangular in shape; apex not prolonged into marked lobe; stylus moderately large, apical in position (Fig. 117). Spermathecal gland round (Fig. 116). Male. Paired sternite of abdominal sternum VIII lacking distinct basal projections (as in Fig. 97f). Aedeagus moderately robust; in lateral view more or less evenly arcuate throughout length; apex not spatulate (Figs. 177c-196c). Internal sac various, with median dorsal pocket low to high; apical and dorsal median pocket individually distinct or not; various lobes present (Figs.

177a,b-196a,b). Apical sclerite, complex present; individual sclerites distinctly scythe-like, simple, lacking median projection.

Comparisons.--- *Cleonidius* species will prove most difficult to distinguish from various *Lixus* species. As noted elsewhere, some Old World species presently regarded as *Lixus*, but not examined by me, may prove to be *Cleonidius*. In North America, north of Mexico, the area where most species of *Cleonidius* are found, no *Lixus* species has a rounded pronotal postocular lobe; at most they possess a variously developed, but usually small, acute postocular projection or else have the anterolateral margin of the pronotum straight to slightly sinuate (Fig. 4). Postocular vibrissae in most New World *Lixus* are of unequal length, long, and have their greatest length behind the base of the eye (Fig. 4). All New World *Lixus* I have examined which possess a variously developed postocular lobe and that have postocular vibrissae of more or less uniform length have the inner margin of the femora variously dentate. Femora are not dentate in any *Cleonidius* species.

Unfortunately there are species of *Cleonidius* which lack or have only slightly developed postocular lobes and have postocular vibrissae as in most *Lixus* species (Figs. 145a, 152a-155a). These can be distinguished from *Lixus* by the relative lengths of the antennal articles, by having a more robust rostrum, and by having the ventral tarsal

pilosity variously reduced in extent.

Although there are exceptions (Fig. 141a), world *Lixus* species in general have a very elongate-narrow rostrum (Fig. 4) whereas *Cleonidius* species have a more robust rostrum (Figs. 137a-140a, 142a-156a). Eyes are rounded and slightly convex in many *Lixus*, otherwise are oval to elongate-oval as in all *Cleonidius*. Whereas characters used herein to separate *Lixus* from *Cleonidius* work for the New World fauna, the structural diversity of *Lixus* species in other geographic areas does not permit separation on the basis of these same characters. In the Old World, some *Lixus* species examined are separable from *Cleonidius* species only by the lack of the internal tubercle at the base of abdominal sternum VII in females, the presence of which is, by definition, universally diagnostic for *Cleonidius*. Such is the situation with *Cylindropterus luxeri* Chevrolat, *Lixus* (*Lixoglyptus*) *spartii* Olivier, and *Lixus* (*Lixoglyptus*) *circumcinctus* Boheman, the taxa chosen as the out-groups for the phylogenetic analysis of relationships of species of *Cleonidius*.

*Apleurus* (*Gibbostethus*) *hystrix* Fall, because of its more elongate-narrow body form (Fig. 24), may be confused with some *Cleonidius* species, but the tumescent mesosternal process (Fig. 58), robust and deeply punctate rostrum, and prosternal swellings immediately in front of the prosternal impressions (Fig. 7) in the former will readily separate the two.

**Checklist of included species.**--- Nineteen species are recognized in the New World from Nicaragua north to southern Canada. The genus also occurs in the Palearctic Region where it is represented by at least, species formerly placed in the subgenus *Lixestus* Reitter of *Lixus* Fabricius. Of species placed in *Lixestus*, I have examined, and therefore herein include only *Lixus (Lixestus) vibex* (Pallas), the type species of that subgenus by monotypy. Subsequently, aside from species regarded as synonyms of *L. vibex*, Csiki (1934) included *L. pallasi* Faust, originally described as a variation of *L. vibex* and therefore likely a *Cleonidius*, and Ter-Minasyan (1978) further included *L. meles* Boheman. Inclusion of *L. meles* in *Lixestus* by Ter-Minasyan (1978) is supported by Petri (1905) wherein *L. vibex* and *L. meles* key out adjacent to one another. A key to separate the three species included in *Lixestus* is given by Ter-Minasyan (1978).

The twenty species herein placed in *Cleonidius* are as follows:

*Cleonidius poricollis* species group

1. *C. poricollis* (Mannerheim)
2. *C. eustictorrhinus* Anderson
3. *C. modestus* (Mannerheim)
4. *C. subcylindricus* Casey
5. *C. longinatus* Anderson



6. *C. texanus* (LeConte)

*Cleonidius americanus* species group

7. *C. americanus* Csiki
8. *C. frontalis* (LeConte)
9. *C. canescens* (LeConte)
10. *C. infrequens* Anderson
11. *C. puberulus* (LeConte)
12. *C. collaris* (LeConte)
13. *C. notolomus* Anderson

*Cleonidius lobigerinus* species group

14. *C. lobigerinus* Casey
15. *C. calandroides* (Randall)

*Cleonidius boucardi* species group

16. *C. boucardi* (Chevrolat)
17. *C. trivittatus* (Say)
18. *C. placidus* Csiki
19. *C. quadrilineatus* (Chevrolat)

*Incertae sedis*

20. *C. vibex* (Pallas)

Phylogenetic relationships.--- See "Phylogenetic analysis" section.

#### 4.5.1 Key to species of adult *Cleonidius*

1. Tarsal claws widely divergent, lacking basal internal flange (as in Fig. 66).....  
.....*C. Vibex* (Pallas)
- 1'. Tarsal claws at most only slightly divergent, basal internal flange well-developed (as in Fig. 67)....2
- 2(1'). Head above eye with deep straight sulcus extended posteroventrally from above dorsal margin of eye to area under postocular lobe (Fig. 139). Elytra with apices produced and acuminate (Fig. 120) .....  
.....*C. modestus* (Mannerheim)
- 2'. Head above eye lacking deep straight sulcus. Elytra with apices rounded to at most only slightly acuminate and slightly produced .....3
- 3(2'). Pronotum extremely irregularly elevated; markedly constricted dorsolaterally near apical margin, deeply and broadly impressed medially at base and longitudinally along lateral margins; with larger punctures on disk very sparse, shallow and indistinct. Metathoracic wings short (approximately equal to one-half length of elytra).....  
.....*C. collaris* (LeConte)
- 3'. Pronotum more regular in elevation and sculpture;

- only shallowly to moderately deeply impressed medially at base (also shallowly longitudinally along lateral margins in very few specimens); with larger punctures on disk denser, deeper and more distinct. Metathoracic wings short (approximately equal to or less than one-half length of elytra) to long (approximately equal to or greater than length of elytra).....4
- 1(3'). Pronotum with postocular lobes moderately to well-developed and with postocular vibrissae short to moderately long (maximum length equal to or less than one-half width of eye in lateral view) (Figs. 137a-144a, 146a-151a). Prosternum in lateral view with anterior flat portion one-third as long as, to approximately equal in length to, more posterior angulate portion (Figs. 137a-144a, 146a-151a).....5
- 1'. Pronotum with postocular lobes lacking to at most moderately developed and with postocular vibrissae long (maximum length equal to or greater than one-half width of eye in lateral view) (Figs. 145a, 152a-155a). Prosternum in lateral view with anterior flat portion no more than one-half as long as more posterior angulate portion (Fig. 145a, 152a-155a).....19
- 5(4). Elytra with sutural intervals lacking scales throughout the greater part of their length, with at most scattered very fine erect hair-like scales

- (Fig. 118); interval 2 with scales contrastingly dense. ....*C. porcollis* (Mannerheim)
- 5'. Elytra with sutural interval with scales uniformly distributed throughout their length (although sparsely so or abraded in some specimens); interval 2 with scales similar in density to those of sutural interval.....6
- 6(5)'. Head behind eye with moderately deep curved sulcus extended posteroventrally from upper one-half of posterior margin of eye to area under postocular lobe (largely covered in many specimens by postocular lobe) (Fig. 150).....7
- 6'. Head behind eye lacking distinct curved sulcus, with at most one or more wrinkles of the cuticle... ..8
- 7(6)'. Distributed in the northern Atlantic coastal states of the United States and Atlantic coastal provinces of Canada (Fig. 229). Dorsal surface of pronotum and elytra with suberect vestiture short, indistinct. Elytral scale pattern of most specimens not distinctly vittate; scales on interval 2 and intervals 6 to 8 not markedly smaller than scales on adjacent intervals; underlying dark cuticle on interval 2 and intervals 6 to 8 largely obscured by overlying white scales (Fig. 132).....  
 ....*C. calandroides* (Randall) (in part)
- 7'. Distributed in Mexico, and/or Canada and the United

States west of 85° W longitude (Fig. 228). Dorsal surface of pronotum and elytra with erect vestiture short to moderately long, distinct. Elytral scale pattern distinctly vittate; scales on interval 2 and intervals 6 to 8 markedly smaller than scales on adjacent intervals; underlying dark cuticle on interval 2 and intervals 6 to 8 not largely obscured by overlying white scales (Fig. 131).....

.....*C. lobigerinus* Casey (in part)

8(6'). Elytra distinctly laterally margined with white scales (especially in basal one-half); intervals 9 to 11 with dense moderately elongate broad white scales; otherwise moderately densely uniformly covered with fine white scales which are smaller than those on intervals 9 to 11 (Figs. 119, 121-123).....9

8'. Elytra not laterally margined, either vittate (Figs. 130-131), more or less uniformly densely scaled (Fig. 129), or mottled with irregularly distributed patches of larger and denser scales (Figs. 124-125); intervals other than 9 to 11 with scales equally large to those on intervals 9 to 11 .....12

9(8). Head, pronotum and elytra lacking dorsal erect or suberect vestiture (Fig. 140a). Postocular lobes only moderately developed (Fig. 140a).....

.....*C. subcylindricus* Casey

9'. Head, pronotum and elytra with dorsal erect or suberect vestiture short to long (Figs. 138a, 141a-142a). Postocular lobes well-developed (Figs. 138a, 141a-142a).....10

10(9'). Rostrum with large deep dense punctures; medially longitudinally tumescent and with median carina as fine shiny line at crest of median tumescence; robust (width at apex greater than 0.58 times length) (Fig. 138). .....  
.....*C. eustictorrhinus* Anderson

10'. Rostrum with small shallow moderately dense punctures; not medially longitudinally tumescent, carina lacking to as variously developed fine shiny line; robust to elongate-narrow (width at apex less than 0.62 times length) (Figs. 141-142). .....11

11(10'). Rostrum (especially in females) very long and narrow (width at apex less than 0.50 times length) (Fig. 141a). Head, pronotum and elytral declivity (and to a lesser extent elytral disk) with dorsal erect vestiture moderately long to very long (lacking from a very few specimens) (Fig. 141a). Associated with Leguminosae.....  
.....*C. longinasus* Anderson

11'. Rostrum (of both sexes) shorter and more robust (width at apex greater than 0.50 times length) (Fig. 142). Head, pronotum and elytra with dorsal suberect to erect vestiture short to moderately

- long (Fig. 142a). Associated primarily with Rosaceae and Rhamnaceae, rarely Leguminosae at lower altitudes ..... *C. texanus* (LeConte)
- 12(8'). Dorsum (especially of head, pronotum and elytral declivity) with erect vestiture long to very long (Fig. 143a). Rostrum in dorsal view slightly expanded laterally at midlength (Fig. 143b); in lateral view with prementum swollen ventrally (Fig. 143a)..... *C. americanus* Csiki
- 12'. Dorsum with suberect to erect vestiture either lacking or short variously on head, pronotum, and elytra (Figs. 144a, 146a-147a, 149a-152a). Rostrum in lateral view with prementum flat to slightly swollen ventrally (Fig. 144a); in dorsal view not to slightly expanded laterally at midlength (Fig. 144a). ..... 13
- 13(12'). Distributed in the northern Atlantic coastal states of the United States and Atlantic coastal provinces of Canada (Fig. 229)..... *C. calandroides* (Randall)
- 13'. Distributed in Mexico, and/or Canada and the United States west of 85° W. longitude..... 14
- 14(12'). All tarsi with similar and extensive ventral pilose vestiture, articles 1 to 3 with large elongate-oval ventral pilose pads covering virtually the entire ventral surface of each article (except basal one-third to one-half of article 1) ..... 15
- 14'. Tarsi with ventral pilose pads small, decreased in

- extent on more posterior tarsi, either with articles 1 and 2 with small elongate-narrow ventral pilose pads, apical tufts of pilosity, or, with pads lacking; article 3 with moderately large pads not covering more than apical two-third of ventral surface ..... 16
- 15(14). Rostrum not dorsally medially tumescent; base in lateral view appearing continuous with frons (Fig. 152a). Elytra with interval 3 (in some specimens also intervals 4 and 5), and intervals 9 to 11 with dense white scales, otherwise with scales very sparse or lacking (Fig. 133).....  
.....*C. boucardi* (Chevrolat) (in part)<sup>9</sup>
- 15'. Rostrum dorsally medially longitudinally tumescent; base in lateral view separated from frons by distinct transverse impression (Fig. 150a). Elytra with scales white, fine and sparse, densest on sutural interval, intervals 3 to 7, and 9 to 11....  
.....*C. Jobigerinus* Casey (in part)
- 16(14'). Rostrum very short and robust (width at apex greater than 0.70 times length) (Figs. 146-147). ..  
..... 17
- 16'. Rostrum more elongate-narrow (width at apex less than 0.70 times length) (Figs. 144, 149)..... 18
- 17(16). Rostrum with median carina very distinct, sharp and elevated, also with faint lateral carinae at lateral margins (Fig. 146b); rostrum flat from



- immediately laterad of median carina to lateral margins; rostrum more or less quadrate in cross-section, the lateral margins sharp. Articles of tarsus of hind leg with ventral pilose pads moderately large. Metathoracic wings long (greater than length of elytra) to short (approximately one-half or less than length of elytra).....
- .....*C. infrequens* Anderson
- 17'. Rostrum with median carina indistinct, low and only slightly elevated, without any trace of lateral carinae (Fig. 146b); rostrum declivous from immediately laterad of median carina to lateral margins; rostrum less quadrate in cross-section, the lateral margins rounded. Articles of tarsus of hind leg with ventral pilose pads small.
- Metathoracic wings short (less than one-half length of elytra).....*C. puberulus* (LeConte)
- 18(16'). Elytra elongate compared to length pronotum (length elytra less than 0.40 times length pronotum) (Fig. 125). Metathoracic wings long (greater than elytra in length). Rostrum more or less circular in cross-section, lateral margins rounded; in dorsal view, slightly swollen laterally at midlength; in lateral view with prementum slightly swollen ventrally. Fore-tibia of most females with inner margin with very large prominent denticles (Fig. 144c).....*C. frontalis* (LeConte)

- 18'. Elytra more robust compared to length pronotum (length elytra greater than 0.40 times length pronotum) (Fig. 130). Metathoracic wings present, variable in length from short (approximately equal to one-half length of elytra) to long (greater than elytra in length). Rostrum more or less quadrate in cross-section, lateral margins sharp; in dorsal view, not swollen laterally at midlength; in lateral view with prementum flat. Fore-tibia of female with inner margin with slightly developed, small denticles.....*C. notolomus* Anderson
- 19(4'). Metathoracic wings short, approximately equal to one-half length of elytra. Eye oval (width greater than 0.60 times length) (Fig. 145a).....  
.....*C. canescens* (LeConte)
- 19'. Metathoracic wings long (approximately equal to or greater than length of elytra). Eye elongate-oval (width less than 0.65 times length) (Figs. 152a-155a).....20
- 20(19'). Elytra, except sutural interval and portions of interval 2, uniformly covered with dense white scales (Fig. 135).....*C. placidus* Csiki
- 20'. Elytra distinctly vittate, intervals 6 to 8 with scales lacking or very sparse compared to intervals 3 to 5 and 9 to 11 (Figs. 133-134, 136) .....21
- 21(20'). Rostrum not or only slightly dorsally medially tumescent, not distinctly carinate (very slightly

- so in few specimens) (Fig. 152a). Metatarsus with articles 2 and 3 with ventral pilose pads large, covering virtually the entire ventral surface; article 1 with pads elongate-broad, covering the apical two-thirds of the ventral surface.....
- .....*C. boucardi* (Chevrolat) (in part)
- 21'. Rostrum dorsally medially tumescent, with low rounded carina (Figs. 153a, 155a). Metatarsus with article 3 with ventral pilose pads moderately large, covering the apical one-half to two-thirds of ventral surface; article 2 with pads elongate-narrow, covering the apical two-thirds of the ventral surface; article 1 with pads either present as apical tufts, or elongate and very narrow, covering not more than the apical one-half of the ventral surface.....22
- 22(21'). Elytra elongate-narrow (width less than 0.55 times length) (Fig. 136). Pronotal disk with moderately long erect hair-like scales (Fig. 155a). Associated primarily with Rosaceae and Rhamnaceae, rarely Leguminosae.....
- .....*C. quadrilineatus* (Chevrolat)
- 22'. Elytra more robust (width greater than 0.55 times length) (Fig. 134). Pronotal disk with at most short, suberect, indistinct hair-like scales (Fig. 153a). Associated with Leguminosae.....
- .....*C. trivittatus* (Say)

#### 4.5.2 Species accounts

##### 4.5.2.1 *Cleonidius poricollis* species group

**Diagnosis.**--- Size small to moderate for *Cleonidius* (Fig. 197). Dorsal erect or suberect vestiture absent (Fig. 140a) to present, dense and very long (Fig. 141a). Rostrum very elongate-narrow (Fig. 141a) to moderately robust (Fig. 138a), not to markedly medially tumescent, not to variously carinate; lateral margins rounded. Pronotum with postocular lobes moderately (Fig. 140a) to well-developed (Fig. 141a); postocular vibrissae uniformly short to more or less uniformly moderately long. Elytra with marginal band of scales, only intervals 9 to 11 with white scales large and broad, otherwise with intervals with scales small and fine to absent (Figs. 118-123). Tarsus with ventral pilose vestiture extensive (covering almost entire ventral surface of each article) to slightly reduced in extent (covering not less than apical one-half of articles 2 and 3). Wings present, of variable length. Female with abdominal sternum VIII with basal arm short; lateral arms more or less straight and divergent throughout length (Figs. 157-162). Male with aedeagus with internal sac with apex of dorsal median pocket rounded (Figs. 177b-182b); lobes B, D, and E absent in some species (Figs. 180a,b-182a,b); dorsal median

pocket low to high in lateral view.

**Phylogenetic relationships.**--- Monophyly of this species group is based only on distribution of the apotypic states of the secondary characters of the length of the basal arm and shape of the lateral arms of abdominal sternum VIII in females. The *C. poricollis* group is hypothesized to be the sister group to the remainder of *Cleonidius* (Figs. 236-237).

4.5.2.1.1 *Cleonidius poricollis* (Mannerheim)

(Figs. 118, 137, 157, 177, 219)

*Lixus poricollis* Mannerheim 1843:291. Holotype (not examined; depository unknown but I suspect types are in the Zoological Museum, Moscow State University). Type locality, California. Gemminger and von Harold 1871 (catal.). LeConte 1876a,b. Henshaw 1885 (check.).

*Cleonus (Cleonidius) poricollis*; Casey 1891 (key, misinterp. as synonym of *Cleonidius vittatus*). Csiki 1934 (catal.).

*Cleonus (Cleonidius) virgatus*; Casey 1891 (key, misident.).

*Cleonus (Cleonidius) vittatus*; Casey 1891 (key, misident.).

Hatch 1971 (key, redesc., misident.).

*Cleonus erysimi* Fall 1901:261. New synonymy. Lectotype (here designated), male, one of an undetermined number of syntypes, labelled "'male symbol'", "Redondo/ Cal. 4.7.94.", "Type/ *erysimi*", "M.C.Z./ Type/ 25193", "H.C. FALL/ COLLECTION" and with my designation label

"*Cleonus erysimi*/ Fall LECTOTYPE/ desig. Anderson"  
 (MCZC). Type locality, Redondo, California. Leng 1920  
 (catal.).

*Cleonus poricollis*; Leng 1920 (catal., misinterp. as synonym  
 of *Cleonidius vittatus*).

*Cleonus (Cleonidius) erysimi*; Csiki 1934 (catal.).

*Cleonis erysimi*; O'Brien and Wibmer 1982 (catal., distn.).

*Cleonis poricollis*; O'Brien and Wibmer 1982 (catal.,  
 distn.).

*Cleonis virgatus*; O'Brien and Wibmer 1982 (catal., distn.,  
 misident.).

*Cleonis vittatus*; O'Brien and Wibmer 1982 (catal., distn.,  
 misident.).

Notes about synonymy.--- Type material of *C. poricollis* has  
 not been examined. I suspect that it is deposited in the  
 Zoological Museum, Moscow State University. B.A. Korotyaev  
 (Zoological Institute, Leningrad) has informed me that there  
 are types of two species of *Lixus* from California in that  
 collection. I have been unable to borrow this material but  
 from the description of Mannerheim (1843) I regard it to be  
 this species.

Like *C. poricollis*, *Cleonus erysimi* Fall also has the  
 sutural interval lacking distinct scales.

Problems in recognition.--- Individuals of this species are

easily recognized by the sutural interval of the elytra lacking scales, with at most only scattered very fine erect hair-like scales; interval 2 has scales that are contrastingly dense (Fig. 118). Only some individuals of *C. modestus* are superficially similarly patterned however close examination of the sutural interval reveals there are small scales present. Regardless, *C. modestus* individuals are easily recognized by the deep straight sulcus above each eye (Fig. 139).

Individuals of *C. eustictorrhinus* and *C. texanus*, especially the former, may also prove difficult to separate from those of *C. poricollis* should the elytra be largely abraded of vestiture. *Cleonidius texanus* specimens have a slightly more elongate-narrow rostrum (Fig. 142) and the length of the elytra as compared to the length of the pronotum is greater (see Figs. 197, 200). *Cleonidius eustictorrhinus* and *C. poricollis* individuals will prove most difficult to separate, if abraded, although most *C. poricollis* are brachypterous, whereas all *C. eustictorrhinus* are macropterous.

**Description.**--- Specimens examined. 129 males, 179 females. Data about variation in LR, WF, WRA, LP, WPB, WE1H, WE1M, LE1, LEy, WEy, WEy/LEy, WPB/LP, WRA/LR, WRA/WF, WE1M/LE1, and LP/LE1 are presented in Tables 29 and 30. Size. Length, male, 7.2-11.3 mm; female, 6.1-12.0 mm. Width, male, 2.2-4.1 mm; female, 2.1-4.5 mm. Head. Eye elongate-oval. Area behind

eye with numerous wrinkles of cuticle. Frons with sparse to moderately dense, small, shallow to moderately deep punctures, some longitudinally confluent and irregularly impressed immediately above eyes, punctures sparser medially; interspersed with minute punctures. Frons also with moderately dense to dense, elongate-narrow erect hair-like scales immediately above eyes; otherwise with scattered erect hair-like scales and moderately dense to dense, elongate-narrow appressed white scales, sparse to absent medially. Frons separated from base of rostrum by moderately deep to deep transverse impression (Fig. 137a). Base of median tumescence or carina of rostrum with small moderately deep fovea. Width of frons greater than or subequal to width at apex of rostrum. Rostrum. Moderately robust, slightly more so in males (width at apex  $0.569-0.800$  times length) (Fig. 137). In lateral view straight to very slightly curved downward. In dorsal view with postgenae not to very slightly expanded laterally, antennal scrobes very slightly visible. Slightly to markedly medially tumescent from point of antennal insertion to middle of frons (more markedly so basally), with median carina variously developed as a rounded to sharp, moderately elevated, fine to broad, glabrous shiny line at crest of median tumescence. Dorsal and lateral punctation moderately dense to dense, small to moderately large, shallow to deep (longitudinally confluent in many specimens), smaller and not longitudinally confluent apically, dorsally interspersed with minute punctures.



Dorsally (especially laterally) with sparse to moderately dense, moderately long erect hair-like scales, and laterally with sparse to moderately dense, elongate-narrow white appressed to recumbent scales, medially with vestiture absent to very sparse, appressed, elongate-fine. Epistoma not swollen. Pronotum. Median carina variously developed in anterior one-half to three-quarters as slightly to moderately elevated glabrous shiny broad line. Dorsal punctation moderately large to large, moderately dense to dense, deep; smaller and shallower apically; shallower on flanks; areas between large dorsal punctures with minute regularly impressed punctures. Dorsally with moderately dense, moderately long erect hair-like scales each situated in large puncture (Fig. 137a). Scales dorsally absent along midline, moderately dense, elongate-fine, appressed in pair of paramedian apically slightly narrowed stripes; laterally, absent to very small and sparse. Lateral margins with appressed white scales elongate-fine to elongate-narrow, moderately dense in moderately broad stripe. Flanks with scales smaller and slightly less dense than along lateral margins. Median basal area of disk very shallowly to moderately deeply impressed. Anterolateral margin with pronotal postocular lobes moderately to well-developed; postocular vibrissae short, of uniform length to very slightly longer immediately behind base of eye (approximately one-half width of eye or less) (Fig. 137a). In dorsal view with lateral margins very slightly arcuate

and convergent from base to apex; subapical constriction very slight and indistinct. **Prosternum.** With moderately large, moderately deep to deep impression, and moderately to well-developed rounded swelling anterior to each procoxal cavity. In lateral view with anterior flat to slightly angulate portion long, approximately one-half to two-thirds length of posterior more angulate portion, anterior and posterior portions not to slightly differentiated by shallow transverse impression (Fig. 137a). **Elytra.** Elongate-narrow in general form (width at midlength 0.454-0.546 times length in males; 0.478-0.529 in females) (Fig. 118). In dorsal view with lateral margins straight to very slightly arcuate or very slightly sinuate, slightly convergent from apical one-third to humerus; slightly to moderately and evenly arcuate from apical one-third to apex. Elytral apices slightly produced, rounded to sub-acuminate, not to slightly divergent. Humerus indistinct, rounded. Dorsally with sparse to moderately dense, suberect to erect, short hair-like scales (abraded in many specimens). Sutural interval with scales lacking throughout greater part of length, with sparse fine scales only at very base; intervals 2 to 7 with scales uniformly moderately dense, various in size from elongate-fine to elongate-narrow on alternate intervals in many specimens; interval 8 with scales absent to very small, fine and sparse; intervals 9 to 11 with scales uniformly moderately dense to dense, moderately large and elongate-narrow, sparser and smaller apically and at base of

interval 11 (Fig. 118). Punctures of elytral striae small, individually distinct, larger and less individually distinct basally, arranged in regular rows. **Wings.** Long (greater than length of elytra [9%, N=29]) to short (less than one-half length of elytra [91%, N=279]). **Legs.** Foretibia of female with inner margin with small denticles in apical three-quarters; subapical tooth small, indistinct to distinct from tibial denticles. Foretibial and mesotibial unci of both sexes moderately large, metatibial uncus small. Tarsal claws connate in basal one-third to one-half, not to slightly divergent, each with basal internal flange well-developed. Ventral tarsal pilose vestiture of all tarsi of male and female present as elongate-narrow pads covering apical one-half to three-quarters of article 1, as rounded pads covering apical one-half of article 2, and as large rounded pads covering apical two-thirds to more or less the entire ventral surface of article 3. **Abdomen.** Ventral surface with sparse to dense appressed elongate-narrow to very elongate-narrow white scales (abraded in part in many specimens). Abdominal sterna III and IV laterally with individually indistinct glabrous patches, each with large puncture and single elongate recumbent to suberect hair-like scale situated in or near the center. Base of abdominal sternum VII of female internally (dorsally) with moderately developed median shiny tubercle. **Genitalia.** Female (four examined). Abdominal sternum VIII with basal arm short; lateral arms straight and divergent from base to apical.

one-third; markedly inwardly arcuate at apical one-third and convergent to apex, moderately expanded at apex (Fig. 157). Stylus moderately large in length compared to length gonocoxite II. Male (four examined). Internal sac (Figs. 177a, b) elongate and low; median dorsal pocket low, with single elongate moderately broad dorsally directed basal lobe (lobe A), with small median dorsally directed lobe at midlength on dorsal surface (lobe B), with small paired dorsally directed lobe at basal one-third on dorsal surface (lobe C), with moderately large paired dorsolaterally directed lobe at midlength near dorsal margin (lobe D), with moderately large paired dorsolaterally directed lobe at apical one-third at dorsal margin (lobe E), and with small ventrally directed paired lobe near base near ventral margin (lobe F). Apex of median dorsal pocket rounded in dorsal view. Apical pocket individually distinct from median dorsal pocket. Ventral median pocket moderately large. Apical pocket with paired small ventrally directed lobe near apical one-third at ventral margin. Eversible apical sclerite complex with paired simple scythe-like sclerite; adjacent ventral surface of apical pocket with pair of large, dark, longitudinal sclerites.

**Geographic variation.**--- None noted.

**Geographic distribution.**--- This species is found from southern British Columbia and Alberta, east to Manitoba,

south to southern California, Arizona and extreme western Texas (Fig. 219).

**Natural history.**--- This species is associated with grassland, desert-grassland transitional habitats, Pacific semi-desert; and Great Basin desert. Adults of this species have been collected on a variety of plants (but especially Cruciferae) as follows; *Cleome* sp. (Capparaceae); *Aplopappus ericoides* (Less.) H. and A., *Senecio* sp. (both Compositae); *Brassica* sp., *Brassica oleracea* L. (cabbage), *Dithyrea californica* Harv., *Erysimum* sp., *E. suffrutescens* (Abrams) G. Rosseb., *Raphanus sativus* L. (radish), *Stanleya pinnata* (Pursh.) Britton var. *inyoensis* (all Cruciferae); *Adenostoma fasciculatum* H. and A., *Prunus persica* (L.) Batsch. (peach) (Rosaceae). The type series of *Cleonus erysimi* Fall was collected from flowers of *Erysimum capitatum* (Dougl.) Greene on sandhills at Redondo Beach, California (Fall 1901). Adults have been reared from larvae found in crowns of *Alyssum* sp. (Cruciferae) at Harris Ranch, Dos Alamos, Santa Barbara Co., California.

Adults of the species have been frequently collected in sand dune habitats in coastal California and in other sandy habitats throughout the species range. Adults have been collected throughout the year at altitudes of 15-2359 m (X=767 m, N=40) (Fig. 233).

**Chorological relationships.**--- Table 67.

**Phylogenetic relationships.**--- Primary characters allow only for placement of this species as a member of an unresolved pentachotomy that also includes *C. modestus*, *C. eustictorrhinus*, the *C. subcylindricus*-*C. longinasus*-*C. texanus* lineage, and the remainder of *Cleonidius* (Fig. 236). Secondary characters allow for further resolution and result in the hypothesis that *C. poricollis* is the sister species of *C. eustictorrhinus*, the two species forming a lineage that is a member of an unresolved trichotomy including *C. modestus* and the *C. subcylindricus*-*C. longinasus*-*C. texanus* lineage (Fig. 237).

Table 29. Descriptive statistics for *C. porticollis*  
based on eight males from Redondo, California

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.68-2.16	1.925	0.279	0.131
WF	1.24-1.56	1.410	0.160	0.075
WRA	1.24-1.56	1.390	0.184	0.087
LP	3.04-3.68	3.400	0.454	0.214
WPB	2.96-4.16	3.510	0.603	0.284
WE1H	3.28-4.64	3.930	0.721	0.340
WE1M	3.60-4.96	4.300	0.780	0.367
LE1	7.04-9.60	8.480	1.405	0.662
LEy	0.92-1.20	1.060	0.143	0.068
WEy	0.56-0.68	0.625	0.078	0.037
B. Proportions.				
WEy/LEy	0.57-0.61	0.590	0.026	0.012
WPB/LP	0.97-1.13	1.031	0.077	0.036
WRA/LR	0.69-0.76	0.723	0.039	0.019
WRA/WF	0.94-1.05	0.986	0.054	0.026
WE1M/LE1	0.49-0.53	0.507	0.021	0.010
LP/LE1	0.38-0.43	0.402	0.022	0.010

Table 30. Descriptive statistics for *C. parvicollis*  
based on eight females from Redondo, California

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.84-2.32	2.050	0.252	0.119
WF	1.28-1.60	1.420	0.154	0.072
WRA	1.20-1.36	1.310	0.095	0.045
LP	3.12-3.92	3.510	0.365	0.172
WPB	3.04-3.76	3.530	0.398	0.187
WE1H	3.36-4.40	3.960	0.476	0.224
WE1M	3.92-4.80	4.420	0.409	0.193
LE1	7.52-9.76	8.720	1.073	0.506
LEy	0.88-1.12	1.035	0.113	0.053
WEy	0.52-0.72	0.640	0.101	0.048
B. Proportions.				
WEy/LEy	0.58-0.67	0.618	0.047	0.022
WPB/LP	0.96-1.07	1.006	0.057	0.027
WRA/LR	0.57-0.68	0.641	0.054	0.026
WRA/WF	0.85-1.03	0.925	0.074	0.035
WE1M/LE1	0.49-0.53	0.508	0.020	0.009
LP/LE1	0.38-0.42	0.403	0.023	0.011



4.5.2.1.2 *Cleonidius eustictorrhinus* Anderson, new species

(Figs. 119, 138, 158, 178, 216)

Type material.--- Holotype, male, with a red label "HOLOTYPE", "SACRAMENTO/ CALIF/ XII-13-1949", "D.GIULIANI/ COLLECTOR", "Derham Giuliani/ Collection/ Calif. Acad. Sci./ Accession 1967", "Collection of the/ CALIFORNIA ACADEMY/ OF SCIENCES, San/ Francisco Calif.", "HOLOTYPE/ *Cleonidius/ eustictorrhinus/* Anderson" and with abdomen on card and genitalia in microvial attached to pin (CASC). Allotype, female, with a red label "ALLOTYPE", "Orange Vale/ Sacto.Co./ July 31, 1938/ Quentin Tornich", "ALLOTYPE/ *Cleonidius/ eustictorrhinus/* Anderson" and with abdomen on card and genitalia in microvial attached to pin (UCBC). Type locality, Sacramento, California.

Paratypes. 11 males, 12 females. UNITED STATES OF AMERICA: California: Orange Vale, 31.VII.38, Q. Tornish, 1F (UCBC); Lancaster, 19.V.37, E.P. Van Duzee, 1M (CASC); 4mi. w. Meadow Valley, 13.V.49, Middlekauff, 1F (UCBC); Madera County, Bates, 25.III.17, 1F (AMNH); Antioch, 1.IX.37, M. Cazier, 1M (AMNH), 26.IV.68, Monroe, 1F (CWOB); Irvine, Davis, 1M (CNCI); San Diego County, 26.IV.91, Blaisdell, 1M (CASC); Stanford, III.04, 1M (LACM), 11.III.06, 1F (LACM); Fresno County, Waltham Canyon, 26.III.37, Blum, 1F (LACM); Baldwin Hills, 11.VII.53, Menke, 1F (LACM); Vine Hill, 23.V.65, Johnson, 1F (CWOB); Atascadero, 18.VI.46, Mansfield, 1M (CASC); Laguna Beach, 30.XII.49, 1M (CWOB); Idyllwild, 12.V.34, Stone, 1F (OSUC); Laguna, 20.VI.30, 1F

(OSUC), 30.VI.30, 1M (CNCI); Irvine, 1M (OSUC); Yosemite, 17.VI.31, Essig, 1F (UCBC); Stanislaus, XII.35, 1M (CFDA); Carmichael, 17.V.58, Wilkey, 1M (CDFA); Napa County, Pope Valley, 9.V.55, Raven, 1F (CASC).

**Derivation of specific epithet.**--- From the Greek "eu" meaning very or exceedingly; "stiktos" meaning punctured; and, "rhinos" meaning nose. This name is used in reference to the markedly punctured rostrum of members of this species.

**Problems in recognition.**--- Individuals of this species are likely to be confused only with *C. texanus*, *C. subcylindricus*, and *C. poricollis*. Individuals of *C. eustictorrhinus* are easily separated from sympatric individuals of *C. poricollis* by the sutural interval lacking scales in the latter (Fig. 118) and by most *C. poricollis* being brachypterous whereas all *C. eustictorrhinus* are macropterous. Individuals of *C. eustictorrhinus* are easily separated from *C. subcylindricus* by geographic distribution, the former being restricted to California (Fig. 216), the latter to the Atlantic Coastal Plain from Florida north to New York (Fig. 215). *C. subcylindricus* individuals also lack suberect or erect dorsal vestiture (Fig. 140a) whereas *C. eustictorrhinus* have short but distinct erect dorsal vestiture (Fig. 138a). Specimens of *C. eustictorrhinus* will however prove most difficult to separate from sympatric

individuals of *C. texanus*. The rostrum is slightly more robust in *C. eustictorrhinus* (Fig. 138) than in *C. texanus* (Fig. 142), *C. eustictorrhinus* tend to be larger in size than *C. texanus*, and the length of the elytra as compared to the length of the pronotum tends to be greater in *C. texanus*, than in *C. eustictorrhinus* (see Figs. 197, 200). In *C. eustictorrhinus* individuals (Fig. 138), the rostrum is medially tumescent, distinctly carinate and has large deep punctures; in *C. texanus* specimens (Fig. 142), the rostrum is not medially tumescent or carinate and has only small shallow punctures.

**Description.**--- Data about variation in LR, WF, WRA, LP, WPB, WE1H, WE1M, LE1, LEy, WEy, WEy/LEy, WPB/LP, WRA/LR, WRA/WF, WE1M/LE1, and LP/LE1 are presented in Tables 31 and 32. **Size.** Length, male, 8.8-10.4 mm; female, 8.8-11.0 mm. Width, male, 3.2-3.9 mm; female, 3.2-4.0 mm. **Head.** Eye < elongate-oval. Area behind eye with numerous wrinkles of cuticle. Frons with moderately dense, small to moderately large, shallow punctures; longitudinally confluent and very irregularly impressed immediately above eyes; larger punctures interspersed with sparse minute punctures. Frons also with scales sparse to moderately dense elongate-narrow, erect immediately above eyes; medially and laterally with (sparse to moderately dense elongate-narrow appressed to suberect white scales; laterally with moderately dense, short to moderately long fine erect hair-like scales. Frons

either continuous with base of rostrum, or slightly separated by a shallow transverse impression (Fig. 138a). Base of median tumescence or carina of rostrum with small shallow to moderately deep fovea. Width of frons greater than or subequal to width at apex of rostrum. **Rostrum.** Moderately robust, slightly more so in males (width at apex 0.564-0.686 times length) (Fig. 138). In lateral view slightly curved downward. In dorsal view with postgenae not to slightly laterally expanded, antennal scrobes slightly visible. Moderately to markedly medially tumescent from point of antennal insertion to middle of frons, with median carina variously developed as a rounded, slightly to moderately elevated, broad, glabrous, shiny line at crest of median tumescence. Dorsal and lateral punctation moderately dense to dense, moderately large to large, shallow to deep (longitudinally confluent in many specimens), smaller and shallower apically, dorsally interspersed with few minute punctures. Dorsally (especially laterally) with sparse to moderately dense, moderately long, suberect to erect hair-like scales, and laterally and medially with sparse to moderately dense, elongate-fine to elongate-narrow, appressed to recumbent, white scales. Epistoma not to very slightly transversely swollen at base. **Pronotum.** Median carina variously developed in anterior one-half as slightly elevated glabrous narrow to broad line. Dorsal punctation small, dense, shallow; smaller and sparser apically; sparser on flanks; areas between dorsal punctures with dense minute

regularly impressed punctures. Dorsally with moderately dense, short erect hair-like scales each situated in larger puncture (Fig. 138a). Scales dorsally absent to sparse, small and fine along midline; moderately dense to dense, elongate-fine, appressed laterad of median line, but not as distinct stripes; laterally, absent to sparse, small and fine. Lateral margins with scales moderately dense to dense, elongate-narrow to moderately robust, appressed to recumbent in a moderately broad stripe. Flanks with scales moderately dense to dense, elongate-fine. Median basal area of disk very shallowly, broadly impressed. Anterolateral margin with pronotal postocular lobes well-developed; postocular vibrissae uniformly short (Fig. 138a). In dorsal view with lateral margins uniformly moderately arcuate and convergent from base to apex; subapical constriction at apical one-fifth indistinct or slight, then straight and convergent to apex (Fig. 119). Prosternum. With shallow to moderately deep impression and moderately to well-developed rounded swelling anterior to each procoxal cavity. In lateral view with anterior flat to slightly angulate portion moderately long, approximately one-half length of posterior more angulate portion; anterior and posterior portions not or slightly differentiated by shallow transverse impression. Elytra. Elongate-narrow in general form (width at midlength 0.491-0.577 times length in males; 0.491-0.553 in females) (Fig. 119). In dorsal view with lateral margins very slightly sinuate, slightly convergent from apical one-third

to humerus; slightly to moderately and evenly arcuate from apical one-third to apex. Elytral apices slightly produced, rounded to sub-acuminate. Humerus distinct. Dorsally with sparse to moderately dense, suberect to erect, very short to short, hair-like scales. Sutural interval to interval 7 with uniformly moderately dense, fine to elongate-fine, white scales and with scattered patches of larger white scales; interval 8 with scales absent to moderately dense, small and fine basally to elongate-fine apically; intervals 9 to 11 with scales moderately dense to dense, moderately large and elongate-narrow to robust, smaller and finer apically in most specimens (Fig. 119). Punctures of elytral striae small, individually distinct, arranged in regular rows. Wings. Long (greater than elytra in length). Legs. Foretibia of female with inner margin with small denticles in apical three-quarters; subapical tooth not distinct from tibial denticles. Foretibial and mesotibial unci of both sexes moderately large, metatibial uncus small. Tarsal claws connate in basal one-half, not to slightly divergent, each with basal internal flange well-developed. Ventral tarsal pilose vestiture of all tarsi of male and female present as elongate-narrow pads covering apical one-half to three-quarters of article 1, as rounded pads covering apical one-half of article 2, and as large rounded pads covering apical two-thirds to more or less the entire ventral surface of article 3. Abdomen. Ventral surface with moderately dense to dense elongate-narrow appressed white scales. Abdominal

sterna III to VI (especially III and IV) with individually indistinct glabrous patches, each with large puncture and single suberect to erect moderately long hair-like scale situated in or near the center. Base of abdominal sternum VII of female internally with slightly developed median shiny tubercle. Genitalia. Female (two examined). Abdominal sternum VIII with basal arm short; lateral arms straight and divergent from base to apical one-third; markedly abruptly inwardly arcuate at apical one-third; apical one-half to one-third markedly expanded (Fig. 158). Stylus moderately large in length compared to length gonocoxite II. Male (two examined). Internal sac (Figs. 178a, b) elongate and low; median dorsal pocket low, with single elongate moderately broad dorsally directed basal lobe (lobe A), with small median dorsally directed lobe at midlength on dorsal surface (lobe B), with moderately large paired dorsally directed lobe near midlength on dorsal surface (lobe C), with moderately large paired dorsolaterally directed lobe near midlength near dorsal margin (lobe D), with large paired anterodorsolaterally directed lobe at apical one-third at dorsal margin (lobe E), and with small ventrally directed paired lobe near base near ventral margin (lobe F). Apex of median dorsal pocket rounded in dorsal view. Apical pocket, individually distinct from median dorsal pocket. Ventral median pocket moderately large. Apical pocket with paired moderately large ventrally directed lobe near apical one-third at ventral margin. Eversible apical sclerite

complex with paired simple scythe-like sclerite; adjacent ventral surface of apical pocket with pair of large, dark, longitudinal sclerites.

Geographic variation.--- None noted.

Geographic distribution.--- This species is known only from California (Fig. 216).

Natural history.--- This species is found only in the Pacific semi-desert region. Adults of *C. eustictorrhinus* have been collected only on *Arctostaphylos* sp. (Ericaceae); *Eriogonum* sp. (Polygonaceae); and beans (Leguminosae). Definite host records are not known. Adults have been collected from March to July, and December at altitudes of 9-2002 m ( $X=596$  m,  $N=8$ ) (Fig. 233).

Chorological relationships.--- Table 67.

Phylogenetic relationships.--- Primary characters allow only for placement of this species as a member of an unresolved pentachotomy that also includes *C. modestus*, *C. poricollis*, the *C. subcylindricus*-*C. longinasus*-*C. texanus* lineage, and the remainder of *Cleonidius* (Fig. 236). Secondary characters allow for further resolution and result in the hypothesis that *C. eustictorrhinus* is the sister species of *C. poricollis*, the two species forming a lineage that a member



of an unresolved trichotomy also including *C. modestus* and the *C. subcylindricus*-*C. longinasus*-*C. texanus* lineage (Fig. 237).

Table 31. Descriptive statistics for *C. eustictorrhinus* based on 13 males from throughout the species range

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.80-2.20	2.000	0.171	0.063
WF	0.96-1.48	1.329	0.238	0.088
WRA	1.08-1.40	1.255	0.142	0.052
LP	2.64-3.60	3.360	0.421	0.156
WPB	2.80-3.92	3.502	0.439	0.162
WE1H	3.44-4.64	4.148	0.547	0.203
WE1M	3.68-4.96	4.437	0.558	0.206
LE1	6.88-9.44	8.369	1.057	0.391
LEy	0.96-1.16	1.068	0.096	0.036
WEy	0.52-0.72	0.637	0.075	0.028
B. Proportions.				
WEy/LEy	0.54-0.67	0.597	0.062	0.023
WPB/LP	1.02-1.09	1.043	0.029	0.011
WRA/LR	0.59-0.69	0.627	0.038	0.014
WRA/WF	0.88-1.13	0.952	0.116	0.043
WE1M/LE1	0.49-0.56	0.530	0.028	0.010
LP/LE1	0.38-0.43	0.402	0.026	0.010

Table 32. Descriptive statistics for *C. eustictorrhinus*  
based on 14 females from throughout the species range

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.80-2.20	2.011	0.200	0.071
WF	1.08-1.52	1.331	0.170	0.061
WRA	1.12-1.36	1.214	0.109	0.039
LP	2.88-3.76	3.406	0.329	0.117
WPB	3.12-4.00	3.543	0.422	0.151
WE1H	3.60-4.64	4.126	0.509	0.181
WE1M	3.92-5.04	4.423	0.497	0.177
LE1	0.75-9.44	8.434	0.962	0.343
LEy	1.00-1.20	1.089	0.113	0.040
WEy	0.56-0.68	0.620	0.056	0.020
B. Proportions.				
WEy/LEy	0.52-0.64	0.571	0.056	0.020
WPB/LP	0.98-1.14	1.040	0.059	0.021
WRA/LR	0.56-0.65	0.604	0.033	0.012
WRA/WF	0.87-1.11	0.916	0.092	0.033
WE1M/LE1	0.49-0.55	0.525	0.024	0.009
LP/LE1	0.38-0.43	0.404	0.026	0.009

4.5.2.1.3 *Cleonidius modestus* (Mannerheim)

(Figs. 120, 139, 159, 179, 217)

*Lixus modestus* Mannerheim 1843:291. Holotype (not examined; depository unknown but I suspect types are in the Zoological Museum, Moscow State University). Type locality, California. Gemminger and von Harold 1871 (catal.). LeConte 1876a,b. Henshaw 1885 (check.).

*Lixus californicus* Motschulsky 1845:378. Holotype (not examined, depository unknown). Type locality, California. Gemminger and von Harold 1871 (catal.). LeConte 1876a,b.

*Lixus pleuralis* LeConte 1858:78. Holotype (examined), female, labelled with a gold circle (=California), "1062", "Type/ 5177", "L./ pleuralis/ Lec." (MCZC). Type locality, on Colorado River below the Gila River, California. Gemminger and von Harold 1871 (catal.). LeConte 1876a (key, redesc.). Henshaw 1881-1882 (check.). Henshaw 1885 (check.). Wickham 1889. Horn 1894 (check.).

*Cleonus (Cleonidius) californicus*; Casey 1891 (as synonym).

*Cleonus (Cleonidius) modestus*; Casey 1891 (key). Wickham 1896 (check.). Fall 1901 (check.). Leng 1920 (catal.). Csiki 1934 (catal.). Blackwelder 1947 (check.). Hatch 1971 (key, misident. of *Cleonidius longinasus*).

*Cleonus (Cleonidius) pleuralis*; Casey 1891 (as synonym). Fall 1901 (check.).

*Cleonus modestus*; Wickham 1896 (check.). Fall 1901 (check.).

Leng 1920 (catal.). Blackwelder 1947 (check.).

*Cleonis californicus*; O'Brien and Wibmer 1982 (catal.,  
distn., as synonym).

*Cleonis modestus*; O'Brien and Wibmer 1982 (catal., distn.).

*Cleonis pleuralis*; O'Brien and Wibmer 1982 (catal., distn.,  
as synonym).

**Notes about synonymy.**--- Type material of *Lixus modestus* and *Lixus californicus* has not been examined. I suspect that type material of the former is deposited in the Zoological Museum, Moscow State University. B.A. Korotyaev (Zoological Institute, Leningrad) has informed me that there are types of two species of *Lixus* from California in that collection. I have been unable to borrow this material but from the description of Mannerheim (1843), and following LeConte (1876a), I regard this species, and *L. californicus* Motschulsky, as conspecific with *Lixus pleuralis*.

**Problems in recognition.**--- Adults of this species are very easily recognized by the deep straight sulcus above each eye (Fig. 139) and the produced and acuminate elytral apices (Fig. 120). Most individuals of *C. lobigerinus* and some *C. calandroides* possess a shallow curved sulcus behind the eye (covered by the pronotal postocular lobe in some specimens and difficult to see) (Fig. 150a), but this should not be confused with the deep straight sulcus in members of *C.*

*modestus*.

**Description.**--- Specimens examined. 89 males, 90 females. Data about variation in LR, WF, WRA, LP, WPB, WE1H, WE1M, LE1, LEy, WEy, WEy/LEy, WPB/LP, WRA/LR, WRA/WF, WE1M/LE1, and LP/LE1 are presented in Tables 33 and 34. **Size.** Length, male, 7.2-10.0 mm; female, 7.3-10.8 mm. Width, male, 2.1-3.1 mm; female, 2.2-3.5 mm. **Head.** Eye elongate-oval. Vertex, above eye with deep straight sulcus extended posteriorly from dorsal margin of eye to area under pronotal postocular lobe. Frons with scattered small shallow punctures and moderately dense minute punctures; also with moderately dense erect hair-like scales above eyes and moderately dense elongate fine appressed white scales, slightly sparser medially in most specimens. Frons continuous with base of rostrum, not separated by transverse impression (Fig. 139a). Most specimens with a shallow median fovea at base of median tumescence or carina of rostrum. Width of frons greater than or subequal to width at apex of rostrum. **Rostrum.** Elongate-narrow, slightly more so in females (width at apex 0.440-0.574 times length) (Fig. 139). In lateral view straight to very slightly curved downward. In dorsal view slightly narrowed apically, postgenae not laterally expanded, antennal scrobes not distinctly visible. Slightly medially tumescent with median carina in most specimens very low, glabrous shiny line at crest of median tumescence. Dorsal and lateral punctation sparse, small and shallow,

interspersed with moderately dense minute punctures. Dorsally (especially laterally) with moderately dense erect fine hair-like scales and with moderately dense fine elongate white appressed scales smaller, finer and sparser apical of point of antennal insertion. Epistoma not swollen. Pronotum. Median carina lacking in most specimens, indicated in very few specimens by low, glabrous shiny line in anterior one-half. Dorsal and lateral punctation moderately large, moderately dense, deep; smaller, sparser and shallower apically and medially, the areas between large dorsal punctures with minute, very shallow punctures. Dorsally with moderately dense, short to moderately long, fine erect hair-like scales each situated in large puncture (Fig. 139a). Scales dorsally very sparse to absent along median line, moderately dense, moderately large and elongate in pair of broad paramedian apically slightly narrowed stripes; laterally, small and fine. Lateral margins with appressed white scales large and elongate-narrow in moderately broad stripe. Flanks with scales moderately dense, small and fine. Median basal area of disk deeply and broadly impressed. Anterolateral margin with pronotal postocular lobes well-developed; postocular vibrissae very short, of uniform length (Fig. 139a). In dorsal view with lateral margins slightly arcuate and slightly convergent apically from base to more or less apical one-fifth; gradually to abruptly constricted at apical one-fifth, convergent to apex (Fig. 120). Prosternum. With moderately

deep impression, and moderately to well-developed rounded swelling anterior to each procoxal cavity. In lateral view with anterior flat to slightly angulate portion short, not distinctly differentiated from longer posterior more angulate portion. Elytra. Very elongate-narrow in general form (width at midlength 0.402-0.459 times length in males; 0.389-0.472 in females) (Fig. 120). In dorsal view with lateral margins straight, subparallel to slightly divergent from apical one-third to humerus; slightly and evenly arcuate from apical one-third to apex. Elytral apices produced, acuminate and divergent. Humerus distinct. Dorsally with moderately dense, erect, short to moderately long, fine hair-like scales. Scale pattern various; sutural interval to interval 8 with uniformly moderately dense, elongate fine appressed white scales to moderately dense, elongate fine on intervals 2 to 5 and moderately dense but smaller and finer on sutural interval and intervals 6 to 8 (interval 8 with scales very sparse to lacking in some specimens); intervals 9 to 11 with scales more or less uniformly moderately dense, moderately large and elongate-narrow (Fig. 120). Punctures of elytral striae small, individually distinct, arranged in regular rows. Wings. Long (greater than elytra in length). Legs. Foretibia of female with inner margin with small denticles in apical three-quarters; subapical tooth small, indistinct from tibial denticles. Foretibial uncus of both sexes moderately large, mesotibial and metatibial unci small. Tarsal claws



connate in basal one-third to one-half, not divergent, basal internal flange lacking. Ventral tarsal pilose vestiture of all tarsi of male and female present as elongate-narrow pads covering apical one-half of article 1, as rounded pads covering apical one-half of article 2, and as large rounded pads covering apical three-quarters of article 3. **Abdomen.** Ventral surface with moderately dense to dense appressed very elongate-narrow white scales, sparser along midline. Abdominal sterna III to VI (especially III and IV) with moderately large individually distinct to indistinct glabrous patches, each with large puncture and single elongate appressed to suberect hair-like scale situated in or near the center. Base of abdominal sternum VII of female internally (dorsally) with slightly developed, median, shiny tubercle. **Genitalia.** Female (four examined). Abdominal sternum VIII with basal arm very short, indistinct; lateral arms straight and divergent from base to apical one-quarter, markedly inwardly arcuate at apical one-quarter and convergent to apex, slightly expanded at apex (Fig. 159). Stylus moderately large in length compared to length gonocoxite II. Male (three examined). Internal sac (Figs. 179a,b) elongate and low; median dorsal pocket low, with single short broad elongate dorsally directed basal lobe (lobe A), with small median dorsally directed lobe at midlength on dorsal surface (lobe B), with moderately large paired dorsally directed lobe at basal one-third on dorsal surface (lobe C), with moderately large paired dorsally

directed lobe at midlength near dorsal margin (lobe D), with moderately large paired dorsolaterally directed lobe at apical one-third at dorsal margin (lobe E), and with small ventrally directed paired lobe near base near ventral margin (lobe F). Apex of median dorsal pocket truncate in dorsal view. Apical pocket individually distinct from median dorsal pocket. Ventral median pocket moderately large. Apical pocket with paired moderately large ventrally directed lobe near apical one-third at ventral margin. Eversible apical sclerite complex with paired simple scythe-like sclerite; adjacent ventral surface of apical pocket with pair of large, dark, longitudinal sclerites.

**Geographic variation.** None noted.

**Geographic distribution.**--- Members of this species are found in the southwestern United States of America and adjacent northern Mexico from California, southern Nevada, southern Utah, central New Mexico and extreme western Texas, south to the Mexican border, except in the west where they are found south into Baja California Norte, Mexico (Fig. 217).

**Natural history.**--- This species is associated with desert-grassland transitional habitat, and Chihuahuan, Mojave and Sonoran deserts. Adults of this species have been collected in various sandy habitats such as dunes and dry

stream beds or desert washes on *Ephedra californica* Wats. (Ephedraceae); *Chrysothamnus nauseosus* (Pall.) Britton, *C. paniculatus* (Gray) H.M. Hall, *Hymenoclea* sp., *H. salsola* Torr. and Gray, (all Compositae); *Asclepias* sp. (Asclepiadaceae); *Atriplex polycarpa* (Torr.) Wats. (Chenopodiaceae); *Medicago sativa* L. (Leguminosae); and *Brassica?* sp., *Lepidium* sp. (Cruciferae). Definite hosts are not known.

Adults have been collected from February to October at altitudes ranging from -96-1747 m (X=886 m, N=26) (Fig. 233).

**Chorological relationships.**--- Table 67.

**Phylogenetic relationships.**--- Primary characters allow only for placement of this species as a member of an unresolved pentachotomy that also includes *C. poricollis*, *C. eustictorrhinus*, the *C. subcylindricus*-*C. longinasus*-*C. texanus* lineage, and the remainder of *Cleonidius* (Fig. 236). Secondary characters allow for further resolution and result in the hypothesis that this species is a member of an unresolved trichotomy that also includes the *C. poricollis*-*C. eustictorrhinus* lineage and the *C. subcylindricus*-*C. longinasus*-*C. texanus* lineage (Fig. 237).

Table 33. Descriptive statistics for *C. modestus* based on eight males from Palm Springs, California

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.72-1.96	1.840	0.140	0.066
WF	0.96-1.12	1.040	0.096	0.045
WRA	0.80-1.00	0.935	0.090	0.043
LP	2.72-3.04	2.900	0.154	0.072
WPB	2.40-3.04	2.800	0.308	0.145
WE1H	3.04-3.68	3.460	0.378	0.178
WE1M	3.20-3.60	3.430	0.235	0.111
LE1	7.44-8.72	8.040	0.700	0.330
LEy	0.84-1.00	0.955	0.081	0.038
WEy	0.44-0.56	0.495	0.064	0.030
B. Proportions.				
WEy/LEy	0.46-0.58	0.519	0.068	0.032
WPB/LP	0.88-1.03	0.965	0.065	0.031
WRA/LR	0.47-0.56	0.509	0.049	0.023
WRA/WF	0.83-1.00	0.900	0.086	0.041
WE1M/LE1	0.40-0.46	0.427	0.031	0.014
LP/LE1	0.35-0.38	0.361	0.023	0.011

Table 34. Descriptive statistics for *C. modestus* based on eight females from Palm Springs, California

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.48-1.92	1.710	0.258	0.122
WF	0.88-1.08	0.990	0.110	0.052
WRA	0.72-1.00	0.865	0.140	0.066
LP	2.40-3.04	2.720	0.385	0.181
WPB	2.24-3.04	2.650	0.442	0.208
WE1H	2.80-3.68	3.230	0.540	0.255
WE1M	2.72-3.68	3.110	0.546	0.257
LE1	5.76-8.96	7.410	1.556	0.733
LEy	0.76-1.00	0.915	0.122	0.057
WEy	0.40-0.56	0.485	0.081	0.038
B. Proportions.				
WEy/LEy	0.46-0.57	0.530	0.053	0.025
WPB/LP	0.93-1.03	0.973	0.060	0.029
WRA/LR	0.46-0.53	0.506	0.040	0.019
WRA/WF	0.78-0.93	0.873	0.078	0.037
WE1M/LE1	0.39-0.47	0.422	0.038	0.018
LP/LE1	0.33-0.42	0.370	0.043	0.020

4.5.2.1.4 *Cleonidius subcylindricus* Casey

(Figs. 121, 140, 160, 180, 215)

*Cleonus (Cleonidius) subcylindricus* Casey 1891:193. Holotype (examined), female, labelled "Fla", "CASEY/ bequest/ 1925", "TYPE USNM/ 37283", "C.Cl./ subcylindricus/ Cas" (USNM). Type locality, Florida. Csiki 1934 (catal.).

*Cleonus (Cleonidius) graniferus* Casey 1891:194. New synonymy. Holotype (examined), female, labelled "Ga.", "CASEY/ bequest/ 1925", "TYPE USNM/ 37284", "C.Cl./ graniferus/ Cas." (USNM). Type locality, Georgia. Csiki 1934 (catal.).

*Lixus lupinus* Blatchley 1914:248. Lectotype (designated by Blatchley [1930:38, error as female] examined), male, labelled "TYPE", "Dunedin, Fla./ W.S.B. Coll./ 1.24.1913", "Purdue/ Blatchley/ collection" and with lectotype designation label of Blatchley (Purdue). Type locality, Dunedin, Florida. Blatchley and Leng 1916 (key, redesc.). Leng 1920 (catal.). Blatchley 1930.

*Cleonus subcylindricus*; Blatchley and Leng 1916 (key, redesc.). Leng 1920 (catal.).

*Cleonus graniferus*; Blatchley and Leng 1916 (key, redesc.). Leng 1920 (catal.).

*Cleonus lupinus*; Kingsolver 1972.

*Cleonis graniferus*; O'Brien and Wibmer 1982 (catal., distn.).

*Cleonis lupinus*; O'Brien and Wibmer 1982 (catal., distn., as synonym).

*Cleonis subcylindricus*; O'Brien and Wibmer 1982 (catal.,  
distn.).

*lupinus*, *Incertae sedis*; Csiki 1934 (catal.).

**Notes about synonymy.**--- *Cleonus graniferus* Casey is a developmentally abnormal female of *C. subcylindricus* with a short robust rostrum, legs and antennae. Similar developmental abnormalities are also known in other species of *Cleonidius*. *Lixus lupinus* Blatchley was placed in synonymy with *Cleonus subcylindricus* Casey by Kingsolver (1972).

**Problems in recognition.**--- This species is one of only two species of *Cleonidius* restricted in their distribution to eastern North America (Figs. 215, 229). These two species differ markedly in many structural features and there should be no problem in separating individuals of the two. *C. subcylindricus* individuals are easily distinguished from other members of the *C. texanus* group, with which they may be confused, by the only slightly developed pronotal postocular lobes and by lack of dorsal suberect or erect vestiture (Fig. 140a).

Problems in separation of *C. subcylindricus* from eastern species of *Lixus* are resolved by *Lixus* species, among other characters, lacking pronotal postocular lobes and instead possessing at most a small acute postocular

projection and postocular vibrissae that are not more or less uniform in length, but rather variable in length and distinctly longest at some point behind the eye (Fig. 4).

**Description.** --- Specimens examined. 11 males, 18 females. Data about variation in LR, WF, WRA, LP, WPB, WE1H, WE1M, LE1, LEy, WEy, WEy/LEy, WPB/LP, WRA/LR, WRA/WF, WE1M/LE1, and LP/LE1 are presented in Tables 35 and 36. **Size.** Length, male, 8.9-10.5 mm; female, 8.5-11.3 mm. Width, male, 2.9-3.5 mm; female, 2.8-3.9 mm. **Head.** Eye elongate-oval. Area behind eye with numerous wrinkles of cuticle. Frons with very dense, small, shallow punctures, some confluent in some specimens. Frons lacking suberect or erect vestiture, with moderately dense, small to moderately large, very fine and elongate appressed white scales (scales abraded medially in most specimens). Frons separated from base of rostrum by shallow transverse impression (Fig. 140a). Base of rostral carina with small shallow fovea. Width of frons less than width at apex of rostrum. **Rostrum.** Moderately robust, very slightly more so in males (width at apex 0.561-0.690 times length) (Fig. 140). In lateral view slightly curved downward. In dorsal view with postgenae slightly expanded laterally, antennal scrobes partially visible. Slightly medially tumescent from point of antennal insertion to base of frons, with distinct, low, glabrous median carina from point of antennal insertion to middle of frons. Dorsal and lateral punctation very dense, small, shallow (many



punctures longitudinally confluent in some specimens). Dorsally lacking suberect or erect vestiture; with moderately dense elongate-fine appressed white scales, scales smaller medially. Epistoma slightly elevated and swollen. Pronotum. Median carina lacking to variously developed as a low, glabrous, shiny line. Dorsal punctation small, dense and shallow; larger and sparser on flanks; areas between larger dorsal punctures with very dense shallow minute punctures, the areas between them shiny and irregularly elevated. Dorsally lacking suberect or erect vestiture (Fig. 140a). Scales dorsally absent to moderately dense and very small along midline; moderately dense, small and fine, appressed in a pair of broad paramedian apically narrowed stripes; laterally absent to moderately dense and very small. Lateral margins with appressed white scales large and robust, moderately dense in a moderately broad stripe. Flanks with scales small and fine. Median basal area of disk very shallowly and very broadly impressed. Anterolateral margin with rounded pronotal postocular lobes moderately developed; postocular vibrissae of approximately uniform length (slightly longer behind basal portion of eye in some specimens which have less well-developed postocular lobes). In dorsal view with lateral margins straight, slightly convergent from base to apical one-quarter, then straight and more markedly convergent to apex; subapical constriction indistinct (Fig. 121). Prosternum. With shallow to moderately deep transverse impression and with prosternal

swelling lacking to slightly developed anterior to each procoxal cavity. In lateral view with anterior flat to slightly angulate portion short, not distinctly differentiated from posterior longer more angulate portion.

**Elytra.** Elongate-narrow in general form (width at midlength 0.449-0.500 times length in males; 0.434-0.533 in females) (Fig. 121). In dorsal view with lateral margins subparallel from apical one-third to humerus; moderately evenly arcuate from apical one-third to apex. Elytral apices not produced, sub-acuminate, not divergent. Humerus distinct. Dorsally lacking suberect or erect vestiture. Sutural interval to interval 8 with scales uniformly moderately dense, fine and elongate, white, with scattered patches of slightly larger white scales; intervals 9 to 11 with scales uniformly moderately dense, large, robust and white (Fig. 121). Punctures of elytral striae small, individually distinct, larger and less individually distinct towards base, arranged in regular rows. Wings. Long (greater than elytra in length). Legs. Foretibia of female with inner margin with small denticles in apical three-quarters; subapical tooth small, indistinct from tibial denticles. Foretibial, mesotibial, and metatibial unci of both sexes moderately large. Tarsal claws connate in basal one-half, not divergent, each with basal internal flange well-developed. Ventral tarsal pilose vestiture of all tarsi of male and female extensive, present as rounded pads which cover more or less entire ventral surface of all articles, with

exception of basal portion of article 1. **Abdomen.** Ventral surface with moderately dense appressed very elongate white scales. Abdominal sterna III and IV (especially laterally), with distinct moderately large glabrous patches, each with large puncture and single elongate appressed scale situated in or near the center; lacking-suberect or erect vestiture except for small, scattered, sparse hair-like scales on sternum VII of male. Base of abdominal sternum VII of female internally (dorsally) with slightly developed median shiny tubercle. **Genitalia.** Female (four examined). Abdominal sternum VIII with basal arm very short; lateral arms straight to very slightly sinuate and divergent from base to apex, apices not expanded (Fig. 160). Stylus large in length compared to length gonocoxite II. Male (two examined). Internal sac (Figs. 180a,b) elongate, moderately high; median dorsal pocket moderately high, with single narrow moderately elongate dorsally directed basal lobe (lobe A), with moderately large paired dorsally directed lobe at basal one-third on dorsal surface (lobe C), and with small ventrally directed paired lobe near base near ventral margin (lobe F); lobes B, D, and E lacking. Apex of median dorsal pocket rounded in dorsal view. Apical pocket individually distinct from median dorsal pocket. Ventral median pocket moderately large. Apical pocket with paired moderately large ventrally directed lobe near apical one-third at ventral margin. Eversible apical sclerite complex with paired simple scythe-like sclerite; adjacent ventral surface of apical

pocket with pair of large, dark, longitudinal sclerites.

**Geographic variation.**--- None noted.

**Geographic distribution.**--- This species is distributed along the Atlantic Coastal Plain from New York south to Florida (Fig. 215).

**Natural history.**--- This species is found throughout the Atlantic Coastal Plain in the southeastern mesophytic evergreen forest region. This species was described from several specimens collected on flowers of the hoary lupine, *Lupinus diffusus* Nutt. (Leguminosae) near Dunedin, Florida between January 24 and March 18 (Blatchley 1914; Blatchley and Leng 1916). Other adults have been collected in roots of *Lupinus* sp. (Wilmington, North Carolina) and have been reared from *Lupinus cumulicola* Small (Spring Valley, Florida). Adult specimens have been collected from January to May at elevations of sea level to 182 m ( $\bar{X}$ =36 m, N=7) (Fig. 233)

**Chorological relationships.**--- Table 67. In the northern part of its range, this species is sympatric only with *C. calandroides*.

**Phylogenetic relationships.**--- Primary characters allow only for placement of this species as a member of an unresolved

trichotomy that also includes *C. longinasus* and *C. texanus*. This lineage then comprises an unresolved pentachotomy with *C. porlcollis*, *C. eustictorrhinus*, *C. modestus*, and the remainder of *Cleonidius* (Fig. 236). Secondary characters support monophyly of the *C. subcylindricus*-*C. longinasus*-*C. texanus* lineage and result in the hypothesis that this species is the sister species to the *C. longinasus*-*C. texanus* lineage. The *C. subcylindricus*-*C. longinasus*-*C. texanus* lineage is left as a member of an unresolved trichotomy that also includes the *C. porlcollis*-*C. eustictorrhinus* lineage and *C. modestus* (Fig. 237).

Table 35. Descriptive statistics for *C. subcylindricus* based on seven males from throughout the species range

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.68-2.08	1.937	0.227	0.114
WF	1.00-1.16	1.051	0.102	0.052
WRA	1.12-1.36	1.217	0.124	0.062
LP	2.96-3.52	3.291	0.313	0.158
WPB	2.88-3.68	3.303	0.431	0.217
WE1H	3.52-4.32	3.943	0.425	0.214
WE1M	3.68-4.64	4.206	0.571	0.288
LE1	7.68-9.44	8.663	1.077	0.543
LEy	0.92-1.12	1.034	0.117	0.059
WEy	0.56-0.68	0.617	0.068	0.034
B. Proportions.				
WEy/LEy	0.57-0.64	0.597	0.036	0.018
WPB/LP	0.95-1.05	1.002	0.053	0.027
WRA/LR	0.58-0.69	0.630	0.054	0.027
WRA/WF	1.07-1.24	1.158	0.081	0.041
WE1M/LE1	0.45-0.50	0.485	0.028	0.014
LP/LE1	0.36-0.40	0.381	0.019	0.010

Table 36. Descriptive statistics for *C. subcylindricus* based on 17 females from throughout the species range

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.68-2.28	2.049	0.281	0.091
WF	0.92-1.40	1.118	0.168	0.054
WRA	1.12-1.40	1.266	0.134	0.043
LP	2.80-3.92	3.426	0.515	0.166
WPB	2.80-4.00	3.449	0.502	0.162
WE1H	3.28-4.72	4.061	0.643	0.208
WE1M	3.52-5.12	4.348	0.766	0.248
LE1	7.52-10.24	8.951	1.210	0.391
LEy	0.88-1.16	1.052	0.123	0.040
WEy	0.52-0.76	0.609	0.084	0.027
3. Proportions.				
WEy/LEy	0.52-0.68	0.580	0.065	0.021
WPB/LP	0.96-1.07	1.008	0.052	0.017
WRA/LR	0.56-0.69	0.619	0.042	0.013
WRA/WF	1.00-1.26	1.137	0.087	0.028
WE1M/LE1	0.43-0.53	0.485	0.039	0.013
LP/LE1	0.36-0.41	0.383	0.022	0.007

4.5.2.1.5 *Cleonidius longinasus* Anderson, new name

(Figs. 122, 141, 161, 181, 214)

*Lixus mixtus* LeConte 1876a:416 [not Fabricius 1792:417].

Holotype (examined), female, labelled "Col",

"B.D.Smith", "Type/ 5178", "*L. mixtus*/ Lec." (MCZC).

Type locality, Colorado. Henshaw 1881-1882 (check.).

Henshaw 1885 (check.) Leng 1920 (catal.).

*Cleonus* (*Cleonidius*) *modestus*; Hatch 1971 (key, redesc.,  
misident.).*Cleonus mixtus*; Kingsolver 1972.*Cleonis mixtus*; O'Brien and Wibmer 1982 (catal., distn.).*mixtus, incertae sedis*; Csiki 1934 (catal.).

Notes about synonymy.--- The name *Lixus mixtus* LeConte is a junior homonym and is here replaced with the name *C. longinasus* Anderson.

Derivation of specific epithet.--- From the Latin "*longus*" meaning long and "*nasus*" meaning nose. This name is used in reference to the very long rostrum of females of this species (Fig. 141a).

Problems in recognition.--- Most individuals, especially females, of this species are very easily recognized by the very long and narrow rostrum (Fig. 141a). Otherwise, individuals, especially males, are likely only to be



confused with *C. eustictorrhinus*, *C. subcylindricus* and *C. texanus*. Individuals of *C. longinasus* are separated from *C. subcylindricus* by the latter entirely lacking suberect or erect dorsal vestiture and being distributed only along the Atlantic Coastal Plain from New York south to Florida. *C. longinasus* and *C. eustictorrhinus* are sympatric in California but individuals are easily distinguished by the more robust, medially tumescent rostrum of the latter (Fig. 138). On the other hand, although most *C. texanus* and *C. longinasus*, where they are sympatric in California and Oregon, can be separated on the basis of the length and width of the rostrum, length of erect dorsal vestiture, and association with different plant taxa, some specimens, especially males, are very difficult to reliably separate as discussed under the "Problems in recognition" section for *C. texanus*.

A very few individuals of *C. longinasus* have scales on elytral intervals 3 to 5 larger than those on adjacent intervals giving the elytra an indistinctly vittate scale pattern and perhaps resulting in confusion with species with distinctly vittate scale patterns. Specimens from San Francisco, California entirely lack suberect or erect vestiture but have the long narrow rostrum typical of members of this species.

**Description.**--- Specimens examined. 44 males, 80 females.  
Data about variation in LR, WF, WRA, LP, WPB, WE1H, WE1M,

LEl, LEy, WEy, WEy/LEy, WPB/LP, WRA/LR, WRA/WF, WEIM/LEl, and LP/LEl are presented in Tables 37 and 38. Size. Length, male, 7.2-9.8 mm; female, 6.1-11.2 mm. Width, male, 2.2-3.4 mm; female, 2.1-3.8 mm. Head. Eye elongate-oval. Area behind eye with numerous wrinkles of cuticle. Frons with sparse to moderately dense, small, shallow punctures, some longitudinally confluent and irregularly impressed immediately above eyes, interspersed with minute punctures. Frons also with sparse to moderately dense, elongate-fine to elongate-narrow, white, appressed to suberect scales medially, elongate-narrow and erect immediately above eyes. Some specimens also with moderately dense very elongate-fine erect hair-like scales medially and immediately above eyes. Frons continuous with base of rostrum or only very slightly separated by very shallow indistinct transverse impression (Fig. 141a). Base of median carina of rostrum (if present) with small, shallow fovea. Width of frons greater than width at apex of rostrum. Rostrum. Very elongate-narrow, especially so in females (width at apex in male 0.400-0.556 times length; 0.344-0.438 in female) (Fig. 141). In lateral view straight to moderately curved downward (markedly so near apex in some specimens). In dorsal view with postgenae not laterally expanded, antennal scrobes not or only very slightly visible. Rostrum not medially tumescent, with median carina lacking to variously developed as rounded slightly elevated fine glabrous shiny line. Dorsal and lateral punctation dense, small, shallow (longitudinally

confluent in many specimens), smaller and not longitudinally confluent apically, dorsally interspersed with minute punctures. Dorsally with suberect or erect vestiture lacking (few specimens) to with sparse to moderately dense, moderately long to very long, erect hair-like scales in basal one-half, erect vestiture lacking from apical one-half; with sparse to moderately dense appressed to suberect elongate-fine scales in basal one-half, lacking from (perhaps abraded) or very sparse in apical one-half in most specimens. Epistoma not swollen. Pronotum. Median carina lacking to variously developed in anterior one-half as slightly elevated narrow shiny glabrous line. Dorsal punctation moderately large, moderately dense to dense, moderately deep; smaller and shallower apically; sparser on flanks; areas between large dorsal punctures with minute regularly impressed punctures. Dorsally (especially laterally) with sparse to moderately dense, moderately long to very long, erect hair-like scales, each situated in large puncture (Fig. 141a). Scales dorsally absent to moderately dense, small and fine along median line; sparse to dense, elongate-fine to elongate-narrow laterad of median line (forming distinct apically narrowed stripes in many specimens); laterally, absent or sparse to moderately dense, small and fine. Lateral margins with scales moderately dense to dense, elongate-narrow to moderately robust, appressed to recumbent in moderately broad stripe. Flanks with scales moderately dense to dense, elongate-narrow. Median basal

area of disk broadly and shallowly impressed. Anterolateral margin with pronotal postocular lobes well-developed; postocular vibrissae uniformly short. In dorsal view with lateral margins uniformly very slightly arcuate and convergent from base to apex; subapical constriction indistinct. Prosternum. With very shallow, indistinct impression and very slightly to moderately developed rounded swelling anterior to each procoxal cavity. In lateral view with anterior flat to slightly angulate portion long, approximately one-half to two-thirds length of posterior more angulate portion; anterior and posterior portions not to slightly differentiated by shallow transverse impression.

• Elytra. Elongate-narrow in general form (width at midlength 0.481-0.540 times length in males; 0.481-0.580 in females), (Fig. 122). In dorsal view with lateral margins straight to very slightly arcuate or slightly sinuate, slightly convergent from apical one-third to humerus; slightly to moderately and evenly arcuate from apical one-third to apex. Elytral apices slightly produced, rounded to sub-acuminate. Humerus distinct. Dorsally with sparse to moderately dense, short to very long erect hair-like scales (longest and densest on declivity of most specimens; absent on specimens from San Francisco, California). Sutural interval to interval 7 with uniformly moderately dense, fine to elongate-fine scales, scales on intervals 3 to 5 larger than those on sutural interval and interval 2, and intervals 6 and 7 in some specimens; interval 8 with scales small, very

fine to fine, sparse to moderately dense; intervals 9 to 11 with scales moderately dense to dense, elongate-narrow to moderately robust, sparser and smaller apically in some specimens (Fig. 122). Punctures of elytral striae small, individually distinct, arranged in regular rows. **Wings.** Long (greater than elytra in length). **Legs.** Foretibia of female with inner margin with small denticles in apical one-half; subapical tooth small, indistinct from tibial denticles. Foretibial and mesotibial unci of both sexes large to moderately large, metatibial uncus small. Tarsal claws connate in basal one-third to one-half, not divergent, each with basal internal flange well-developed. Ventral tarsal pilose vestiture of all tarsi of male and female present as elongate-narrow pads covering apical one-half to two-thirds of article 1, as large oval pads covering apical three-quarters to more or less the entire ventral surface of article 2, and as large rounded pads covering apical three-quarters to more or less the entire ventral surface of article 3. **Abdomen.** Ventral surface with moderately dense, elongate-narrow appressed white scales. Abdominal sterna III to VI (especially III and IV) with individually indistinct to distinct glabrous patches, each with large puncture and single suberect to erect moderately long hair-like scale situated in or near the center. Base of abdominal sternum VII of female internally (dorsally) with well-developed median shiny tubercle. **Genitalia.** Female (two examined). Abdominal sternum VIII with basal arm short; lateral arms

straight and divergent from base to apical one-quarter, inwardly arcuate at apical one-quarter and convergent to apex, apices slightly expanded (Fig. 161). Stylus moderately large compared to length gonocoxite, II. Male (two examined). Internal sac (Figs. 181a,b) elongate, moderately high; median dorsal pocket moderately high, with single moderately broad elongate dorsally directed basal lobe (lobe A), with large paired dorsally directed lobe at basal one-third on dorsal surface (lobe C), and with small ventrally directed paired lobe near base near ventral margin (lobe F); lobes B, D, and E lacking. Apex of median dorsal pocket rounded in dorsal view. Apical pocket individually distinct from median dorsal pocket. Ventral median pocket moderately large. Apical pocket with paired moderately large ventrally directed lobe near apical one-third at ventral margin. Eversible apical sclerite complex with paired simple scythe-like sclerite; adjacent ventral surface of apical pocket with pair of large, dark, longitudinal sclerites.

**Geographic variation.**--- Specimens from San Francisco, California lack dorsal suberect or erect vestiture but are otherwise typical *C. longinasus*. Some specimens from California and Oregon have less elongate-narrow rostra than typical for this species. This is likely due to hybridization with *C. texanus* which in these areas also occurs on Leguminosae. It does not appear to be host plant induced for all *C. longinasus* are found on Leguminosae.

**Geographic distribution.**--- This species is distributed from southern British Columbia and Washington, south to Idaho, western Montana and Wyoming in the east, south to southern California in the west (Fig. 214).

**Natural history.**--- This species is associated with grassland, Pacific semi-desert and Great Basin desert habitats. Adults of this species have only been found associated with Leguminosae. They have been collected on *Lupinus* spp. and *Astragalus* spp. (both Leguminosae) throughout California. No definite host records are known but it is likely that various annual legumes serve as potential host plants throughout the species range. Adults have been collected from February to December at elevations of from 31-2293 m (X=990 m, N=14) (Fig. 233). Unlike adults of *C. texanus* which are known to occur, albeit rarely, on low altitude Leguminosae, *C. longinasus* are not known from Rosaceae, the presumed primary host of the former species.

**Chorological relationships.**--- Table 67.

**Phylogenetic relationships.**--- Primary characters allow only for placement of this species as a member of an unresolved trichotomy that also includes *C. subcylindricus* and *C. texanus*. This lineage then comprises an unresolved pentachotomy with *C. porlcollis*, *C. eustictorrhinus*, *C. modestus*, and the remainder of *Cleonidius* (Fig. 236).

Secondary characters support monophyly of the *C. subcylindricus-C. longinasus-C. texanus* lineage and allow for further resolution and result in the hypothesis that this species is the sister species to *C. texanus* and that the *C. subcylindricus-C. longinasus-C. texanus* lineage is a member of an unresolved trichotomy that also includes the *C. poricollis-C. eustictorrhinus* lineage and *C. modestus* (Fig. 23).



Table 37. Descriptive statistics for *C. longinasus* based on 10 males from Hemet Reservoir, Herkey Creek, and Idyllwild, California

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.80-2.28	1.980	0.283	0.119
WF	1.04-1.24	1.112	0.089	0.037
WRA	0.84-1.08	0.952	0.101	0.043
LP	2.56-3.44	2.976	0.434	0.183
WPB	2.72-3.44	3.056	0.298	0.126
WE1H	3.20-4.00	3.560	0.345	0.145
WE1M	3.52-4.40	3.936	0.466	0.196
LE1	6.72-8.64	7.744	0.934	0.394
LEy	0.92-1.04	0.960	0.063	0.027
WEy	0.52-0.64	0.556	0.066	0.028
B. Proportions.				
WEy/LEy	0.52-0.65	0.579	0.060	0.025
WPB/LP	0.97-1.13	1.030	0.071	0.030
WRA/ER	0.40-0.56	0.485	0.080	0.034
WRA/WF	0.74-0.93	0.858	0.130	0.043
WE1M/LE1	0.48-0.54	0.509	0.031	0.013
LP/LE1	0.36-0.40	0.384	0.023	0.010

Table 38. Descriptive statistics for *C. longinasus* based on 10 females from Hemet Reservoir, Herkey Creek, and Idyllwild, California

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.92-2.52	2.296	0.261	0.110
WF	1.08-1.20	1.172	0.064	0.027
WRA	0.80-1.00	0.888	0.093	0.039
LP	2.96-3.44	3.184	0.232	0.098
WPB	2.96-3.36	3.264	0.202	0.085
WE1H	3.36-4.08	3.824	0.329	0.139
WE1M	3.84-4.64	4.272	0.432	0.182
LE1	7.52-8.80	8.224	0.744	0.313
LEy	0.92-1.08	0.984	0.064	0.027
WEy	0.52-0.60	0.564	0.034	0.014
B. Proportions.				
WEy/LEy	0.52-0.63	0.574	0.048	0.020
WPB/LP	0.95-1.11	1.027	0.081	0.034
WRA/LR	0.34-0.44	0.388	0.043	0.018
WRA/WF	0.70-0.83	0.758	0.069	0.029
WE1M/LE1	0.48-0.58	0.520	0.043	0.018
LP/LE1	0.37-0.40	0.388	0.015	0.006

4.5.2.1.6 *Cleonidius texanus* (LeConte)

(Figs. 123, 142, 162, 182, 218)

- Lixus texanus* LeConte 1876a:155. Lectotype (here designated), one of two syntypes, male, labelled with a dark red circle (=Texas), "67", "TYPE/ 5250", "L.texanus/ Lec" and with my designation label "*Lixus texanus* LeC./ LECTOTYPE/ desig. Anderson" (MCZC). Type locality, Texas. LeConte 1876a (in part; misident. of *Cleonidius quadrilineatus*, [mixed type Series]). Henshaw 1881-1882 (check.). Henshaw 1885 (check.).
- Cleonus (Cleonidius) circumductus* Casey 1891:192. New synonymy. Holotype (examined), male, labelled "Ari.", "CASEY/ bequest/ 1925", "TYPE USNM/ 37282", "C.Cl./ circumductus/ Cas." (USNM). Type locality, Arizona. Csiki 1934 (catal.).
- Cleonus (Cleonidius) texanus*; Casey 1891 (key). Csiki 1934 (catal.).
- Cleonus circumductus*; Wickham 1902 (check.). Leng 1920 (catal.).
- Cleonus texanus*; Wickham 1902 (check.). Leng 1920 (catal.).
- Cleonus canescens*; Yothers 1916 (misident.). Essig 1958 (biol., misident.).
- Cleonis circumductus*; O'Brien and Wibmer 1982 (catal., distn.).
- Cleonis modestus*; O'Brien and Wibmer 1982 (in part; catal., distn., misident.).
- Cleonis texanus*; O'Brien and Wibmer 1982 (catal., distn.).

Notes about synonymy.--- Casey (1891) distinguished *C. circumductus* Casey from *C. texanus* (LeConte) by the longer, denser and more erect dorsal vestiture. I find these features of the dorsal vestiture to vary throughout the species range and do not consider these differences as warranting separate species status. (See also sections on "Problems in recognition" and "Variation".)

Problems in recognition.--- Individuals of this species are likely to be confused only with individuals of *C. longinasus*, *C. eustictorrhinus*, and *C. subcylindricus*. *C. subcylindricus* individuals are easily separated as they lack suberect or erect dorsal vestiture and are found only throughout the Atlantic Coastal Plain of the eastern United States of America. Separation of *C. eustictorrhinus* and *C. texanus* individuals is more difficult. Both species occur in California but can be separated by the more robust and medially tumescent rostrum of *C. eustictorrhinus* (Fig. 138) as outlined in the key.

Individuals, especially males, of *C. texanus* and *C. longinasus* are perhaps the most difficult species of *Cleonidius* to reliably separate. Both species are sympatric over a wide geographic range but are especially problematical to distinguish in California and Oregon. Most *C. longinasus* are slightly larger in size, have long to very long dorsal erect vestiture (Fig. 141a), a very elongate-narrow rostrum (especially so in females) (Fig.

141a), and are associated with both low and high altitude Leguminosae. Most *C. texanus* on the other hand have generally shorter dorsal erect vestiture (Fig. 142a), a shorter, slightly more robust rostrum (Fig. 142a), and are associated with Rosaceae or Rhamnaceae. All individuals from Rosaceae or Rhamnaceae are structurally typical *C. texanus*, although not all specimens associated with Leguminosae are *C. longinasus*. I have seen both male and female specimens with short rostra and short dorsal erect vestiture, both character states of *C. texanus*, from low altitude Leguminosae. I consider these as *C. texanus*. Various individuals from the area of sympatry, including some females, but especially males and those individuals lacking data on plant associations, are less easy to separate based on these structural differences. The rostrum in these specimens is generally of a length precluding reliable assignment to either species, however, length of dorsal erect vestiture of many specimens allows for a tentative identification. As a consequence of this variation, some may consider these two forms as questionably warranting separate species status, but the moderately extensive distribution of *C. longinasus*, yet the lack of *C. longinasus* or *C. texanus* on Leguminosae from areas (particularly Arizona and Texas) where *C. texanus* is found on Rosaceae and Rhamnaceae, and the lack of individuals with structural character states of *C. longinasus* from Rosaceae or Rhamnaceae yet presence of individuals with structural features of *C. texanus* from

Leguminosae, argues against conspecificity (the structural differences perhaps induced by association with different host plants). I suspect hybridization to be the likely cause of the complexities in variation but because of the noted assymetry in plant associations, do not regard it as warranting conspecificity of the two forms.

**Description.**--- Specimens examined. 156 males, 174 females. Data about variation in LR, WF, WRA, LP, WPB, WE1H, WE1M, LE1, LEy, WEy, WEy/LEy, WPB/LP, WRA/LR, WRA/WF, WE1M/LE1, and LP/LE1 are presented in Tables 39 and 40. **Size.** Length, male, 6.2-8.6 mm; female, 5.3-8.8 mm. Width, male, 1.9-3.2 mm; female, 1.8-3.0 mm. **Head.** Eye elongate-oval. Area behind eye with numerous wrinkles of cuticle. Frons with sparse to moderately dense, small, shallow punctures, some longitudinally confluent and irregularly impressed immediately above eyes, interspersed with minute punctures. Frons also with moderately dense, elongate-fine, white appressed to suberect scales medially, elongate-narrow and erect immediately above eyes. Frons continuous with base of rostrum or only very slightly separated by very shallow indistinct transverse impression (Fig. 142a). Base of median carina of rostrum (if present) with small, shallow fovea. Width of frons greater than or subequal to width at apex of rostrum. **Rostrum.** Elongate-narrow, slightly more so in females (width at apex 0.512-0.676 times length in male; 0.459-0.658 in female) (Fig. 142). In lateral view slightly

curved downward (markedly so near apex in some specimens). In dorsal view with postgenae not to very slightly laterally expanded, antennal scrobes only very slightly visible. Rostrum not to very slightly medially tumescent, with median carina lacking. Dorsal and lateral punctation moderately dense to dense, small, shallow (longitudinally confluent in some specimens), smaller and not longitudinally confluent apically, dorsally interspersed with minute punctures. Dorsally with suberect or erect vestiture lacking (few specimens) to with sparse, short, sub-erect to erect hair-like scales laterally and basally, erect vestiture lacking from apical one-half; with sparse to moderately dense recumbent elongate-very-fine to elongate-fine scales laterally in basal one-half, lacking from (perhaps abraded) or very sparse in apical one-half and medially in most specimens. Epistoma not swollen. Pronotum. Median carina lacking to variously developed in anterior one-half as an indistinct slightly elevated narrow shiny glabrous line. Dorsal punctation moderately large, moderately dense to dense, moderately deep; smaller and shallower apically; sparser on flanks; areas between large dorsal punctures with minute regularly impressed punctures. Dorsally (especially laterally) with sparse to moderately dense, short to moderately long, erect hair-like scales, each situated in large puncture (Fig. 142a). Scales dorsally sparse to moderately dense, small and fine along median line; moderately dense, elongate-fine to elongate-narrow laterad

of median line; laterally, sparse to moderately dense, small and fine. Lateral margins with scales moderately dense to dense, elongate-narrow to moderately robust, appressed to recumbent in moderately broad stripe. Flanks with scales moderately dense, elongate-narrow. Median basal area of disk broadly and shallowly impressed. Anterolateral margin with pronotal postocular lobes well-developed; postocular vibrissae uniformly short. In dorsal view with lateral margins uniformly very slightly arcuate and convergent from base to apex; subapical constriction indistinct, or straight and convergent from base to apical one-quarter, slightly constricted at apical one-quarter then straight and convergent to apex. Prosternum. With shallow to moderately deep impression and very slightly to well-developed rounded swelling anterior to each procoxal cavity. In lateral view with anterior flat to slightly angulate portion long, approximately one-half to two-thirds length of posterior more angulate portion; anterior and posterior portions not to slightly differentiated by shallow transverse impression. Elytra. Elongate-narrow in general form (width at midlength 0.447-0.514 times length in males; 0.443-0.506 in females) (Fig. 123). In dorsal view with lateral margins straight to very slightly arcuate or slightly sinuate, slightly convergent from apical one-third to humerus; slightly to moderately and evenly arcuate from apical one-third to apex. Elytral apices slightly produced, rounded to sub-acuminate. Humerus distinct. Dorsally with sparse to moderately dense,



short suberect to erect hair-like scales (longest and densest on declivity of most specimens). Sutural interval to interval 7 with uniformly moderately dense, fine to elongate-fine scales, interval 8 with scales small, very fine to fine, sparse to moderately dense; intervals 9 to 11 with scales moderately dense to dense, elongate-narrow to moderately robust, sparser and smaller apically in some specimens (Fig. 123). Punctures of elytral striae small, individually distinct, arranged in regular rows. Wings. Long (greater than elytra in length). Legs. Foretibia of female with inner margin with small denticles in apical one-half; subapical tooth small, indistinct from tibial denticles. Foretibial and mesotibial unci of both sexes large to moderately large, metatibial uncus small. Tarsal claws connate in basal one-half, not divergent, each with basal internal flange well-developed. Ventral tarsal pilose vestiture of all tarsi of male and female present as elongate-narrow pads covering apical one-half to two-thirds of article 1, as large oval pads covering apical three-quarters to more or less the entire ventral surface of article 2 and as large rounded pads covering apical three-quarters to more or less the entire ventral surface of article 3. Abdomen. Ventral surface with moderately dense, elongate-narrow appressed white scales. Abdominal sterna III to VI (especially III and IV) with individually indistinct to distinct glabrous patches, each with large puncture and single suberect to erect moderately long hair-like scale

situated in or near the center. Base of abdominal sternum VII of female internally (dorsally) with well-developed median shiny tubercle. Genitalia. Female (four examined). Abdominal sternum VIII with basal arm short; lateral arms straight and divergent from base to apical one-third, inwardly arcuate at apical one-third and convergent to apex, apices slightly expanded (Fig. 162). Stylus moderately large compared to length gonocoxite II. Male (four examined). Internal sac (Figs. 182a,b) elongate, moderately high; median dorsal pocket moderately high, with single moderately broad elongate dorsally directed basal lobe (lobe A), with moderately large paired dorsally directed lobe at basal one-third on dorsal surface (lobe C), and with small ventrally directed paired lobe near base near ventral margin (lobe F); lobes B, D, and E lacking. Apex of median dorsal pocket rounded in dorsal view. Apical pocket individually distinct from median dorsal pocket. Ventral median pocket moderately large. Apical pocket with paired moderately large ventrally directed lobe near apical one-third at ventral margin. Eversible apical sclerite complex with paired simple scythe-like sclerite; adjacent ventral surface of apical pocket with pair of large, dark, longitudinal sclerites.

**Geographic variation.** None noted, but see "Geographic variation" section for *C. longinasus* and "Problems in recognition" section for *C. texanus*.

**Geographic distribution.**--- This species is distributed from Washington and Idaho, east to Colorado, south to southeastern Texas in the east, and southern California in the west (Fig. 218).

**Natural history.**--- This species is associated with the western xerophytic evergreen forest habitat. Adults have been collected primarily on Rosaceae and Rhamnaceae throughout the species range but also on Leguminosae, although only in coastal California and Oregon. Records from Rosaceae and Rhamnaceae are primarily at higher altitudes, regardless of geographic location (although especially in oak-pinyon-juniper woodlands). Records from Leguminosae are known only from lower altitudes. Adults have been collected on the following plants; *Lupinus* sp. (Leguminosae); *Ceanothus* sp., *C. cuneatus* (Hook.) Nutt., *C. leucodermis* Greene, *C. integerrimus* Hook. and Arn. (Rhamnaceae); *Cercocarpus* sp., *C. ledifolius* Nutt., *C. montanus* Raf., *Cowanla mexicana* D. Don., *Prunus amygdalus* Batsch. (almond), *P. persica* Batsch. (peach), *Pyrus communis* L. (pear), *Purshia tridentata* (Pursh.) DC. (all Rosaceae). Adults have been reared only from *Lupinus affinis* J.G. Agardh. (Leguminosae) at Berkeley, California, but I suspect that a variety of Rosaceae and Rhamnaceae serve as hosts at higher altitudes throughout most of the species range. Adults have been collected from January to October at altitudes ranging from 15-2998 m ( $\bar{X}$ =1345 m, N=56) (Fig. 233). I and others

have collected specimens of *C. texanus* and *C. quadrilineatus* together on various Rosaceae and Rhamnaceae in Arizona and Texas.

**Chorological relationships.**--- Table 67.

**Phylogenetic relationships.**--- Primary characters allow only for placement of this species as a member of an unresolved trichotomy that also includes *C. subcylindricus* and *C. longinasus*. This lineage then comprises an unresolved pentachotomy with *C. poricollis*, *C. eustictorrhinus*, *C. modestus*, and the remainder of *Cleonidius* (Fig. 236).

Secondary characters support monophyly of the *C. subcylindricus*-*C. longinasus*-*C. texanus* lineage and allow for further resolution and result in the hypothesis that this species is the sister species to *C. longinasus* and that the *C. subcylindricus*-*C. longinasus*-*C. texanus* lineage is a member of an unresolved trichotomy that also includes the *C. poricollis*-*C. eustictorrhinus* lineage and *C. modestus* (Fig. 237).

Table 39. Descriptive statistics for *C. texanus* based on eight males from Mile 19, Hitchcock Highway, Santa Catalina Mountains, Arizona

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.44-1.64	1.535	0.101	0.048
WF	0.84-0.96	0.900	0.064	0.030
WRA	0.80-0.88	0.830	0.042	0.020
LP	2.16-2.56	2.320	0.192	0.090
WPB	2.48-2.56	2.510	0.062	0.029
WE1H	2.88-3.20	3.030	0.175	0.082
WE1M	3.12-3.36	3.230	0.156	0.052
LE1	6.32-6.96	6.580	0.325	0.153
LEy	0.80-0.88	0.825	0.045	0.021
WEy	0.44-0.52	0.475	0.038	0.018
B. Proportions.				
WEy/LEy	0.52-0.62	0.576	0.051	0.024
WPB/LP	1.00-1.15	1.084	0.072	0.034
WRA/LR	0.51-0.56	0.541	0.025	0.012
WRA/WF	0.87-1.00	0.924	0.073	0.034
WE1M/LE1	0.47-0.51	0.491	0.021	0.010
LP/LE1	0.34-0.38	0.353	0.023	0.011

Table 40. Descriptive statistics for *C. texanus* based on eight females from Mile 19, Hitchcock Highway, Santa Catalina Mountains, Arizona

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.32-1.72	1.515	0.199	0.094
WF	0.84-1.00	0.905	0.078	0.026
WRA	0.68-0.80	0.740	0.064	0.030
LP	2.00-2.56	2.360	0.301	0.142
WPB	2.24-2.72	2.530	0.231	0.109
WE1H	2.56-3.20	2.990	0.307	0.145
WE1M	2.72-3.44	3.210	0.336	0.158
LE1	5.52-7.04	6.530	0.748	0.352
LEy	0.76-0.92	0.830	0.077	0.036
WEy	0.44-0.52	0.480	0.056	0.026
B. Proportions.				
WEy/LEy	0.55-0.62	0.578	0.036	0.017
WPB/LP	1.00-1.12	1.075	0.063	0.030
WRA/LR	0.46-0.55	0.490	0.045	0.021
WRA/WF	0.77-0.87	0.818	0.048	0.023
WE1M/LE1	0.48-0.51	0.492	0.017	0.008
LP/LE1	0.34-0.38	0.361	0.016	0.008

4.5.2.2 *Cleonidius americanus* species group

**Diagnosis.**--- Size small to moderate for *Cleonidius* (Fig. 197). Dorsal erect or suberect vestiture absent to present, dense, short to very long. Rostrum moderately elongate-narrow (Fig. 143) to very robust (Figs. 146-147), slightly to markedly medially tumescent, not to variously carinate; lateral margins rounded to sharp. Pronotum with postocular lobes absent (Fig. 145) to present and well-developed (Fig. 143); postocular vibrissae uniformly short to long, of unequal length, and longest behind base of eye. Elytra with scale pattern either more or less vittate, intervals other than 9 to 11 with white scales equal in size to those on intervals 9 to 11 (Figs. 146, 149), or, with scattered patches of large white scales giving a mottled appearance (Figs. 143-144). Tarsus with ventral pilose vestiture slightly to markedly reduced in extent (covering apical one-half to two-thirds of articles 2 and 3) to absent or nearly so on all articles. Wings present, of variable length. Female with abdominal sternum VIII with basal arm short to long; lateral arms arcuate to more or less straight throughout length. Male with aedeagus with internal sac with apex of dorsal median pocket rounded; lobes A to F present; dorsal median pocket low in lateral view.

**Phylogenetic relationships.**--- Monophyly of this species group is based only on distribution of the apotypic states of the secondary character of the extent of ventral tarsal

pilosity (homoplasious within *Cleonidius* in species of the *C. boucardi* group). This monophyly is further weakly supported by the tendency for individuals of species placed in the *C. americanus* group to be brachypterous more so than individuals of species in other groups and to be the only species group whose members exhibit a mottled elytral scale pattern. These states however cannot be considered synapotypic for they are not shared by all members of the species group and independent evolution of the apotypic state within species of the group is equally parsimonious with consideration of the apotypic state as a groundplan state for the group with subsequent reversions to the plesiotypic state having taken place. The *C. americanus* group is hypothesized to be the sister group to the *C. Jobigerinus* group-*C. boucardi* group lineage, this lineage then sister to the *C. poricollis* group (Figs. 236-237).

#### 4.5.2.2.1 *Cleonidius americanus* Csiki

(Figs. 124, 143, 163, 183, 223)

*Cleonus* (*Cleonidius*) *sparsus*; Casey 1891 (key, misident.).

*Cleonus basalis* Fall 1897:242 [not Chevrolat 1860:80].

Holotype (examined), male, labelled "S.Clemente/ Id.

CAL./ 5.29.97", "TYPE", "basalis/ Fall", "M.C.Z./ Type/

25192", "H.C. FALL/ COLLECTION" (MCZC). Type locality,

San Clemente Island, California.

*Cleonus* (*Cleonidius*) *americanus* Csiki 1934:64. New name for

*Cleonus basalis* Fall. Blackwelder 1939 (check.).



*Cleonis americanus*; O'Brien and Wibmer 1982 (catalog.,  
distribution.).

*Cleonis basalis*; O'Brien and Wibmer 1982 (catalog., distribution., as  
junior homonym).

**Problems in recognition.**--- Unabraded specimens of this species should be easily recognized because of the long dorsal-erect vestiture (Fig. 143a). Only specimens of *C. longinasus* have equally long erect dorsal vestiture, but they possess a different elytral scale pattern and have a more elongate-narrow rostrum that is not laterally expanded at midlength nor has the prementum swollen ventrally, as do members of *C. americanus* (Fig. 143). Indeed, *C. americanus* and *C. frontalis* are the only species of *Cleonidius* in which the rostrum, in lateral view, has the prementum slightly to markedly swollen ventrally, and in dorsal view is variously expanded laterally at midlength (Figs. 143-144). Abraded specimens throughout the species range in California might therefore only be otherwise confused with *C. frontalis*. From *C. frontalis*, individuals of *C. americanus* can be distinguished by their larger, deeper, and denser dorsal pronotal punctation, absence of large denticles on the inner margin of the fore-tibia of females (Fig. 143c), and the rostrum straight, not slightly curved downward.

**Description.**--- Specimens examined. 65 males, 69 females.

Data about variation in LR, WF, WRA, LP, WPB, WE1H, WE1M, LE1, LEy, WEy, WEy/LEy, WPB/LP, WRA/LR, WRA/WF, WE1M/LE1, and LP/LE1 are presented in Tables 41 and 42. Size. Length, male, 4.9-10.9 mm; female, 5.4-11.3 mm. Width, male, 1.9-4.1 mm; female, 2.1-4.1 mm. Head. Eye elongate-oval. Area behind eye with numerous slight wrinkles of cuticle. Frons with sparse to moderately dense, small, shallow punctures, some longitudinally confluent and irregularly impressed immediately above eyes, interspersed with minute punctures. Frons also with sparse, elongate-fine, white appressed to recumbent scales medially, moderately dense, elongate-narrow and erect immediately above eyes; with dense, very long erect hair-like scales laterally, lacking medially. Frons continuous with base of rostrum or only very slightly separated by shallow transverse impression (Fig. 143a). Base of median carina of rostrum (if present) with small, shallow fovea. Width of frons greater than width at apex of rostrum. Rostrum. Moderately, elongate-narrow (width at apex 0.508-0.625 times length in males; 0.432-0.588 in females) (Figs. 143a,b). In lateral view straight. In dorsal view with postgenae not to very slightly laterally expanded, antennal scrobes not distinctly visible. Rostrum not to very slightly medially tumescent; slightly laterally expanded at midlength (Fig. 143b); lateral margins more or less rounded, not sharp; with median carina variously developed as moderately elevated broad shiny glabrous line. Dorsal and lateral punctation sparse to moderately dense, small,

shallow to deep (longitudinally confluent in many specimens), dorsally interspersed with minute punctures. Dorsally with moderately dense to dense, very long, erect hair-like scales (especially laterally); with moderately dense recumbent elongate-fine scales laterally in basal one-half, lacking from (perhaps abraded) or very sparse in apical one-half and medially in most specimens. Epistoma not swollen. Rostrum in lateral view with submentum slightly to markedly swollen ventrally (Fig. 143a). Pronotum. Median carina lacking to variously developed in anterior one-half as low, narrow shiny glabrous line. Dorsal punctation and that of flanks large, dense, deep; smaller, sparser and shallower apically; areas between large dorsal punctures with minute regularly impressed punctures. Dorsally with moderately dense, very long, erect hair-like scales, each situated in large puncture (Fig. 143a). Scales dorsally absent to sparse, small and fine along median line; moderately dense, elongate-fine, appressed to recumbent in pair of paramedian slightly apically narrowed stripes; laterally, absent to very sparse, small and fine. Lateral margins with scales moderately dense, elongate-narrow, appressed in narrow stripe. Flanks with scales uniformly moderately dense, elongate-fine. Median basal area of disk broadly and shallowly to moderately deeply impressed. Anterolateral margin with pronotal postocular lobes moderately-developed; postocular vibrissae uniformly short to of unequal length, moderately long and with greatest

width approximately one-half width of eye, immediately behind base of eye (Fig. 143a). In dorsal view with lateral margins slightly arcuate and convergent from base to apical one-third, slightly constricted at apical one-third then straight and convergent to apex. Prosternum. With small shallow impression and very slightly to well-developed rounded swelling anterior to each procoxal cavity. In lateral view with anterior flat to slightly angulate portion long, approximately two-thirds length of posterior more angulate portion; anterior and posterior portions slightly differentiated by shallow transverse impression. Elytra. Elongate-narrow in general form (width at midlength 0.500-0.600 times length in males; 0.500-0.560 in females) (Fig. 124). In dorsal view with lateral margins straight, slightly convergent from apical one-third to humerus; moderately and evenly arcuate from apical one-third to apex. Elytral apices not produced, rounded, not divergent. Humerus distinct. Dorsally with moderately dense, moderately long erect hair-like scales. Scales variable in color from golden to white; scale pattern various from mottled, with uniformly moderately dense small, fine scales and irregularly distributed patches of denser and larger white scales (Fig. 124); to vittate, with scales uniformly moderately dense, sutural interval and interval 2 (except at base) with golden scales; elongate-very-fine, stria 1 with irregularly distributed patches of larger elongate-narrow white scales, base of interval 2 and intervals 3 to 5 with scales

elongate-narrow, white, intervals 6 to 11 various from with irregularly distributed patches of denser and larger, elongate-narrow white scales interspersed among elongate-very-fine white or golden scales; to uniformly covered (except intervals 6 and 7 at basal one-third) with large elongate-narrow white scales. Punctures of elytral striae small, individually distinct, larger and less individually distinct basally; arranged in regular rows. Wings. Long (greater than elytra in length). Legs. Foretibia of female with inner margin with small denticles in apical three-quarters; subapical tooth small, indistinct from tibial denticles. Foretibial and mesotibial unci of both sexes large to moderately large, metatibial uncus small. Tarsal claws connate in basal one-third, not divergent, each with basal internal flange slightly developed. Ventral tarsal pilose vestiture of foretarsus of male present as elongate-narrow pads covering apical one-third to one-half of article 1, as elongate-oval pads covering apical one-half to two-thirds of article 2, and as large rounded pads covering apical two-thirds to more or less the entire ventral surface of article 3; of foretarsus of female, as apical tufts to small elongate pads covering apical one-quarter of article 1, as small rounded pads covering apical one-third of article 2, as moderately large rounded pads covering apical two-thirds of article 3; of mesotarsus of male, as elongate-narrow pads covering apical one-half of article 1, as elongate-oval pads covering apical one-half to

two-thirds of article 2, as large rounded pads covering apical two-thirds to three-quarters of article 3; of mesotarsus of female, as apical tufts to small elongate pads covering apical one-quarter of article 1, as small rounded pads covering apical one-third of article 2, as moderately large rounded pads covering apical one-half to two-thirds of article 3; of metatarsus of male, as elongate-narrow pads covering apical one-third to one-half of article 1, as elongate-oval pads covering apical one-half to two-thirds of article 2, as large rounded pads covering apical two-thirds to three-quarters of article 3; of metatarsus of female, as apical tufts to small elongate pads covering apical one-quarter of article 1, as small rounded pads covering apical one-third of article 2, as moderately large rounded pads covering apical one-half of article 3. **Abdomen.** Ventral surface with moderately dense, elongate-narrow appressed white scales. Abdominal sterna III to VI (especially III and IV) with individually indistinct to distinct moderately large glabrous patches, each with large puncture and single erect very long hair-like scale situated in or near the center. Base of abdominal sternum VII of female internally (dorsally) with only a very slightly developed median shiny tubercle. **Genitalia.** Female (three examined). Abdominal sternum VIII with basal arm short; lateral arms moderately and evenly inwardly arcuate at point slightly beyond midlength, apices slightly expanded (Fig. 163). Stylus moderately large compared to length gonocoxite II. Male

241

(three examined). Internal sac (Figs. 183a,b) elongate and low; median dorsal pocket low, with single elongate narrow dorsally directed basal lobe (lobe A), with small median dorsally directed lobe at midlength on dorsal surface (lobe B), with moderately large paired dorsally directed lobe at basal one-third on dorsal surface (lobe C), with moderately large paired dorsolaterally directed lobe near midlength near dorsal margin (lobe D), with large paired dorsolaterally directed lobe just beyond midlength at dorsal margin (lobe E), and with small ventrally directed paired lobe near base near ventral margin (lobe F). Apex of median dorsal pocket truncate in dorsal view. Apical pocket individually distinct from median dorsal pocket. Ventral median pocket moderately large. Apical pocket with paired moderately large ventrally directed lobe near apical one-third at ventral margin. Eversible apical sclerite complex with paired simple scythe-like sclerite; adjacent ventral surface and basal portion of sides of apical pocket with pair of large, dark, longitudinal sclerites.

**Geographic variation.**--- None noted.

**Geographic distribution.**--- This species is found only in California, from the San Francisco Bay region south along the coast and central valleys to the Mexican border (Fig. 223).

**Natural history.**--- This species is found only in the Pacific semi-desert habitat. Adults of this species have been collected on *Artemisia tridentata* Nutt., *Chrysothamnus nauseosus* (Pall.) Britton, *Franseria* sp., *Gutierrezia* sp., and *Aplopappus* sp. (all Compositae). No definite hosts are known but it is probable that a variety of Compositae serve as potential hosts. Adults have been collected from March to November at elevations of from 15-1446 m (X=323 m, N=7) (Fig. 233).

**Chorological relationships.**--- Table 67.

**Phylogenetic relationships.**--- Primary characters show this species to be the sister-species of *C. frontalis*, comprising a lineage that is a member of the unresolved heptachotomy also including the *C. infrequens-C. puberulus* lineage, *C. canescens*, *C. notolomus*, *C. collaris*, the *C. lobigerinus* species group, and the *C. boucardi* species group (Fig. 236). Secondary characters support monophyly of the *C. americanus-C. frontalis* lineage which is then considered the sister-group of the rest of the *C. americanus* species group (Fig. 237).



Table 41. Descriptive statistics for *C. americanus*  
based on eight males from Mercy Hot Springs, California

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.28-2.44	1.815	0.555	0.262
WF	0.88-1.44	1.155	0.261	0.123
WRA	0.80-1.24	1.005	0.234	0.111
LP	1.84-3.60	2.670	0.836	0.394
WPB	1.92-3.60	2.760	0.831	0.392
WE1H	2.40-4.56	3.390	1.058	0.499
WE1M	2.88-4.88	3.700	0.985	0.464
LE1	4.80-9.28	6.980	2.135	1.007
LEy	0.72-1.16	0.950	0.239	0.113
WEy	0.40-0.72	0.570	0.156	0.073
B. Proportions.				
WEy/LEy	0.54-0.65	0.600	0.053	0.025
WPB/LP	1.00-1.09	1.035	0.044	0.021
WRA/LR	0.51-0.63	0.559	0.052	0.025
WRA/WF	0.81-0.91	0.870	0.049	0.023
WE1M/LE1	0.50-0.60	0.534	0.044	0.021
LP/LE1	0.37-0.40	0.383	0.013	0.006

Table 42. Descriptive statistics for *C. americanus* based on eight females from Mercy Hot Springs, California

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.36-2.44	1.755	0.504	0.238
WF	0.88-1.52	1.130	0.294	0.138
WRA	0.72-1.28	0.910	0.260	0.123
LP	2.00-3.44	2.480	0.669	0.316
WPB	1.92-3.68	2.610	0.806	0.380
WE1H	2.40-4.48	3.180	0.931	0.439
WE1M	2.72-4.88	3.540	0.981	0.462
LE1	4.96-8.96	6.560	1.796	0.846
LEy	0.72-1.28	0.925	0.251	0.118
WEy	0.40-0.72	0.540	0.136	0.064
B. Proportions.				
WEy/LEy	0.53-0.67	0.586	0.064	0.030
WPB/LP	0.96-1.10	1.048	0.073	0.034
WRA/LR	0.43-0.59	0.521	0.065	0.031
WRA/WF	0.70-0.91	0.806	0.094	0.044
WE1M/LE1	0.50-0.56	0.540	0.029	0.013
LP/LE1	0.36-0.40	0.379	0.020	0.009

4.5.2.2.2 *Cleonidius frontalis* (LeConte)

(Figs. 125, 144, 164, 184, 226)

*Cleonus frontalis* LeConte 1876a:150. Lectotype (here designated), female, one of three syntypes, labelled with a pale green circle (=Wyoming Territory), "Black Hills/ Hammond", "TYPE/ 5245", "C. frontalis/ Lec." and with my designation label "*Cleonus/ frontalis* LeC./ LECTOTYPE/ desig. Anderson" (MCZC). Type locality, Black Hills, South Dakota. Henshaw 1881-1882 (check.). Henshaw 1885 (check.). Wickham 1889. Wickham 1902 (check.). Fall and Cockerell 1907 (check.). Blatchley and Leng 1916 (key, redesc.). Leng 1920 (catal.). Bleasdel 1937 (check.).

*Cleonus sparsus* LeConte 1876a:152 [not Zoubkoff 1833:334].

New synonymy. Holotype (examined), female, labelled "Col", "C./ sparsus/ TYPE Lec.", "Horn Coll/ H8528" and with a red square label (MCZC). Type locality, Colorado. Henshaw 1881-1882 (check.). Henshaw 1885 (check.). Wickham 1902 (check.). Leng 1920 (catal.). Essig 1958 (biol., in part misident. of *Cleonidius lobgerinus*). Arnett et al. 1980 (probable misident. of *Cleonidius lobgerinus*).

*Cleonus (Cleonidius) frontalis*; Casey 1891 (key). Csiki 1934 (catal.).

*Cleonus (Cleonidius) sparsus*; Casey 1891 (key, misident. of *Cleonidius americanus*). Hatch 1971 (key, redesc., in part misident. of *Cleonidius lobgerinus*).

*Cleonus (Cleonidius) stratus* Csiki 1934:65. New synonymy.

New name for *Cleonus sparsus* LeConte. Blackwelder 1939 (check.).

*Cleonis frontalis*; O'Brien and Wibmer 1982 (catal., distn., in part misident. of *Cleonidius lobigerinus* and *Cleonidius puberulus*).

*Cleonis sparsus*; O'Brien and Wibmer 1982 (catal., distn.).

*Cleonis stratus*; O'Brien and Wibmer 1982 (catal., distn.).

**Notes about synonymy.**--- To initially distinguish *C. sparsus* and *C. frontalis* as separate species, LeConte (1876a) emphasized differences in the extent of ventral pilosity and form of hind tarsal article 3. Variation in these characters is continuous and known to occur in a number of other widespread species, thus I consider these two forms as conspecific. *Cleonidius frontalis* was arbitrarily chosen over *C. sparsus* as the valid name for this species.

Casey (1891) incorrectly placed *C. puberulus* as a junior synonym of *C. frontalis* and misidentified individuals of *C. americanus* as *C. sparsus* in his key. Likewise, other authors have frequently confused *C. frontalis*, *C. puberulus*, *C. sparsus*, and less frequently also *C. lobigerinus*.

**Problems in recognition.**--- Individuals of this species are very likely to be confused with those of *C. americanus*, *C. notolomus* and perhaps, *C. lobigerinus* and *C. trivittatus*.

They are easily separated from *C. americanus* by the long to very long erect dorsal vestiture and dense, large and deep pronotal punctures of the latter. Furthermore, only a few *C. frontalis* are known from California (Fig. 226), whereas *C. americanus* is restricted to that state (Fig. 223).

Separation from *C. lobigerinus* is quite simple as members of that species, where the two are sympatric, possess a curved sulcus behind each eye (Fig. 150a). Similarly, separation from *C. trivittatus* is simple, as individuals of that species lack or at most have only very slightly developed postocular lobes (Fig. 153a).

Separation from *C. notolomus* is most difficult and discussed under "Problems in recognition" section for that species.

Most individuals of *Cleonidius* with an elytral scale pattern that is mottled with irregularly distributed patches of larger white scales, and not distinctly vittate (Fig. 125), are this species.

**Description.**--- Specimens examined. 121 males, 115 females. Data about variation in LR, WF, WRA, LP, WPB, WE1H, WE1M, LE1, LEy, WEy, WEy/LEy, WPB/LP, WRA/LR, WRA/WF, WE1M/LE1, and LP/LE1 are presented in Tables 43 and 44. **Size.** Length, male, 5.7-10.5 mm; female, 5.7-10.8 mm. Width, male, 2.2-3.9 mm; female, 2.2-4.1 mm. **Head.** Eye elongate-oval (Fig. 145a). Area behind eye with numerous slight wrinkles of cuticle. **Frons** with sparse to moderately dense, small, shallow

punctures, some longitudinally confluent and irregularly impressed immediately above eyes, interspersed throughout with minute punctures. Frons also with moderately dense, elongate-narrow, white appressed to recumbent scales medially, moderately dense, elongate-narrow and suberect to erect immediately above eyes; otherwise lacking suberect or erect vestiture. Frons slightly separated from base of rostrum by shallow to moderately deep transverse impression (Fig. 144a). Base of median carina of rostrum (if present) with small, shallow fovea. Width of frons greater than, to subequal to, width at apex of rostrum. Rostrum. Moderately robust, slightly less so in females (width at apex 0.561-0.707 times length in male; 0.538-0.650 in female) (Figs. 144a,b). In lateral view straight to very slightly curved downward. In dorsal view with postgenae not to very slightly laterally expanded, antennal scrobes very slightly visible. Rostrum slightly to moderately medially tumescent from point of antennal insertion to middle of frons (more markedly so basally); not distinctly to slightly laterally expanded at midlength; lateral margins more or less rounded, not sharp; with median carina absent to variously developed as low glabrous shiny line at crest of median tumescence. Dorsal and lateral punctation sparse to dense, small, shallow (longitudinally confluent in some specimens); smaller and less dense apically; dorsally interspersed with minute punctures. Dorsally with sparse to moderately dense, short, suberect hair-like scales (especially laterally);

with moderately dense appressed elongate-narrow scales laterally in basal one-half, sparse in apical one-half and medially in most specimens. Epistoma not swollen. Rostrum in lateral view with submentum slightly swollen ventrally (Fig. 144a). Pronotum. Median carina lacking to variously developed in anterior one-half as low, rounded, moderately broad shiny glabrous line. Dorsal punctation moderately large, moderately dense, moderately deep to deep; smaller, sparser and shallower apically and on flanks; areas between large dorsal punctures with minute regularly impressed punctures. Dorsally with moderately dense, short to moderately long, suberect to erect hair-like scales, each situated in large puncture (Fig. 144a). Scales dorsally absent to very sparse, small and fine along median line; moderately dense, elongate-fine, appressed in pair of narrow paramedian stripes; laterally, absent to very sparse, small and fine. Lateral margins with scales moderately dense to dense, elongate-fine to elongate-narrow, appressed to recumbent in moderately broad stripe. Flanks with scales moderately dense, elongate-fine to elongate-narrow. Median basal area of disk broadly and shallowly to moderately deeply impressed. Anterolateral margin with pronotal postocular lobes moderately to well-developed; postocular vibrissae uniformly short (Fig. 144a). In dorsal view with lateral margins slightly to moderately arcuate and slightly convergent from base to apex, or slightly constricted at apical one-third to one-quarter then straight and convergent

to apex. Prosternum. With small shallow to moderately deep impression and slightly to well-developed rounded swelling anterior to each procoxal cavity. In lateral view with anterior flat to slightly angulate portion long, approximately two-thirds length of, to slightly shorter than, length of posterior more angulate portion; anterior and posterior portions not to very slightly differentiated by shallow transverse impression. Elytra. Elongate-narrow in general form (width at midlength 0.500-0.560 times length in males; 0.491-0.575 in females) (Fig. 125). In dorsal view with lateral margins straight to very slightly sinuate, slightly convergent from apical one-third to humerus; moderately and evenly arcuate from apical one-third to apex. Elytral apices not produced, rounded, not to very slightly divergent. Humerus distinct. Dorsally with sparse, indistinct, very short suberect hair-like scales. Scales various in color from golden to white; scale pattern various with sutural interval and interval 2 with golden to white scales sparse to moderately dense, very small and fine to elongate-fine, intervals 3 to 5 with scales moderately dense, elongate-fine to elongate-narrow, white, intervals 6 to 8 with scales sparse to moderately dense, small and fine to elongate-narrow, white, intervals 9 to 11 with scales moderately dense, elongate-fine to elongate narrow, white; striae of many specimens variously with irregularly distributed patches of larger white scales giving mottled appearance (Fig. 125). Punctures of elytral striae small,



individually distinct; arranged in regular rows. Wings. Long (greater than elytra in length). Legs. Foretibia of female with inner margin with very large denticles in apical three-quarters; subapical tooth large, indistinct to distinct from tibial denticles (Fig. 144c). Foretibial and mesotibial unci of both sexes moderately large, metatibial uncus small. Tarsal claws connate in basal one-third, not divergent, each with basal internal flange well-developed. Ventral tarsal pilose vestiture of foretarsus of male present as elongate-oval pads covering apical one-quarter to one-third of article 1, as rounded pads covering apical one-third to one-half of article 2, and as large rounded pads covering apical one-half to two-thirds of article 3; of foretarsus of female, as apical tufts on article 1, as small rounded pads covering apical one-quarter of article 2, as moderately large rounded pads covering apical one-half of article 3; of mesotarsus of male, as elongate-narrow pads covering apical one-third of article 1, as elongate-oval pads covering apical one-third to one-half of article 2, as large rounded pads covering apical one-half to two-thirds of article 3; of mesotarsus of female, as apical tufts on article 1, as small elongate-oval pads covering apical one-third of article 2, as moderately large rounded pads covering apical one-half of article 3; of metatarsus of male, as elongate-narrow pads covering apical one-quarter to one-half of article 1, as elongate-narrow pads covering apical one-third to one-half of article 2, as elongate-oval

to rounded pads covering apical one-half to two-thirds of article 3; of metatarsus of female, as elongate-narrow pads covering apical one-quarter to one-half of article 1, as elongate-oval pads covering apical one-third of article 2, as moderately large rounded pads covering apical one-half of article 3. **Abdomen.** Ventral surface with dense, elongate-narrow to broad appressed white scales, sparser and smaller medially on abdominal sterna V to VII. Abdominal sterna III to VI (especially III and IV) with individually indistinct to distinct large rounded glabrous patches, each with large puncture and single suberect to erect moderately long hair-like scale situated in or near the center. Base of abdominal sternum VII of female internally (dorsally) with moderately to well-developed median shiny tubercle.

**Genitalia.** Female (four examined). Abdominal sternum VIII with basal arm short; lateral arms very slightly and evenly inwardly arcuate from base to apical one-quarter, more markedly inwardly arcuate at apical one-quarter and convergent to apex, apices slightly expanded (Fig. 164).

Stylus moderately large compared to length gonocoxite II.

Male (three examined). Internal sac (Figs. 184a,b) elongate and low; median dorsal pocket low, with single broad short dorsally directed basal lobe (lobe A), with small median dorsally directed lobe at midlength on dorsal surface (lobe B), with moderately large paired dorsally directed lobe at basal one-third on dorsal surface (lobe C), with moderately large paired dorsolaterally directed lobe near midlength

near dorsal margin (lobe D), with large paired dorsolaterally directed lobe just beyond midlength at dorsal margin (lobe E), and with small ventrally directed paired lobe near base near ventral margin (lobe F). Apex of median dorsal pocket truncate in dorsal view. Apical pocket individually distinct from median dorsal pocket. Ventral median pocket moderately large. Apical pocket with paired moderately large ventrally directed lobe near apical one-third at ventral margin. Eversible apical sclerite complex with paired simple scythe-like sclerite; adjacent ventral surface and basal portion of sides of apical pocket with pair of large, dark, longitudinal sclerites.

**Geographic variation.**--- None noted.

**Geographic distribution.**--- This species is distributed from southern Alberta east to southern Manitoba, south to California in the west, east to Nebraska, Kansas and western Texas (Fig. 226).

**Natural history.**--- This species is found in grassland, desert-grassland transitional habitats, and Great Basin desert. Very little is known of the natural history of this species given the large number of specimens collected and examined. Plant associations are very few, the only records known are of adults on *Atriplex* sp., *A. canescens* (Pursh) Nutt., (Chenopodiaceae); and *Melilotus* sp. (Leguminosae). No

definite hosts are known. Adults have been collected from January to November, primarily in grassland habitats, at elevations of from 339-3130 m (X=1595 m, N=38) (Fig. 233).

**Chorological relationships.**--- Table 67.

**Phylogenetic relationships.**--- Primary characters show this species to be the sister-species of *C. americanus*, comprising a lineage that is a member of the unresolved heptachotomy also including the *C. infrequens*-*C. puberulus* lineage, *C. canescens*, *C. notolomus*, *C. collaris*, the *C. lobigeninus* species group, and the *C. boucardi* species group (Fig. 236). Secondary characters support monophyly of the *C. americanus*-*C. frontalis* lineage that is then considered the sister-group of the rest of the *C. americanus* species group (Fig. 237).

Table 43. Descriptive statistics for *C. frontalis* based on eight males from Medicine Hat, Alberta

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.52-2.00	1.745	0.231	0.109
WF	1.00-1.28	1.145	0.150	0.071
WRA	0.92-1.20	1.070	0.163	0.077
LP	2.32-3.04	2.710	0.376	0.177
WPB	2.64-3.44	3.030	0.437	0.206
WE1H	3.20-4.16	3.650	0.544	0.256
WE1M	3.52-4.56	4.020	0.634	0.299
LE1	6.40-8.80	7.540	1.279	0.603
LEy	0.80-1.00	0.925	0.130	0.061
WEy	0.52-0.64	0.565	0.075	0.035
B. Proportions.				
WEy/LEy	0.54-0.65	0.612	0.055	0.026
WPB/LP	1.09-1.14	1.118	0.028	0.013
WRA/LR	0.56-0.71	0.614	0.064	0.030
WRA/WF	0.89-0.97	0.934	0.037	0.017
WE1M/LE1	0.50-0.55	0.534	0.029	0.014
LP/LE1	0.35-0.38	0.360	0.017	0.008

Table 44. Descriptive statistics for *C. frontalis* based on eight females from Medicine Hat, Alberta

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.52-2.04	1.785	0.264	0.125
WF	1.00-1.36	1.170	0.183	0.087
WRA	0.96-1.12	1.055	0.078	0.037
LP	2.40-3.12	2.880	0.406	0.191
WPB	2.64-3.60	3.140	0.424	0.200
WE1H	3.20-4.40	3.830	0.542	0.256
WE1M	3.52-4.88	4.220	0.601	0.283
LE1	6.56-8.48	7.740	1.034	0.487
LEy	0.80-1.12	1.000	0.150	0.071
WEy	0.48-0.60	0.570	0.070	0.033
B. Proportions.				
WEy/LEy	0.52-0.60	0.572	0.052	0.025
WPB/LP	1.03-1.16	1.092	0.074	0.035
WRA/LR	0.55-0.65	0.594	0.055	0.026
WRA/WF	0.82-1.00	0.907	0.090	0.042
WE1M/LE1	0.52-0.58	0.545	0.030	0.014
LP/LE1	0.36-0.39	0.372	0.016	0.007

4.5.2.2.3 *Cleonidius canescens* (LeConte)

(Figs. 126, 145, 165, 185, 227)

*Cleonus canescens* LeConte 1876a:151. Lectotype (here designated), male, one of three syntypes, labelled "Col.", "Type/ 5243", "C. canescens/ Lec." and with my designation label "Cleonus/ canescens/ Lec. LECTOTYPE/ desig. Anderson" (MCZC). Type locality, Colorado. Henshaw 1881-1882 (check.). Henshaw 1885 (check.). Wickham 1902 (check.). Fall and Cockerell 1907 (check.). Yothers 1916 (misident. of *Cleonidius texanus*). Leng 1920 (catal.). Essig 1958 (biol., misident. of *Cleonidius texanus*).

*Cleonus* (*Cleonidius*) *canescens*; Casey 1891 (key).

*Cleonus* (*Cleonidius*) *grandirostris* Casey 1891:189. Lectotype (here designated), male, one of five syntypes, labelled "N.M.", "CASEY/ bequest/ 1925", "TYPE USNM/ 37278", "C.C./ *grandirostris*/ Cas." and with my designation label "Cleonidius/ grandirostris/ Cas./ LECTOTYPE/ desig. Anderson" (USNM). Type locality, New Mexico. Csiki 1934 (catal.).

*Cleonus grandirostris*; Wickham 1896 (check.). Fall and Cockerell 1907 (check.). Leng 1920 (catal.).

*Cleonus* (*Cleonidius*) *coloradensis* Csiki 1934:64. Unjustified replacement name for *Cleonus canescens* LeConte.

Blackwelder 1939 (check.).

*Cleonis canescens*; O'Brien and Wibmer 1982 (catal., distn.).

*Cleonis coloradensis*; O'Brien and Wibmer 1982 (catal.,

distn., as synonym).

*Cleonis grandirostris*; O'Brien and Wibmer 1982 (catal.,  
distn., as synonym).

**Notes about synonymy.**--- O'Brien and Wibmer (1982) have previously synonymized *C. canescens* and *C. grandirostris*. *C. coloradensis* is an unjustified replacement name for *C. canescens*.

**Problems in recognition.**--- This species is easily recognized by the less elongate-oval form of the eye (width greater than 0.60 times length) (Fig. 145a), markedly medially tumescent and robust rostrum (Fig. 145a), anterolateral margin of pronotum lacking postocular lobe (Fig. 145a), and short metathoracic wings.

**Description.**--- **Specimens examined.** 182 males, 216 females. Data about variation in LR, WF, WRA, LP, WPB, WE1H, WE1M, LE1, LEy, WEy, WEy/LEy, WPB/LP, WRA/LR, WRA/WF, WE1M/LE1, and LP/LE1 are presented in Tables 45 and 46. **Size.** Length, male, 6.8-8.8 mm; female, 6.9-10.5 mm. Width, male, 2.5-3.7 mm; female, 2.3-4.5 mm. **Head.** Eye more or less oval (width greater than 0.60 times length). Area behind eye with at most only 2 or 3 slight wrinkles of cuticle. Frons with moderately dense, small, shallow punctures, some longitudinally confluent and irregularly impressed.



immediately above eyes, interspersed throughout with minute punctures. Frons also with dense, elongate-narrow, white suberect scales, slightly smaller medially in some specimens, moderately dense, elongate-narrow and erect immediately above eyes; with moderately dense short suberect hair-like scales. Frons continuous with base of rostrum (Fig. 145a). Base of median tumescence or carina of rostrum (if present) with small, shallow fovea. Width of frons greater than, to subequal to, width at apex of rostrum. **Rostrum.** Robust, (width at apex 0.636-0.844 times length in males; 0.640-0.868 in females) (Fig. 145). In lateral view markedly curved downward. In dorsal view with postgenae moderately laterally expanded, antennal scrobes distinctly visible. Rostrum markedly medially tumescent from point of antennal insertion to middle of frons; not laterally expanded at midlength; lateral margins more or less rounded, not sharp; with median carina absent to variously developed as low glabrous shiny line at crest of median tumescence. Dorsal and lateral punctation moderately dense, small, shallow to deep; dorsally interspersed with minute punctures. Dorsally with moderately dense, short, erect hair-like scales (especially laterally); with dense suberect moderately robust scales, smaller and finer apicad of point of antennal insertion. Epistoma moderately transversely swollen at base. **Pronotum.** Median carina lacking to slightly developed in anterior one-half as low, rounded, shiny narrow glabrous line. Dorsal and lateral punctation small to

moderately large, moderately dense, shallow to deep; smaller apically; areas between large dorsal punctures with minute regularly impressed punctures. Dorsally with moderately dense, very short, fine erect hair-like scales, each situated in large puncture (Fig. 145a). Scales dorsally either uniformly moderately dense medially on disk, or absent along median line; moderately dense, moderately robust, appressed in pair of broad apically slightly narrowed paramedian stripes; laterally, absent to sparse, small and fine. Lateral margins with scales moderately dense, moderately robust, appressed. Flanks with scales moderately dense in dorsal one-half, slightly larger in ventral one-half. elongate-fine to elongate-narrow. Median basal area of disk narrowly and moderately deeply impressed. Anterolateral margin with pronotal postocular lobes lacking; postocular vibrissae of unequal length, long, greatest length (approximately two-thirds width of eye) immediately behind base of eye (Fig. 145a). In dorsal view with lateral margins straight to slightly arcuate and slightly convergent from base to apical one-quarter; slightly constricted at apical one-quarter then straight and convergent to apex. Prosternum. With small shallow impression and at most very slight rounded swelling anterior to each procoxal cavity. In lateral view with anterior flat to slightly angulate portion lacking or at most very short and indistinct from posterior more angulate portion. Elytra. Moderately robust in general form (width at midlength 0.563-0.620 times length in males;

0.536-0.622 in females) (Fig. 126). In dorsal view with lateral margins very slightly arcuate from apical one-third to humerus; moderately and evenly arcuate from apical one-third to apex. Elytral apices not produced, subacuminate, not divergent. Humerus rounded, indistinct. Dorsally with moderately dense, short suberect hair-like scales. Scale pattern various; either more or less mottled, with scales moderately dense, small, elongate-narrow to moderately robust on sutural interval and interval 2, moderately dense, large, moderately robust on intervals 3 to 6, moderately dense, large, but with scattered patches of very much smaller scales on intervals 7 to 11 (Fig. 126); or vittate, with scales moderately dense, small and fine on sutural interval and interval 2, moderately dense, uniformly large, moderately robust on intervals 3 to 5, moderately dense, moderately robust, small on intervals 6 and 7, absent to sparse, small, fine on interval 8, uniformly moderately dense, large, moderately robust on intervals 9 to 11. Punctures of elytral striae small, individually distinct; arranged in regular rows. Wings. Short (approximately equal to one-half length of elytra). Legs. Foretibia of female with inner margin with small denticles in apical two-thirds; subapical tooth small, indistinct from tibial denticles. Foretibial and mesotibial unci of both sexes large to moderately large, metatibial uncus small. Tarsal claws connate in basal one-half, not divergent, each with basal internal flange well-developed. Ventral tarsal pilose

vestiture of foretarsus of male present as elongate-narrow pads covering apical one-quarter of article 1, as elongate-oval pads covering apical one-quarter to one-half of article 2, and as large rounded pads covering apical two-thirds of article 3; of foretarsus of female, as apical tufts on article 1, as small rounded pads at ventral apex of article 2, as moderately large rounded pads covering apical one-half to two-thirds of article 3; of mesotarsus of male, as elongate-narrow pads covering apical one-quarter of article 1, as elongate-oval pads covering apical one-third of article 2, as large rounded pads covering apical one-half to two-thirds of article 3; of mesotarsus of female, as apical tufts on article 1, as small rounded pads at ventral apex of article 2, as moderately large rounded pads covering apical one-half to two-thirds of article 3; of metatarsus of male, as apical tufts on article 1, as small elongate pads at ventral apex of article 2, as large rounded pads covering apical one-half to two-thirds of article 3; of metatarsus of female, as apical tufts on article 1, as small elongate pads at ventral apex of article 2, as moderately large elongate pads covering apical one-half to two-thirds of article 3.

**Abdomen.** Ventral surface with moderately dense to dense, elongate-narrow appressed white scales, sparser and smaller medially on abdominal sterna V to VII. Abdominal sterna III to VI (especially III and IV) with individually indistinct to distinct small to large rounded glabrous patches, each with large puncture and single suberect moderately long

hair-like scale situated in or near the center. Base of abdominal sternum VII of female internally (dorsally) with at most very slightly-developed median shiny tubercle.

**Genitalia.** Female (four examined). Abdominal sternum VIII with basal arm long; lateral arms markedly inwardly arcuate at midlength, apical one-half of each arm laterally expanded (Fig. 165). Stylus moderately large compared to length gonocoxite II. Male (four examined). Internal sac (Figs. 185a,b) elongate and low; median dorsal pocket low, with single moderately broad elongate dorsally directed basal lobe (lobe A), with moderately large median dorsally directed lobe at midlength on dorsal surface (lobe B), with moderately large paired dorsolaterally directed lobe near midlength near dorsal margin (lobe D), with large paired dorsolaterally directed lobe just beyond midlength at dorsal margin (lobe E), and with small ventrally directed paired lobe near base near ventral margin (lobe F); lobe C absent. Apical pocket not distinct from median dorsal pocket. Ventral median pocket moderately large. Apical pocket with paired moderately large ventrally directed lobe near apical one-third at ventral margin. Eversible apical sclerite complex with paired simple scythe-like sclerite; adjacent ventral surface of apical pocket with pair of large, dark, longitudinal sclerites.

**Geographic variation.**--- There is geographic variation in the elytral and abdominal scale patterns and in depth and

size of head, rostral and pronotal punctation. Specimens from the western part of the species range in Arizona, Utah and Idaho have an elytral scale pattern that is of the vittate form. The ventral surface of the abdomen of these individuals has only small indistinct glabrous patches, and head, rostral and pronotal punctures are small and shallow. Specimens from the rest of the species range to the east have an elytral scale pattern that is of the more or less mottled form (Fig. 126), ventral abdominal glabrous patches are large and distinct, and punctation of the head, rostrum and pronotum is slightly larger and deeper.

The rostrum is also more elongate-narrow in the western form than in the eastern form based on measurements of rostral width at apex compared to length of rostrum in both male and female individuals from populations at 6 miles SE. Turkey, Texas; 19 miles SW. Kayenta, Arizona; and Denver, Colorado. Individuals from the population at Denver have the most robust rostra (males,  $X=0.805$   $N=8$ ; females,  $X=0.824$   $N=8$ ), those from 6 mi. SE. Turkey have an intermediate sized rostrum (males,  $X=0.780$   $N=8$ ; females,  $X=0.751$   $N=7$ ), and those from 19 mi. SW. Kayenta have the least robust rostrum (males,  $X=0.673$   $N=12$ ; females,  $X=0.669$   $N=6$ ).

Patterns of variation in all characters in this species parallel those in *A. angularis* wherein gene flow appears to be occurring across the continental divide in southern New Mexico and Arizona, but not to the north in Colorado and Utah.

**Geographic distribution.**--- This species is distributed in the southcentral United States of America from extreme southeastern Idaho east to Nebraska, south in the west to southern Arizona, east to extreme southern Texas (Fig. 227). There is a questionable record from Lassen National Park in northern California.

**Natural history.**--- This appears to be a species of low to high elevation grassland, desert-grassland transitional, Texas semi-desert, and Great Basin desert habitats. Plant associations are few. Adults have been collected only on *Symphoricarpos* sp. (Caprifoliaceae); *Atriplex canescens* (Pursh) Nutt. (Chenopodiaceae); *Medicago sativa* L. (alfalfa) (Leguminosae); and *Mentzelia* sp. (Loasaceae). Definite hosts are not known. Specimens have been collected in all months of the year at elevations of 200-2868 m (X=1678 m, N=56) (Fig. 233).

**Chorological relationships.**--- Table 67.

**Phylogenetic relationships.**--- Primary characters show this species to be a member of the unresolved heptachotomy also including the *C. frontalis*-*C. americanus* lineage, the *C. infrequens*-*C. puberulus* lineage, *C. notolomus*, *C. collaris*, the *C. lobigerinus* species group, and the *C. boucardi* species group (Fig. 236). Secondary characters place the *C. notolomus*-*C. collaris*-*C. infrequens*-*C. puberulus* lineage as

the sister group of this species (Fig. 237).



Table 45. Descriptive statistics for *C. canescens* based on eight males from Denver, Colorado

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.48-2.00	1.860	0.246	0.116
WF	1.28-1.64	1.510	0.156	0.074
WRA	1.24-1.60	0.495	0.163	0.077
LP	2.24-3.20	2.820	0.419	0.198
WPB	2.64-3.76	3.370	0.507	0.239
WE1H	3.12-4.24	3.870	0.497	0.234
WE1M	3.28-4.64	4.190	0.608	0.287
LE1	5.60-7.76	7.100	1.086	0.512
LEy	0.80-1.00	0.965	0.104	0.049
WEy	0.56-0.72	0.645	0.075	0.035
B. Proportions.				
WEy/LEy	0.63-0.72	0.669	0.049	0.023
WPB/LP	1.16-1.25	1.195	0.046	0.022
WRA/LR	0.77-0.84	0.805	0.037	0.017
WRA/WF	0.97-1.03	0.990	0.037	0.017
WE1M/LE1	0.57-0.61	0.591	0.022	0.010
LP/LE1	0.38-0.42	0.398	0.023	0.011

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.52-2.04	1.880	0.276	0.130
WF	1.28-1.72	1.565	0.199	0.094
WRA	1.32-1.72	1.545	0.195	0.092
LP	2.40-3.12	2.880	0.357	0.168
WPB	2.72-3.84	3.390	0.517	0.244
WE1H	3.20-4.40	3.940	0.614	0.290
WE1M	3.36-4.88	4.300	0.740	0.348
LE1	5.76-8.00	7.220	1.220	0.575
LEy	0.88-1.08	0.985	0.124	0.058
WEy	0.56-0.72	0.660	0.085	0.040
B. Proportions.				
WEy/LEy	0.62-0.73	0.671	0.061	0.029
WPB/LP	1.13-1.23	1.176	0.052	0.025
WRA/LR	0.76-0.87	0.824	0.054	0.025
WRA/WF	0.95-1.03	0.988	0.048	0.022
WE1M/LE1	0.57-0.62	0.596	0.028	0.013
LP/LE1	0.38-0.43	0.400	0.026	0.012

Type material.--- Holotype, male, with a red label "HOLOTYPE", "Scott City Kan./5-29-59/H. Willis", "HOLOTYPE/*Cleonidius/ Infrequens/ Anderson*" and with abdomen on card and genitalia in microvial attached to pin (CWOB). Allotype, female, with a red label "ALLOTYPE", "TEX. Muleshoe/Bailey Co./16.IV.1971/G.B. Marshall, "ALLOTYPE/*Cleonidius/ Infrequens/ Anderson*" and with abdomen on card and genitalia in microvial attached to pin (CWOB). Type locality, Scott City, Kansas.

Paratypes. 1 male, 5 females. MEXICO: San Luis Potosi: San Luis Potosi, 17.VI.68, Arsego, 1F (USNM).

UNITED STATES OF AMERICA: New Mexico: Fort Sumner, 19.VIII.51, Robinson, 1M (CWOB); 15mi. n. Las Cruces, 24.VII.79, Richmond, 1F (NMSU). Texas: 4mi. s. Big Springs, 3.VI.72, C.W. O'Brien, 1F (CWOB); D'Hanis, 23.II.63, Fohn, 1F (TAMU); Van Horn, 24.VI.42, Scullen, 1F (OSUC).

Derivation of specific epithet.--- From the Latin "*in*" meaning not and "*frequens*" meaning frequent. This name is used in reference to the rarity of individuals of this species.

Problems in recognition.--- Individuals of this species are likely to be confused only with those of *C. puberulus* and perhaps also *C. notolomus*. From *C. notolomus*, specimens are

differently impressed pronotum. Characters allowing for separation of *C. infrequens* from *C. puberulus* are discussed under "Problems in recognition" section for that species.

**Description.**--- Data about variation in LR, WF, WRA, LP, WPB, WE1H, WE1M, LE1, LEy, WEy, WEy/LEy, WPB/LP, WRA/LR, WRA/WF, WE1M/LE1, and LP/LE1 are presented in Tables 47 and 48. **Size.** Length, male, 8.3-8.8 mm; female, 8.2-10.0 mm. Width, male, 3.1-3.3 mm; female, 3.0-3.8 mm. **Head.** Eye elongate-oval. Area behind eye with numerous slight wrinkles of cuticle. Frons with sparse, small, shallow punctures, some longitudinally confluent and irregularly impressed immediately above eyes, interspersed throughout with minute punctures. Frons also with moderately dense to dense, elongate-fine to elongate-narrow, white appressed scales medially, suberect to erect immediately above eyes; otherwise lacking suberect or erect vestiture. Frons distinctly separated from base of rostrum by deep transverse impression (Fig. 146a). Base of median carina of rostrum with small, shallow fovea. Width of frons less than to subequal to width at apex of rostrum. **Rostrum.** Robust, (width at apex 0.780-0.791 times length in males; 0.745-0.917 in females) (Fig. 146). In lateral view straight to very slightly curved downward. In dorsal view with postgenae slightly laterally expanded, antennal scrobes

laterally expanded at midlength; lateral margins sharp, rostrum flat from immediately laterad of median carina to lateral margins; with median carina present, very distinct, as sharp, elevated, glabrous shiny line at crest of median tumescence; also with indistinct low lateral carina at lateral margin in most specimens (Fig. 146b). Dorsal and lateral punctation moderately dense, moderately large to large, moderately deep and longitudinally confluent (especially so immediately laterad of median carina); dorsally interspersed with sparse minute punctures. Dorsally lacking suberect or erect vestiture; with sparse to moderately dense elongate-narrow to robust appressed white scales. Epistoma transversely swollen at base. Pronotum. Median carina lacking to variously developed in anterior one-half as very low, indistinct, broad glabrous line. Dorsal and lateral punctation small to moderately large, moderately dense to dense, moderately deep, not confluent; smaller sparser and shallower apically; areas between large dorsal punctures with minute regularly impressed punctures; punctation sparser and shallower on flanks. Dorsally with moderately dense, short, suberect hair-like scales, each situated in large puncture (Fig. 146a). Scales dorsally absent to sparse, small and fine along median line: dense, elongate-fine to elongate-narrow, appressed in pair of apically slightly narrowed paramedian stripes; laterally,

stripe. Flanks with scales moderately dense to dense, elongate-fine to elongate-narrow. Median basal area of disk broadly and moderately deeply impressed. Anterolateral margin with pronotal postocular lobes well-developed; postocular vibrissae of unequal length, short, greatest length immediately behind base of eye (Fig. 146a). In dorsal view with lateral margins straight and subparallel to slightly and evenly arcuate from base to apical one-third to one-quarter; constricted at apical one-third to one-quarter then straight and convergent to apex, Prosternum. With small shallow impression and very slightly to moderately developed rounded swelling anterior to each procoxal cavity. In lateral view with anterior flat to slightly angulate portion long, approximately one-half to two-thirds length of posterior more angulate portion; posterior and anterior portions not or only slightly separated by shallow transverse impression. Elytra. Moderately robust in general form (width at midlength 0.554-0.605 times length in males; 0.522-0.602 in females) (Fig. 127). In dorsal view with lateral margins very slightly sinuate and convergent from apical one-third to humerus; moderately and evenly arcuate from apical one-third to apex. Elytral apices not produced, subacuminate, not divergent. Humerus rounded, indistinct. Dorsally with very sparse, irregularly distributed very short suberect hair-like scales; with sutural interval and

and fine to elongate-fine white scales; stria 1 with irregularly distributed patches of larger and denser white scales; intervals 3 to 5 with uniformly dense, elongate-narrow to moderately robust white scales; intervals 6 to 8 with scales absent to very sparse, very small and fine; stria 6 and 7 with irregularly distributed patches of larger and denser white scales; intervals 9 to 11 with scales uniformly dense, white, elongate-narrow to robust (Fig. 127). Punctures of elytral striae small, individually distinct; arranged in regular rows. Wings. Long (greater than length of elytra [62%, N=5]) to short (approximately equal to one-half length of elytra or less [38%, N=3]). Legs. Foretibia of female with inner margin with small to large denticles in apical three-quarters; distinct subapical tooth present. Foretibial and mesotibial unci of both sexes moderately large, metatibial uncus small. Tarsal claws connate in basal one-half, slightly divergent, each with basal internal flange well-developed. Ventral tarsal pilose vestiture of foretarsus of both sexes present as elongate pad covering apical one-third on article 1, as large rounded pad covering apical one-half on article 2 and apical three quarters on article 3; of mesotarsus of both sexes as elongate pad covering apical one-quarter to one-third on article 1, as large rounded pad covering apical one-half on article 2 and apical three quarters on article 3; and of metatarsus of both sexes as elongate pad covering apical

elongate pad covering apical one-half on article 2 and as elongate-oval pad covering apical one-half to two-thirds on article 3. Abdomen. Ventral surface with moderately dense to dense, elongate-narrow to moderately robust appressed white scales, sparser and smaller medially on abdominal sternum VII of female. Abdominal sterna III to VI (especially III and IV) with individually indistinct to distinct moderately large rounded glabrous patches, each with large puncture and single suberect to erect, short to moderately long hair-like scale situated in or near the center. Base of abdominal sternum VII of female internally (dorsally) with a slightly-developed median shiny tubercle. Genitalia. Female (five examined). Abdominal sternum VIII with basal arm long; lateral arms very slightly inwardly arcuate throughout length, expanded abruptly at apex (Fig. 166). Stylus moderately large compared to length gonocoxite II. Male (two examined). Internal sac (Figs. 186a,b) elongate and low; median dorsal pocket low, with single moderately broad elongate dorsally directed basal lobe (lobe A), with small median dorsally directed lobe at midlength on dorsal surface (lobe B), with moderately large paired dorsally directed lobe at basal one-third on dorsal surface (lobe C), with moderately large paired dorsolaterally directed lobe near midlength near dorsal margin (lobe D), with large paired dorsolaterally directed lobe just beyond midlength at dorsal margin (lobe E), and with small ventrally directed paired



and median dorsal pocket individually distinct; apex of dorsal median pocket rounded in dorsal view. Ventral median pocket moderately large. Apical pocket with paired moderately large ventrally directed lobe near apical one-third at ventral margin. Eversible apical sclerite complex with paired simple scythe-like sclerite; adjacent ventral surface and basal portion of sides of apical pocket with pair of large, dark, longitudinal sclerites.

Geographic variation.--- None noted.

Geographic distribution.--- Very few individuals of this species are known from western Kansas, southern New Mexico, and western and central Texas, south to San Luis Potosi, Mexico (Fig. 222).

Natural history.--- No definite hosts or host plant associations are known for this species. Adults have been collected in grassland and desert-grassland transitional habitats from February to August at elevations of 873-2242 m ( $\bar{X}$ =1418 m, N= 7) (Fig. 233).

Chorological relationships.--- Table 67.

Phylogenetic relationships.--- Primary characters show this species to be the sister-species of *C. puberulus*. This

including the *C. frontalis*-*C. americanus* lineage, the *C. infrequens*-*C. puberulus* lineage, *C. notolomus*, *C. collaris*, the *C. lobigerinus* species group, and the *C. boucardi* species group (Fig. 236). Secondary characters place the *C. notolomus*-*C. collaris* lineage as the sister group of this lineage (Fig. 237).

based on two males from throughout the species range

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.64-1.72	1.680	-	-
WF	1.24-1.24	1.240	-	-
WRA	1.28-1.36	1.320	-	-
LP	2.96-3.20	3.080	-	-
WPB	2.96-3.44	3.200	-	-
WE1H	3.52-3.92	3.720	-	-
WE1M	4.08-4.16	4.120	-	-
LE1	6.88-7.36	7.120	-	-
LEy	0.88-1.00	0.940	-	-
WEy	0.52-0.60	0.560	-	-
B. Proportions.				
WEy/LEy	0.59-0.60	0.596	-	-
WPB/LP	1.00-1.08	1.038	-	-
WRA/LR	0.78-0.79	0.786	-	-
WRA/WF	1.03-1.10	1.065	-	-
WE1M/LE1	0.55-0.61	0.580	-	-
LP/LE1	0.43-0.44	0.433	-	-

based on six females from throughout the species range

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.44-1.88	1.667	0.239	0.130
WF	1.24-1.40	1.327	0.080	0.043
WRA	1.32-1.44	1.373	0.090	0.049
LP	3.04-3.52	3.267	0.308	0.167
WPB	3.12-3.60	3.373	0.278	0.151
WE1H	3.60-4.32	4.040	0.460	0.251
WE1M	3.76-4.72	4.400	0.563	0.307
LE1	7.20-8.48	7.840	0.759	0.413
LEy	0.96-1.16	1.067	0.105	0.057
WEy	0.52-0.60	0.573	0.049	0.027
B. Proportions.				
WEy/LEy	0.52-0.58	0.538	0.032	0.018
WPB/LP	1.00-1.05	1.033	0.031	0.017
WRA/LR	0.75-0.92	0.828	0.088	0.048
WRA/WF	1.00-1.09	1.036	0.061	0.033
WE1M/LE1	0.52-0.60	0.561	0.046	0.025
LP/LE1	0.40-0.45	0.417	0.027	0.015

4.3.2.2.5 *Cleonidius puberulus* (LeConte)

(Figs. 115, 128, 147, 167, 187, 225)

*Cleonus puberulus* LeConte 1876a:151. Lectotype (here designated), female, one of seven syntypes, labelled "N", "Type 5242", "*C. puberulus*/ Lec." and with my designation label "*Cleonus/ puberulus*/ LeC. LECTOTYPE/ desig. Anderson" (MCZC). Type locality, Nevada. Henshaw 1881-1882 (check.). Henshaw 1885 (check.). Leng 1920 (catal., misinterp. as synonym *Cleonidius frontalis*).

*Cleonus (Cleonidius) puberulus*; Casey 1891 (misinterp. as synonym *Cleonidius frontalis*). Csiki 1934 (catal., misinterp. as synonym *Cleonidius frontalis*).

*Cleonus (Cleonidius) lobigerinus*; Champion 1902-1906 (in part, distn., misident.).

*Cleonis puberulus*; O'Brien and Wibmer 1982 (catal., distn., misinterp. as synonym *Cleonidius frontalis*).

**Notes about synonymy.**--- The type locality for this species, given as Nevada (LeConte 1876a), lies outside the geographic range as indicated by specimens I have examined. Perhaps LeConte (1876a) was in error in that the type locality may have been Nebraska, not Nevada (presumably an interpretation made by LeConte based on the "N" on the specimen labels).

**Problems in recognition.**--- Individuals of this species are likely to be confused only with those of *C. infrequens*.

Members of both species possess a very broad rostrum not found in other *Cleonidius* species (Figs. 146-147). In *C. Infrequens* however, the median rostral carina is sharply elevated and distinct, the lateral margins of the rostrum are sharp, the dorsal surface of the rostrum flat from immediately laterad of the median carina to the lateral margins, the rostral punctures are deep and large, there are very slight low lateral rostral carinae, ventral pilose tarsal vestiture is moderately extensive, and both macropterous and brachypterous forms are known. All of these character states differ in *C. puberulus* individuals. Geographic distributions of the two species (Figs. 222, 225) are allopatric although there may be sympatry in Colorado or Kansas.

Specimens of *C. puberulus* may also be confused with small *C. lobigerinus*, *C. notolomus*, and *C. frontalis* but can be distinguished by their more robust rostrum (Fig. 147) and lack of the various diagnostic character states of these other species.

**Description.**--- Specimens examined. 10 males, 26 females. Data about variation in LR, WF, WRA, LP, WPB, WE1H, WE1M, LE1, LEy, WEy, WEy/LEy, WPB/LP, WRA/LR, WRA/WF, WE1M/LE1, and LP/LE1 are presented in Tables 49 and 50. Size. Length, male, 5.1-8.3 mm; female, 6.4-8.4 mm. Width, male, 2.1-3.1 mm; female, 2.6-3.3 mm. Head. Eye elongate-oval. Area behind eye with numerous slight wrinkles of cuticle. Frons with

sparse, small, shallow punctures; some longitudinally confluent and irregularly impressed immediately above eyes, interspersed throughout with minute punctures. Frons also with moderately dense to dense, elongate-fine to elongate-narrow, white appressed scales medially/ suberect to erect immediately above eyes; otherwise lacking suberect or erect vestiture. Frons separated from base of rostrum by shallow to moderately-deep transverse impression (Fig. 147a). Base of median carina of rostrum with small, shallow fovea. Width of frons greater than to subequal to width at apex of rostrum. Rostrum. Robust, (width at apex 0.707-0.865 times length in males; 0.744-0.818 in females) (Fig. 147). In lateral view very slightly curved downward. In dorsal view with postgenae slightly laterally expanded, antennal scrobes partially visible. Rostrum slightly medially tumescent from point of antennal insertion to middle of frons; not laterally expanded at midlength; lateral margins rounded, rostrum declivous from immediately laterad of median carina to lateral margins; with median carina present as variously developed slightly elevated, narrow line, more or less obscured in most specimens by overlying scales. Dorsal and lateral punctation moderately dense, small to moderately large, shallow, not longitudinally confluent; dorsally interspersed with dense minute punctures. Dorsally lacking suberect or erect vestiture to with sparse short suberect hair-like scales; moderately dense to dense, elongate-fine to robust appressed white scales. Epistoma

glabrous line. Dorsal and lateral punctation small, moderately dense to dense, shallow, not confluent; smaller sparser and shallower apically; areas between large dorsal punctures with minute regularly impressed punctures; punctation sparser and shallower on flanks. Dorsally with sparse to moderately dense, very short, suberect hair-like scales, each situated in large puncture (Fig. 147a). Scales dorsally absent to sparse, small and fine along median line; dense, elongate-fine to elongate-narrow, appressed in pair of apically slightly narrowed paramedian stripes; laterally, absent to sparse, small and fine. Lateral margins with scales dense, elongate-narrow, appressed in moderately broad stripe. Flanks with scales moderately dense to dense, elongate-fine to elongate-narrow. Median basal area of disk broadly and moderately deeply impressed. Anterolateral margin with pronotal postocular lobes well-developed; postocular vibrissae of unequal length, short, greatest length immediately behind base of eye (Fig. 147a). In dorsal view with lateral margins straight and subparallel to slightly and evenly arcuate from base to apical one-third to one-quarter; constricted at apical one-third to one-quarter then straight and convergent to apex. Prosternum. With small shallow impression and a very slightly developed rounded swelling anterior to each procoxal cavity. In lateral view with anterior flat to slightly angulate portion long,



separated by shallow to moderately deep transverse impression. Elytra. Moderately robust in general form (width at midlength 0.531-0.611 times length in males; 0.543-0.597 in females) (Fig. 128). In dorsal view with lateral margins very slightly sinuate and convergent from apical one-third to humerus; moderately and evenly arcuate from apical one-third to apex. Elytral apices not produced, rounded to subacuminate, not to very slightly divergent. Humerus rounded, indistinct. Dorsally with sparse to moderately dense, very short suberect hair-like scales; with sutural interval and interval 2 with very sparse to moderately dense, very small and fine to elongate-fine white scales; stria 1 with irregularly distributed patches of larger and denser white scales; intervals 3 to 5 with uniformly dense, elongate-narrow to moderately robust white scales; intervals 6 to 8 with scales absent to dense, very small and fine to moderately robust; stria 6 and 7 with irregularly distributed patches of larger and denser white scales in those specimens with scales of intervals 6 to 8 small and fine; intervals 9 to 11 with scales uniformly dense, white, elongate-narrow to robust (Fig. 148). Punctures of elytral striae small, individually distinct; arranged in regular rows. Wings. Very short (approximately one-quarter to one-third length of elytra). Legs. Foretibia of female with inner margin with small denticles in apical three-quarters;

uncus small. Tarsal claws connate in basal one-quarter to one-third, slightly divergent, each with basal internal flange moderately to well-developed. Ventral tarsal pilose vestiture of foretarsus of male present as small elongate pad covering apical one-quarter to one-third on article 1, as small rounded pad covering apical one-half on article 2, and as large round pad covering apical two-thirds on article 3; of female as apical tuft on article 1, as small rounded pad covering apical one-quarter on article 2, and as large rounded pad covering apical one-half on article 3; of mesotarsus of male as apical tuft to small elongate pad covering apical one-quarter on article 1, as apical tuft to small elongate pad covering apical one-third on article 2, and as small rounded pad covering apical one-third on article 3; of female as apical tuft on articles 1 and 2, and as small rounded pad covering apical one-third on article 3; of metatarsus of male lacking from, to as small elongate pad covering apical one-quarter on article 1, as apical tuft to small elongate pad covering apical one-third on article 2, and as small elongate pad covering apical one-third on article 3; of female lacking from, to as apical tuft on article 1, as apical tuft on article 2, and a small elongate pad covering apical one-quarter on article 3. **Abdomen.**

Ventral surface with moderately dense to dense, elongate-narrow to moderately robust appressed white scales,

female. Abdominal sternum III to VI (especially III and IV) with individually indistinct to distinct moderately large rounded glabrous patches, each with large puncture and single suberect, short hair-like scale situated in or near the center. Base of abdominal sternum VII of female internally (dorsally) with slightly to moderately developed median shiny tubercle. Genitalia. Female (four examined). Abdominal sternum VIII with basal arm moderately long; lateral arms slightly inwardly arcuate at mid-length, very slightly expanded at apex (Fig. 167). Stylus moderately large compared to length gonocoxite II. Male (three examined). Internal sac (Figs. 187a,b) elongate and low; median dorsal pocket low; with single moderately broad elongate dorsally directed basal lobe (lobe A), with small median dorsally directed lobe at midlength on dorsal surface (lobe B), with moderately large paired dorsally directed lobe at basal one-third on dorsal surface (lobe C), with moderately large paired dorsolaterally directed lobe near midlength near dorsal margin (lobe D), with large paired dorsolaterally directed lobe just beyond midlength at dorsal margin (lobe E), and with small ventrally directed paired lobe near base near ventral margin (lobe F). Apical pocket and median dorsal pocket individually distinct; apex of dorsal median pocket rounded in dorsal view. Ventral median pocket moderately large. Apical pocket with paired moderately large ventrally directed lobe near apical

complex with paired simple scythe-like sclerite; adjacent ventral surface of sides of apical pocket with pair of large, dark, longitudinal sclerites.

**Geographic variation.**--- None noted.

**Geographic distribution.**--- This species is known only from extreme southern Alberta and Manitoba, south to southern Wyoming and Colorado (Fig. 225).

**Natural history.**--- A single adult has been collected on *Lesquerella ludoviciana* (Nutt.) Wats. (Cruciferae) in Kansas. Definite hosts are not known. Adults have been collected from April to October in grassland habitats at elevations of 1369-2607 m (X=2186 m, N=4) (Fig. 233).

**Chorological relationships.**--- Table 67. This species is allopatric with its sister-species *C. infrequens*.

**Phylogenetic relationships.**--- Primary characters show this species to be the sister-species of *C. infrequens*. This lineage is then a member of the unresolved heptachotomy also including the *C. frontalis*-*C. americanus* lineage, the *C. infrequens*-*C. puberulus* lineage, *C. notolomus*, *C. collaris*, the *C. jobligerinus* species group, and the *C. boucardi* species group (Fig. 236). Secondary characters place the *C.*

lineage (Fig. 237).

on six males from Colorado Springs, Colorado

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.44-1.64	1.513	0.128	0.070
WF	1.24-1.40	1.313	0.110	0.060
WRA	1.16-1.28	1.213	0.082	0.045
LP	2.48-3.04	2.627	0.317	0.173
WPB	2.64-2.96	2.760	0.197	0.107
WE1H	3.04-3.44	3.200	0.201	0.109
WE1M	3.36-3.60	3.493	0.145	0.079
LE1	5.76-6.48	6.067	0.425	0.232
LEy	0.84-1.04	0.900	0.106	0.058
WEy	0.52-0.56	0.533	0.031	0.017
B. Proportions.				
WEy/LEy	0.50-0.67	0.600	0.085	0.046
WPB/LP	0.97-1.13	1.054	0.086	0.047
WRA/LR	0.71-0.87	0.804	0.083	0.045
WRA/WF	0.86-1.00	0.926	0.080	0.044
WE1M/LE1	0.53-0.61	0.577	0.039	0.021
LP/LE1	0.40-0.47	0.433	0.033	0.018

on 10 females from Colorado Springs, Colorado

Character	Range	Mean	1.5SD	2SE
-----------	-------	------	-------	-----

A. Measurements, in mm.

LR	1.28-1.76	1.548	0.255	0.107
WF	1.20-1.48	1.356	0.161	0.068
WRA	1.00-1.36	1.196	0.178	0.075
LP	2.24-3.04	2.648	0.454	0.192
WPB	2.24-3.36	2.792	0.520	0.219
WE1H	2.64-3.84	3.280	0.640	0.270
WE1M	2.96-4.08	3.608	0.597	0.252
LE1	4.96-7.52	6.328	1.253	0.529
LEy	0.76-1.04	0.916	0.159	0.067
WEy	0.48-0.60	0.552	0.079	0.033

B. Proportions.

WEy/LEy	0.58-0.63	0.604	0.035	0.015
WPB/LP	0.97-1.11	1.054	0.067	0.028
WRA/LR	0.74-0.82	0.774	0.037	0.015
WRA/WF	0.83-0.94	0.881	0.050	0.021
WE1M/LE1	0.54-0.60	0.572	0.031	0.013
LP/LE1	0.40-0.45	0.420	0.021	0.009

(Figs. 129, 148, 168, 188, 223)

*Cleonus collaris* LeConte 1876a:149. Holotype (examined), male, labelled "Col", "*Cleonus/ collaris/* TYPE LeC.", "Horn Coll/ H 8517" and with red square label (MCZC). Type locality, Colorado. Henshaw 1881-1882 (check.). Henshaw 1885 (check.). Wickham 1902 (check.). Leng 1920 (catal.).

*Cleonus (Cleonidius) collaris*; Casey 1891 (key). Csiki 1934 (catal.).

*Cleonis collaris*; O'Brien and Wibmer 1982 (catal., distn.).

**Problems in recognition.**--- The markedly irregularly elevated and sculptured pronotum with only sparse shallow and indistinct large punctures is characteristic of members this species and should serve for reliable identification. The pronotum of some specimens of *C. notolomus* is similarly broadly and deeply medially impressed at the base and has shallow longitudinal impressions but is otherwise not as irregularly sculptured as that of *C. collaris*; it also has distinct moderately large, moderately deep dorsal punctation.

**Description.**--- **Specimens examined.** 7 males, 6 females. Data about variation in LR, WF, WRA, LP, WPB, WELH, WELM, LEL, LEY, WEY, WEY/LEY, WPB/LP, WRA/LR, WRA/WF, WELM/LEL, and



male, 8.4-10.4 mm; female, 9.3-10.9 mm. Width, male, 3.3-4.4 mm; female, 4.0-4.6 mm. Head. Eye elongate-oval. Area behind eye with numerous slight wrinkles of cuticle. Frons with only dense minute punctures. Frons also with dense, moderately robust, pale brownish appressed scales medially, suberect to erect immediately above eyes; otherwise lacking suberect or erect vestiture. Frons very slightly separated from base of rostrum by very shallow transverse impression (Fig. 148a). Width of frons greater than width at apex of rostrum. Rostrum. Moderately robust (width at apex 0.612-0.725 times length in male; 0.607-0.690 in female) (Fig. 148). In lateral view very slightly curved downward. In dorsal view with postgenae slightly to moderately laterally expanded, antennal scrobes partially visible. Rostrum slightly medially tumescent from point of antennal insertion to middle of frons (more markedly so basally); not laterally expanded at midlength; lateral margins sharp; with median carina present as variously developed slightly elevated, glabrous, shiny narrow line at crest of median tumescence. Dorsal and lateral punctation sparse, small, shallow, not longitudinally confluent; dorsally interspersed with dense minute punctures. Dorsally lacking suberect or erect vestiture; with dense moderately robust to robust appressed white to pale brown scales. Epistoma moderately transversely swollen at base. Pronotum. Extremely irregularly elevated, markedly constricted dorsolaterally at apical

one quarter, deeply and broadly impressed medially at base and longitudinally along lateral margins. Median carina variously developed in anterior one-half from absent to as low, rounded moderately broad glabrous line. Dorsal and lateral punctation and that of flanks small, very sparse, shallow and indistinct; areas between larger dorsal punctures with dense minute regularly impressed punctures. Dorsolaterally with sparse, short, suberect hair-like scales, each situated in large puncture; medially lacking suberect or erect vestiture (Fig. 148a). Scales dorsally absent to sparse, moderately robust along median line; very dense, moderately robust, appressed in pair of broad apically slightly narrowed paramedian stripes; laterally, absent to sparse, small and elongate-fine. Lateral margins with scales very dense, moderately robust, appressed to recumbent. Flanks with scales dense to very dense, moderately robust, appressed. Anterolateral margin with pronotal postocular lobes well-developed; postocular vibrissae of unequal length, moderately long, greatest length (approximately one-half width of eye) immediately behind base of eye (Fig. 148a). In dorsal view with lateral margins straight, sinuate or slightly arcuate from base to apical one-quarter; markedly constricted at apical one-quarter then straight and convergent to apex.

**Prosternum.** With small shallow impression and slightly developed rounded swelling anterior to each procoxal cavity. In lateral view with anterior flat to slightly angulate

portion long, approximately one-half to two-thirds length of posterior more angulate portion; posterior and anterior portions slightly to markedly separated by shallow to moderately deep transverse impression. Elytra. Moderately robust in general form (width at midlength 0.589-0.644 times length in males; 0.596-0.645 in females) (Fig. 129). In dorsal view with lateral margins very slightly and evenly arcuate, sinuate at basal one-quarter and convergent from apical one-third to humerus; moderately and evenly arcuate from apical one-third to apex. Elytral apices not produced, rounded, not to very slightly divergent. Humerus distinct. Dorsally lacking suberect or erect vestiture; with uniformly dense to very dense, moderately robust appressed white to pale brown scales, absent to sparse and small in scattered patches on striae 1 and 2, and intervals 9 to 11 (Fig. 129). Punctures of elytral striae individually indistinct, obscured by overlying scales. Wings. Short (approximately one-half length of elytra). Legs. Foretibia of female with inner margin with small denticles in apical one-half; no distinct subapical tooth present. Foretibial and mesotibial unci of both sexes moderately large, metatibial unci small. Tarsal claws connate in basal one-third, not divergent, each with basal internal flange slightly developed. Ventral tarsal pilose vestiture of foretarsus of male present as elongate pad covering apical one-third on article 1, as large rounded pad covering apical one-half on article 2, and as large round pad covering apical three-quarters on article

3; of female as small elongate pad at apex on article 1, as small rounded pad at apex on article 2, and as large round pad covering apical one-half on article 3; of mesotarsus of male as small elongate pad covering apical one-quarter to one-third on article 1, as moderately large elongate pad covering apical one-third to one-half on article 2, and as large elongate-oval pad covering apical two-thirds on article 3; of female as small elongate pad at apex on articles 1 and 2, and as large rounded pad covering apical one-half on article 3; of metatarsus of male as small elongate tuft at apex to covering apical one-quarter on article 1, as small elongate pad covering apical one-third to one-half on article 2, and as large elongate-oval pad covering apical one-half on article 3; of female lacking from, to as apical tuft on article 1, as apical tuft on article 2, and as elongate-oval pad covering apical one-half on article 3. **Abdomen.** Ventral surface with moderately dense to dense, moderately robust appressed white to pale-brown scales, sparser and smaller medially on abdominal sterna V to VII. Abdominal sterna III to VI (especially III and IV laterally) with individually distinct large rounded glabrous patches, each with large puncture and single recumbent to suberect, moderately long hair-like scale situated in or near the center. Base of abdominal sternum VII of female internally (dorsally) with markedly developed median shiny tubercle. **Genitalia.** Female (two examined). Abdominal sternum VIII with basal arm moderately long; lateral arms

slightly inwardly arcuate throughout length, very slightly expanded at apex (Fig. 168). Stylus moderately large compared to length gonocoxite II. Male (four examined). Internal sac (Figs. 188a,b) elongate and low; median dorsal pocket low, with single moderately broad elongate dorsally directed basal lobe (lobe A), with small median dorsally directed lobe at midlength on dorsal surface (lobe B), with moderately large paired dorsally directed lobe at basal one-third on dorsal surface (lobe C), with small paired dorsolaterally directed lobe near midlength near dorsal margin (lobe D), with large paired dorsolaterally directed lobe just beyond midlength at dorsal margin (lobe E), and with small ventrally directed paired lobe near base near ventral margin (lobe F). Apical pocket and median dorsal pocket individually distinct; apex of dorsal median pocket truncate in dorsal view. Ventral median pocket moderately large. Apical pocket with paired moderately large ventrally directed lobe near apical one-third at ventral margin. Eversible apical sclerite complex with paired simple scythe-like sclerite; adjacent ventral surface of sides of apical pocket with pair of large, dark, longitudinal sclerites.

**Geographic variation.**--- None noted.

**Geographic distribution.**--- This species is known only from Colorado and adjacent southern Wyoming, western Kansas and

Colorado was collected on *Lepidium* sp. (pepper-grass; Cruciferae). The very few adults known have been collected in grassland habitats from March to May at elevations of 1697-2608 m (X=2005 m, N=3) (Fig. 233).

**Chorological relationships.**--- Table 67.

**Phylogenetic relationships.**--- Primary characters show this species to be a member of the unresolved heptachotomy also including the *C. frontalis*-*C. americanus* lineage, the *C. infrequens*-*C. puberulus* lineage, *C. canescens*, *C. notolomus*, the *C. lobigerinus* species group, and the *C. boucardi* species group (Fig. 236). Secondary characters indicate *C. collaris* is the sister-species of *C. notolomus*, these two species forming a lineage that is sister to the *C. infrequens*-*C. puberulus* lineage within the *C. americanus* species group (Fig. 237).

Character	Range	Mean	1.5SD	2SE
-----------	-------	------	-------	-----

A. Measurements, in mm.

LR	1.60-2.20	1.971	0.333	0.168
WF	1.24-1.72	1.486	0.248	0.125
WRA	1.16-1.44	1.280	0.162	0.082
LP	2.72-3.84	3.211	0.633	0.319
WPB	3.04-4.24	3.611	0.673	0.339
WE1H	3.68-5.20	4.446	0.840	0.423
WE1M	4.24-5.60	4.937	0.758	0.382
LE1	6.72-9.12	7.909	1.412	0.712
LEy	0.88-1.24	1.011	0.176	0.089
WEy	0.56-0.72	0.629	0.090	0.045

B. Proportions.

WEy/LEy				
WPB/LP	1.07-1.21	1.126	0.075	0.038
WRA/LR	0.61-0.73	0.652	0.062	0.031
WRA/WF	0.83-0.94	0.864	0.065	0.033
WE1M/LE1	0.59-0.64	0.625	0.028	0.014
LP/LE1	0.40-0.42	0.406	0.017	0.008

Character	Range	Mean	1.5SD	2SE
-----------	-------	------	-------	-----

A. Measurements, in mm.

LR	1.68-2.24	1.940	0.321	0.175
WF	1.32-1.72	1.513	0.219	0.119
WRA	1.16-1.36	1.240	0.126	0.069
LP	2.80-3.68	3.173	0.561	0.306
WPB	3.28-4.16	3.747	0.534	0.291
WE1H	3.84-5.12	4.507	0.798	0.435
WE1M	4.40-5.76	5.080	0.861	0.469
LE1	7.20-8.96	8.133	1.034	0.563
LEy	1.00-1.08	1.053	0.049	0.027
WEy	0.56-0.72	0.633	0.110	0.060

B. Proportions.

WEy/LEy	0.52-0.67	0.601	0.094	0.051
WPB/LP	1.13-1.24	1.183	0.060	0.033
WRA/LR	0.61-0.69	0.642	0.051	0.028
WRA/WF	0.79-0.88	0.822	0.058	0.031
WE1M/LE1	0.60-0.65	0.623	0.028	0.015
LP/LE1	0.37-0.41	0.390	0.025	0.014



*Cleonis frontalis*; O'Brien and Wibmer 1982 (in part; catal.,  
distn., misident.).

Type material.--- Holotype, male, with a red label  
"HOLOTYPE", "Grant Co NM/ X 1932", "RT Kellogg", "HOLOTYPE/  
*Cleonidius/ notolomus/* Anderson" and with abdomen on card  
and genitalia in microvial attached to pin (USNM). Allotype,  
female, with a red label "ALLOTYPE", "New Mex:  
Hidalgo/Co., 1Mi.W.Rodeo/VII-31-1971, J. Doyen & Tschinkel",  
"J.Doyen Lot/71G20", "ALLOTYPE/ *Cleonidius/ notolomus/*  
Anderson" and with abdomen on card and genitalia in  
microvial attached to pin (UCBC). Type locality, Grant Co.,  
New Mexico.

Paratypes. 15 males, 16 females. MEXICO: Durango: 26mi.  
w. Durango, 13.VII.75, L.E. Watrous, 1M (CWOB). Mexico:  
Chapingo, 5.VII.55, Gonzalez, 1F (USNM).

UNITED STATES OF AMERICA: Arizona: Cochise County,  
Guadalupe Canyon, 31.VII.75, S. McCleve, 1F (SMC); 25mi. n.  
Douglas, 10.VII.74, S. McCleve, 1M (SMC); Tucson, XII.27,  
Downe, 1F (UAT); Cochise, 5.VIII.54, F. Werner, 1F (UAT);  
Douglas, 23.VIII.27, Jones, 1M (LACM); Chiricahua Mountains,  
Shake Gulch, 8.VI.68, Menke, 1F (USNM); Sabino Canyon,  
2.VII.52, Kelsey, 1F (UAT); Aguirre Lake (10mi. ne. Sasabe),  
2.VIII.78, Hetz, 1M (UAT); Hereford, 3.III.41, Jones, 1M

Cochise, 25.V.51, Statham, 1M (USNM); Sierra Vista,  
4.VIII.79, A.E. Lewis, 1M (CWOB). New Mexico: Grant County,  
27.IX.35, Kellogg, 1F (CASC); Silver City, VI.33, Kellogg,  
2F (OSUC); White Sands National Monument, Lake Lucero,  
2.VIII.74, D. Chandler, 1M (CWOB); White Sands National  
Monument, R. Lenczy, 1F (USNM), VIII.78, R. Lenczy, 1M  
(USNM); 5mi. n. Carlsbad, 21.IX.56, MacSwain, 1F (UCBC);  
Catron County, Quemado, 24.III.64, Hogg, 1F (NMSU). Texas:  
El Paso, VII, H.F. Wickham, 2M, 4F (USNM); Van Horn,  
24.VI.42, Scullen, 1M (OSUC). Utah: Callao, 8.VI.22,  
Spalding, 1M (USNM).

**Derivation of specific epithet.**--- From the Greek "*notos*"  
meaning south and "*tomus*" meaning border. This name is used  
in reference to the distribution of individuals of this  
species near the southern border of the United States of  
America.

**Problems in recognition.**--- Adult individuals of this  
species are most difficult to separate from those of *C.*  
*frontalis*. Individuals of both sexes of *C. frontalis*  
generally have a proportionately longer pronotum compared to  
elytra (see Fig. 200), a more elongate-narrow elytra (see  
Fig. 197), and more rounded lateral margins of the rostrum.

large denticles along the inner margin of the fore-tibia (Fig. 144c); these denticles are small in all *C. notolomus*. Most *C. frontalis* individuals have the rostrum very slightly laterally swollen at midlength and have the submentum slightly ventrally swollen (Figs. 144a,b); this is not so for *C. notolomus*. All *C. frontalis* are also macropterous, whereas *C. notolomus* are either macropterous or brachypterous. The two species also differ largely in their distributions although there is a zone of sympatry in eastern New Mexico and western Texas.

Individuals of *C. notolomus* may also be confused with those of *C. collaris*, *C. infrequens*, *C. puberulus*, *C. lobigerinus*, and *C. trivittatus*. *Cleonidius infrequens* and *C. puberulus* individuals are distinguished by their more robust rostra (Figs. 146-147); *C. collaris* by the much more markedly irregularly sculptured and shallowly and indistinctly punctate pronotum, and not distinctly vittate scale pattern; *C. trivittatus* by the at most only slightly developed postocular lobes (Fig. 153a); *C. lobigerinus* by the curved sulcus behind the eye and the more elongate erect dorsal vestiture (Fig. 150a).

**Description.**--- Data about variation in LR, WF, WRA, LP, WPB, WE1H, WE1M, LE1, LEy, WEy, WEy/LEy, WPB/LP, WRA/LR, WRA/WF, WE1M/LE1, and LP/LE1 are presented in Tables 53 and

elongate-oval. Area behind eye with numerous slight wrinkles of cuticle. Frons with sparse to moderately dense, small shallow punctures, interspersed with minute punctures. Frons also with moderately dense, elongate-narrow appressed white scales medially, suberect to erect immediately above eyes; otherwise lacking suberect or erect vestiture. Frons slightly separated from base of rostrum by shallow transverse impression (Fig. 149a). Base of median carina of rostrum with small, shallow fovea. Width of frons greater than width at apex of rostrum. Rostrum. Moderately robust, (width at apex 0.604-0.767 times length in male; 0.585-0.680 in female) (Fig. 149). In lateral view slightly curved downward. In dorsal view with postgenae at most slightly laterally expanded, antennal scrobes partially visible. Rostrum slightly medially tumescent from point of antennal insertion to middle of frons; not laterally expanded at midlength; lateral margins sharp; with median carina as narrow low sharp and distinct shiny line at crest of median tumescence. Dorsal and lateral punctation moderately dense to dense, small to moderately large, shallow to moderately deep; smaller and less dense apicad of point of antennal insertion; dorsally interspersed with dense minute punctures. Dorsally (especially laterally) with sparse to moderately dense, very short, suberect hair-like scales; also with moderately dense, elongate-narrow, appressed white

low, rounded moderately broad glabrous line. Dorsal and lateral punctation small to moderately large, moderately dense, moderately deep, not confluent; smaller, sparser and shallower apically and on flanks; areas between larger dorsal punctures with minute regularly impressed punctures. Dorsally with moderately dense, short, suberect hair-like scales, each situated in large puncture (Fig. 149a). Scales dorsally absent along median line; dense, elongate-narrow, appressed in pair of broad apically slightly narrowed paramedian stripes; laterally, absent to sparse, small and elongate-fine. Lateral margins with scales dense, elongate-narrow, appressed to recumbent in moderately broad stripe. Flanks with scales elongate-narrow, slightly less dense than on lateral margins. Median basal portion of disk deeply and broadly impressed; also longitudinally shallowly impressed along lateral margins in most specimens. Anterolateral margin with pronotal postocular lobes well-developed; postocular vibrissae of unequal length, moderately long, greatest length (approximately one-half width of eye) immediately behind base of eye (Fig. 149a). In dorsal view with lateral margins slightly to moderately arcuate and convergent to apex; in some specimens, slightly constricted at apical one-quarter then straight and convergent to apex. Prosternum. With small shallow impression and slightly to moderately developed rounded

approximately one-half to subequal in length to posterior more angulate portion; posterior and anterior portions not or slightly separated by a shallow to moderately deep transverse impression. Elytra. Moderately robust in general form (width at midlength 0.561-0.643 times length in males; 0.576-0.638 in females) (Fig. 130). In dorsal view with lateral margins straight to very slightly sinuate to slightly and evenly arcuate and convergent from apical one-third to humerus; moderately and evenly arcuate from apical one-third to apex. Elytral apices not produced, rounded, very slightly divergent. Humerus distinct to indistinct. Dorsally with moderately dense, short, suberect to erect hair-like scales; with sutural interval and interval 2 with very sparse to moderately dense, very small and fine to elongate-fine, golden to white scales; intervals 3 to 5 with uniformly dense, elongate-narrow to moderately robust white scales; intervals 6 to 8 with scales lacking to very sparse, very small and very fine, golden to white; intervals 9 to 11 with uniformly dense, elongate-narrow to moderately robust white scales; striae 1, 6 and 7 with irregularly distributed patches of larger and denser white scales than on adjacent intervals (Fig. 130). Punctures of elytral striae small, individually distinct, arranged in regular rows. Wings. Long (greater than elytra in length [21%, N=7]), to short (approximately one-half length of

small subapical tooth present, slightly distinct from tibial denticles. Foretibial and mesotibial unci of both sexes moderately large, metatibial uncus small. Tarsal claws connate in basal one-third, slightly divergent, each with basal internal flange well-developed. Ventral tarsal pilose vestiture of foretarsus of male and female present as elongate-narrow pad covering apical one-third to one-half on article 1, as moderately large elongate pad covering apical one-third to one-half on article 2, and as large rounded pad covering apical two-thirds on article 3; of mesotarsus of male and female as small elongate pad at apex on article 1, as small rounded pad covering apical one-quarter to one-third on article 2, and as moderately large rounded pad covering apical one-third to one-half on article 3; of metatarsus of male and female as elongate-narrow inner pad covering apical one-half and as outer apical tuft on article 1, as elongate-narrow pad covering apical one-half on article 2, and as moderately large elongate-oval pad covering apical one-half to two-thirds on article 3.

**Abdomen.** Ventral surface with dense, moderately robust appressed white scales, sparser and smaller medially on abdominal sternum VII of female. Abdominal sterna III to VI (especially III and IV) with individually indistinct to distinct moderately large rounded glabrous patches, each with large puncture and single suberect to erect, short to

(dorsally) with slightly to markedly developed median shiny tubercle. Genitalia. Female (five examined). Abdominal sternum VIII with basal arm moderately long; lateral arms slightly to moderately inwardly arcuate at midlength; slightly expanded at apex (Fig. 169). Stylus moderately large compared to length gonocoxite II. Male (two examined). Internal sac (Figs. 189a,b) elongate and low; median dorsal pocket low, with single moderately broad elongate dorsally directed basal lobe (lobe A), with small median dorsally directed lobe at midlength on dorsal surface (lobe B), with moderately large paired dorsally directed lobe at basal one-third on dorsal surface (lobe C), with small paired dorsolaterally directed lobe near midlength near dorsal margin (lobe D), with large paired dorsolaterally directed lobe just beyond midlength at dorsal margin (lobe E), and with small ventrally directed paired lobe near base near ventral margin (lobe F). Apical pocket and median dorsal pocket individually distinct; apex of dorsal median pocket truncate in dorsal view. Ventral median pocket moderately large. Apical pocket with paired moderately large ventrally directed lobe near apical one-third at ventral margin. Eversible apical sclerite complex with paired simple scythe-like sclerite; adjacent ventral surface and basal portion of sides of apical pocket with pair of large, dark, longitudinal sclerites.



**Geographic distribution.**--- This species is known only from southeastern Arizona, southern New Mexico, and extreme western Texas, south to central Mexico (Fig. 225).

**Natural history.**--- This species appears to be associated with desert-grassland transitional habitats. Very little is known of the natural history of this species. There is only a single record of one adult on *Atriplex canescens* (Pursh) Nutt. (Chenopodiaceae). No definite hosts are known. Two individuals have been collected at ultraviolet lights. Adults have been collected from March to December at elevations of 870-2161 m ( $X=1448$  m,  $N=14$ ) (Fig. 233).

**Chorological relationships.**--- Table 67.

**Phylogenetic relationships.**--- Primary characters show this species to be a member of the unresolved heptachotomy also including the *C. frontalis*-*C. americanus* lineage, the *C. infrequens*-*C. puberulus* lineage, *C. canescens*, *C. collaris*, the *C. lobigerinus* species group, and the *C. boucardi* species group (Fig. 236). Secondary characters indicate *C. notolomus* is the sister-species of *C. collaris*, these two species forming a lineage that is sister to the *C. infrequens*-*C. puberulus* lineage within the *C. americanus* species group (Fig. 237).

on 11 males from throughout the species range

Character	Range	Mean	1.5SD	2SE
-----------	-------	------	-------	-----

A. Measurements, in mm.

LR	1.56-2.24	2.015	0.298	0.120
WF	1.20-1.52	1.407	0.144	0.058
WRA	1.16-1.40	1.313	0.096	0.039
LP	2.48-3.68	3.389	0.518	0.208
WPB	2.88-4.00	3.615	0.425	0.171
WE1H	3.36-4.64	4.327	0.541	0.217
WE1M	3.68-5.20	4.822	0.666	0.268
LE1	5.92-9.12	8.044	1.337	0.537
LEy	1.096-1.16	1.058	0.102	0.041
WEy	0.56-0.72	0.636	0.068	0.027

B. Proportions.

WEy/LEy	0.55-0.65	0.602	0.057	0.023
WPB/LP	1.02-1.16	1.070	0.061	0.025
WRA/LR	0.60-0.77	0.656	0.081	0.033
WRA/WF	0.84-0.97	0.935	0.057	0.023
WE1M/LE1	0.56-0.64	0.601	0.038	0.015
LP/LE1	0.40-0.44	0.422	0.024	0.010

on 13 females from throughout the species range

Character	Range	Mean	1.5SD	2SE
-----------	-------	------	-------	-----

A. Measurements, in mm.

LR	1.80-2.32	2.059	0.242	0.090
WF	1.24-1.60	1.437	0.177	0.066
WRA	1.16-1.48	1.302	0.162	0.060
LP	3.12-4.16	3.575	0.466	0.172
WPB	3.20-4.40	3.809	0.555	0.206
WE1H	3.76-5.12	4.517	0.675	0.250
WE1M	4.24-5.92	5.083	0.760	0.281
LE1	7.04-9.44	8.455	1.159	0.429
LEy	0.96-1.20	1.083	0.119	0.044
WEy	0.56-0.72	0.646	0.073	0.027

B. Proportions.

WEy/LEy	0.54-0.65	0.597	0.051	0.019
WPB/LP	1.00-1.12	1.065	0.051	0.019
WRA/LR	0.59-0.68	0.633	0.049	0.018
WRA/WF	0.85-0.95	0.906	0.042	0.015
WE1M/LE1	0.58-0.64	0.601	0.030	0.011
LP/LE1	0.40-0.45	0.423	0.022	0.008

dense and moderately long (Fig. 150a). Head behind eye with variously developed curved sulcus extended from above eye to area under postocular lobe (Fig. 150a). Rostrum moderately robust, moderately to markedly medially tumescent, not to very slightly carinate; lateral margins rounded. Pronotum with postocular lobes well-developed; postocular vibrissae uniformly short (Figs. 150a-151a). Elytra with scale pattern more or less vittate, intervals other than 9 to 11 with white scales equal in size to those on intervals 9 to 11 (Figs. 131-132). Tarsus with ventral pilose vestiture extensive (covering greater part of ventral surface of each article) to slightly reduced in extent (covering not less than apical one-half of articles 2 and 3). Wings present, of variable length (short in only few specimens). Female with abdominal sternum VIII with basal arm short; lateral arms arcuate (Figs. 170-171). Male with aedeagus with internal sac with apex of dorsal median pocket rounded; lobes A to F present; dorsal median pocket low in lateral view (Figs. 190-191).

**Phylogenetic relationships.**--- Monophyly of this species group is based only on distribution of the apotypic state of the primary characters of the presence of a curved sulcus behind the eye. Primary characters show this species group

(all belonging to the *C. americanus* group), and the *C. boucardi* species group (Fig. 236). Secondary characters indicate that the *C. lobigerinus* group is the sister-group to the *C. boucardi* species group, this lineage sister to the *C. americanus* species group (Fig. 237).

4.5.2.3.1 *Cleonidius lobigerinus* Casey

(Figs. 131, 150, 170, 190, 228)

*Cleonis vittatus* Kirby 1837:199 [not Zoubkoff 1829:163].

Lectotype (here designated), male, one of two syntypes, labelled "Co-type", "namer", "*Cleonus/ vittatus/ Kirby/ Co-type*" and with my designation label "LECTOTYPE/  
*Cleonis vittatus/ Kirby desig./ Anderson*" (BMNH). Type locality, North America. O'Brien and Wibmer 1982 (catal., distn., misident. of *Cleonis porlicollis* as synonym).

*Cleonus vittatus*; Melsheimer 1853 (check.). Gemminger and von Harold 1871 (catal.). LeConte 1876a (key). Henshaw 1885 (check.). Fall 1901 (check.). Fletcher 1906. Fall and Cockerell 1907 (check.). Gibson 1914. Leng 1920 (catal.).

*Cleonus virgatus* LeConte 1876a:150 [not Gyllenhal 1834:242].

Holotype (examined), male, labelled "Cal", "C./  
*virgatus/ TYPE Lec.*", "Horn Coll/ H8521" and a red

*Cleonus (Cleonidius) lobigerinus* Casey 1891:191. Lectotype (here designated), female, one of eight syntypes, labelled "Ari", "CASEY/ bequest/ 1925", "TYPE USNM/ 37281", "C.Cl./ lobigerinus/ Cas" and with my designation label "*Cleonus/ lobigerinus* Csy./ LECTOTYPE/ desig. Anderson" (USNM). Type locality, Peach Springs, Arizona. Champion 1902-1906 (in part; distn., misident. of *Cleonidius puberulus*). Csiki 1934 (catal.). Hatch 1971 (key, redesc.).

*Cleonus (Cleonidius) kirbyi* Casey 1891:188 (in key). **New synonymy.** Holotype (examined), female, labelled "Vic./ Vanc.I.", "CASEY/ bequest/ 1925", "TYPE USNM/ 37299", "C.Cl./ kirbyi/ Cas." (USNM). Type locality, Victoria, British Columbia. Csiki 1934 (catal.). Hatch 1971 (key, redesc.).

*Cleonus (Cleonidius) virgatus*; Casey 1891 (key, misident. of *Cleonidius poricollis*). Csiki 1934 (catal., misinterp. as synonym of *Cleonus poricollis*).

*Cleonus (Cleonidius) vittatus*; Casey 1891 (key, misident. of *Cleonidius poricollis*). Hatch 1971 (key, redesc., misident. of *Cleonidius poricollis*).

*Cleonus lobigerinus*; Wickham 1896 (check.). Fall 1901 (check.). Yothers 1916. Leng 1920 (catal.). Tanner 1966 (redesc., biol.).

Leng 1920 (catal.).  
*Cleonidius vittatus*; Faust 1904 (desig. as type species of  
*Cleonidius*).  
*Cleonus quadrilineatus*; Anderson 1914 (misident.).  
*Cleonus sparsus*; Essig 1958 (in part; biol., misident.).  
Arnett-et al. 1980 (probable misident.).  
*Cleonus (Cleonidius) quadrilineatus*; Hatch 1971 (key,  
redesc., misident.).  
*Cleonus (Cleonidius) sparsus*; Hatch 1971 (in part; key,  
redesc., misident.).  
*Cleonis frontalis*; O'Brien and Wibmer 1982 (in part; catal.,  
distn., misident.).  
*Cleonis kirbyi*; O'Brien and Wibmer 1982 (catal., distn.).  
*Cleonis lobigerinus*; O'Brien and Wibmer 1982 (catal.,  
distn.).  
*Cleonis virgatus*; O'Brien and Wibmer 1982 (catal., distn.,  
misident. of *Cleonis poricollis*, as synonym).  
*vittatus, Incertae sedis*; Chevrolat 1873.

**Notes about synonymy,---** There has been much confusion regarding this species and consequently, numerous misidentifications in the literature. *Cleonidius lobigerinus*, as considered here, includes all those western

sulcus well to only slightly developed, to absent. These sympatric specimens are otherwise indistinguishable and are considered herein to be conspecific. Primary types of *Cleonis vittatus* Kirby, *Cleonus virgatus* LeConte, *Cleonus* (*Cleonidius*) *lobigerinus* Casey, and *Cleonus* (*Cleonidius*) *kirbyi* Casey each possess a curved sulcus behind the eye.

*Cleonus* (*Cleonidius*) *kirbyi* Casey was initially proposed by Casey (1891) as a replacement name for *Cleonus vittatus* LeConte, however LeConte (1876a) did not describe a new species, *Cleonus vittatus*, but rather was referring (correctly I might add, contrary to assertions of Casey [1891]) to *Cleonis vittatus* Kirby. Thus *Cleonus* (*Cleonidius*) *kirbyi* Casey should be considered a new species description and not a replacement name. Casey (1891), based only on the brief and inadequate published descriptions of *Cleonis vittatus* Kirby and *Lixus porlicollis* Mannerheim, incorrectly considered *Cleonis vittatus* Kirby as a distinct species from *Cleonus vittatus* Kirby, *sensu* LeConte (1876a), and as conspecific with *Cleonus porlicollis* (Mannerheim).

Both *Cleonis vittatus* Kirby and *Cleonus virgatus* LeConte are junior homonyms. I have chosen *C. lobigerinus* as the valid name over *C. kirbyi* as the former is the more widely used and familiar name.



Faust and labelled in his distinctive handwriting. This indicates that his designation was based on a correct interpretation of *Cleonis vittatus* Kirby as conspecific with *C. lobigerinus* Casey.

Problems in recognition.--- Most members of this widespread species are easily recognized by the presence of a variously developed (usually deep) curved sulcus behind each eye (Fig. 150a). In many specimens the sulcus is present but largely to completely covered by the postocular lobe. In specimens in which the postocular lobe covers the sulcus, reorientation of the head will expose the sulcus to view. If this is not done, such individuals will otherwise prove difficult to separate from other species, especially *C. notolomus* and *C. frontalis*. Specimens lacking a sulcus or with only a slightly developed sulcus appear confined to Oregon and British Columbia (perhaps also Washington, although no such specimens have been seen from that area) and are separated from other species of *Cleonidius* by the sparse scales of the elytra (forming a faintly vittate pattern), possession of large ventral tarsal pilose pads, and rostrum with distinct median tumescence, the base of the rostrum distinctly separated from the frons by a transverse impression (Fig. 150a).

scale pattern (intervals 6 to 8 with scales as large as those on intervals 9 to 11) (Fig. 132) in the former. The sulcus is variously developed in individuals of both species but is moderately deep and more distinct in most *C. lobigerinus* and in only few *C. calandroides*.

**Description.**--- Specimens examined. 387 males, 459 females. Data about variation in LR, WF, WRA, LP, WPB, WELH, WELM, LE1, LEy, WEy, WEy/LEy, WPB/LP, WRA/LR, WRA/WF, WELM/LE1, and LP/LE1 are presented in Tables 55 and 56. Size. Length, male, 4.3-9.3 mm; female, 5.6-10.3 mm. Width, male, 1.8-3.6 mm; female, 2.0-4.0 mm. Head. Eye elongate-oval. Area behind eye with numerous slight wrinkles of cuticle or with variously developed (moderately deeply in most specimens) curved sulcus extended posteroventrally from upper one-half of posterior margin of eye to area under postocular lobe (sulci continue ventrally and meet at gular suture but are covered by prothorax and are not visible) (Fig. 150a). Frons with sparse to moderately dense, small shallow punctures, interspersed with minute punctures. Frons also with very sparse to moderately dense, short to moderately long, suberect to erect hair-like scales, moderately dense elongate-fine appressed white scales medially, suberect to erect immediately above eyes. Frons distinctly separated

with small, shallow to moderately deep rovea. width of frons less than, to subequal to, width at apex of rostrum.

Rostrum. Moderately robust, (width at apex 0.571-0.711 times length in male; 0.568-0.758 in female) (Fig. 150). In

lateral view straight to very slightly curved downward. In

dorsal view with postgenae moderately to markedly laterally expanded, antennal scrobes distinctly visible. Rostrum

moderately to markedly medially tumescent from point of antennal insertion to middle of frons; not laterally

expanded at midlength; lateral margins rounded; with median carina variously developed as narrow low glabrous shiny line

at crest of median tumescence. Dorsal and lateral punctation moderately dense to dense, small, shallow (many

longitudinally confluent); dorsally interspersed with dense minute punctures. Dorsally with very sparse to moderately

dense, short to moderately long, suberect to erect hair-like scales; also with moderately dense, elongate-fine, appressed

to recumbent white scales, sparser medially. Epistoma slightly swollen. Pronotum. Median carina variously

developed in anterior one-half from absent to as low, rounded moderately broad glabrous line. Dorsal and lateral

punctation moderately large, moderately dense to dense, moderately deep to deep, not confluent; smaller, sparser and

shallower apically and on flanks; areas between larger dorsal punctures with or without minute regularly impressed

(Fig. 150a). Scales dorsally absent to small fine and sparse along median line; moderately dense to dense, elongate-fine to elongate-narrow, appressed in pair of broad apically slightly narrowed paramedian stripes; laterally, absent to sparse, small and elongate-fine. Lateral margins with scales moderately dense to dense, elongate-fine to elongate-narrow, appressed to recumbent in moderately broad stripe. Flanks with scales elongate-fine to elongate-narrow, sparse to moderately dense. Median basal portion of disk shallowly to deeply and narrowly impressed. Anterolateral margin with pronotal postocular lobes well-developed; postocular vibrissae uniformly short (Fig. 150a). In dorsal view with lateral margins slightly arcuate and convergent from base to apical one-quarter, slightly constricted at apical one-quarter then straight and convergent to apex.

**Prosternum.** With small shallow to moderately deep impression and moderately to markedly developed rounded swelling anterior to each procoxal cavity. In lateral view with anterior flat to slightly angulate portion long, approximately two-thirds to subequal in length to, posterior more angulate portion; posterior and anterior portions not or slightly separated by shallow transverse impression.

**Elytra.** Elongate-narrow in general form (width at midlength 0.500-0.634 times length in males; 0.500-0.641 in females) (Fig. 131). In dorsal view with lateral margins straight to

apical one-third to apex. Elytral apices not produced, rounded, slightly divergent. Humerus distinct. Dorsally with suberect or erect vestiture lacking to dense, moderately long, erect hair-like scales; with sutural interval with dense, very small and fine to elongate-fine, golden to white scales, interspersed with scattered patches of larger white scales, to with scales uniformly white, dense, small and elongate-fine; intervals 2 with scales golden to white, dense, very small and fine, with scattered patches of larger white scales; intervals 3 to 5 with dense, small and fine to elongate-narrow white scales; intervals 6 to 8 with scales dense, very small and very fine, golden to white, with scattered patches of larger white scales; intervals 9 to 11 with uniformly dense, small to large, elongate-fine to elongate-narrow white scales (Fig. 131). Punctures of elytral striae small, individually distinct, arranged in regular rows. Wings. Long (greater than elytra in length [99%, N=838]), to short (approximately one-half length of elytra or less [1%, N=8]). Legs. Foretibia of female with inner margin with small denticles in apical one-half; subapical tooth indistinct from tibial denticles. Foretibial uncus of both sexes moderately large, mesotibial and metatibial unci small. Tarsal claws connate in basal one-third, not divergent, each with basal internal flange slightly developed. Ventral tarsal pilose vestiture of

three-quarters on article 1, as moderately large rounded pad covering apical one-half on article 2, and as large round pad covering apical three-quarters on article 3 in male, apical one-half to three-quarters in female. **Abdomen.**

Ventral surface with moderately dense, elongate-narrow to moderately robust appressed white scales. Abdominal sterna III to VI (especially III and IV) with individually indistinct to distinct moderately large rounded glabrous patches, each with large puncture and single suberect to erect, moderately long hair-like scale situated in or near the center. Base of abdominal sternum VII of female internally (dorsally) with slightly to well-developed median shiny tubercle. **Genitalia.** Female (four examined). Abdominal sternum VIII with basal arm short; lateral arms slightly inwardly arcuate throughout length; very slightly expanded at apex (Fig. 170). Stylus moderately large compared to length gonocoxite II. Male (four examined). Internal sac (Figs. 190a,b) elongate and low; median dorsal pocket low, with single moderately broad elongate dorsally directed basal lobe (lobe A), with small median dorsally directed lobe at midlength on dorsal surface (lobe B), with moderately large paired dorsally directed lobe at basal one-third on dorsal surface (lobe C), with moderately large paired dorsolaterally directed lobe near midlength near dorsal margin (lobe D), with large paired dorsolaterally

near ventral margin (lobe F). Apical pocket and median dorsal pocket individually distinct; apex of dorsal median pocket rounded in dorsal view. Ventral median pocket moderately large. Apical pocket with paired moderately large ventrally directed lobe near apical one-third at ventral margin. Eversible apical sclerite complex with paired simple scythe-like sclerite; adjacent ventral surface of sides of apical pocket with pair of large, dark, longitudinal sclerites.

Geographic variation.--- Specimens from coastal localities in Oregon and British Columbia (likely also Washington) differ in a number of respects from individuals from throughout the rest of the species range. These individuals have generally slightly larger prosternal swellings, generally deeper and denser punctation of head, rostrum and pronotum, have smaller elytral scales, the pattern only faintly vittate, lack a sulcus behind the eye, lack or else have very short indistinct dorsal suberect or erect vestiture, and have broad foretarsal articles with more extensive ventral tarsal pilosity. Not all individuals however possess all of these character states; mixed series are known in which some individuals possess various combinations of the above character states but otherwise possess the typical *C. lobigerinus* states. As such, no

distinct taxonomic entity. A large number of these specimens have been reared from *Erysimum* sp. (wallflower; Cruciferae) at Victoria, British Columbia, but other typical *C. lobigerinus* have also been reared from various Cruciferae throughout the species range. *C. calandroides*, sister-species of *C. lobigerinus* also occurs on Cruciferae.

Individuals from a series of specimens from San Francisco, California, have the elytra uniformly covered with moderately dense, elongate-fine white scales; the elytra do not appear distinctly vittate. Otherwise these specimens are typical *C. lobigerinus* and in my view do not warrant separate status.

**Geographic distribution.**--- This is the most widely distributed species of *Cleonidius*. It is found throughout western North America across southern British Columbia, Alberta and Saskatchewan in Canada into the adjacent United States of America east to Michigan in the north, south along the west coast to California in the west and south to western Texas and central Mexico in the east (Fig. 228).

Smith (1957) discusses taxa found in prairie habitats in Michigan that are disjunct with the western prairie. The distribution of *C. lobigerinus* in the northern United States appears to fit this pattern.



transitional, Pacific semi-desert, Great Basin desert, Sonoran desert, Mojave desert, and western xerophytic evergreen forest. Adults of this species have been collected on a wide taxonomic range of plants as follows; *Asclepias* sp. (Asclepiadaceae); *Atriplex* sp., *A. nuttallii* Wats., *A. rosea* L., *Beta vulgaris* L. (beet), *Eurotia lanata* (Pursh) Moq., *Salsola kali* L. (all Chenopodiaceae); *Ambrosia* sp., *Franseria confertiflora* (DC) Rydb., *Chrysothamnus* sp., *C. nauseosus* (Pall.) Britton, *C. viscidiflorus* (Hook.) Nutt. var. *typicus*, *Gutierrezia californica* (DC.) Torr. and Gray, *Aplopappus acradenius* (Greene) Blake, *A. venetus* Blake ssp. *vernonioides* (Nutt.) Hall, *Hymenoclea monogyra* T. and G., *Zinnia* sp. (all Compositae); *Sisymbrium altissimum* L., "S. pestifer", *Erysimum* sp. (wallflower), *Descurainia sophia* (L.) Webb., *Brassica campestris* L. (mustard), *B. oleracea* L. (broccoli) (all Cruciferae); *Salvia* sp. (Labiatae); *Lupinus* sp., *Trifolium* sp., *Melilotus albus* Desr., *Medicago sativa* L. (all Leguminosae); *Fragaria* sp., *Rubus* sp., *Purshia tridentata* (Pursh) DC. (all Rosaceae). Adults have been reared or collected from the roots of *Chrysothamnus* sp., *Gutierrezia* sp., *G. microcephala* (DC.) Gray, *Aplopappus* sp., *A. venetus* (Hbk.) Blake (all Compositae); *Brassica* sp., *Raphanus sativus* L. (radish), *Erysimum* sp., *Descurainia pinnata* (Walt.) Britton (all Cruciferae). Tanner (1966) reports collection of adults from a *Grayia-Lycium* community,

radishes at Corvallis, Oregon (as *Cleonus sparsus*) and as attacking wallflower roots on a seed farm on Vancouver Island, British Columbia (as *Cleonus kirbyi*).

This species has been collected on various types of vegetation primarily in dry washes and stream beds throughout the species range. Individuals have also been collected on sand dunes or other sandy habitats.

I have seen a single specimen collected from the stomach of *Bufo* sp. (Amphibia).

Adults have been collected from June to November at elevations from sea level to 4004 m ( $X=1469$  m,  $N=132$ ) (Fig. 233).

**Chorological relationships.**--- Table 67. *C. lobigerinus* is allopatric to its sister-species *C. calandroides*.

**Phylogenetic relationships.**--- Primary characters show this species to be the sister-species of *C. calandroides*, the only other species in the *C. lobigerinus* species group (Fig. 236).

BASED ON EIGHT MEASUREMENTS FROM THE N. PACIFIC OAK, CALIFORNIA

Character	Range	Mean	1.5SD	2SE
-----------	-------	------	-------	-----

A. Measurements, in mm.

LR	1.52-1.88	1.700	0.195	0.092
WF	0.88-1.08	0.950	0.131	0.062
WRA	0.96-1.12	1.045	0.087	0.041
LP	2.24-2.72	2.500	0.238	0.112
WPB	2.48-3.04	2.730	0.268	0.126
WE1H	2.96-3.68	3.320	0.345	0.163
WE1M	3.20-3.76	3.530	0.311	0.146
LE1	6.40-7.36	6.880	0.602	0.284
LEy	0.84-1.00	0.920	0.072	0.034
WEy	0.44-0.52	0.475	0.038	0.018

B. Proportions.

WEy/LEy	0.48-0.55	0.517	0.038	0.018
WPB/LP	1.03-1.15	1.093	0.061	0.029
WRA/LR	0.59-0.68	0.616	0.049	0.023
WRA/WF	1.00-1.18	1.104	0.083	0.039
WE1M/LE1	0.50-0.53	0.513	0.015	0.007
LP/LE1	0.34-0.39	0.364	0.027	0.013

based on eight females from 1mi. N. Little Lake, California

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.52-1.92	1.735	0.208	0.098
WF	0.88-1.12	0.980	0.120	0.057
WRA	1.00-1.24	1.075	0.122	0.057
LP	2.24-2.96	2.530	0.345	0.163
WPB	2.56-3.28	2.860	0.394	0.186
WE1H	2.96-4.16	3.430	0.582	0.274
WE1M	3.36-4.56	3.750	0.600	0.283
LE1	6.40-8.64	7.260	1.096	0.516
LEy	0.84-1.04	0.940	0.091	0.043
WEy	0.44-0.56	0.500	0.072	0.034
B. Proportions.				
WEy/LEy	0.50-0.58	0.532	0.047	0.022
WPB/LP	1.09-1.25	1.132	0.079	0.037
WRA/LR	0.57-0.71	0.621	0.070	0.033
WRA/WF	1.04-1.14	1.098	0.059	0.028
WE1M/LE1	0.50-0.53	0.516	0.015	0.007
LP/LE1	0.32-0.36	0.349	0.024	0.011

(Figs. 132, 151, 171, 191, 229)

*Lixus calandroides* Randall 1838:42. Type(s) lost. Gemminger and von Harold 1871 (catal.). Sprague and Austin 1875. LeConte 1876a (misinterpret., as synonym of *Lixus musculus*).

*Cleonus calandroides*; LeConte 1876b (diagnosis). Henshaw 1885 (check.). Casey 1891 (key). Ely 1913 (biol.). Blatchley and Leng 1916 (key, redesc., biol.). Leng 1920 (catal.). Leonard 1926 (check., biol.). Csiki 1934 (catal.). Bruhn 1947 (morphol.). Sanders 1960 (morphol.). Kissinger 1964 (biol.).

*Cleonus (Cleonidius) calandroides*; Casey 1891 (key). Csiki 1934 (catal.).

*Cleonis calandroides*; O'Brien and Wibmer 1982 (catal., distn.).

**Notes about synonymy.**--- *Lixus calandroides* was described from an unspecified number of specimens from Chelsea Beach, Massachusetts (Randall 1838). Unfortunately type material of Randall has been lost (Sprague and Austin 1875). Following recommendations of the International Code of Zoological Nomenclature, no neotype designation is required, because there has been no recent confusion regarding identity of this species. After identification of specimens as this species by Sprague and Austin (1875), LeConte (1876b), based

transferred the species to *Cleonus* and removed it from his previously proposed synonymy with *Lixus musculus* Say (LeConte 1876a).

**Problems in recognition.**--- This is one of only two species of *Cleonidius* restricted to eastern North America (Figs. 215, 229). These two differ markedly in structural features and should not be confused. The eastern *C. calandroides* is distinguished from the western *C. lobigerinus* by the short suberect dorsal vestiture (Fig. 151a) and less distinctly vittate elytral scale pattern (intervals 6 to 8 with scales as large as those on intervals 9 to 11) (Fig. 132) in the former. The sulcus behind the eye is variously developed in individuals of both species but is moderately deep and distinct in most *C. lobigerinus* and in only few *C. calandroides* (see descriptions).

**Description.**--- Specimens examined. 65 males, 63 females. Data about variation in LR, WF, WRA, LP, WPB, WELH, WELM, LE1, LEy, WEy, WEy/LEy, WPB/LP, WRA/LR, WRA/WF, WELM/LE1, and LP/LE1 are presented in Tables 57 and 58. Size. Length, male, 8.0-9.3 mm; female, 5.5-9.6 mm. Width, male, 3.0-3.6 mm; female, 2.1-3.9 mm. Head. Eye elongate-oval. Area behind eye with numerous slight wrinkles of cuticle or with variously developed (moderately deeply in only few specimens) curved sulcus extended posteroventrally from

postocular lobe (sulci continue ventrally and meet at gular suture but are covered by prothoracic structures and not visible) (as in Fig. 150a). Frons with only scattered small shallow punctures, interspersed with minute punctures. Frons also with moderately dense, short, suberect hair-like scales, sparse to moderately dense elongate-fine appressed white scales medially, suberect to erect immediately above eyes. Frons distinctly separated from base of rostrum by moderately deep to deep transverse impression (Fig. 151a). Base of median tumescence of rostrum with small, shallow to moderately deep fovea. Width of frons greater than to subequal to width at apex of rostrum. Rostrum. Moderately robust, slightly less so in females (width at apex 0.596-0.675 times length in male; 0.558-0.653 in female) (Fig. 151). In lateral view straight to very slightly curved downward. In dorsal view with postgenae moderately laterally expanded, antennal scrobes partially visible. Rostrum medially tumescent from point of antennal insertion to middle of frons; not laterally expanded at midlength; lateral margins rounded; with median carina variously developed as narrow low glabrous shiny line at crest of median tumescence. Dorsal and lateral punctation moderately dense, small, shallow (many longitudinally confluent); dorsally interspersed with dense minute punctures. Dorsally with very sparse to moderately dense, short, suberect hair-like scales (especially laterally); also with

scales, sparser medially. Epistoma slightly swollen.

**Pronotum.** Median carina variously developed in anterior one-half from absent to as low, rounded narrow glabrous line. Dorsal and lateral punctation moderately large, moderately dense to dense, shallow, not confluent; smaller apically and on flanks; areas between larger dorsal punctures with or without minute regularly impressed punctures. Dorsally with moderately dense, very short, suberect hair-like scales, each situated in large puncture (Fig. 151a). Scales dorsally absent to small fine and sparse along median line; moderately dense, elongate-narrow, appressed in pair of broad apically slightly narrowed paramedian stripes; laterally, absent to sparse, small and elongate-fine. Lateral margins with scales dense, elongate-narrow, appressed in moderately broad stripe. Flanks with scales dense, elongate-fine to elongate-narrow. Median basal portion of disk shallowly to deeply and narrowly impressed. Anterolateral margin with pronotal postocular lobes well-developed; postocular vibrissae uniformly short (Fig. 151a). In dorsal view with lateral margins slightly arcuate and slightly convergent from base to apical one-quarter, slightly constricted at apical one-quarter then straight and convergent to apex.

**Prosternum.** With small shallow impression and slight to moderately developed rounded swelling anterior to each procoxal cavity. In lateral view with anterior flat to



slightly angulate portion long, subequal in length to posterior more angulate portion; posterior and anterior portions slightly separated by shallow transverse impression. Elytra. Moderately robust in general form (width at midlength 0.567-0.596 times length in males; 0.550-0.630 in females) (Fig. 132). In dorsal view with lateral margins straight, subparallel to slightly convergent from apical one-third to humerus; moderately and evenly arcuate from apical one-third to apex. Elytral apices not produced, rounded, slightly divergent. Humerus distinct. Dorsally with moderately dense, very short, suberect, indistinct hair-like scales; scales more or less uniformly dense, white; with sutural interval and interval 2 with scales small and fine; intervals 3 to 5 and 9 to 11 with scales large and elongate-narrow; scales small and fine to large and elongate-narrow on intervals 6 to 8 (Fig. 132). Punctures of elytral striae small, individually distinct, arranged in regular rows. Wings. Long (greater than elytra in length). Legs. Foretibia of female with inner margin with small denticles in apical one-half; subapical tooth indistinct from tibial denticles. Foretibial and mesotibial unci of both sexes moderately large, metatibial uncus small. Tarsal claws connate in basal one-half, slightly divergent, each with basal internal flange slightly developed. Ventral tarsal pilose vestiture of foretarsus of male present as large pads covering more or less the entire ventral surface of articles 2 and 3 and apical one-half of article 1; of

remale, as elongate pad covering apical one-half on article 1, as large oval pad covering apical three-quarters on article 2, and as large round pad covering more or less the entire ventral surface of article 3; of mesotarsus of male and female as small elongate-narrow pad covering apical one-half on article 1, as large elongate pad covering more or less the entire ventral surface on articles 2 and 3; of metatarsus of male and female as elongate pad covering apical three-quarters on article 1, as large elongate-oval pad covering apical one-half on article 2, and as large oval pad covering apical three-quarters on article 3. **Abdomen.** Ventral surface with moderately dense to dense, elongate-narrow appressed white scales. Abdominal sterna III to VI (especially III and IV) with individually indistinct to distinct moderately large rounded glabrous patches, each with large puncture and single appressed hair-like scale situated in or near the center. Base of abdominal sternum VII of female internally (dorsally) with moderately developed median shiny tubercle. **Genitalia.** Female (two examined). Abdominal sternum VIII with basal arm short; lateral arms slightly inwardly arcuate throughout length; very slightly expanded at apex (Fig. 171). Stylus moderately large compared to length gonocoxite II. Male (four examined). Internal sac (Figs. 191a,b) elongate and low; median dorsal pocket low, with single moderately broad elongate dorsally directed basal lobe (lobe A), with small median dorsally directed lobe at midlength on dorsal surface

(lobe B), with moderately large paired dorsally directed lobe at basal one-third on dorsal surface (lobe C), with moderately large paired dorsolaterally directed lobe near midlength near dorsal margin (lobe D), with large paired dorsolaterally directed lobe just beyond midlength at dorsal margin (lobe E), and with small ventrally directed paired lobe near base near ventral margin (lobe F). Apical pocket and median dorsal pocket individually distinct; apex of dorsal median pocket rounded in dorsal view. Ventral median pocket moderately large. Apical pocket with paired moderately large ventrally directed lobe near apical one-third at ventral margin. Eversible apical sclerite complex with paired simple scythe-like sclerite; adjacent ventral surface of sides of apical pocket with pair of large, dark, longitudinal sclerites.

**Geographic variation.**--- None noted.

**Geographic distribution.**--- This species is distributed along the northern Atlantic Coast from Long Island, New York, north to New Hampshire and New Brunswick (Fig. 229).

**Natural history.**--- This species breeds in the sea-rocket, *Cakile edentula* (Bigel.) Hook. (Cruciferae). Ely (1913) collected various life stages from larvae to adults in the roots of this plant in mid-August. He reports that larvae live entirely within the root and that the portion of the

root lying within 2 or 3 inches of the surface of the ground is most commonly attacked. Pupation takes place in a cocoon constructed of small shreds of plant material built up in a longitudinal excavation in one side of the root. He further states that the beetles must be very common as virtually all plants examined showed evidence of feeding by larvae.

Blatchley and Leng (1916) also state that larvae breed in *Cakile edentula* and other specimens I have examined bear label data indicating they were reared from this plant species. No other plant associations are known. Adults have been collected only on sea beaches from April to October.

**Chorological relationships.**--- Table 67. *Cleonidius calandroides* is sympatric only with northern *C. subcylindricus*.

**Phylogenetic relationships.**--- Primary characters show this species to be the sister-species of *C. lobigerinus*, the only other species in the *C. lobigerinus* species group (Fig. 236).

Table 57. Descriptive statistics for *C. calandroides* based on eight males from Hampton, New Hampshire

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.60-1.96	1.850	0.166	0.078
WF	1.08-1.24	1.180	0.085	0.040
WRA	1.08-1.24	1.170	0.089	0.042
LP	2.56-3.12	2.960	0.280	0.132
WPB	2.80-3.44	3.180	0.338	0.159
WE1H	3.20-4.08	3.740	0.399	0.188
WE1M	3.68-4.56	4.210	0.430	0.203
LE1	6.40-7.84	7.240	0.638	0.301
LEy	0.84-0.96	0.910	0.053	0.025
WEy	0.44-0.52	0.485	0.038	0.018
B. Proportions.				
WEy/LEy	0.52-0.57	0.533	0.024	0.011
WPB/LP	1.00-1.13	1.075	0.064	0.030
WRA/LR	0.60-0.68	0.633	0.044	0.021
WRA/WF	0.94-1.04	0.992	0.053	0.025
WE1M/LE1	0.57-0.60	0.581	0.016	0.007
LP/LE1	0.39-0.43	0.409	0.021	0.010

Table 58. Descriptive statistics for *C. calandroides*  
based on eight females from Hampton, New Hampshire

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.92-2.08	1.975	0.078	0.037
WF	1.20-1.28	1.235	0.050	0.024
WRA	1.12-1.28	1.190	0.077	0.036
LP	3.04-3.20	3.100	0.085	0.040
WPB	3.20-3.60	3.370	0.197	0.093
WE1H	3.84-4.24	4.030	0.202	0.095
WE1M	4.24-4.72	4.510	0.256	0.121
LE1	7.36-8.16	7.820	0.394	0.186
LEy	0.88-1.08	0.955	0.093	0.044
WEy	0.48-0.56	0.530	0.042	0.020
B. Proportions.				
WEy/LEy	0.52-0.64	0.556	0.055	0.026
WPB/LP	1.03-1.18	1.087	0.070	0.033
WRA/LR	0.56-0.65	0.603	0.046	0.021
WRA/WF	0.93-1.00	0.964	0.040	0.019
WE1M/LE1	0.55-0.63	0.577	0.038	0.018
LP/LE1	0.38-0.41	0.397	0.015	0.007

#### 4.5.2.4 *Cleonidius boucardi* species group

**Diagnosis.**--- Size moderately large to large for *Cleonidius* (Fig. 197). Dorsal erect or suberect vestiture absent (Fig. 152a) to present, moderately dense and moderately long (Fig. 155a). Rostrum moderately robust, not (Fig. 152a) to moderately (Fig. 155a) medially tumescent, not to variously carinate; lateral margins rounded. Pronotum with postocular lobes absent (Fig. 155a) to very slightly developed (Fig. 152a); postocular vibrissae moderately long to long, of unequal length and longest behind base of eye. Elytra with scale pattern more or less vittate, intervals other than 9 to 11 with white scales equal in size to those on intervals 9 to 11 (Figs. 133-136). Tarsus with ventral pilose vestiture extensive (covering greater part of ventral surface of each article) to slightly reduced in extent (covering not less than apical one-half of articles 2 and 3). Wings present, long. Female with abdominal sternum VIII with basal arm short to moderate in length; lateral arms arcuate (Figs. 172-175). Male with aedeagus with internal sac with apex of dorsal median pocket variously emarginate or sinuate (Figs. 192a,b-195a,b); lobes A to F present; dorsal median pocket low in lateral view.

**Phylogenetic relationships.**--- Primary characters show this species group to be monophyletic and a member of the unresolved heptachotomy also including the *C. frontalis*-*C. americanus* lineage, the *C. infrequens*-*C. puberulus* lineage,

monophyly of the group and indicate that the *C. boucardi* group is the sister-group to the *C. lobgerlingus* species group, this lineage sister to the *C. americanus* species group (Fig. 237).

4.5.2.4.1 *Cleonidius boucardi* (Chevrolat)

(Figs. 133, 152, 172, 192, 221)

*Apleurus boucardi* Chevrolat 1873:79. Holotype (examined), male, labelled "Puebla", "[letter illegible] Boucard", pale blue square label, "Typus", "41", "471/ 85", "Riksmuseum/ Stockholm" and with my label "*Apleurus boucardi*/ Chevrolat/ HOLOTYPE/ lab. Anderson" (NRS). Type locality, Puebla, Mexico.

*Cleonus carinicornis* LeConte 1876a:152 [not Gyllenhal 1834:241]. New synonymy. Holotype (examined), female, labelled with a pale green circle (=Wyoming Territory), "Type/ 524.1", "*C. carinicornis*/ Lec." (MCZC). Type locality, Colorado. Henshaw 1881-1882 (check.). Henshaw 1885 (check.). Wickham 1902 (check.). Blatchley and Leng 1916 (key, redesc.). Leng 1920 (catal.). Brimley 1938 (check.). Arnett *et al.* 1980.

*Cleonus (Cleonidius) boucardi*; Casey 1891 (key). Champion 1902-1906 (in part; distn., misident of *Cleonidius lobgerlingus*). Csiki 1934 (catal.).



Holotype (examined), female, labelled "Ari", "CASEY/  
bequest/ 1925", "TYPE USNM/ 37280", "C.CI./ lecontel/  
Cas." (USNM). Type locality, Arizona. Csiki 1934  
(catal.).

*Cleonus lecontel*; Leng 1920 (catal.).

*Cleonus (Cleonidius) lecontellus* Csiki 1934:64. New  
synonymy. New name for *Cleonus carinicolllis* LeConte.  
Blackwelder 1939 (check.).

*Cleonus boucardi*; Blackwelder 1947 (check.).

*Cleonis boucardi*; O'Brien and Wibmer 1982 (catal., distn.).

*Cleonis carinicolllis*; O'Brien and Wibmer 1982 (catal.,  
distn., as jr. homonym).

*Cleonis lecontel*; O'Brien and Wibmer 1982 (catal., distn.).

*Cleonis lecontellus*; O'Brien and Wibmer 1982 (catal.,  
distn.).

**Notes about synonymy.**--- Chevrolat (1873), in describing *C.*  
*boucardi* noted the similarity to *C. trivittatus* (Say).

Although most individuals of the two species are quite  
distinct, some individuals are difficult to reliably assign  
as belonging to either species. One such specimen is the  
holotype of *Cleonus carinicolllis* LeConte, which has dense,  
extensive ventral tarsal pilose pads, moderately developed

that of most *C. boucardi* (the latter, all states of *C. trivittatus*). The type locality is "Colorado Territory" and therefore within the ranges of both species. I have chosen to regard this individual as conspecific with *C. boucardi* and accordingly emphasize the taxonomic significance of the characters of extent of ventral tarsal vestiture and development of postocular lobes as diagnostic for these two species.

*Cleonidius lecontellus* Csiki is a typical *C. boucardi*, otherwise differing from the latter, as noted by Casey (1891), only in details of the elytral scale pattern.

**Problems in recognition.**--- Most individuals of this species are likely to be confused only with *C. trivittatus* and *C. quadrilineatus*. They are however, easily separated by the rostrum not or only slightly dorsally medially tumescent (Fig. 152a), postocular lobes slightly to moderately developed (Fig. 152a), and with all tarsal articles with extensive ventral pilose pads (covering more or less the entire ventral surface of each article) in *C. boucardi* specimens. In both *C. trivittatus* and *C. quadrilineatus* the rostrum is medially dorsally tumescent and carinate (Figs. 153a, 155a), the postocular lobes are not to moderately developed (but only in few specimens of *C. trivittatus*,

and Texas are very similar to *C. trivittatus* in elytral scale pattern and in rostral characters but can be separated by their extensive ventral tarsal pilosity and moderately developed postocular lobes.

*Cleonidius boucardi* may also be confused with some *C. lobigerinus* that have a similar faintly vittate elytral scale pattern and lack the curved sulcus behind the eye. Although *C. boucardi* and *C. lobigerinus* are sympatric over part of their ranges, those individuals of *C. lobigerinus* that are most likely to be confused with *C. boucardi* are found only in the Pacific Northwest; *C. boucardi* does not occur in this or adjacent areas.

This is the only species of *Cleonidius* that has been extensively collected in Mexico (Fig. 221).

**Description.**--- Specimens examined. 73 males, 82 females. Data about variation in LR, WF, WRA, LP, WPB, WE1H, WE1M, LE1, LEy, WEy, WEy/LEy, WPB/LP, WRA/LR, WRA/WF, WE1M/LE1, and LP/LE1 are presented in Tables 59 and 60. **Size.** Length, male, 7.5-14.7 mm; female, 8.4-18.6 mm. Width, male, 3.2-5.9 mm; female, 3.4-7.0 mm. **Head.** Eye elongate-oval. Area behind eye with numerous slight wrinkles of cuticle. Frons with sparse to moderately dense, small to large, shallow to deep punctures, largest, deepest and longitudinally confluent in

elongate-fine, appressed to recumbent white scales medially, suberect to erect immediately above eyes. Frons continuous with base of rostrum or only very slightly separated from base of rostrum by very shallow transverse impression (Fig. 152a). Base of median tumescence of rostrum with small, shallow to moderately deep fovea. Width of frons subequal to width at apex of rostrum. Rostrum. Moderately robust, (width at apex 0.574-0.674 times length in male; 0.593-0.658 in female) (Fig. 152). In lateral view straight to very slightly curved downward. In dorsal view with postgenae slightly laterally expanded, antennal scrobes distinctly visible. Rostrum not to very slightly medially tumescent from point of antennal insertion to middle of frons; not laterally expanded at midlength; lateral margins rounded; with median carina variously developed as narrow to broad low glabrous shiny line. Dorsal and lateral punctation sparse to dense, small to large, moderately deep to deep (dense and large in most specimens with many punctures longitudinally confluent); punctures smaller and shallower apically; dorsally interspersed with dense minute punctures. Dorsally with scattered, short, suberect hair-like scales in some specimens; also with sparse to moderately dense, elongate-fine to elongate-narrow, appressed to recumbent white scales. Epistoma very slightly swollen. Pronotum.

elevated sharp narrow glabrous shiny line, well-developed and distinct in most specimens. Dorsal and lateral punctation small, moderately dense, shallow, not confluent; smaller apically and on flanks; areas between larger dorsal punctures with dense minute regularly impressed punctures. Lacking dorsal suberect or erect vestiture (Fig. 152a). Scales dorsally absent in moderately broad median longitudinal line; moderately dense, elongate-fine to elongate-narrow, appressed in pair of narrow to broad apically slightly narrowed paramedian stripes; laterally, scales absent to very sparse, small and fine. Lateral margins with scales moderately dense, elongate-narrow, appressed in moderately broad stripe. Flanks with scales sparse to moderately dense, elongate-fine. Median basal portion of disk shallowly to moderately deeply and broadly impressed. Anterolateral margin with pronotal postocular lobes very slightly to moderately developed; postocular vibrissae of unequal length, moderately long to long, greatest length (from one-half to two-thirds width of eye) behind base of eye (Fig. 152a). In dorsal view with lateral margins uniformly slightly arcuate and convergent from base to apex, or slightly constricted at apical one-quarter, then straight and convergent to apex. Prosternum. With small very shallow indistinct impression and lacking or with at most only very slight rounded swelling anterior to each procoxal

length of posterior more angulate portion; posterior and anterior portions not to slightly differentiated by shallow transverse impression. Elytra. Moderately robust in general form (width at midlength 0.567-0.620 times length in males; 0.537-0.605 in females) (Fig. 133). In dorsal view with lateral margins straight, to very slightly sinuate and slightly convergent from apical one-third to humerus; moderately and evenly arcuate from apical one-third to apex. Elytral apices not to slightly produced, rounded to subacuminate, not to slightly divergent. Humerus distinct. Dorsally lacking suberect or erect dorsal vestiture; sutural interval with moderately dense, small fine white scales; interval 2 with scales absent to sparse small and fine; interval 3 with scales moderately dense large elongate-narrow; intervals 4 and 5 with scales absent to moderately dense, moderately large and elongate-fine; intervals 6 to 8 with scales absent to sparse small and fine; intervals 9 to 11 with scales moderately dense, large elongate-narrow; striae 1, 6 and 7 with irregularly distributed patches of larger and denser white scales (Fig. 133). Punctures of elytral striae small, individually distinct, arranged in regular rows. Wings. Long (greater than elytra in length). Legs. Foretibia of female with inner margin with small to large denticles in apical three-quarters; subapical tooth large, distinct from apical

connate in basal one-third to one-half, not to slightly divergent, each with basal internal flange well-developed. Ventral tarsal pilose vestiture of all tarsal articles present as large pads covering more or less entire ventral surface of articles 2 and 3 and apical one-half to three-quarters of article 1. **Abdomen.** Ventral surface with sparse to moderately dense, elongate-fine to elongate-narrow appressed white scales. Abdominal sterna III to VI lacking glabrous patches or with sterna III and IV with individually indistinct small rounded glabrous patches, each with large puncture and single appressed hair-like scale situated in or near the center. Base of abdominal sternum VII of female internally (dorsally) with very slightly developed median shiny tubercle. **Genitalia.** Female (four examined). Abdominal sternum VIII with basal arm short; lateral arms moderately inwardly arcuate throughout length; slightly expanded at apex (Fig. 172). Stylus moderately large compared to length gonocoxite II. Male (three examined). Internal sac (Figs. 192a,b) elongate and low; median dorsal pocket low, with single narrow elongate dorsally directed basal lobe (lobe A), with small median dorsally directed lobe at midlength on dorsal surface (lobe B), with moderately large paired dorsally directed lobe at basal one-third on dorsal surface (lobe C), with moderately large paired markedly dorsolaterally directed lobe near midlength near dorsal

with small ventrally directed paired lobe near base near ventral margin (lobe F). Apical pocket and median dorsal pocket individually distinct; apex of dorsal median pocket slightly emarginate medially in dorsal view (Fig. 192b). Ventral median pocket large. Apical pocket with paired moderately large ventrally directed lobe near apical one-third at ventral margin. Eversible apical sclerite complex with paired simple scythe-like sclerite; adjacent ventral surface and basal portion of sides of apical pocket with pair of large, dark, longitudinal sclerites.

**Geographic variation.**--- Specimens from Kansas and Texas in the northern part of the species range have a rostrum that is slightly medially tumescent with smaller dorsal punctation, and an elytral scale pattern that is more distinctly vittate.

**Geographic distribution.**--- This species is found from extreme southern California and Arizona in the west, and Kansas and northern Texas in the east, south through central Mexico to Honduras (Fig. 221).

**Natural history.**--- This species is found in mesquite-grassland and grassland habitat throughout Mexico and the southern United States. There are no plant



255-3385 m (X=1900 m, N=29) (Fig. 233).

Chorological relationships.--- Table 67.

Phylogenetic relationships.--- Primary characters indicate only that this species is a member of an unresolved quadrichotomy composed of the four species in the *C. boucardi* group (Fig. 236). Distribution of the apotypic states of the secondary characters of extent of development of postocular lobes and extent of ventral tarsal pilosity place this species as the sister-species to the *C. trivittatus*-*C. quadrilineatus*-*C. placidus* lineage in the *C. boucardi* group (Fig. 237).

Character	Range	Mean	1.5SD	2SE
<b>A. Measurements, in mm.</b>				
LR	1.80-2.52	2.065	0.386	0.182
WF	1.08-1.48	1.280	0.190	0.089
WRA	1.08-1.68	1.315	0.307	0.145
LP	2.96-4.40	3.570	0.714	0.337
WPB	3.12-4.80	3.810	0.791	0.373
WE1H	3.76-5.68	4.550	0.972	0.458
WE1M	4.24-6.16	5.040	0.981	0.462
LE1	7.20-10.72	8.62	1.900	0.896
LEy	0.96-1.32	1.095	0.192	0.091
WEy	0.60-0.76	0.665	0.106	0.050
<b>B. Proportions.</b>				
WEy/LEy	0.58-0.64	0.608	0.036	0.017
WPB/LP	1.04-1.09	1.067	0.022	0.010
WRA/LR	0.57-0.67	0.635	0.051	0.024
WRA/WF	0.87-1.14	1.024	0.113	0.053
WE1M/LE1	0.57-0.62	0.586	0.024	0.011
LP/LE1	0.40-0.44	0.415	0.018	0.009

On eight females from 9 mi. W. Tepic, Jalisco, Mexico

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.52-2.40	2.115	0.501	0.236
WF	0.96-1.52	1.320	0.308	0.145
WRA	1.00-1.48	1.320	0.264	0.125
LP	2.56-4.16	3.560	0.846	0.399
WPB	2.48-4.56	3.760	1.073	0.506
WE1H	3.04-5.44	4.550	1.284	0.605
WE1M	3.68-5.76	5.03	1.177	0.555
LE1	6.08-10.72	9.020	2.461	1.160
LEy	0.84-1.32	1.105	0.217	0.102
WEy	0.56-0.80	0.650	0.110	0.052
B. Proportions.				
WEy/LEy	0.54-0.67	0.591	0.061	0.029
WPB/LP	0.97-1.13	1.051	0.075	0.036
WRA/LR	0.59-0.66	0.627	0.031	0.014
WRA/LE1	0.95-1.06	1.004	0.068	0.032
WE1M/LE1	0.54-0.61	0.560	0.032	0.015
LP/LE1	0.38-0.42	0.397	0.021	0.010

*Cleonus trivittatus* Say 1831:10. Neotype (here designated, from the LeConte Collection), female, labelled with a pale green circle (=Wyoming Territory), "1847", "J. LECONTE/ COLLECTION", "*Cl./ trivittatus/ Say*" and with my designation label "*Cleonus trivittatus/ Say* NEOTYPE/ desig. Anderson" (MCZC). Type locality, Colorado. Melsheimer 1853 (check.). LeConte 1859b (error as *C. vittatus* Say). Gemminger and Harold 1871 (catal.). LeConte 1876a (key). Henshaw 1885 (check.). Wickham 1902 (check.). Fall and Cockerell 1907 (check.). Leng 1920 (catal.). Blackwelder 1947 (check.). Essig 1958 (biol.). Kumar et al. 1976 (in part; biol.).

*Rhynchophorus praepotens* Say 1831:21. Type destroyed.

*Lixus praepotens*; Boheman 1836 (redescr.).

*Apleurus trivittatus*; Chevrolat 1873.

*Cleonus inornatus* LeConte 1876a:149. New synonymy. Holotype (examined), female, labelled "Cal", "*C./ inornatus/ TYPE LeC*", "Horn Coll/ H 8519" (MCZC). Type locality, Owen's Valley, California. Henshaw 1881-1882 (check.). Henshaw 1885 (check.). Fall 1901 (check.). Leng 1920 (catal.).

*Cleonus (Cleonidius) bicarinatus* Casey 1894:190 [not Gebler 1830:158]. New synonymy. Holotype (examined), male, labelled "Tex", "CASEY/ bequest/ 1925", "TYPE USNM/ 37279", "*C.Cl./ bicarinatus/ Cas*" (USNM). Type

- Cleonus (Cleonidius) carinicolllis*; Casey 1891 (key, misident.).
- Cleonus (Cleonidius) inornatus*; Casey 1891 (key). Csiki 1934 (catal.).
- Cleonus (Cleonidius) praepotens*; Casey 1891 (as synonym).
- Cleonus (Cleonidius) trivittatus*; Casey 1891 (key). Csiki 1934 (catal.).
- Cleonus bicarinatus*; Leng 1920 (catal.).
- Cleonus quadrilineatus*; Pierce 1907 (biol., misident.). Chittenden 1911 (biol., misident.). Blatchley and Leng 1916 (figure, misident.). Yothers 1916 (misident., in part). Essig 1958 (biol., misident.). Arnett *et al.* 1980 (key, redesc., biol., misident.).
- Cleonis bicarinatus*; O'Brien and Wibmer 1982 (catal., distn.).
- Cleonis inornatus*; O'Brien and Wibmer 1982 (catal., distn.).
- Cleonis praepotens*; O'Brien and Wibmer 1982 (catal., distn. as synonym).
- Cleonis trivittatus*; O'Brien and Wibmer 1982 (catal., distn.).

**Notes about synonymy.**--- The type series of this species has been lost. I have selected and designated as neotype, a specimen from the LeConte collection, because it is generally considered that LeConte compared his material with

that of Say before the loss of the latter. This collection thus represents one of the best sources of reliably identified specimens of Say species. The type locality of the original type series is given as "Arkansaw" (Say 1831) but it is later stated that he (Say) "obtained two or three specimens near the Rocky Mountains". The neotype selected is from "Colorado Territory" which probably refers to the same general area as that intended by the use of "Arkansaw" by Say (1831), which, as indicated by the reference to the Rocky Mountains, undoubtedly refers to more than what is presently the state of Arkansas.

*Rhynchophorus praepotens* Say (type lost) was regarded by LeConte (1859b) as conspecific with *Cleonus trivittatus* Say. This synonymy has since been generally followed and thus no neotype designation is required. LeConte (1859b) also states that *Lixus praepotens* (Say) of Boheman (1836) is evidently this same species. *Lixus praepotens* (Say) of Boheman (1836) therefore represents only a reassignment from *Rhynchophorus* to *Lixus* and a species redescription, not description of a new species as indicated by Csiki (1934) and O'Brien and Wibmer (1982).

The holotype of *Cleonus Inornatus* LeConte from Owen's Valley, California, as noted under "Notes about synonymy" section for *C. placidus*, although possessing a similar elytral scale pattern to *C. placidus* individuals, is here considered conspecific with *C. trivittatus*. This decision is based on the character states of the reduced extent of

tarsal vestiture, presence of slight postocular lobes, and short dorsal erect vestiture on the pronotum shared with *C. trivittatus* and not *C. placidus*. *Cleonus (Cleonidius) bicarinatus* Casey is a typical, but abraded *C. trivittatus*.

**Problems in recognition.**--- Specimens of *C. trivittatus* are most likely to be confused with those of *C. quadrilineatus*, *C. placidus* and *C. boucardi*. Discussions of characters allowing for separation of the former are given under the "Problems in recognition" section for each of the latter three species.

**Description.**--- **Specimens examined.** 128 males, 159 females. Data about variation in LR, WF, WRA, LP, WPB, WE1H, WE1M, LE1, LEy, WEy, WEy/LEy, WPB/LP, WRA/LR, WRA/WF, WE1M/LE1, and LP/LE1 are presented in Tables 61 and 62. **Size.** Length, male, 9.4-17.3 mm; female, 8.2-18.4 mm. Width, male, 3.8-9.4 mm; female, 3.0-8.2 mm. **Head.** Eye elongate-oval. Area behind eye with numerous slight wrinkles of cuticle. Frons with punctures sparse to moderately dense, small, shallow, some large and longitudinally confluent immediately above eyes; interspersed with dense minute punctures. Frons also with sparse to moderately dense, elongate-narrow appressed to recumbent white scales medially, erect immediately above eyes. Frons separated from base of rostrum by shallow to moderately deep transverse impression (Fig. 153a). Base of median tumescence of rostrum with small, shallow fovea.

Width of frons greater than to subequal to width at apex of rostrum. Rostrum. Moderately robust, (width at apex 0.578-0.732 times length in male; 0.567-0.796 in female) (Fig. 153). In lateral view very slightly curved downward. In dorsal view with postgenae slightly laterally expanded, antennal scrobes partially visible. Rostrum slightly to markedly medially tumescent from point of antennal insertion to middle of frons (especially basally) (Fig. 153a); not laterally expanded at midlength; lateral margins rounded; with median carina variously developed as narrow low glabrous shiny line at crest of median tumescence. Dorsal and lateral punctation sparse to dense, small, shallow; dorsally interspersed with dense minute punctures. Dorsally either lacking suberect or erect vestiture or with at most sparse to moderately dense, short, suberect hair-like scales; also with moderately dense to dense, elongate-narrow, appressed to recumbent white scales laterally. Scales sparser medially and apically. Epistoma not to very slightly transversely swollen at base. Pronotum. Median carina variously developed in anterior one-half from as minutely punctate, slightly elevated broad glabrous line to slightly elevated narrow glabrous shiny line. Dorsal and lateral punctation small to moderately large, sparse to moderately dense, shallow to moderately deep; smaller and shallower apically; sparser and shallower on flanks; areas between larger dorsal punctures with dense minute regularly impressed punctures. Lacking dorsal suberect or erect



vestiture or with sparse to moderately dense, short, suberect hair-like scales each situated in large puncture (Fig. 153a). Scales dorsally absent to sparse small and fine, along median line; moderately dense to dense, elongate-narrow, appressed in pair of narrow to broad apically slightly narrowed paramedian stripes; laterally, scales absent to sparse, small and fine. Lateral margins with scales moderately dense to dense, elongate-narrow, appressed in moderately broad stripe. Flanks with scales moderately dense to dense, elongate-narrow. Median basal portion of disk shallowly to moderately deeply and broadly impressed. Anterolateral margin with pronotal postocular lobes lacking (most specimens) to moderately developed (few specimens); postocular vibrissae of unequal length, moderately long to very long, greatest length (from one-half to slightly greater than two-thirds width of eye) behind base of eye (Fig. 153a). In dorsal view with lateral margins uniformly slightly arcuate to straight and convergent from base to apical one-quarter to one-third, slightly constricted at apical one-quarter to one-third, then straight and convergent to apex. Prosternum. With small shallow to moderately deep impression and lacking or with at most only very slight rounded swelling anterior to each procoxal cavity. In lateral view with anterior flat to slightly angulate portion very short to short, approximately one-half length of posterior more angulate portion or less; posterior and anterior portions not to slightly

differentiated by a shallow transverse impression. Elytra. Moderately robust in general form (width at midlength 0.547-0.617 times length in males; 0.524-0.608 in females) (Fig. 134). In dorsal view with lateral margins very slightly sinuate to moderately and evenly arcuate and slightly convergent from apical one-third to humerus; moderately and evenly arcuate from apical one-third to apex. Elytral apices not to slightly produced, rounded to subacuminate, not to slightly divergent. Humerus distinct. Dorsally lacking suberect or erect dorsal vestiture or with at most sparse, short suberect hair-like scales on declivity; sutural interval with dense, small, fine to elongate-fine white (rarely golden) scales; interval 2 with scales very sparse to dense small and fine; intervals 3 to 5 with scales dense large elongate-narrow; intervals 6 to 8 with scales absent to moderately dense small and fine to elongate-fine; intervals 9 to 11 with scales dense, large elongate-narrow; striae 1, 6 and 7 with irregularly distributed patches of larger and denser white scales (Fig. 134). Punctures of elytral striae small, individually distinct, arranged in regular rows. Wings. Long (greater than elytra in length). Legs. Foretibia of female with inner margin with small to large denticles in apical three-quarters; subapical tooth small to large, not to distinct from tibial denticles. Foretibial and mesotibial unci of both sexes moderately large, metatibial unci small. Tarsal claws connate in basal one-third, slightly divergent,

each with basal internal flange well-developed. Ventral tarsal pilose vestiture of foretarsus and mesotarsus of male present as small elongate pads covering apical one-third of article 1, as rounded pads covering apical one-half of article 2, and as large rounded pads covering apical three-quarters of article 3; of foretarsus and mesotarsus of female as apical tufts to small elongate-narrow pads covering apical one-third of article 1, as rounded pads covering apical one-quarter to one-half of article 2, and as large rounded pads covering apical one-half to two-thirds of article 3; of metatarsus of male as elongate-narrow pad covering apical one-quarter on inner lobe and as apical tuft on outer lobe of article 1, as elongate pads covering apical one-half of article 2, as moderately large elongate-oval pads covering apical one-half to three-quarters on article 3; of metatarsus of female as apical tuft to elongate-narrow pad covering apical one-quarter on inner lobe and as elongate-narrow pad covering apical one-half on outer lobe of article 1, as elongate pads covering apical one-quarter to three-quarters of article 2, as elongate-oval pads covering apical one-third to one-half on article 3. **Abdomen.** Ventral surface with moderately dense, elongate-narrow to moderately robust appressed white scales. Abdominal sterna III to VI (especially sterna III and IV) with distinct rounded glabrous patches, each with large puncture and single appressed to suberect moderately long hair-like scale situated in or near the center. Base of abdominal sternum

VII of female internally (dorsally) with very slightly developed median shiny tubercle. Genitalia. Female (six examined). Abdominal sternum VIII with basal arm short; lateral arms moderately inwardly arcuate throughout length; not to slightly expanded at apex (Fig. 173). Stylus moderately large compared to length gonocoxite II. Male (six examined). Internal sac (Figs. 193a,b) elongate and low; median dorsal pocket low, with single moderately broad dorsally directed basal lobe (lobe A), with small median dorsally directed lobe at midlength on dorsal surface (lobe B), with moderately large paired dorsally directed lobe at basal one-third on dorsal surface (lobe C), with moderately large paired markedly dorsolaterally directed lobe near midlength near dorsal margin (lobe D), with small paired dorsolaterally directed lobe just beyond midlength at dorsal margin (lobe E), and with small ventrally directed paired lobe near base near ventral margin (lobe F). Apical pocket and median dorsal pocket individually distinct; apex of dorsal median pocket slightly emarginate medially in dorsal view (Fig. 193b). Ventral median pocket large. Apical pocket with paired moderately large ventrally directed lobe near apical one-third at ventral margin. Eversible apical sclerite complex with paired simple scythe-like sclerite; adjacent ventral surface and basal portion of sides of apical pocket with pair of large, dark, longitudinal sclerites.

**Geographic variation.**--- Individuals from the northern part of the species range (especially Alberta, Canada) possess moderately developed postocular lobes and correspondingly shorter postocular vibrissae. Throughout the rest of the species range, lobes are lacking or only slightly developed and postocular vibrissae are longer.

In the western part of the species range in California and Nevada (especially the former), individuals possess short suberect dorsal vestiture on the head and pronotum. Throughout the rest of the species range, dorsal suberect or erect dorsal vestiture is lacking or at most very short and indistinct.

**Geographic distribution.**--- This species is widespread in the western United States of America and Canada from extreme southern British Columbia south to southern California in the west, and from southern Manitoba, south through eastern South Dakota, Kansas to western Texas (Fig. 220).

**Natural history.**--- This species appears to be associated primarily with grassland and desert-grassland transitional habitats throughout the species range. Adults have been collected exclusively on Leguminosae as follows; *Astragalus* spp. (as "loco" or "locoweed"), *A. bisulcatus* (Hook.) Gray var. *haydenianus*, *A. earlei* Greene ex. Rydb., *A. flavus* Nutt., *A. utahensis* Torr. nad Gray, *A. wootoni* Sheldon, *Oxytropis lambertii* Pursh, and *O. sericea* Nutt.. Adults have

been reared from *Astragalus* sp. (as "locoweed"). Adults have been collected throughout the year at elevations of from 131-3640 m (X=1768 m, N=42) (Fig. 233).

A single specimen was found in the stomach contents of *Bufo* sp. (Amphibia).

**Chorological relationships.**--- Table 67.

**Phylogenetic relationships.**--- Primary characters indicate only that this species is a member of an unresolved quadrichotomy composed of the four species in the *C. boucardi* group (Fig. 236). Distribution of the apotypic state of the secondary character of length of dorsal vestiture place this species as the sister-species to the *C. quadrilineatus*-*C. placidus* lineage in the *C. boucardi* group (Fig. 237).

Table 61. Descriptive statistics for *C. trivittatus*  
based on eight males from Albuquerque, New Mexico

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	2.32-2.72	2.565	0.253	0.119
WF	1.60-1.88	1.740	0.190	0.089
WRA	1.44-1.76	1.585	0.178	0.084
LP	3.92-4.72	4.270	0.374	0.176
WPB	4.32-5.52	4.670	0.577	0.272
WE1H	4.80-6.08	5.330	0.608	0.287
WE1M	5.44-6.64	5.900	0.604	0.285
LE1	9.28-11.68	10.400	1.126	0.531
LEy	1.08-1.52	1.270	0.212	0.100
WEy	0.60-0.84	0.735	0.120	0.056
B. Proportions.				
WEy/LEy	0.55-0.63	0.579	0.035	0.017
WPB/LP	1.06-1.17	1.093	0.055	0.026
WRA/LR	0.58-0.65	0.618	0.031	0.015
WRA/WF	0.87-0.98	0.911	0.051	0.024
WE1M/LE1	0.55-0.59	0.568	0.018	0.009
LP/LE1	0.39-0.42	0.411	0.017	0.008

Table 62. Descriptive statistics for *C. trivittatus*  
based on eight females from Albuquerque, New Mexico

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.92-2.96	2.450	0.495	0.350
WF	1.32-1.84	1.670	0.279	0.132
WRA	1.16-1.68	1.500	0.246	0.116
LP	3.28-4.56	4.120	0.714	0.337
WPB	3.36-5.04	4.440	0.889	0.419
WE1H	3.76-6.16	5.170	1.202	0.566
WE1M	4.32-6.64	5.700	1.191	0.561
LE1	7.68-11.20	9.940	1.789	0.843
LEy	0.96-1.44	1.225	0.231	0.109
WEy	0.56-0.80	0.710	0.119	0.056
B. Proportions.				
WEy/LEy	0.56-0.61	0.581	0.030	0.014
WPB/LP	1.02-1.12	1.076	0.047	0.022
WRA/LR	0.57-0.68	0.615	0.061	0.029
WRA/WF	0.86-0.95	0.899	0.040	0.019
WE1M/LE1	0.55-0.59	0.572	0.024	0.011
LP/LE1	0.41-0.43	0.415	0.016	0.007



4.5.2.4.3 *Cleonidius placidus* Csiki

(Figs. 135, 154, 174, 194, 222)

*Cleonus pacificus* Fall 1901:260 [not Olivier 1807:268].

Holotype (examined), male, labelled "Redondo/  
Cal.4.7.98", "TYPE/ *pacificus*", "M.C.Z./ Type/ 25194",  
"H.C.FALL/ COLLECTION", "*Cleonus/ pacificus/ Fall*"  
(MCZC). Type locality, Redondo, California. Leng 1920  
(catal.).

*Cleonus (Cleonidius) placidus* Csiki 1934:65. New name for  
*Cleonus pacificus* Fall. Blackwelder 1939 (check.).

*Cleonis pacificus*; O'Brien and Wibmer 1982 (catal., distn.,  
as jr. homonym).

*Cleonis placidus*; O'Brien and Wibmer 1982 (catal., distn.).

**Notes about synonymy.** --- This species is commonly known as  
*C. pacificus* (Fall). This name is a junior homonym and *C.*  
*placidus* Csiki is the valid name. The holotype of *Cleonus*  
*Inornatus* LeConte from Owen's Valley, California has an  
elytral scale pattern similar to that of *C. placidus* (Fig.  
135) but differs in other structural features (possession of  
short dorsal suberect vestiture of the head and pronotum,  
slightly developed postocular lobes, and less extensive  
ventral tarsal pilosity) that are found in Californian *C.*  
*trivittatus*. It is herein regarded as a synonym of *C.*  
*trivittatus* (see also "Notes about synonymy" section under  
*C. trivittatus*).

Problems in recognition.--- Most individuals of this species are easily recognized by the lack of, or at most only very slightly developed postocular lobes (Fig. 154a), moderately long dorsal erect vestiture of the head and pronotum (Fig. 154a), and distinctive elytral scale pattern (Fig. 135). From *C. trivittatus*, most individuals can be separated by the elytral scale pattern and also by the presence of moderately long dorsal erect vestiture on the head and pronotum in *C. placidus*; in *C. trivittatus*, dorsal erect or suberect vestiture is lacking or at most only suberect and short (Fig. 153a). The two species are also largely allopatric in their distributions; *C. placidus* is known only from California (Fig. 222), whereas *C. trivittatus* is widely distributed in western North America, but known from only a few localities in California (Fig. 220). These few *C. trivittatus* from California are difficult to separate from *C. placidus* however most of them possess short erect dorsal vestiture, whereas this vestiture is slightly longer on the head and pronotum of most *C. placidus*; they also possess less extensive ventral tarsal pilosity, slightly developed postocular lobes, and lack the distinctive elytral scale pattern of *C. placidus*. Both species are known only from Leguminosae.

*Cleonidius placidus* individuals may also be confused with individuals of *C. quadrilineatus*, but as for *C. trivittatus* specimens, the distinctive scale pattern of *C. placidus* should serve to separate the two (Fig. 135).

are only found in California, whereas *C. quadrilineatus* are known from Rosaceae and Rhamnaceae (rarely from Leguminosae) and are widespread in southwestern North America (including California). Most individuals of *C. placidus* are also slightly larger than are most *C. quadrilineatus* (see Fig. 197).

**Description.** --- **Specimens examined.** 57 males, 49 females. Data about variation in LR, WF, WRA, LP, WPB, WE1H, WE1M, LE1, LEy, WEy, WEy/LEy, WPB/LP, WRA/LR, WRA/WF, WE1M/LE1, and LP/LE1 are presented in Tables 63 and 64. **Size.** Length, male, 9.9-17.1 mm; female, 12.6-17.6 mm. Width, male, 3.7-6.2 mm; female, 4.3-6.7 mm. **Head.** Eye elongate-oval. Area behind eye with numerous slight wrinkles of cuticle. **Frons** with punctures sparse to moderately dense, small to moderately large, shallow, some longitudinally confluent immediately above eyes; interspersed with dense minute punctures. Frons also with moderately dense, moderately long, erect hair-like scales and sparse to dense, elongate-narrow appressed white scales medially, erect immediately above eyes. Frons continuous with base of rostrum or only slightly separated by shallow transverse impression (Fig. 154a). Base of median tumescence of rostrum with small, shallow fovea. Width of frons greater than width at apex of rostrum. **Rostrum.** Moderately robust, (width at apex 0.586-0.645 times length in male; 0.568-0.618 in

downward. In dorsal view with postgenae slightly laterally expanded, antennal scrobes visible. Rostrum slightly to moderately medially tumescent from point of antennal insertion to middle of frons (Fig. 154a); not laterally expanded at midlength; lateral margins rounded; with median carina variously developed as rounded to sharp, narrow low glabrous shiny line at crest of median tumescence. Dorsal and lateral punctation sparse to moderately dense, small to moderately large, shallow to moderately deep; punctures smaller and irregularly impressed apically, in some specimens dorsally interspersed with dense minute punctures. Dorsally with moderately dense, moderately long to long, erect hair-like scales; also with sparse, elongate-narrow, recumbent white scales laterally, scales sparser medially and apically. Epistoma not to very slightly transversely swollen at base. Pronotum. Median carina variously developed in anterior one-half from as minutely punctate, slightly elevated broad glabrous line to slightly elevated narrow glabrous shiny line, well-developed and distinct in most specimens. Dorsal and lateral punctation moderately large, moderately dense to dense, moderately deep; smaller and shallower apically; sparser, smaller and shallower on flanks; areas between larger dorsal punctures with dense minute regularly impressed punctures. Dorsally, laterally and medially with moderately dense, moderately long to long, erect hair-like scales each situated in large puncture (Fig.

moderately dense to dense, elongate-narrow, appressed in pair of narrow apically convergent and slightly narrowed paramedian stripes; laterally, scales absent. Lateral margins and flanks with scales moderately dense to dense, elongate-narrow. Median basal portion of disk moderately deeply to deeply and broadly impressed. Anterolateral margin with pronotal postocular lobes lacking (most specimens) to only very slightly developed (few specimens); postocular vibrissae of unequal length, very long, greatest length (slightly greater than two-thirds width of eye) behind base of eye (Fig. 154a). In dorsal view with lateral margins uniformly very slightly arcuate and convergent from base to apical one-quarter, slightly constricted at apical one-quarter, then straight and convergent to apex.

**Prosternum.** With small shallow to moderately deep impression and lacking or with at most only very slight rounded swelling anterior to each procoxal cavity. In lateral view with anterior flat to slightly angulate portion very short, indistinct. **Elytra.** Moderately elongate-narrow in general form (width at midlength 0.508-0.555 times length in males; 0.500-0.567 in females) (Fig. 135). In dorsal view with lateral margins straight to very slightly sinuate and slightly convergent from apical one-third to humerus; moderately and evenly arcuate from apical one-third to apex. Elytral apices not to slightly produced, rounded to subacuminate, not to slightly divergent. Humerus distinct.

evident on declivity); sutural interval with sparse to moderately dense, small fine white scales; interval 2 with irregularly distributed patches of dense, elongate-fine white scales, otherwise with scales absent to sparse, small and fine; intervals 3 to 11 with scales uniformly moderately dense to dense, elongate-fine to elongate-narrow (some specimens with irregularly distributed patches of even larger denser, white scales, especially on intervals 9 to 11) (Fig. 135). Punctures of elytral striae small, individually distinct, arranged in regular rows. Wings. Long (greater than elytra in length). Legs. Foretibia of female with inner margin with small denticles in apical three-quarters; subapical tooth not distinct from tibial denticles. Foretibial and mesotibial unci of both sexes moderately large, metatibial unculus small. Tarsal claws connate in basal one-third, slightly divergent, each with basal internal flange well-developed. Ventral tarsal pilose vestiture of foretarsus and mesotarsus of male present as elongate pads covering apical two-thirds of article 1, as large rounded pads covering apical two-thirds of article 2, and as large rounded pads covering more or less entire ventral surface of article 3; of foretarsus of female as small elongate pads covering apical one-third of article 1, as rounded pads covering apical one-half of article 2, as large rounded pads covering apical three-quarters of article 3; of mesotarsus of female as small elongate pads covering

pads covering apical one-half of article 2, as large rounded pads covering apical two-thirds of article 3; of metatarsus of male as elongate pads covering apical one-half to two-thirds of article 1, as large rounded pads covering apical two-thirds of article 2, as large rounded pads covering more or less entire ventral surface of article 3; of metatarsus of female as elongate-narrow pads covering apical one-quarter to two-thirds of article 1, as elongate pads covering apical one-half to two-thirds of article 2, as large rounded pads covering apical two-thirds of article 3.

**Abdomen.** Ventral surface with moderately dense to dense, elongate-narrow to moderately robust appressed white scales; scales sparser along midline on sternum VII of female.

Abdominal sterna III to VI (especially sterna III and IV) with distinct large rounded glabrous patches, each with large puncture and single recumbent to suberect moderately long hair-like scale situated in or near the center. Base of abdominal sternum VII of female internally (dorsally) with very slightly developed median shiny tubercle. **Genitalia.**

Female (two examined). Abdominal sternum VIII with basal arm moderate in length; lateral arms moderately inwardly arcuate at midlength; slightly expanded at apex (Fig. 174). Stylus moderately large compared to length gonocoxite II. Male (two examined). Internal sac (Figs. 194a,b) elongate and low; median dorsal pocket low, with single moderately broad dorsally directed basal lobe (lobe A), with small median

B), with moderately large paired dorsally directed lobe at basal one-third on dorsal surface (lobe C), with moderately large paired dorsolaterally directed lobe near midlength near dorsal margin (lobe D), with moderately large paired dorsolaterally directed lobe just beyond midlength at dorsal margin (lobe E), and with small ventrally directed paired lobe near base near ventral margin (lobe F). Apical pocket and median dorsal pocket individually distinct; apex of dorsal median pocket bisinuate medially in dorsal view (Fig. 194b). Ventral median pocket moderately large. Apical pocket with paired moderately large ventrally directed lobe near apical one-third at ventral margin. Eversible apical sclerite complex with paired simple scythe-like sclerite; adjacent ventral surface and basal portion of sides of apical pocket with pair of large, dark, longitudinal sclerites.

**Geographic variation.**--- None noted.

**Geographic distribution.**--- This species is known only from California in the coastal and central valleys from San Francisco south to San Diego (Fig. 222).

**Natural history.**--- This species is found only in the Pacific semi-desert region. Adults have been collected almost exclusively from *Astragalus* spp. (Leguminosae). A



L. (Umbelliferae). Adults have also been reared from roots of *Astragalus* sp.. Adults have been collected throughout the year at elevations of from 9-1747 m ( $X=327$  m,  $N=10$ ) (Fig. 233).

Chorological relationships.--- Table 67.

Phylogenetic relationships.--- Primary characters indicate only that this species is a member of an unresolved quadrichotomy composed of the four species in the *C. boucardi* group (Fig. 236). Distribution of the apotypic state of the secondary character of length of dorsal vestiture place this species as the sister-species of *C. quadrilineatus*, the two species forming a lineage that is the sister group of *C. trivittatus* (Fig. 237).

on 10 males from Panoche, California

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	2.40-2.80	2.592	0.240	0.101
WF	1.56-1.84	1.692	0.174	0.074
WRA	1.48-1.72	1.596	0.134	0.057
LP	3.60-4.32	4.008	0.364	0.154
WPB	4.00-4.88	4.408	0.465	0.196
WE1H	4.80-5.76	5.304	0.463	0.195
WE1M	5.20-6.48	5.776	0.699	0.295
LE1	9.92-11.68	10.800	1.104	0.466
LEy	1.28-1.48	1.364	0.125	0.053
WEy	0.72-0.84	0.764	0.052	0.022
B. Proportions.				
WEy/LEy	0.53-0.59	0.561	0.032	0.013
WPB/LP	1.04-1.18	1.100	0.072	0.030
WRA/LR	0.59-0.65	0.616	0.022	0.009
WRA/WF	0.91-1.00	0.945	0.048	0.020
WE1M/LE1	0.51-0.56	0.535	0.024	0.010
LP/LE1	0.35-0.40	0.372	0.021	0.009

on eight females from Panoche, California

Character	Range	Mean	1.5SD	2SE
-----------	-------	------	-------	-----

A. Measurements, in mm.

LR	2.12-2.96	2.565	0.439	0.207
WF	1.32-2.00	1.725	0.307	0.145
WRA	1.24-1.68	1.510	0.228	0.108
LP	3.12-4.56	4.020	0.669	0.315
WPB	3.36-4.96	4.290	0.793	0.374
WE1H	4.16-6.08	5.250	0.916	0.432
WE1M	4.64-6.80	5.760	0.996	0.469
LE1	8.80-12.48	11.000	1.799	0.848
LEY	1.08-1.48	1.325	0.191	0.090
WEy	0.68-0.80	0.740	0.079	0.037

B. Proportions.

WEy/LEY	0.52-0.63	0.561	0.060	0.028
WPB/LP	1.00-1.11	1.066	0.056	0.026
WRA/LR	0.57-0.62	0.590	0.024	0.011
WRA/WF	0.84-0.94	0.878	0.060	0.028
WE1M/LE1	0.50-0.57	0.524	0.030	0.014
LP/LE1	0.35-0.39	0.366	0.024	0.011

(Figs. 136, 155, 175, 195, 224)

*Apleurus quadrilineatus* Chevrolat 1873:80. Holotype (examined), male, labelled "66", "ex coll./ Sallè", "MUSEUM PARIS/ 1952/ COLL R OBERTHUR", "*apleurus/ quadrilineatus/ Chev. type texas*", and with an inverted label "*Lixus/ praepotens/ Boh. Texas*", and with a label indicating it as the holotype "*Apleurus/ quadrilineatus/ Chev. HOLOTYPE/ labelled Anderson*" (MNHP). Type locality, Texas.

*Cleonus quadrilineatus*; LeConte 1876a (key, diag.). Henshaw 1885 (check.). Wickham 1889. Wickham 1896 (check.). Fall 1897. Fall 1901. Wickham 1902 (check.). Fall and Cockerell 1907 (check.). Pierce 1907 (biol., misident. of *Cleonidius trivittatus*). Chittenden 1911 (biol., misident. of *Cleonidius trivittatus*). Anderson 1914 (misident. of *Cleonidius lobigerinus*). Blatchley and Leng 1916 (figure, misident. of *Cleonidius trivittatus*). Yothers 1916 (prob. misident. of *Cleonidius trivittatus*, in part). Leng 1920 (catal.). Essig 1958 (biol., misident. of *Cleonidius trivittatus*). Arnett et al. 1980 (key, redesc., biol., misident. of *Cleonidius trivittatus*).

*Lixus texanus*; LeConte 1876a (in part, mixed type series).

*Cleonus (Cleonidius) quadrilineatus*; Casey 1891 (key). Csiki 1934 (catal.). Hatch 1971 (key, redesc., misident. of *Cleonidius lobigerinus*).

distn.).

**Notes about synonymy.**--- The holotype of *C. quadrilineatus* is the typical elongate-narrow Rosaceae form. Many literature references to *C. trivittatus* refer to this species.

**Problems in recognition.**--- Specimens of *C. quadrilineatus* are very likely to be commonly confused with those of the broadly sympatric *C. trivittatus*. *Cleonidius quadrilineatus* specimens can be separated from the latter by their more elongate-narrow elytral form (Fig. 136) and by presence of moderately long erect hair-like scales on the pronotal disk (Fig. 155a), which in *C. trivittatus* are at most only short, suberect and indistinct (Fig. 153a). Most *C. quadrilineatus* are also associated with Rosaceae or Rhamnaceae, although rarely also with Leguminosae, whereas all *C. trivittatus* are associated with Leguminosae. Some *C. trivittatus* have slightly to moderately developed postocular lobes (especially from the northern part of the range) whereas all *C. quadrilineatus* lack postocular lobes entirely (Fig. 155a).

In California, specimens of *C. quadrilineatus* may be confused with *C. placidus*, especially specimens of the latter in which the distinctive elytral scale pattern (Fig.

long, erect hair-like scales on the pronotum (Fig. 154a), as do *C. quadrilineatus*; but are associated exclusively with Leguminosae. *C. placidus* individuals are also slightly larger than are those of *C. quadrilineatus* (Fig. 197) and have slightly more extensive ventral tarsal pilosity.

Description.--- Specimens examined. 108 males, 107 females. Data about variation in LR, WF, WRA, LP, WPB, WE1H, WE1M, LE1, LEy, WEy, WEy/LEy, WPB/LP, WRA/LR, WRA/WF, WE1M/LE1, and LP/LE1 are presented in Tables 65 and 66. Size. Length, male, 8.3-14.3 mm; female, 9.3-13.1 mm. Width, male, 2.9-5.3 mm; female, 3.2-4.8 mm. Head. Eye elongate-oval. Area behind eye with numerous slight wrinkles of cuticle. Frons with punctures sparse to moderately dense, small to moderately large, shallow; some large and longitudinally confluent immediately above eyes; interspersed with minute punctures. Frons also with sparse to moderately dense, elongate-narrow appressed white scales medially, erect immediately above eyes. Frons continuous with base of rostrum or separated from base of rostrum by shallow transverse impression (Fig. 155a). Base of median tumescence of rostrum with small, shallow fovea. Width of frons greater than to subequal to width at apex of rostrum. Rostrum. Moderately robust, (width at apex 0.604-0.703 times length in male; 0.582-0.690 in female) (Fig. 155). In lateral view very slightly curved downward. In dorsal view with postgenae slightly laterally

moderately medially tumescent from point of antennal insertion to middle of frons (especially basally) (Fig. 155a); not laterally expanded at midlength; lateral margins rounded; with median carina variously developed as narrow low glabrous shiny line at crest of median tumescence. Dorsal and lateral punctation sparse to dense, small to moderately large, shallow; punctures smaller apically; dorsally interspersed with dense minute punctures. Dorsally with moderately dense, moderately long, suberect to erect, hair-like scales (especially laterally); also with moderately dense, elongate-narrow, recumbent white scales laterally, scales absent to sparse medially and apically. Epistoma not to very slightly transversely swollen at base. Pronotum. Median carina variously developed in anterior one-half from as minutely punctate, slightly elevated broad glabrous line to slightly elevated narrow glabrous shiny distinct line. Dorsal and lateral punctation moderately large, moderately dense, moderately deep; smaller and shallower apically; sparser and shallower on flanks; areas between larger dorsal punctures with dense minute regularly impressed punctures. Dorsally, laterally and medially, with moderately dense, moderately long, erect hair-like scales each situated in large puncture (Fig. 155a). Scales dorsally absent to sparse small and fine along median line; moderately dense, elongate-fine to elongate-narrow, appressed in pair of narrow apically slightly narrowed

small and fine. Lateral margins with scales moderately dense to dense, elongate-narrow to moderately robust, appressed in moderately broad stripe. Flanks with scales moderately dense, elongate-fine to elongate-narrow. Median basal portion of disk shallowly to moderately deeply and broadly impressed. Anterolateral margin with pronotal postocular lobes lacking; postocular vibrissae of unequal length, very long, greatest length (greater than two-thirds width of eye) behind base of eye (Fig. 155a). In dorsal view with lateral margins uniformly slightly arcuate and convergent from base to apex, to straight and convergent from base to apical one-quarter to one-third, slightly constricted at apical one-quarter to one-third, then straight and convergent to apex. Prosternum. With small shallow to moderately deep impression and slight rounded swelling anterior to each procoxal cavity. In lateral view with anterior flat to slightly angulate portion very short, indistinct. Elytra. Elongate-narrow in general form (width at midlength 0.482-0.545 times length in males; 0.477-0.524 in females) (Fig. 136). In dorsal view with lateral margins straight to very slightly sinuate slightly convergent from apical one-third to humerus; moderately and evenly arcuate from apical one-third to apex. Elytral apices not to slightly produced, rounded to subacuminate, not to slightly divergent. Humerus distinct. Dorsally with sparse, short suberect hair-like scales (most evident on declivity);



scales; interval 2 with scales very sparse to dense small and fine, white; intervals 3 to 5 with scales dense large elongate-narrow, white; intervals 6 to 8 with scales sparse to moderately dense, small and fine to elongate-fine, white; intervals 9 to 11 with scales dense, large elongate-narrow, white; striae 1, 6 and 7 with irregularly distributed patches of larger and denser white scales (Fig. 136).

Punctures of elytral striae small, individually distinct, arranged in regular rows. Wings. Long (greater than elytra in length). Legs. Foretibia of female with inner margin with small to moderately large denticles in apical three-quarters; subapical tooth indistinct from tibial denticles. Foretibial and mesotibial unci of both sexes moderately large, metatibial uncus small. Tarsal claws connate in basal one-third, slightly divergent, each with basal internal flange well-developed. Ventral tarsal pilose vestiture of foretarsus and mesotarsus of male present as small elongate pads covering apical one-half of article 1, as large, elongate-oval pads covering apical two-thirds of article 2, and as large rounded pads covering apical three-quarters of article 3; of foretarsus and mesotarsus of female as as small elongate-narrow pads covering apical one-quarter of article 1, as small rounded pads covering apical one-third of article 2, and as moderately large rounded pads covering apical one-half to two-thirds of article 3; of metatarsus of male as elongate-narrow pad.

on outer lobe of article 1, as elongate pads covering apical one-half to two-thirds of article 2, as large rounded pads covering apical three-quarters on article 3; of metatarsus of female as elongate-very-narrow pad covering apical one-half on inner lobe and apical one-third on outer lobe of article 1, as elongate pads covering apical one-third to one-half of article 2, as moderately large elongate-oval pads covering apical one-half to two-thirds on article 3.

**Abdomen.** Ventral surface with moderately dense, elongate-narrow to moderately robust appressed white scales. Abdominal sterna III to VI (especially sterna III and IV) with distinct rounded glabrous patches, each with large puncture and single appressed to erect moderately long hair-like scale situated in or near the center. Base of abdominal sternum VII of female internally (dorsally) with slightly to moderately developed median shiny tubercle.

**Genitalia.** Female (four examined). Abdominal sternum VIII with basal arm short; lateral arms moderately inwardly arcuate throughout length; not to slightly expanded at apex (Fig. 175). Stylus moderately large compared to length gonocoxite II. Male (four examined). Internal sac (Figs. 195a,b) elongate and low; median dorsal pocket low, with single moderately broad dorsally directed basal lobe (lobe A), with small median dorsally directed lobe at midlength on dorsal surface (lobe B), with moderately large paired dorsally directed lobe at basal one-third on dorsal surface

dorsolaterally directed lobe near midlength near dorsal margin (lobe D), with small paired dorsolaterally directed lobe just beyond midlength at dorsal margin (lobe E), and with small ventrally directed paired lobe near base near ventral margin (lobe F). Apical pocket and median dorsal pocket individually distinct; apex of dorsal median pocket slightly emarginate medially in dorsal view (Fig. 195b). Ventral median pocket large. Apical pocket with paired moderately large ventrally directed lobe near apical one-third at ventral margin. Eversible apical sclerite complex with paired simple scythe-like sclerite; adjacent ventral surface and basal portion of sides of apical pocket with pair of large, dark, longitudinal sclerites.

**Geographic variation.**--- None noted.

**Geographic distribution.**--- This species is found from California east to eastern Nebraska, Oklahoma, and central Texas (Fig. 224). There is a single Mexican record from near Zacatecas.

**Natural history.**--- This species is found primarily in the western xerophytic evergreen forest region but also in desert-grassland transitional habitats. Adults of this species have been collected primarily on Rosaceae or Rhamnaceae, especially in oak-pinyon-juniper woodlands

Arizona. Adults have been collected on *Astragalus wootoni* Sheldon, *Lupinus* sp. (Leguminosae); *Ceanothus* sp. (Rhamnaceae); *Cercocarpus montanus* Raf., *Rosa* sp. (Rosaceae). Adults have been reared only from *Astragalus* sp. roots at Sulphur Springs Valley, Arizona. I suspect that a variety of Rosaceae or Rhamnaceae serve as hosts at higher altitudes throughout the species range. Adults have been collected from January to October at elevations from 15-4186 m ( $\bar{X}=1729$  m,  $N=63$ ) (Fig. 233).

Adults have been found in the stomach contents of *Geococcyx californicus* (roadrunner; Aves) at Carlsbad, New Mexico.

Myself and others have collected this species and *C. texanus* together on various Rosaceae and Rhamnaceae in Texas and Arizona.

Chorological relationships.--- Table 67.

Phylogenetic relationships.--- Primary characters indicate only that this species is a member of an unresolved quadrichotomy composed of the four species in the *C. boucardi* group (Fig. 236). Distribution of the apotypic state of the secondary character of length of dorsal vestiture place this species as the sister-species of *C. placidus*, the two species forming a lineage that is the sister group of *C. trivittatus* (Fig. 237).

based on eight males from southern Arizona localities

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.84-2.36	2.055	0.270	0.127
WF	1.24-1.52	1.365	0.185	0.087
WRA	1.16-1.48	1.320	0.181	0.086
LP	3.04-3.76	3.380	0.434	0.204
WPB	3.28-4.08	3.620	0.496	0.234
WE1H	3.84-5.04	4.360	0.609	0.287
WE1M	4.32-5.76	4.850	0.714	0.337
LE1	8.48-10.56	9.500	1.065	0.502
LEy	1.00-1.28	1.140	0.128	0.060
WEy	0.60-0.76	0.670	0.077	0.036
B. Proportions.				
WEy/LEy	0.53-0.61	0.588	0.038	0.018
WPB/LP	1.05-1.09	1.071	0.022	0.010
WRA/LR	0.60-0.67	0.643	0.035	0.017
WRA/WF	0.91-1.07	0.968	0.084	0.040
WE1M/LE1	0.48-0.55	0.510	0.027	0.013
LP/LE1	0.34-0.37	0.356	0.014	0.007

based on eight females from southern Arizona localities

Character	Range	Mean	1.5SD	2SE
A. Measurements, in mm.				
LR	1.68-2.20	1.985	0.353	0.166
WF	1.04-1.52	1.325	0.269	0.127
WRA	1.00-1.40	1.230	0.207	0.098
LP	2.48-3.68	3.250	0.672	0.317
WPB	2.72-3.92	3.440	0.670	0.316
WE1H	3.20-4.88	4.190	0.918	0.433
WE1M	3.60-5.36	4.670	0.985	0.464
LE1	7.20-10.40	9.220	1.809	0.853
LEy	0.92-1.20	1.110	0.153	0.072
WEy	0.56-0.72	0.655	0.084	0.040
B. Proportions.				
WEy/LEy	0.55-0.64	0.591	0.046	0.022
WPB/LP	1.02-1.10	1.060	0.043	0.020
WRA/LR	0.58-0.69	0.621	0.050	0.023
WRA/WF	0.84-1.00	0.932	0.076	0.036
WE1M/LE1	0.49-0.52	0.506	0.019	0.009
LP/LE1	0.34-0.37	0.352	0.015	0.007

This category is used herein for a single species of uncertain placement, *Cleonidius vibex* (Pallas). Although adults of *C. vibex* have a vittate elytral scale pattern, lack extensive ventral tarsal pilose pads, possess a moderately broad rostrum, and are brachypterous, all of which suggest inclusion in the *C. americanus* group, the nature of formation of the vittate scale pattern is fundamentally different from that in other *Cleonidius* and thus not homologous. In other *Cleonidius* the vittate pattern is produced as a result of differing sizes and densities of scales on the various elytral intervals; in *C. vibex*, the pattern is produced as a result of presence or absence of white pigment in the individual scales. This suggests independent derivation of a vittate elytral pattern and perhaps of states of other characters as well. Furthermore autapotypic states of lack of larger punctures on the rostrum and head, and widely divergent tarsal claws suggests a primitive position in the genus. Placement of *C. vibex* as sister to the remainder of *Cleonidius* can thus only be substantiated by autapotypic features in that species and not by synapotypy in the remainder of *Cleonidius*, and by the adaptive argument concerning the fundamentally different vittate elytral scale pattern.

(Figs. 156, 176, 196)

*Curculio vibex* Pallas 1781:32. Types not examined. Herbst  
1795.

*Lixus denudatus* Zoubkoff 1833:319. Types not examined. Faust  
1890. Petri 1912.

*Lixus karelini* Boheman 1836:22. Types not examined. Faust  
1890. Petri 1912.

*Lixus vibex*; Boheman 1836. Capiomont and Leprieur 1874.  
Faust 1890. Petri 1912. Lukjanovitsh 1926.

*Lixus vibex scutellaris* Petri 1905:103. Types not examined.

*Lixus (Lixestus) vibex*; Reitter 1916. Csiki 1934.

*Lixus (Lixesthus) vibex*; Ter-Minasyan 1978.

**Notes about synonymy.**--- No types have been examined by me  
of this species or its synonyms. Synonymy presented here  
follows Petri (1912) and Csiki (1934).

Two additional species have been placed in *Lixus*  
(*Lixestus*) by Ter-Minasyan (1978). These are *L. pallas*  
Faust and *L. meles* Boheman. These species will likely also  
prove to be *Cleonidius* but representatives have not been  
examined by me.

**Problems in recognition.**--- Individuals of this species are  
easily distinguished from Nearctic *Cleonidius* by the widely  
divergent tarsal claws each lacking a basal internal flange,



The rostrum of this species has uniformly very dense and small punctures, but no or else very few larger punctures typical of Nearctic species of *Cleonidius*. Individuals also differ from those of Nearctic species in that the contrasting dark and light areas of the pronotum and elytra are a result of apparently having white scales, such that the underlying dark cuticle is obscured, or unpigmented, such that the dark cuticle is not obscured. Individuals of Nearctic species on the other hand, have a vittate elytral scale pattern and a pronotal scale pattern that is a result of differing sizes of scales; small scales do not obscure the dark cuticle, whereas larger scales do.

The robust, downwardly curved rostrum and elytral scale pattern of individuals of this species result in superficial resemblance to *C. canescens*.

**Description.**--- Specimens examined. 1 male, 1 female. **Size.** Length, male, 7.3 mm; female, 7.1 mm. Width, male, 2.7 mm; female, 2.6 mm. **Head.** Eye elongate-oval. Area behind eye with numerous slight wrinkles of cuticle. Frons with only uniformly very dense minute punctures. Frons also with very dense, truncate grey-brown appressed scales. Frons continuous with base of rostrum (Fig. 156a). Base of median tumescence or carina of rostrum (if present) with small, shallow fovea. Width of frons greater than width at apex of rostrum. **Rostrum.** Robust, (width at apex 0.73 times length

moderately curved downward. In dorsal view with postgenae very slightly laterally expanded, antennal scrobes distinctly visible. Rostrum markedly medially tumescent from point of antennal insertion to middle of frons (Fig. 156a); not laterally expanded at midlength; lateral margins more or less rounded, not sharp; with median carina absent. Dorsal and lateral punctation very sparse, small, shallow; sparser apically and medially; dorsally interspersed with minute punctures. Dorsally lacking suberect or erect vestiture, with only uniformly very dense, appressed truncate grey-brown scales (almost completely obscuring punctation), sparser apically and medially. Epistoma not swollen.

Pronotum. Median carina lacking. Dorsal and lateral punctation sparse, small, shallow, interspersed with very dense minute regularly impressed punctures. Suberect or erect vestiture lacking. Scales dorsally uniformly dense small truncate appressed medially; laterally transparent in pair of narrow stripes (appearing black). Lateral margins and flanks with scales uniformly dense, small, truncate and appressed, almost completely obscuring punctation. Median basal area of disk broadly and very shallowly impressed. Anterolateral margin with pronotal postocular lobes well-developed; postocular vibrissae short, of uniform length (Fig. 156a). In dorsal view with lateral margins straight and convergent from apical one-third to base; slightly arcuate from apical one-third to apex; subapical

moderately deep impression anterior to each procoxal cavity; swelling lacking. In lateral view with anterior flat to slightly angulate portion moderately long, approximately one-half length of posterior more angulate portion; anterior and posterior portions slightly differentiated by shallow transverse impression. Elytra. Moderately elongate-narrow in general form (width at midlength 0.52 times length in male; 0.53 in female). In dorsal view with lateral margins very slightly sinuate and slightly convergent from apical one-third to humerus; moderately arcuate at apical one-third then straight and convergent to apex. Elytral apices not produced, rounded, slightly divergent. Humerus rounded, indistinct. Dorsally lacking suberect or erect vestiture. Scales uniformly very dense, small, robust, appressed grey-brown on sutural interval and intervals 2 to 5 and 8 to 11; intervals 6 and 7 with scales truncate, transparent, the underlying dark cuticle visible through scales. Punctures of elytral striae small, individually distinct; arranged in regular rows. Wings. Moderate (slightly shorter than length of elytra). Legs. Foretibia of female with inner margin with small denticles in apical one-half; no distinct subapical tooth present. Foretibial and mesotibial unci of both sexes very large, metatibial uncus moderately large. Tarsal claws widely divergent, each lacking distinct basal internal flange. Ventral tarsal pilose vestiture of foretarsus and mesotarsus of both sexes lacking from articles 1 and 2,

metatarsus of both sexes, lacking from articles 1 and 2, present as minute apical tuft on article 3. **Abdomen.** Ventral surface with dense, truncate, small appressed grey-brown scales. Abdominal sterna III to VI sparse moderately large shallow punctures, no distinct glabrous patches or suberect or erect vestiture present. Base of abdominal sternum VII of female internally (dorsally) with at most moderately-developed median shiny tubercle. **Genitalia.**

Female (one examined). Abdominal sternum VIII with basal arm long; lateral arms straight and divergent from base to approximately apical one-third; moderately inwardly arcuate at apical one-third and slightly convergent to apex; apices moderately expanded (Fig. 176). Stylus moderately large compared to length gonocoxite II. Male (one examined).

Internal sac (Figs. 196a,b) elongate and low; median dorsal pocket low, with single moderately broad elongate dorsally directed basal lobe (lobe A), with small median dorsally directed lobe at midlength on dorsal surface (lobe B), with moderately large-paired dorsally directed lobe at basal one-third at dorsal margin (lobe C), with moderately large paired dorsolaterally directed lobe near midlength near dorsal margin (lobe D), with large paired dorsolaterally directed lobe just beyond midlength at dorsal margin (lobe E), and with small ventrally directed paired lobe near base near ventral margin (lobe F). Apical pocket individually distinct from median dorsal pocket; apex of median dorsal

Apical pocket with paired moderately large ventrally directed lobe near apical one-third at ventral margin. Eversible apical sclerite complex with paired simple scythe-like sclerite; adjacent ventral surface of apical pocket with pair of large, dark, longitudinal sclerites.

**Geographic variation.**--- Only two specimens of this species have been examined.

**Geographic distribution.**--- Southern Russia, Persia, Siberia and Mongolia (Csiki 1934).

**Natural history.**--- No information is available about the natural history of this species. Based on its general geographic distribution I suspect that it is a grassland or semi-desert inhabiting species.

**Chorological relationships.**--- This is the only Palearctic species of *Cleonidius* I have seen.

**Phylogenetic relationships.**--- See "*Incertae sedis*" section preceding.

During the course of study, individuals of two adventitious species were encountered from North American localities. These are *Stephanocleonus glaucus* labelled "S.C." (1), and "Aames, Iowa" (1); and, an unidentified species of *Tetragonothorax* from "Mobile, Alabama" (1). These species are considered not to be established in North America.

#### 4.7 Extant taxa erroneously placed in Cleoninae

*Cleonis chilensis* Blanchard 1851:326 is listed in both *Cleonus (Incertae Sedis)* and *Adloristus* in the Coleopterorum Catalogus (Csiki 1934; Schenkling and Marshall 1931) and also in Blackwelder (1947). Kuschel (1949:31) subsequently placed it in *Scotoeborus* (Leptopiinae [now Entiminae], Cyliindrorhini). I have examined the type of this species (deposited in MHNP) and agree that it is not Cleoninae.

*ArgentInorhynchus breyeri* Brèthes 1910:211 was described from Argentina and originally noted as "Cleonidarum". It has since been regarded as Cleoninae by (Csiki 1934) who placed the genus as a subgenus of *Cleonus*. Likewise, and probably following Csiki, Blackwelder (1947) lists the species in *Cleonus*. Kuschel (1950:112) subsequently returned *ArgentInorhynchus* to generic status and placed it in Eriirhininae. The holotype of this species

The genus *Lepyrus* is also not Cleoninae and has been removed to Hylobiinae (Solari 1941; Ter-Minasyan 1963).

#### 4.8 Fossil material

Various Tertiary coleopterous fossils have been identified as Cleoninae however, preservation is such that most of these are actually not reliably identifiable to subfamily. Kingsolver (1961) has examined much of the material of Scudder (1893) and Wickham (1911, 1912) deposited at the Museum of Comparative Zoology, Harvard University, and states this to be the case. Matthews (1977) also notes a *Cleonus* sp. from the Tertiary of Banks Island in the Canadian Arctic but this identification is based only on elytral fragments. I have not examined any of these specimens.

Sleeper (1969) described two species of *Cleonidius* (as *Cleonus*) from Pleistocene Rita Blanca Lake deposits in Texas. Unfortunately I have not seen the specimens, and descriptions and figures do not provide information permitting meaningful comparison with extant species.

Sleeper (1969) states that *Cleonidius ritablancensis* is near *Cleonidius sparsus* and that *Cleonidius channingensis* is near *Cleonidius circumductus*, however, the history of

Unfortunately location of the deposition of the material was not stated by Sleeper (1969).

Fossils of species identified as the genus *Cleonus* are frequently encountered in late Pleistocene deposits in northern North America (Matthews 1974, 1975; Hughes et al. 1981; Mott et al. 1981; Morlan and Matthews 1983). Some of these have been tentatively identified as *Cleonus plumbeus*, but as noted by Morgan et al. (1985), they may actually represent one or more Palearctic species or even one of the herein-newly-described Nearctic species. All of this northern Pleistocene material actually belongs in the genus *Stephanocleonus*, and the present revision of North American members of that genus reveals that taxa traditionally regarded as *S. plumbeus* are a complex of six species, four of which are newly described herein. All of this fossil material is deposited with John V. Matthews, Jr., Geological Survey of Canada, Ottawa; none of it has been examined by me.



## 5.1 Introductory comments

*My profession embodies one theme even more inclusive than evolution--the nature and meaning of history. History employs evolution to structure biological events in time. History subverts the stereotype of science as a precise, heartless enterprise that strips the uniqueness from any complexity and reduces everything to timeless, repeatable, controlled experiments in a laboratory. Historical sciences are different, not lesser. Their methods are comparative, not always experimental; they explain, but usually do not try to predict; they recognize the irreducible quirkiness that history entails, and acknowledge the limited power of present circumstances to impose or elicit optimal solutions.-----Steven Jay Gould, Prologue to "The Flamingo's Smile", 1985.*

Inquiry in historical science proceeds primarily by investigation of relationships between pattern and process. Factual data are assembled and patterns are searched for and recognized in these data. Recognition of a pattern, especially if repeated within the study group or in other groups, begs explanation by proposal of a common causal mechanism in the form of a testable hypothesis. Proposal and subsequent testing of this hypothesis, the former inductive, the latter hypothetico-deductive, are the cornerstones of science in general and systematic research in particular. Comparison of these approaches in systematics is adequately treated by Wiley (1981) and application of the hypothetico-deductive method to systematics by Gaffney

hypothetico-deductive science will be employed and applied. Although some recent authors advocate only the hypothetico-deductive approach, I believe that much useful information and advances in biological knowledge have been and should continue to be, achieved through use of both approaches, but that they operate best at different stages in the process of scientific inquiry; induction for hypothesis formulation, deduction for hypothesis testing.

Herein, an inductive approach will be used primarily to attempt to formulate explanations of details of the evolutionary and biogeographic history of the weevil subfamily Cleoninae, especially species of the genera *Cleonidius* and *Apleurus* in North America. From these analyses, evident patterns will be identified and more general hypotheses concerning biogeographic patterns and evolutionary trends in arid-land-adapted organisms will be proposed, so as to be available for subsequent testing in future studies for their applicability and predictive value. In addition, some competing and previously proposed hypotheses concerning general evolutionary or biogeographic patterns will be tested by application of the appropriate results of this study of North American Cleoninae.

### 5.2.1 Methods of phylogeny reconstruction

Since the publication of Hennig (1966), there has been an expanding emphasis on both theoretical and practical aspects of phylogenetic and biogeographic studies. While the pages of *Systematic Zoology* clearly document an intense interest in theoretical aspects, increased attention to practical aspects and applications is evidenced simply by the numbers of revisionary works that deal extensively with reconstruction of phylogeny and its applicability to character evolution, classification and biogeography. The appeal of the phylogenetic approach (Hennig 1966) to phylogeny reconstruction over various other approaches (Wiley 1981) lies primarily in the potential for critical assessment of resultant hypotheses of relationship, their logical predictive nature and their general applicability. The approach I have taken herein is phylogenetic (Hennig 1966). Character states are determined as apotypic (derived) or plesiotypic (primitive) and only distributions of apotypic states are used in the analysis.

Of paramount importance in reconstructing phylogeny are one, determination of polarity of character states; and two, recognition of homoplasy or simply, determination of which characters actually reflect descent from a common ancestor, not other processes.

For secondary characters, out-group criteria were employed where possible although in-group or local parsimony (Maddison et al. 1984), or the functional out-group of Watrous and Wheeler (1981), were also used, especially in instances where more than one character state occurs in the out-groups or where, although both out-groups have the same state, there is incompatibility with the reconstructed phylogeny as determined by primary characters.

Character weighting was employed. Characters used in the phylogenetic analysis of species in *Cleonidius* and *Apleurus* are considered at two levels as indicated above. Primary characters are those which are deemed of high weight because there is no ambiguity in determining polarity of the various character states based on an out-group analysis (Watrous and Wheeler 1981; Maddison et al. 1984), and the apotypic state, so determined, is not widely distributed elsewhere in Cleoninae. Characters meeting these criteria are given further support as primary characters should they, as a group, result in a reconstructed phylogeny of the genus in question in which the distributions of apotypic states of the various characters are compatible, with the result that there is minimal homoplasy. Characters that exhibit more than a little homoplasy elsewhere in Cleoninae may be regarded as primary should the distribution of their apotypic states be entirely compatible with the distribution

lesser weight because of widespread presence of the apotypic state elsewhere in Cleoninae. Lack of complete compatibility with primary characters, and/or problems in determining polarity of the character states based strictly on an out-group analysis necessitating reference to in-group relationships previously resolved on the basis of the distribution of states of primary characters also results in such characters being considered at a secondary level. Distribution of apotypic states of secondary characters may or may not be compatible with distributions of apotypic states of primary or other secondary characters with the result that there is generally more homoplasy in characters included in this more resolved reconstructed phylogeny of the genus is question.

Characters examined but not included in the phylogenetic analysis are those that exhibit extensive homoplasy elsewhere in Cleoninae and are not compatible with, or, when compared with other potential secondary characters, exhibit minimal compatibility with, distributions of apotypic states of primary characters.

Primary characters are the characters used to establish a basic set of relationships that may or may not be totally dichotomously resolved, but in which confidence can generally be regarded as high. This set of relationships is then accepted *a priori* and the distribution of states of

be used to reinforce previously established relationships or to further resolve the reconstructed phylogeny by providing additional characters in which confidence is low.

I do not use a complex system of weighting characters such as that of Hecht and Edwards (1976, 1977). Instead, I use this simple two level weighting system which depends solely upon the distributions of character states, and makes no inferences or assumptions about complexity, function or adaptive value which may not have a logically sound basis. As such, I think this system better reflects the confidence level for a given set of relationships especially when these are depicted on successively more resolved cladograms.

#### 5.2.2 Phylogenetic position of Cleoninae

The detailed phylogenetic position of Cleoninae is unclear although relationships at some level appear to be with Hylobiinae. Taxonomy of these two subfamilies has a history of repeated misclassification and a repeated concern for adequate distinction (Aslam 1963; Marshall 1932; Solari 1941; Ter-Minasyan 1963). In addition, Aslam (1961) has noted that among Curculionidae, the Hylobiinae and Cleoninae are unique in their possession of free-lobed testes, and, along with a few other subfamilies, both have the vasa efferentia free from the testicular lobes and each of the

primitive Curculionoidea (Morimoto 1962) and cannot be considered evidence for a sister-group relationship but rather, as will be noted forthwith, for a primitive position for Cleoninae with respect to other Curculionidae. Although unjustified by synapotypy as sister-taxon to Cleoninae, Hylobiinae are employed herein as the out-group for the purpose of tentative polarization of character states in genera of Cleoninae.

Evidence that Cleoninae are a very primitive lineage within Curculionidae is extensive. Indeed they may prove sister to the rest of the Curculionidae based on the large number of plesiotypic character states that they possess.

Buchner (1933, 1953) and Scheinert (1933) discuss at length the mechanism by which adult female Cleoninae pass on symbiotic bacteria to the egg and what the site of localization of these symbionts is in the larvae. In Cleoninae, symbiont pouches are paired and attach to the vagina near the base of gonocoxite II. Bacteria develop in these pouches embedded in a secretion originating from the epithelial cells of the pouch. As eggs pass through the vagina they are coated with bacteria which are then ingested by young larvae. In the larvae, there are four variously shaped evaginations in the anterior part of the midgut where the bacteria are localized. Both of these states are unique within Curculionidae to Cleoninae and are very similar to

pouches to those in adult female Cleoninae also occur in various Chrysomelidae, a constituent taxon of the presumed sister-lineage of Curculionoidea; site of localization in larvae was not examined. Although Curculionoidea generally regarded to be primitive have not been examined for these characters, the widespread distribution of the states in related groups suggests that these states are plesiotypic in Curculionidae and that Cleoninae is a very primitive lineage within that family.

Wing structure is also primitive (Kingsolver 1961), because in at least some Cleoninae, the branches of the second anal vein are joined at the base and the basal one-half of the first anal vein is present. These features appear to be unique to Cleoninae within Curculionidae and are undoubtedly plesiotypic because they are found in primitive Curculionoidea.

Ting (1936) has also noted that in Curculionidae, Cleoninae are the only subfamily in which adults have the labial palpi ventral in their site of attachment to the labium. Although Ting (1936) suggests this to be the autapotypic state, the fact that the same state is found in primitive Curculionoidea suggests that it is plesiotypic.

Further evidence for a primitive phylogenetic position for the subfamily comes from the structure of the testes (as noted previously), and generalized stem or root mining



labial palpi, which, although described as of one article (Aslam 1963; Ter-Minasyan 1963), are of three very small and telescoped articles (Figs. 61-63). This character state appears to be unique to Cleoninae within Curculionoidea. Most other Curculionoidea possess larger three-segmented labial palpi; some lack palpi entirely or have the palpi variously modified, but these latter states are clearly autapotypic. At present, this is the only autapotypic character state supporting monophyly of Cleoninae.

### 5.2.3 Phylogenetic relationships among tribes of Cleoninae

Within Cleoninae, Csiki (1934) recognized four tribes; Lepyriini, Cleonini, Rhinocyllini, and Lixini. Lepyriini, has since properly been removed from Cleoninae and placed in Hylobiinae (Solari 1941; Ter-Minasyan 1963). Since then, Aslam (1963) and Ter-Minasyan (1978) have recognized only two tribes, Cleonini and Lixini; O'Brien and Wibmer (1982) continue to recognize three tribes, as above, excluding Lepyriini.

In general, the tribe Lixini are all those Cleoninae with a more elongate-narrow cylindrical rostrum and elongate-narrow body form. The tribe Cleonini on the other hand, are taxa with a more robust rostrum and body form. Cleonini are generally associated with arid habitats;

are otherwise placed as Lixini.

No satisfactory suprageneric classification yet exists for Cleoninae, because the phylogenetic relationships of the world genera must be assessed before such can be realized. No tribal placement is therefore herein attempted although generic relationships of North American taxa, traditionally placed as Cleonini, are discussed and represent a start at a reconstructed phylogeny of genera and revised tribal classification.

#### 5.2.4 Phylogenetic relationships of genera of New World

##### Cleonini

In an attempt to determine the relative position of *Apleurus* and *Cleonidius*, a survey of representatives of 50 genera of world Cleoninae was undertaken (Appendix 3). In instances where type species have not been designated for a given genus, the species considered representative of that genus is one originally included and which could subsequently be designated as the type species. Where possible, relationships were inferred on the basis of shared, presumably derived character states, preferably unique to those taxa or found in few other Cleoninae (preferably those taxa, which for reasons of distribution of other more widespread and presumably apotypic character

corroborated by examination of distribution of either apotypic or plesiotypic states of other characters (with more widespread distributions) for concordance.

Confidence in phylogenetic relationships of *Apleurus* is high because a large number of representatives of genus-group names in Cleonini were examined and considered in the search for out-groups. Unfortunately, confidence in out-group relationships of *Cleonidius* is low, largely because few representatives of genus-group names of Lixini were examined. Although *Cleonidius* have traditionally been considered as Cleonini (the initial search for an out-group accepted that premise), indications are that *Cleonidius* (and *Cylindropterus*) are more closely related to Lixini than to any traditional Cleonini. In fact, as noted elsewhere, some *Lixus* species proved to belong in *Cleonidius*. A revision of this large and structurally diverse genus is required on a world basis to better formulate the phylogenetic relationships of *Cleonidius*.

For *Apleurus*, out-groups *Cnemodontus* and *Chromoderus*, together with *Apleurus* itself, are inferred to compose a monophyletic group based primarily upon the presence of a variously developed acute pronotal postocular projection fringed with anteriorly directed, long, postocular vibrissae of unequal length (Figs. 81-88). This state is otherwise found only in species of *Lixus* (Fig. 4), *Lixocleonus* and

considered closely related to *Apleurus*.

Recognition of this group as monophyletic is supported by the distributions of the following character states, which are shared, except as noted, by members of the three genera and in most instances, by a variable number of other genera as well. Polarity of the states of these characters may or may not have been determined.

1. Body form more or less robust. The three taxa under consideration possess this state as do numerous other Cleoninae. Hylobiinae are also more or less robust in body form; therefore, this state is likely plesiotypic.

2. Pronotum with scale pattern with large white scales present in a lateral stripe of variable width, small and fine in a moderately broad to very broad apically narrowed median stripe; median area largely black in color, the underlying dark cuticle not obscured by the overlying scales (Figs. 24, 68-80). This state is present in *Apleurus* and *Chromoderus*, as well as numerous other genera of Cleoninae. A distinct pronotal scale pattern is not evident in *Cnemodontus*. Polarity of the states of this character has not been determined.

3. Eye like an inverted, elongate-teardrop in shape. This state is found in the three genera in question as well as in numerous other genera of Cleoninae. Polarity of the states of this character has not been determined.

Polarity of the states of this character has not been determined.

5. Female with abdominal sternum VIII with basal arm absent (Figs. 56e-57e, 89a-96a). This state is found in all three genera as well as in numerous other genera of Cleoninae. Hylobiinae possess a long basal arm and thus lack of the arm is likely apotypic but also homoplasious.

6. Tarsi with articles moderately wide, not markedly elongate-narrow. This state is found in *Chromoderus*, *Cnemodontus* and some *Apleurus* species as well as numerous other Cleoninae. Hylobiinae also possess this state and thus it is likely plesiotypic.

7. Tarsi with ventral pilose pads present, large and distinctly delimited. This state is found in *Chromoderus*, *Cnemodontus* and some *Apleurus* species as well as numerous other Cleoninae. Hylobiinae also possess this state and thus it is likely plesiotypic.

8. Pronotum with dorsal surface punctate, not tuberculate. This state is found in *Chromoderus*, *Cnemodontus* and *Apleurus* species as well as numerous other Cleoninae. Polarity of the states of this character has not been determined.

9. Female with spermathecal gland spherical, not variously elongate and cylindrical. This state is found in

10. Rostrum moderately elongate-narrow to robust. This state is found in the three taxa in question as well as various other Cleoninae. Polarity of the states of this character has not been determined.

11. Tibiae with corbel ridge sharp (as in Fig. 6). This state is found in *Chromoderus*, *Cnemodontus* and *Apleurus* species as well as numerous other Cleoninae. Polarity of the states of this character has not been determined.

12. Ventral surface of male lacking extensive long erect vestiture. This state is found in *Chromoderus*, *Cnemodontus* and *Apleurus* species as well as numerous other Cleoninae. Hylobiinae also possess this state and thus it is likely plesiotypic.

13. Female with gonocoxite II triangular in shape (Figs. 89b-96b), apex not prolonged into a marked lobe (Fig. 54). This state is found in *Chromoderus*, *Cnemodontus* and *Apleurus* species as well as numerous other Cleoninae. Hylobiinae also possess this state and thus it is likely plesiotypic.

14. Antenna with apical three articles with placoidal sensillae. This state is found in *Chromoderus*, *Cnemodontus* and *Apleurus* species as well as various other Cleoninae. Polarity of the states of this character has not been determined and not all Cleoninae have been examined in

resemblance to *Cleonidius* species, yet lack of the internal swelling at the base of abdominal sternum VII in the female which characterizes the latter. No synapomorphies were identified to unite the three taxa as a monophyletic group and indeed, the group may prove to be paraphyletic. Distributions of the states of the following characters support a close but unresolved relationship among these three taxa.

1. Body form more or less elongate-narrow, cylindrical. The three taxa under consideration possess this state as do *Mecaspis*, *Lixocleonus*, and most *Lixus*. Hylobiinae are more robust in body form therefore elongate-narrow form is likely apotypic.

2. Pronotum with scale pattern slightly to distinctly vittate, with white scales largest and/or densest immediately laterad of midline, smaller and/or sparser laterally; largely whitish in color, the underlying dark cuticle largely obscured by the white scales (Figs. 118-136). This state is present in *Lixoglyptus* and *Cleonidius* species as well as *Pseudocleonus*, *Mecaspis*, and various other species of *Lixus*. A distinct pronotal scale pattern is lacking from many genera including *Cylindropterus*.

character has not been determined.

4. Eye more or less flat in dorsal view. This state is found in the three taxa in question as well as numerous other Cleoninae. Polarity of the states of this character has not been determined.

5. Tibiae with corbel ridge rounded (as in Fig. 5). This state is present in the three taxa in question, *Lixocleonus*, *Menocleonus*, *Epirhynchus*, and all *Lixus* species. Polarity of the states of this character have not been determined.

6. Tarsi with articles moderately wide, not markedly elongate-narrow. This state is found in the three taxa in question as well as numerous other Cleoninae. Hylobiinae also possess this state and thus it is likely plesiotypic.

7. Tarsi with ventral pilose pads present, large and distinctly delimited. This state is found in the three taxa in question as well as numerous other Cleoninae. Hylobiinae also possess this state and thus it is likely plesiotypic.

8. Pronotum with dorsal surface punctate, not tuberculate. This state is found in the taxa in question as well as numerous other Cleoninae. Polarity of the states of this character has not been determined.

9. Female with spermathecal gland spherical, not elongate and cylindrical. This state is found in the three



10. Rostrum moderately to very elongate-narrow. This state is found in the three taxa in question (although not all species of *Cleonidius*) as well as various other Cleoninae. Polarity of the states of this character has not been determined.

11. Pronotum with postocular lobes present. This state is found in the three taxa in question (although not in all species of *Cleonidius*) as well as numerous other Cleoninae. Hylobiinae also possess this state and thus it is likely plesiotypic.

12. Female with abdominal sternum VIII with basal arm present, short to long (Figs. 113-114, 157-176). This state is found in the three taxa in question as well as in numerous other genera of Cleoninae. Hylobiinae possess a long basal arm and thus this state is likely plesiotypic with a successively shorter arm increasingly apotypic.

13. Antenna with apical three articles with placoidal sensillae lacking or else very few in number. This state is found in the three genera in question as well as *Lixocleonus* and *Lixus* species. Polarity of states of this character has not been determined and not all Cleoninae have been examined in detail for this character.

14. Prosternum with swelling in front of fore-coxal cavity. This state is found in the three taxa in question

determined.

#### 5.2.4.1 Genus *Apleurus* Chevrolat

##### 5.2.4.1.1 Character evolution

###### Primary characters.

Character 1. Epistoma, apical margin.--- Two states of this character are recognized:

1. emarginate (Fig. 81b);
2. rounded and produced (Figs. 82b-88b).

Out-groups, most Cleoninae, and Hylobiinae possess an emarginate epistoma. This state is considered plesiotypic. An apically rounded epistoma is otherwise known only in *Gonocleonus*, *Leucochromus*, *Eurycleonus*, and *Koeniglus*.

Character 2. Prementum, setae.--- Two arbitrarily defined states of this character are recognized:

1. lacking or with only one large seta per side (Figs. 61-62);
2. with two or more large setae per side (Fig. 63).

Out-groups *Cnemodontus* and *Chromoderus* lack and possess one large seta per side respectively. Two or more setae per side is considered apotypic. Other Cleoninae were not

setae on each side of the prementum.

**Character 3. Pronotum, tubercles, development and shape of anterior angles.**--- Three states are recognized:

1. absent (Figs. 24, 68-69);
2. present, anterior angles obtuse (Figs. 75-80);
3. present, anterior angles acute (Figs. 70-74).

Out-groups lack distinct tubercles but the pronota are slightly constricted at the apical one-third to one-quarter, the position of the tubercles if developed. This and lack of acutely-angled tubercles in other Cleoninae which possess variously developed pronotal tubercles, justify recognition of obtusely angled tubercles as plesiotypic.

**Character 4. Tarsus, ventral pilose vestiture.**--- Three arbitrarily defined states are recognized:

1. extensive;
2. moderately reduced in extent;
3. markedly reduced in extent, or absent.

Out-groups each possess extensive ventral tarsal pilose vestiture. This state is considered plesiotypic and successive stages in reduced extent of pilose vestiture increasingly apotypic. Hylobiinae possess extensive ventral pilose vestiture. Extent of ventral tarsal pilose vestiture varies extensively in Cleoninae and on that basis alone this

states of this character with numerous other primary characters results in increased confidence that it also be considered primary.

**Character 5. Female, stylus, size.**--- Three states are recognized:

1. large (Figs. 89b-91b);
2. small (Figs. 92b-94b);
3. absent (Figs. 95b-96b).

Out-groups and all Cleoninae examined (except for *Brachycleonus* in which a stylus is absent) possess a moderately large stylus. Presence of a stylus is considered plesiotypic and decreased size is considered increasingly apotypic.

**Character 6. Female, stylus, position.**--- Three states are recognized:

1. apical (Figs. 89b-91b);
2. anteapical (Figs. 92b-94b);
3. absent, anteapical sclerotized ridge present (Figs. 95b-96b).

Out-groups and all Cleoninae examined in which a stylus is present, possess an apically situated stylus. This state is considered plesiotypic. The anteapical sclerotized ridge is regarded as positionally homologous with the stylus in

Character 7. Male, abdominal sternum VII, dorsally directed tooth at apical margin.--- Two states are recognized:

1. absent;
2. present.

Out-groups and all Cleoninae except *Brachycleonus* and *Centrocleonus* lack a dorsally directed tooth on the apical margin of abdominal sternum VII. Lack of a tooth is considered plesiotypic.

Character 8. Male, abdominal sternum VIII, interior angles of each sternite.--- Three states are recognized:

1. basal projection absent (Fig. 97f);
2. basal projection slightly to moderately developed (Fig. 101e);
3. basal projection present, markedly developed (Fig. 103e).

Both out-groups lack a basal projection. Thus, this state is considered plesiotypic. Extent of development from slight to marked represents successively apotypic states. Other Cleoninae have not been examined in detail for this character although a distinct basal projection is lacking from both *Cleonidius* and *Stephanocleonus*.

**Secondary characters.**

1.  $X < 7.0$  mm;
2.  $7.0 \text{ mm} < X < 9.0$  mm;
3.  $X > 9.0$  mm.

Large size (state 3) is widespread in Cleoninae but the out-groups *Chromoderus* and *Cnemodontus* possess state 1 and state 2 respectively. Small size of *Chromoderus* and primitive *Apleurus* species, however, suggests that the common ancestor of *Chromoderus* and *Apleurus* was also of small size, and that medium to large sizes (states 2 and 3) in *Apleurus* are successively apotypic. Whether the medium size of *Chromoderus* is then primitive or represents an independent increase in size is not known.

**Character 10. Eye, prominence and convexity.**--- Three arbitrarily defined states are recognized:

1. markedly convex and prominent;
2. slightly to moderately prominent and convex;
3. flat.

Flat, non-prominent eyes are known in virtually all Cleoninae with the exception of *Lixocleonus*, *Microlarinus*, and various *Lixus* species. Out-groups *Cnemodontus* and *Chromoderus* have markedly convex and flat eyes respectively. As for character 9, occurrence of markedly convex eyes in one of the out-groups and in primitive species of *Apleurus*, suggests that this state is plesiotypic. Decreased

Character 11. Suberect or erect vestiture, length.--- Three states are recognized:

1. absent or indistinct;
2. short;
3. long.

Out-groups lack distinct erect vestiture. Thus, this state is considered plesiotypic. Presence and subsequently increased length of vestiture are considered successively apotypic. Long suberect or erect vestiture is known in numerous genera of Cleoninae and other Curculionidae (e.g. *Trigonoscuta*, *Miloderes*, etc.) and appears correlated with life in sand substrates.

Character 12. Rostrum, carina.--- Two states are recognized:

1. present (Figs. 84-86);
2. absent (Figs. 81-83, 87-88).

Both out-groups possess a rostral carina. However, primitive *Apleurus* species lack a carina. This suggests that lack of a carina is apotypic for *Apleurus* but that presence of a carina in certain species of *Apleurus* is also apotypic.

Character 13. Rostrum, declivity at apex.--- Two states are recognized:

1. flat to at most slightly declivous (Figs. 81a-83a);

apex, thus this state is considered plesiotypic. A rostrum with a steeply declivous apex also occurs in many other Cleoninae.

**Character 14. Maxillary palpus, setae.**--- Two states are recognized:

1. one large seta on palpifer, one on stipes (Fig. 64);
2. one large seta on palpifer (Fig. 65).

Out-groups possess large setae on both palpifer and stipes. Thus, lack of a seta from the stipes is considered apotypic. Whether this seta is lacking or simply reduced in size was not determined. Other Cleoninae were not examined in detail for this character.

**Character 15. Pronotum, width.**--- Two states are recognized (see Fig. 109):

1. width greatest at base to width at base more or less subequal to width at tubercles;
2. width distinctly greatest at tubercles.

Out-groups have pronota that are widest at the base. Thus, this state is considered plesiotypic. Pronota that are distinctly widest at the tubercles are also known in *Tetragonothorax* and *Gonocleonus*.

**Character 16. Pronotum, punctation.**--- Two states are



2. large and deep.

Out-groups have pronota that have small shallow punctures. Thus, a pronotum with large and deep punctures is considered apotypic. No other Cleoninae examined have similarly large and deep pronotal punctures.

\*  
Character 17. Pronotum, median carina.--- Two states are recognized:

1. absent;
2. present, of variable extent.

Out-groups lack a pronotal carina. Thus, presence of a carina is considered apotypic. Presence of a carina is widespread and common in most genera of Cleoninae

Character 18. Prosternum, impressions and swellings.--- Four arbitrarily defined states are recognized in this lineage:

1. impression anterolaterad of fore-coxal cavity present; no swelling present;
2. impression anterolaterad of fore-coxal cavity present; swelling immediately anterior to impression present, low and rounded (Figs. 82a-83a);
3. impression anterolaterad of fore-coxal cavity present; swelling immediately anterior to impression present, high (Figs. 81a, 84a-88a);
4. impression anterolaterad of fore-coxal cavity present;

front of fore-coxal cavity.

All Cleoninae possess an impression, although various in depth and extent, immediately anterolaterad of each fore-coxal cavity. Swellings, if present, are found in two locations, one, immediately in front of the fore-coxal cavity, adjacent to the impression, or two, immediately in front of the impression. Out-group *Cnemodontus* lacks swellings and *Chromoderus* is unique in Cleoninae examined in that it possesses swellings in front of both the fore-coxal cavity and the impression. Primitive *Apleurus* possess low rounded swellings anterior to the impression, thus this state is judged plesiotypic for *Apleurus*; high swellings in front of the prosternal impressions are considered apotypic. Swellings in front of the impression are widespread in Cleoninae but are generally low and rounded except in *Stephanocleonus* and *Conorhynchus*. They are however absent in most genera. *Cosmogaster*, *Mecaspis*, *Lixocleonus*, *Microcleonus*, *Lixus* and *Cleonidius* all have a swelling of variable extent in front of the fore-coxal cavity.

Character 19. Elytra, elevation of intervals.--- Two states are recognized;

1. all intervals equally flat or only slightly elevated and convex;
2. sutural interval and intervals 3, 5, 7, and 9 slightly to

Out-groups possess state 1 which is therefore considered plesiotypic. The apotypic state occurs in many genera of Cleoninae and other Curculionidae. Development of elevated and convex elytral intervals appears to be correlated with reduction in size or absence of the metathoracic wings.

**Character 20. Fore-tibia of female, apex.**--- Two states are recognized:

1. uncus and large subapical second spur present;
2. uncus and at most only small subapical second spur present.

Out-groups each possess a large second spur in addition to the uncus thus this state is considered plesiotypic. Both states are widespread in Cleoninae.

**Character 21. Fore-tibia, inner margins.**--- Two states are recognized:

1. large denticles present;
2. with at most only small denticles present.

*Cnemodontus* individuals possess large fore-tibial denticles; *Chromoderus* individuals, only small denticles. Both states are widespread in other Cleoninae. Presence of large denticles is considered plesiotypic.

**Character 22. Female abdominal sternum VIII, lateral**

1. more or less straight throughout length (Figs. 49e-50e, 90b-91b, 95b-96b);
2. arcuate (Figs. 89b, 92b-94b).

Abdominal sternum VIII in females of both out-groups has lateral arms that are more or less straight. This state is considered plesiotypic. Both states are widespread in Cleoninae.

**Character 23. Female abdominal tergum VII, median longitudinal carina.**--- Three states are recognized:

1. absent;
2. slightly developed;
3. markedly developed.

*Cnemodontus* lacks a longitudinal carina whereas a markedly developed carina is present in *Chromoderus*. Other Cleoninae examined lack a carina. Thus, this state is considered plesiotypic.

**Character 24. Female abdominal tergum VIII, median longitudinal carina.**--- Three states are recognized:

1. absent;
2. slightly developed;
3. markedly developed.

*Cnemodontus* lacks a longitudinal carina whereas a slightly developed carina is present in *Chromoderus*. Other Cleoninae examined lack a carina. Thus, this state is

Character 25. Male, aedeagus, curvature.--- Two states are recognized:

1. markedly arcuate at base (Figs. 56a-57a);
2. more or less evenly arcuate throughout length (Figs. 97d, 98c-104c).

Both out-groups possess an aedeagus that is markedly arcuate at the base. Thus, this state is considered plesiotypic. Both states occur in *Stephanocleonus* (Ter-Minasyan 1979). The aedeagus of all *Cleonidius* species, *Lixoglyptus* and *Cylindropterus* is evenly arcuate throughout its length. Other Cleoninae have not been examined in detail for this character.

Character 26. Male, aedeagus, apex.--- Two states are recognized:

1. spatulate (Figs. 56a-57a);
2. not spatulate (Figs. 97d, 98c-104c).

Both out-groups possess an aedeagus with a spatulate apex. Thus, this state is considered plesiotypic. Both states are widely distributed in Cleoninae.

Character 27. Male, aedeagus, internal sac, paired lateral sclerite at midlength.--- Two states are recognized:

1. present (Fig. 57b);
2. absent (Figs. 56b, 97a-104a).

sclerite at the midlength of the internal sac; *Chromoderus* males lack such sclerites. Lack of the sclerites is considered apotypic. *Stephanocleonus*, *Prionorhinus*, and *Lixomorphus* males also possess such sclerites, and another unpaired sclerite, situated in a ventral position closer to the apex of the internal sac, is present in *Menocleonus*, *Pleurocleonus*, and *Xanthochelus*. This latter sclerite, however, is questionably homologous with the paired sclerite at midlength.

Character 28. Male, aedeagus, internal sac, shape.--- Two states are recognized:

1. dorsal median pocket low and with two pairs of lobes (Figs. 56b-57b, 97a-99a);
2. dorsal median pocket high and with one pair of lobes (Figs. 100a-104a).

*Cnemodontus* males possess an internal sac with state 1; *Chromoderus* males possess an internal sac with state 2. The shape of the internal sac varies in other Cleoninae examined, but only in some species of *Cleonidius* is it similarly high and possessing of only a single pair of lobes. State 1 is thus considered plesiotypic.

Character 29. Male, aedeagus, internal sac, lobe A.--- Two states are recognized:

1. present (Figs. 56b,c-57b,c);

Males of each out-group possess lobe A. Thus, this state is plesiotypic. This character has not been examined in detail in other Cleoninae. However, lobe A is present in at least *Cleonidius*, *Lixoglyptus*, *Cylindropterus*, *Adosomus*, *Cleonis* and *Cyphocleonus*. It is absent from those *Stephanocleonus* examined.

Character 30. Male, aedeagus, internal sac, apical sclerite complex, median projection.--- Two states are recognized:

1. long (Figs. 56d-57d, 103d-104d);
2. absent or present, but short and indistinct (Figs. 97c, 98d-101d).

Both *Cnemodontus* and *Chromoderus* males possess apical sclerite complexes with a well-developed median projection. Presence of state 2 in primitive *Apleurus* species suggests presence of state 1 within *Apleurus* is secondarily apotypic. This character has not been examined in detail in other Cleoninae.

Character 31. Male, aedeagus, internal sac, apical sclerite complex, sclerotization of base and sides of the apical pocket.--- Two states are recognized:

1. well-sclerotized, dark brown in color;
2. lightly-sclerotized, pale brown in color.

Males of each out-group have the base and sides of the apical pocket markedly sclerotized thus this state is

in detail in other Cleoninae.

#### 5.2.4.1.2 Phylogenetic relationships of included species

Confidence in phylogenetic relationships of species of *Apleurus* (Figs. 234-235) is high; most primary characters exhibit very little homoplasy elsewhere in Cleoninae and the distribution of apotypic states is concordant with distributions of apotypic states in most secondary characters. Furthermore, primary characters are employed to define most major lineages and are more general in their applicability (Fig. 234) (i.e. not all primary characters are used to define the same or few lineages). Because of this, confidence in recognizing homoplasy in secondary characters is enhanced and their applicability amplified.

Relationships among *Chromoderus*, *Cnemodontus* and *Apleurus* based on primary characters are unclear (Fig. 234). Apotypic states of secondary characters of size of female fore-tibial denticles, and presence or absence of a paired lateral sclerite at midlength in the internal sac of the aedeagus of males, suggests *Chromoderus* is the sister group to *Apleurus*, with *Cnemodontus* as sister to the *Chromoderus-Apleurus* lineage (Fig. 235). Flat eyes in *Chromoderus* and most other Cleoninae, and shared presence of swollen and convex eyes in *Cnemodontus* and most *Apleurus* is the only evidence that *Cnemodontus* is sister to *Apleurus*. However, I favour consideration of flat eyes as



eyes as plesiotypic for the lineage.

Monophyly of *Apleurus* is based upon distribution of apotypic states of a number of characters, both primary and secondary. All *Apleurus* have ventral tarsal pilose vestiture moderately to markedly reduced in extent, the aedeagus of males is evenly arcuate throughout its length, the apex of the aedeagus is not spatulate, and the internal sac of the aedeagus lacks lobe A. These apotypic states all occur elsewhere in Cleoninae but not in those taxa believed closely related to *Apleurus*. Although there is homoplasy within *Apleurus*, distributions of apotypic states of other characters also support monophyly of the genus. These states are lack of a rostral carina (secondarily derived in *A. angularis*-*A. jacobinus*-*A. porosus* lineage), fore-tibia of female with small subapical second spur (secondarily derived in *A. aztecus*), and internal sac of aedeagus of male with the median projection of the apical sclerite complex absent or short and indistinct (secondarily derived in *A. albovestitus*-*A. saginatus* lineage).

*Apleurus (Gibbostethus) hystrix* is hypothesized to be the most primitive species of *Apleurus* (Figs. 234-235). It is the only species of *Apleurus* which has an emarginate epistoma; all other *Apleurus* species have the epistoma rounded. *Apleurus hystrix* is autapotypic in a number of characters (see species treatment of *A. hystrix*), and also shares, with various other species of *Apleurus*, a number of

apotypic states which are best regarded as homoplasious. All of these apotypic states are of secondary characters and are otherwise found only in the *A. angularis*-*A. jacobinus*-*A. porosus* lineage or the common ancestor of this lineage and the *A. albovestitus*-*A. saginatus* lineage. This suggests that *A. hystrix* might be better placed with these species, but distribution of apotypic states of all primary characters in the analysis argues against this proposal (Fig. 234).

*Apleurus aztecus* is the next most primitive species and is sister to the remainder of the subgenus *Apleurus* (Figs. 234-235). Phylogenetic position of this species is based upon possession of a rounded epistoma, lack of a dorsally directed tooth at the apical margin of abdominal sternum VII in males, and lack of basal projections on the interior angles of each sternite of abdominal sternum VIII also in males.

*Apleurus lutulentus* is sister to the ancestor of the *A. angularis*-*A. jacobinus*-*A. porosus* and *A. albovestitus*-*A. saginatus* lineages (Figs. 234-235). This position is due to distributions of apotypic states of numerous primary as well as secondary characters. These latter five species comprise a monophyletic group divisible into two lineages as noted above. Monophyly of the *A. angularis*-*A. jacobinus*-*A. porosus* lineage is based upon possession of acute-angled pronotal tubercles, presence of rostral carina, maxillary palpus with only one large seta on palpifer, large and deep pronotal punctation, alternate elytral intervals elevated and convex,

female abdominal sternum VIII with slight median longitudinal carina, and internal sac of aedeagus of male with sides and base of apical pocket only lightly sclerotized. Within this group of species, long erect dorsal vestiture and relative width of the basal and apical portions of the pronotum suggests an *A. angularis*-*A. jacobinus* sister species relationship. Widely divergent tarsal claws in all *A. angularis* and some *A. porosus* are likely a result of independent evolution or of hybridization.

The sister species status of *A. albovestitus* and *A. saginatus* (Figs. 234-235) is strongly supported by the lack of a stylus on gonocoxite II of the female and by markedly developed basal projections on the interior angles of each sternite of abdominal sternum VIII in males. Distribution of apotypic states of numerous secondary characters supports this relationship.

#### 5.2.4.2 Genus *Cleonidius* Casey

##### 5.2.4.2.1 Character evolution

#### Primary characters.

Character 1. Submentum, degree of ventral swelling.--- Two states of this character are recognized:

1. flat, not ventrally swollen (Figs. 137a-142a, 145a-156a);

4. swollen ventrally (Figs. 143a-144a).

Out-groups and all Cleoninae examined possess a submentum that is not swollen ventrally. This state is considered plesiotypic.

**Character 2. Rostrum, ratio of length to width in females.**--- Four arbitrarily defined states are recognized (see Fig. 198):

- 1. very elongate,  $X < 0.45$  (Fig. 141);
1. moderately elongate,  $0.45 < X < 0.65$  (Figs. 137-140, 142-144, 148-155);
2. short, robust,  $0.65 < X < 0.77$  (Figs. 145, 156);
3. very short, very robust,  $X > 0.77$  (Figs. 146-147).

Out-groups possess a rostrum that is moderately elongate. This state is deemed plesiotypic and successively shorter and more robust rostra are considered increasingly apotypic. A very elongate rostrum is also considered apotypic. States of this character are widely distributed in Cleoninae although in general members of the tribe Lixini possess a more elongate, narrow rostrum than do members of the Cleonini. Rostral length in Hylobiinae varies from moderately to very elongate.

**Character 3. Rostrum, degree of lateral swelling.**--- Two states are recognized:

1. not swollen at midlength in dorsal view (Figs. 137b-142b, 145b-156b);

2. swollen at midlength in dorsal view (Figs. 1430-1440).

Out-groups possess state 1 which is accordingly considered plesiotypic. No other Cleoninae or Hylobiinae examined possess a similarly laterally expanded rostrum.

**Character 4. Frons/vertex, surface sculpture.**--- Three states of this character are recognized:

1. area behind eye with wrinkles of cuticle;
2. area behind eye with a curved sulcus (Fig. 150a);
3. area behind eye with a deep, straight sulcus (Fig. 139).

Other than various punctures, out-groups and most Cleoninae possess at most only a few wrinkles of the cuticle behind the eye. Some African *Lixus* possess a very deep elongate impression above the eye. State 1 is considered plesiotypic for *Cleonidius*.

**Character 5. Elytra, scale pattern.**--- Three states of this character are recognized:

1. marginal; intervals 9 to 11 with dense, moderately large, white scales, otherwise moderately densely and uniformly covered with fine white scales which are smaller than those on intervals 9 to 11 (Figs. 118-123);
2. vittate; intervals other than 9 to 11 with scales as large as those on intervals 9 to 11 (Figs. 127-128, 130-136);
3. mottled; with irregularly scattered and sized patches of large white scales (Figs. 124-126, 129).

Of the out-groups, *Cylindropterus* lacks a distinct scale pattern whereas *Lixoglyptus* species both have elytral scale patterns that are of the marginal type. Most *Lixus* species possess uniformly dense small fine scales on the elytra although elytra of individuals of a few species are distinctly vittate. Members of the Cleonini have various elytral scale patterns, but these are not recognizable as any of the states discussed herein for *Cleonidius*. State 1 is considered plesiotypic based largely on its occurrence in *Lixoglyptus* species. This state is assumed to represent an intermediate stage between the uniformly distributed small fine scales of most *Lixus* and the vittate state of certain *Cleonidius* species. The mottled state is then assumed to be a result of decreased scale size at various positions of the vittate pattern and is homoplasious within *Cleonidius* although restricted to a single species group.

**Character 6. Abdomen of female, ornamentation of abdominal sternum VII.**--- Two states of this character are recognized:

1. basal margin uniformly flat;
2. basal margin at middle internally (dorsally) with variously developed glabrous shiny tubercle.

Out-groups and all other Cleoninae examined possess state 1. Within *Cleonidius* there is variation within and between species in the degree of development of the tubercle. Very slightly developed tubercles or glabrous patches are also present at the basal margin of abdominal

sterna V and VI in some specimens, but in these individuals, the most markedly developed tubercle is that on sternum VII.

**Character 7. Male, aedeagus, internal sac, height of dorsal median pocket.**--- Two states of this character are recognized:

1. low (Figs. 111a-112a, 177a-179a, 183a-196a);
2. high (Figs. 180a-182a).

Out-groups possess a low dorsal median pocket. Thus, this state is considered plesiotypic. Both states occur elsewhere in Cleoninae but the character has not been examined in detail.

**Character 8. Male, aedeagus, internal sac, apex of dorsal median pocket.**--- Two states of this character are recognized:

1. rounded or truncate in dorsal view (Figs. 111b-112b, 177b-191b, 196b);
2. emarginate or sinuate in dorsal view (Figs. 192b-195b).

Out-groups possess a dorsal median pocket with a rounded to truncate apical margin. Thus, this state is considered plesiotypic. Other Cleoninae have not been examined in detail for this character.

**Character 9. Male, aedeagus, internal sac, lobe B.**--- Two states of this character are recognized:

1. present (Figs. 111a-112a, 177a-179a, 183a-196a);

2. absent (Figs. 180a-182a).

Out-groups possess a lobe B on the internal sac. Thus, this state is plesiotypic. Other Cleoninae have not been examined in detail for this character although both states are found in *Apleurus* species, *Cleonis* and *Cnemodontus* possess a lobe B, and the lobe is lacking from *Stephanocleonus* species examined and *Chromoderus*.

Character 10. Male, aedeagus, internal sac, lobe D.--- Two states of this character are recognized:

1. present (Figs. 111a-112a, 177a-179a, 183a-196a);
2. absent (Figs. 180a-182a).

The internal sac of each out-group possesses a lobe D; however, the lobe is small in *L. spartii*. Lobe D is present in all species of *Apleurus*, *Cleonis*, *Stephanocleonus*, *Chromoderus*, and *Cnemodontus* examined. Absence of the lobe is considered apotypic in *Cleonidius*.

#### Secondary characters.

Character 11. Size, length of elytra.--- Two arbitrarily defined states of this character are recognized (see Fig. 197):

1. large,  $X > 9\text{mm}$ ;
2. small,  $X < 9\text{mm}$ .

Individuals of both out-groups are large in size but distribution of states of primary characters within



Character 12. Rostrum, punctation.--- Two states are recognized:

1. apical portion with punctures moderately dense and moderately deep;
2. apical portion with punctures sparse, small and shallow.

Out-groups each possess state 1. Thus, this state is considered plesiotypic. State 2 occurs in most species of *Lixus* species and appears associated with an elongate-narrow rostrum. The apotypic state of this character is homoplasious within *Cleonidius*.

Character 13. Rostrum, median tumescence.--- Three states are recognized:

1. absent (Figs. 139b, 141b-142b, 152b);
2. present, slightly developed (Figs. 140b, 143b-144b, 146b-149b);
3. present, markedly developed (Figs. 137b-138b, 145b, 150b-151b, 153b-156b).

Out-groups each possess state 1. Thus, this state is considered plesiotypic for *Cleonidius*. Increased degree of development of the median tumescence is considered increasingly apotypic. There is extensive homoplasy in this character.

2. sharp.

Out-groups each possess state 1. Thus, this state is considered plesiotypic. Species of *Lixus* and *Larinus* examined possess rounded lateral margins; species of Cleonini have either state.

Character 15. Pronotum, anterolateral margins.--- Three states are recognized:

1. postocular lobe rounded, well-developed (Figs. 137a-139a, 141a-144a, 146a-151a);
2. postocular lobe rounded, slightly developed (Figs. 140a, 152a);
3. postocular lobe absent, margin straight (Figs. 145a, 153a-155a).

Out-groups and most other Cleoninae possess well-developed postocular lobes. This state is thus considered plesiotypic and successively less markedly developed postocular lobes as increasingly apotypic. Acute pronotal postocular projections, although present in some Cleoninae which lack rounded postocular lobes (e.g. *Lixus*, *Apleurus*), are not present in those *Cleonidius* species that lack a postocular lobe. This character is homoplasious in *Cleonidius*.

1. basal median area shallowly to moderately deeply impressed;
2. basal median area deeply impressed, lateral margins variously longitudinally impressed.

State 1 occurs in both out-groups and is regarded as plesiotypic. Similar sculpture is not present in other Cleoninae examined.

**Character 17. Suberect or erect vestiture, length.**--- Three states are recognized:

1. absent (Figs. 140a, 152a, 156a);
2. present, short to moderate in length (Figs. 138a-139a, 145a-149a, 151a, 153a);
3. present, long (Figs. 137a, 141a-144a, 150a, 154a-155a).

Both out-groups and many *LIXUS* lack suberect or erect vestiture. Thus, this state is considered plesiotypic. Increased length of vestiture is considered as increasingly apotypic. There is extensive homoplasy in this character in *Cleonidius* and in Cleoninae in general.

**Character 18. Tarsus, ventral pilose vestiture.**--- Three states are recognized:

1. extensive;
2. moderately reduced in extent;
3. markedly reduced in extent, to absent.

vestiture. Thus, this state is considered plesiotypic.

Successive stages in reduction of vestiture are considered as increasingly apotypic. Reduced ventral pilose vestiture is otherwise confined to members of Cleonini. This character is homoplasious in *Cleonidius*.

Character 19. Foretibia of female, inner margin.--- Two states are recognized:

1. large denticles present (as in Fig. 144c);
2. only small denticles present (as in Fig. 143c).

Both out-groups possess large denticles on the inner margin of the foretibia. Thus, this state is considered plesiotypic for *Cleonidius*. Distribution of states of primary characters within *Cleonidius* suggests presence of large denticles in some species of *Cleonidius* is secondarily apotypic and homoplasious. Both states of this character occur in other Cleoninae.

Character 20. Female abdominal sternum VIII, length of basal arm.--- Two states are recognized:

1. long (Figs. 165, 167-169, 174);
2. short (Figs. 157-164, 166, 170-174, 175-176).

Out-groups each possess a long basal arm. Thus, this state is considered plesiotypic. Both states of this character are widely distributed in Cleoninae, but as

*Cleonidius*.

Character 21. Female abdominal sternum VIII, shape of lateral arms.--- Two states are recognized:

1. arcuate (Figs. 163, 165-175);
2. more or less straight, divergent (Figs. 157-162, 164, 176).

Out-groups each possess state 1. Thus, this state is considered plesiotypic. This character is homoplasious in *Cleonidius*. Both states occur in other Cleoninae.

Character 22. Male, aedeagus, internal sac, lobe E.--- Two states are recognized:

1. present (Figs. 111a, 177a-179a, 183a-196a);
2. absent (Figs. 112a, 180a-182a).

The internal sac of *Cylindropterus* possesses a lobe E; that of *Lixoglyptus* lacks lobe E. Lobe E is present in species of *Cleonis*, *Cnemodontus*, some *Apleurus* and at least some *Stephanocleonus*. Distribution of states of primary characters within *Cleonidius* suggests presence of lobe E be considered plesiotypic.

relationships in *Cleonidius* is low, due to the low numbers of primary characters as compared to the number of species in the genus, but especially the level at which the primary characters are applicable. Seven of the ten primary characters apply to lineages of only 2 or 3 species; only 3 apply at more general levels, one of which is used to define the monophyly of *Cleonidius* (Fig. 236). This leaves a large number of relationships subject to resolution only on the basis of secondary characters (Fig. 237). Unfortunately, secondary characters containing any phylogenetic information are also not numerous, few are applicable at a general level, and many, especially the latter, are subject to marked homoplasy within *Cleonidius* (Fig. 237).

Basal relationships among *Cleonidius*, *Cylindropterus* and *Lixoglyptus* are very unclear and, as discussed elsewhere (see "Phylogenetic relationships of genera of New World Cleonini") are left as unresolved (Fig. 237). Monophyly of *Cleonidius* however, is strongly supported by the presence of a variously internally (dorsally) developed glabrous shiny median tubercle at the base of abdominal sternum VII of the female. All secondary characters used in support of monophyly of *Cleonidius* involve homoplasy within *Cleonidius* (Fig. 237).

Relationships of the Palearctic species component of *Cleonidius* are herein left as unresolved because of

potentially *Cleonidius*. Further examination of Palearctic taxa should resolve the placement of the Palearctic species component of *Cleonidius*.

Among species groups, monophyly of each of the *C. boucardi* and *C. lobigerinus* groups, is substantiated by a single primary character (Fig. 236) and also in the *C. boucardi* group by an additional two secondary characters (Fig. 237). Other species groups are defined as monophyletic strictly on the basis of secondary characters, some of which involve extensive homoplasy. Monophyly of the *C. lobigerinus* group, which contains only two species, is based upon the primary character of presence or absence of a curved sulcus behind the eye. The apotypic state is found in both *C. lobigerinus* and *C. calandroides*, however, not in all individuals of each of the species. Monophyly of the *C. boucardi* group is based upon the emarginate or sinuate apex of the dorsal median pocket of the internal sac of males, larger size (secondarily reduced in *C. quadrilineatus*), and reduced pronotal postocular lobes (homoplasious in *C. canescens*). In the *C. americanus* group, monophyly is based upon the ventral tarsal pilose pads being reduced in extent (homoplasious in other *Cleonidius* and other Cleoninae). High incidence of brachyptery and of a mottled elytral scale pattern may be further evidence in support of monophyly of this group. Monophyly of the *C. poricollis* group is based

the basal arm short. Both of these characters exhibit homoplasy elsewhere in *Cleonidius*.

Within the *C. poricollis* group, three primary characters attest to the fact that *C. subcylindricus*-*C. texanus*-*C. longinasus* comprise a monophyletic group. Relationships of this lineage, however, to the rest of the species group is unclear (Fig. 237). *Cleonidius poricollis* and *C. eustictorrhinus* are considered sister-species, and this lineage along with *C. modestus* and the *C. subcylindricus*-*C. texanus*-*C. longinasus* lineage, are left as elements of an unresolved trichotomy. *Cleonidius texanus* and *C. longinasus* are regarded as sister-species because of lack of large punctures apically on the rostrum and presence of long dorsal erect vestiture. *Cleonidius subcylindricus* is therefore sister to the *C. texanus*-*C. longinasus* lineage.

In the *C. americanus* group, primary characters support only the sister-species status of each of *C. americanus* and *C. frontalis*, and *C. infrequens* and *C. puberulus* (Fig. 236); all other relationships are based on secondary characters (Fig. 237). *Cleonidius notolomus* and *C. collaris* are regarded as sister-species based upon pronotal sculpture although the state in *C. collaris* is much more extreme in its irregularity than that in *C. notolomus*. A moderately broad rostrum is used to propose monophyly of the *C. canescens*-*C. infrequens*-*C. puberulus*-*C. notolomus*-*C.*



and *C. notolomus*-*C. collaris* lineages. Acceptance of the *C. notolomus*-*C. collaris* lineage as having a secondarily plesiotypic, more elongate-narrow rostrum and of *C. puberulus* having secondarily rounded lateral rostral margins is necessitated by the sister-species status of *C. puberulus* and *C. infrequens*. Other interpretations require that these two not be sister-species and are in my view less parsimonious than that presented above. *Cleonidius canescens* is thus the sister-species of the *C. infrequens*-*C. puberulus*-*C. notolomus*-*C. collaris* lineage and the *C. americanus*-*C. frontalis* lineage sister to the remainder of species in the *C. americanus* group.

In the *C. boucardi* group, *C. boucardi* is considered sister to the remaining three species in the group based upon the ventral tarsal pilose pads being reduced in extent and the pronotal postocular lobes also reduced (both homoplasious in *Cleonidius*) in the latter three species. *C. quadrilineatus* and *C. placidus* are considered sister-species, with *C. trivittatus* as sister to that lineage, because of long dorsal erect vestiture in the former two species.

In the reconstructed phylogeny of *Cleonidius*, great emphasis is placed on scale pattern because this is the only primary character applicable in a broad or general sense within this genus (Fig. 236). As a result, misinterpretation

one character. If this character is misinterpreted, radical changes in the reconstructed phylogeny would result.

No alternative interpretations of this reconstructed phylogeny clearly stand as worthy of individual discussion. They would not likely, however, be concerned as much with relationships of species in groups, as with relationships of the species groups themselves.

### 5.3 Biogeography

#### 5.3.1 Methods of biogeographic analysis

Historical and ecological factors combine to produce the observed distributions of plants and animals, the explanation of which is the goal of the science of biogeography. Unfortunately many past studies of biogeography attempt to explain the distributions observed in terms of only one of ecology or history (Endler 1982). Distinguishing the relative roles of these factors must at least be considered although most agree it is very difficult to separate the effects of the two and to estimate their relative roles (Endler 1982; Vuilleumier and Simberloff 1980). The present study is no exception and my approach, although attempting to encompass aspects of both historical

Knowledge of distribution and aspects of natural history combine with knowledge of phylogenetic relationships and of earth history to allow for discussion of the historical biogeography and ultimately, the evolutionary history of the taxon in question. Ecological biogeography on the other hand does not require knowledge of phylogeny and earth history, but rather ecological theory and principles. Systematists and ecologists are the proponents of the historical and ecological approaches respectively. Rarely are the two approaches combined in a single study and relative roles assessed. This is brought about simply by the nature of the data bases because systematists frequently have not carried out the detailed sampling necessary for the application of ecological theory. Similarly, more often than not, ecologists do not have the knowledge of phylogeny that predicates all historical biogeographic analyses.

In ecological biogeography, the units selected for study are usually communities or various guilds of taxa that share association with a particular resource, be it habitat, food, geographic area, etc. (Cody and Diamond 1975). Ecological biogeographers are primarily concerned with proximal factors that determine the composition of the species assemblage but not the origins of these same factors. Little concern or attention is expressed or given to the phylogenetic relationships of the constituent taxa

only be developed in terms of detailed knowledge of phylogeny. Less frequently, ecological determinants of distributions of constituents of monophyletic groups, say a genus or family, have been studied, although rarely if at all with specific reference to detailed phylogenetic relationships.

Much of ecological biogeography addresses questions of dynamics and proximal origins of diversity, best applicable to islands or insular situations. Diversity patterns on continents are much more complex and are not so easily studied. Unfortunately, knowledge of population dynamics and detailed data about natural history attributes, permitting determination of niche breadths, are required for much meaningful ecological biogeography to be conducted (Cody 1975; Pianka 1975). Because of this, the approach I will take herein is only one of examination of natural history attributes and distributions, with specific reference to present environmental parameters.

Historical biogeographers, on the other hand, recognize and stress the need for knowledge of phylogeny, but generally at the expense of ignoring the role ecology might play in determining distributions. Indeed, to many, knowledge of phylogeny is essential prior to attempts at any meaningful historical biogeography. Wiley (1981) has noted three schools of historical biogeographers: 1, evolutionary;

knowledge of phylogeny and strictly monophyletic groups are the units of study for adherents of phylogenetic and vicariance biogeography and stress vicariance and allopatric speciation. Evolutionary biogeography however, emphasizes the role of dispersal with little regard for speciation mechanism or for details of phylogeny. This latter is not employed by many contemporary biogeographers for it is the approach resulting in the narratives and "scenarios" frowned upon for their *ad hoc* nature, and lack of applicability and rigour. Heuristic advantages of the phylogenetic and vicariance schools of biogeography are consistent with the advantages of a phylogenetic approach to phylogeny reconstruction and are discussed in more detail by Wiley (1981), Patterson (1981), and Platnick and Nelson (1978).

Ball (1975) recognizes three phases through which biogeographic studies pass: 1, descriptive or empirical phase; 2, narrative phase; and 3, analytical phase. These phases, as noted elsewhere herein and by Patterson (1981), apply equally well to systematic studies, especially methods of phylogeny reconstruction. Descriptive or empirical biogeography is simply the data base. What are the extant distributions and natural history attributes of the taxa in question? What habitats do they occupy? What do they eat? When are they active?

biogeographic studies. Narrative explanations, or general evolutionary "scenarios" as they have been called by Eldredge (1979), have frequently been criticized as pseudo-science or bad science because of their unfalsifiable and *ad hoc* nature, and for their explanatory value, which is, more often than not, concerned only with particulars as they apply to the taxon in question (Goudge 1961; Hull 1974; Ball 1975; Eldredge 1979; Eldredge and Cracraft 1980; Brown and Gibson 1983). General applicability is not usually a concern of narrative explanations, although there is no reason why it should not be.

Consequently, Eldredge (1979) notes two ways of improving these "scenarios". Firstly, base them more explicitly on knowledge of phylogenetic relationships, and secondly, eliminate some if not all of the more purely speculative elements. To these suggestions, I add a third; formulate explanations as hypotheses in terms that are testable, predictive, and have potential for more general applicability than just the taxon in question. With these improvements in mind, "scenarios" are capable of producing ideas that can be subsequently rigorously tested in a hypothetico-deductive manner for acceptance as general scientific facts, ideally of widespread applicability and significance. In essence, this is the analytical phase of Ball (1975).

In my analysis of historical biogeography I employ a phylogenetic approach to place aspects of the phylogeny of North American Cleoninae in a geographic perspective and thus come to some degree of understanding of the geographic history of that specific taxon. Resultant potential generalizations and predictions based on these results are enumerated, discussed and formulated as hypotheses of general potential utility.

A vicariance approach, examining distribution patterns of other North American arid land taxa and the plotting of generalized tracks will not be attempted herein but should be carried out as more studies of such taxa, that include a phylogenetic basis, become available.

### 5.3.2 Genus *Apleurus* Chevrolat

#### 5.3.2.1 Descriptive biogeography

##### 5.3.2.1.1 Diversity patterns

The genus *Apleurus* is represented in arid land habitats of southwestern North America and Mexico by eight species placed in two subgenera. Maximum species diversity is in southern California, northwestern Mexico (including Baja California Norte), southern Arizona and southern New Mexico (Fig. 230). Diversity decreases beyond this area especially

to the east and the north. Species of *Apleurus* do not occur north of 45°N latitude, east of 95°W longitude, or south of 15°N latitude (Fig. 230).

Five of the eight species are variously restricted in their distributions and of these five, only two, *A. jacobinus* and *A. hystrix* are macrosympatric. The remaining three species, *A. lutulentus*, *A. angularis*, and *A. albovestitus* are widespread and broadly sympatric with each other and various of the five geographically restricted species. Gross geographic range overlap is relatively low with 12 of 28 possible geographic distributions of species pairs resulting in at least partial sympatry (i.e. on average, each species of *Apleurus* is sympatric with 42.8% of the genus *Apleurus*, or 3.424 species). Nine of the 12 sympatric species pairs exhibit overlapping distributions; three are nested (the area of distribution of one species occurring entirely within the area of distribution of the other).

#### 5.3.2.1.2 Habitat associations

Habitat associations of the species are varied, ranging from grasslands to desert-grassland transitional habitats or deserts in eurytopic species (three of the eight); others, are more stenotopic, found only in grasslands, desert-grassland transitional habitats, or in deserts (five of the eight) (Fig. 238).



*Apleurus aztecus* is a mesquite-grassland inhabitant in central Mexico. *Apleurus lutulentus*, although also found in desert-grassland transitional habitats, is primarily a mesquite-grassland and grassland inhabitant as evidenced by its more easterly distribution into eastern Texas, Oklahoma and Kansas and disjunct southerly distribution into central Mexico, but not in the Mexican portion of the Chihuahuan desert (Fig. 207). *Apleurus angularis* is also found in grasslands, however, in contrast to *A. lutulentus*, does not extend as far east or south as does that species. Rather, *A. angularis* extends further west and north into Great Basin desert. *Apleurus jacobinus* and *A. porosus* are semi-desert and semi-desert to desert inhabiting species respectively. *Apleurus albovestitus* is distributed even more to the west than is *A. angularis*. It occurs in most western semi-desert and desert, and desert-grassland transitional habitats. *Apleurus saginatus* is a desert-grassland inhabitant bordering the eastern Sonoran Desert region (Fig. 206). *Apleurus hystrix* is a sand dune inhabiting species in the California semi-desert habitat.

Relative measures of association of species with particular types of habitats reveals a slightly higher level of association with desert-grassland transitional habitats than with others. These measures were obtained by summing the total number of species of *Apleurus* occupying a particular habitat type scoring one for a marked association (solid box in Fig. 238) and one-half for a less marked

association (half box in Fig. 238). This number is then divided by the number of possible associations with that habitat type to give a relative measure of the degree to which species of the genus occur in the various habitat types. Measures of association with the four desert habitats is 0.156; with semi-deserts, 0.250; with desert-grassland transition habitat, 0.375; and, with mesquite-grasslands and grasslands, 0.313.

#### 5.3.2.1.3 Plant associations

Details of host plant associations are largely unknown for all species and so all plant associations, many likely accidental, have been used as indicative of the probable taxonomic range of host plants (Fig. 239). Adults of all species are found primarily or exclusively on Compositae although in some species, especially *A. albovestitus*, adults are found sporadically on a variety of other types of plants as well. Other than Compositae, Chenopodiaceae and Leguminosae constitute the bulk of the remainder of the associations. Clear associations with taxa lower than the hierarchical level of family are not evident.

#### 5.3.2.1.4 Altitudinal zonation

Altitudinal ranges are shown as a Hubbs-Hubbs diagram (Fig. 232). Although species may be represented over an extensive range of altitudes, most members of each species are found over a narrower part of this range. This is

indicated by the empty rectangles which indicate 1.5 standard deviations on either side of the mean. *Apleurus hystrix*, *A. jacobinus* and *A. porosus*, species restricted to the Pacific coastal areas, are found over a much narrower range of altitudes than are more widespread species. Mean values for altitude of samples of these three coastal species do not significantly differ as evidenced by the overlap of two standard errors on either side of the sample mean. Among the remaining species, only *A. lutulentus* and *A. aztecus* have sample mean values which do not significantly differ. The three remaining species, *A. angularis*, *A. albovestitus*, and *A. saginatus*, have sample mean values which significantly differ from each other, and from any of the species in the genus. On average, the 95% probability limits of the mean value of altitude for a species of *Apleurus*, as defined by two standard errors on either side of the sample mean, is not significantly different from the mean of another species in 14.2% of all possible combinations of species pairs. There is also a trend towards occupation of lower altitudes by increasingly derived species in the genus.

#### 5.3.2.1.5 Seasonal occurrence

Patterns of adult seasonal occurrence are broad with *A. angularis*, *A. jacobinus*, *A. albovestitus* and *A. saginatus* found throughout the year. *Apleurus porosus* adults are found from January to November and are likely active throughout

the entire year. On the other hand, *A. lutulentus* adults are known from March to October; *A. aztecus* from June to August; and *A. hystrix* from February to July. Few records are available, however, for these last two species. Detailed life histories are unknown for all species.

#### 5.3.2.1.6 Co-existence

Consideration of these various niche dimensions as a group reveals that no two species of *Apleurus* share the same or largely overlapping ranges in all niche dimensions. Statistical measures aside, inspection shows species differences along the dimensions of altitude, geographic distribution, and habitat association. All species share a plant association with Compositae. Macrosympatric species differ in various aspects of altitudinal range and habitat association. Species with overlapping altitude ranges are allopatric or differ in habitat association. In general, niche overlap between species of *Apleurus* is low.

#### 5.3.2.1.7 Wing length polymorphism

At least some adult members of most *Apleurus* species lack wings; 97.9% of *A. lutulentus*, 4.5% of *A. angularis*, 99.8% of *A. albovestitus*, and all *A. saginatus* are the only macropterous individuals known. Adults examined of species of restricted distribution such as *A. hystrix*, *A. aztecus*, *A. jacobinus* and *A. porosus* are all apterous. The most widespread species, *A. albovestitus*, *A. lutulentus* and *A.*

*angularis* have variable percentages of macropterous individuals. These individuals are scattered throughout the species ranges and are not restricted in their distributions.

### 5.3.3 Genus *Cleonidius* Chevrolat

#### 5.3.3.1 Descriptive biogeography

##### 5.3.3.1.1 Diversity patterns

The genus *Cleonidius* is represented in North and Central America by 19 species; a single species has so far been recognized from the Palearctic Region but is not considered further in the biogeographic discussions following. Maximum species diversity is in the southwestern United States, especially southern and central California, southern Nevada, Arizona, southern Utah, New Mexico, Colorado and western Texas (Fig. 231). Diversity decreases slightly to the north in northern Utah and Wyoming into eastern Washington and western Montana, but more markedly into Oregon and western Washington to the west, into eastern Texas, Oklahoma and Kansas to the east, and especially to the south into Mexico (Fig. 231) (although this may prove an artifact of insufficient collecting). Species of the genus extend north into southern Canada and south as far as

Honduras; two species occur along the east coast of North America.

Species of *Cleonidius* tend to be more or less widespread in their geographic distributions. Only five of the nineteen species are what I consider restricted in their geographic distributions. Three of these five, *C. americanus*, *C. placidus*, and *C. eustictorrhinus* are macrosympatric and found only in California; *C. collaris* is found only in eastern Colorado and nearby localities in adjacent states; and *C. calandroides* is found only along the northern Atlantic coast. Three other species (*C. puberulus*, *C. notolomus*, and *C. infrequens*) are slightly less restricted in their distributions. Of the remaining 11 species, 10 are moderately to very widespread in the southwestern United States and Mexico, or western North America and are largely macrosympatric with each other and with various of the species with restricted distributions. One species, *C. subcylindricus*, is widespread along the Atlantic Coastal Plain. One of the six species in the *C. poricollis* group, one of two in the *C. lobigerinus* group, one of four in the *C. boucardi* group, and five of seven in the *C. americanus* group have more or less restricted distributions.

Gross geographic range overlap is more extensive than in *Apleurus*. In the *C. poricollis* group, eight of 15 possible geographic distributions of species pairs result in at least partial sympatry (three overlapping, five nested);

two of six comparisons in the *C. boucardi* group (two overlapping) and 11 of 21 in the *C. americanus* group (seven overlapping, four nested) result in at least partial sympatry. Species in the *C. lobigerinus* group are allopatric. For the genus *Cleonidius* as a whole, 87 of 171 possible combinations result in at least some sympatry (62 overlapping, 25 nested). In other words, a single species is on average macrosympatric with 50.9% of the genus, or 9.67 other species of *Cleonidius*.

#### 5.3.3.1.2 Habitat associations

Habitat associations are varied in species of the genus, ranging, as in *Apleurus* species, from grasslands to deserts in some, to principally deserts, to principally grasslands in others (Fig. 240). Measures of primary habitat associations (calculated as for *Apleurus* species), differ slightly from those in *Apleurus* however, because in *Cleonidius*, association with desert habitats is low (0.099), increasing in semi-desert habitats to 0.171, and desert-grassland transitional habitats to 0.316. The highest association is with mesquite-grasslands and grasslands (0.474). Association with western xerophytic evergreen forest is 0.132 whereas no *Apleurus* were associated with this habitat type. Among the species groups, the *C. poricollis* group is relatively well represented in deserts (0.125), also in semi-desert (0.250), but less so in desert-grassland (0.167) and markedly less so in

mesquite-grasslands and grasslands (0.083). In contrast, the *C. americanus* group has values of 0.054 for deserts, 0.107 for semi-desert, 0.429 for desert-grassland, and 0.714 for mesquite-grasslands and grasslands. The *C. boucardi* group has values of 0.000, 0.125, 0.250, and 0.500 respectively for these same habitats. The only southwestern species in the *C. lobigerinus* group is found in a variety of habitats. With the exception of this species, *C. lobigerinus*, other *Cleonidius* species are restricted to one or only a few habitat types.

#### 5.3.3.1.3 Plant associations

Unlike in *Apleurus* species, which are all associated primarily with Compositae, plant associations in *Cleonidius* species are much more varied and span a much broader taxonomic range of plant families (Fig. 241). A number of species are associated with Compositae, Rosaceae, Cruciferae, and Leguminosae. There is slight association with Chenopodiaceae and scattered records of associations with members of various other plant families. In the *C. parvicollis* group, associations are varied, with one species on each of Compositae, Rosaceae and Rhamnaceae, and Cruciferae, and two on Leguminosae. Plant associations are not well known for *C. eustictorrhinus*. In the *C. boucardi* group, two species are associated with Leguminosae, one species with Rosaceae and Rhamnaceae, but also Leguminosae; plant associations are not known for *C. boucardi*.



Associations are primarily with Compositae and Cruciferae in *C. lobigerinus*; exclusively with Cruciferae in *C. calandroides*. Finally, plant associations are few for most species in the *C. americanus* group with the exception of *C. americanus* which appears to be associated with Compositae.

#### 5.3.3.1.4 Altitudinal zonation

Altitudinal ranges are shown as a Hubbs-Hubbs diagram (Fig. 233). In general, each species is found over a wide range of altitudes which overlap extensively with ranges of many other species. An altitudinal overlap value of 67 out of 171 or 0.392, was derived for *Cleonidius*. This measure was based on comparison of the 95% probability limits of the species mean as defined by 2.0 standard errors on either side of the sample mean. This value indicates that each species of *Cleonidius* has on average, a mean value for altitude which is not significantly different from the same values for all other species of *Cleonidius* in 39.2% of all species pair combinations. This value contrasts well with *Apleurus*, where the value was only 14.2%, and demonstrates that *Cleonidius* species are not markedly altitudinally segregated as are *Apleurus* species. This marked overlap also exists in the species groups (0.667 for the *C. poricollis* group, 0.286 for the *C. americanus* group [perhaps artificially low because of few collection localities for *C. puberulus* and *C. collaris*], 0.500 for the *C. boucardi* group). *Cleonidius lobigerinus*, and three members of the *C.*

*boucardi* group, *C. boucardi*, *C. trivittatus* and *C. quadrilineatus*, have the largest altitudinal ranges of species in *Cleonidius*. Individuals of these species have been recorded at maximum altitudes ranging from 3276 to 4004 m.

#### 5.3.3.1.5 Seasonal occurrence

Patterns of adult seasonal occurrence are broad with most species found throughout the greater part of the year from early spring to late winter months.

#### 5.3.3.1.6 Co-existence

In contrast to species of *Apleurus*, examination by inspection of niche dimensions as a group shows the majority of *Cleonidius* species differ most in terms of host plant (Fig. 241) and habitat associations (Fig. 240). Species geographic range overlap is generally high, as is the measure of altitudinal range overlap and as is overlap in seasonal occurrence. Within the *C. poricollis* group, extent of sympatry and altitudinal range overlap are high, but different species in the group are associated with different habitat types and with different plant families. In the *C. americanus* group, extent of sympatry and altitudinal range overlap are low and habitat associations overlap extensively; plant associations are mostly unknown. In the *C. boucardi* group, sympatry is low, habitat associations differ, host plant associations differ slightly, but

altitudinal range overlap is high.

The genus *Cleonidius* is more diverse than *Apleurus* and is more extensive in its distribution. Whereas *Apleurus* species appear to be more associated with deserts and less so with grasslands, *Cleonidius* is very well-diversified in grasslands and desert-grassland transitional habitats and is not associated with deserts to any significant degree (except for species in the *C. porticollis* group). Only in *Cleonidius* are some species associated with the western xerophytic evergreen forest. Patterns in host plant association differ markedly, for species of *Apleurus* are almost exclusively associated with Compositae, whereas species of *Cleonidius* are associated to various degrees with one of Compositae, Cruciferae, Leguminosae, Rhamnaceae, Rosaceae and to a lesser extent Chenopodiaceae.

#### 5.3.3.1.7 Wing length polymorphism

All individuals examined of the *C. boucardi* group are macropterous; in the *C. lobgerinus* group 99% of *C. lobgerinus* and all *C. calandroides* are macropterous; and, in the *C. porticollis* group, only in *C. porticollis* are brachypterous individuals known (91%). In contrast, in the *C. americanus* group, all *C. canescens*, *C. puberulus* and *C. collaris*, and 79% of *C. notolomus* and 38% of *C. infrequens*, are brachypterous.

### 5.3.4 Historical biogeography of New World *Apleurus* and *Cleonidius*

#### 5.3.4.0.1 Introductory Comments--A Warning

The evolutionary history of plant and animal taxa that presently inhabit the southwestern United States and adjacent areas has been very complex. Not surprisingly, determination of what this history has been is also a very difficult task and based on various assumptions. Changing, locally variable climates, mainly a trend during the Tertiary of increasing aridity (but here and there punctuated by pluvial periods of various duration) resulting in various and ephemeral habitat types, continual but localized orogenic activity, and locally variable topography, all result in a complex framework for speciation events to be understood and even approximately dated. Adding to this is perhaps the most significant problem, that of the Quaternary, for it is known that distributions of many organisms changed radically during this time period (Coope 1979). Owing to these presumptive changes, many of the distribution patterns may reflect only Quaternary distributional shifts and not original speciation events, these perhaps having occurred elsewhere and under very different ecological conditions. In other words, the present distributions may be the result of secondary, not primary contact zones (Endler 1977). In the absence of fossils, the heuristic value of historical biogeography may therefore be

questioned because in order to overcome this problem, assumptions about habitat associations, other ecological attributes, and especially distributions of the species at different times, or of their ancestors, are required, based on these features in extant species. These can only be made with reference to a reconstructed phylogeny and as such the claim of phylogeny before biogeography is in large part valid for analyses to be maximally scientific. Whether or not these assumptions should be accepted is a moot point.

Because biogeographic statements are so dependent on a reconstructed phylogeny, another problem is that of confidence in the reconstructed phylogeny. A weak reconstructed phylogeny results in weak biogeographic statements that may prove misleading should they be uncritically accepted as facts.

In *Apleurus*, only the possibility of changed distributions is a problem, for confidence in the reconstructed phylogeny is high. On the other hand, both are — problems in *Cleonidius*. Confidence in the fully resolved reconstructed phylogeny is low due to the large number of secondary characters used in the analysis and concomitant homoplasy in these characters. Some biogeographic statements about details of species origins in *Cleonidius* must therefore be cautiously accepted, and alternatives, some of which will be discussed, must also be considered.

A further complication in *Cleonidius* is recognition of a Palearctic species component which is not herein

considered in the reconstructed phylogeny. The single species examined from the Palearctic region can however be placed either as sister to the Nearctic *Cleonidius*, or as a member of the *C. americanus* group. Biogeographic significance of both placements will be discussed.

#### 5.3.4.0.2 Historical Explanations

In *Apleurus*, diversity gradients do not appear to be easily explainable in terms of latitude, elevation, or any other climatic factor with the possible exception of regional degree of variation in aridity. As noted elsewhere, evolution of *Apleurus* appears to parallel evolution of increasingly arid habitats. This is supported by comparison not only of habitat associations with the reconstructed phylogeny (Fig. 238), but also of altitudinal ranges with the reconstructed phylogeny which trend toward occupation of lower, and thus likely more arid altitudes. Thus, if any ecological factor might prove important in explaining why southern California, Arizona, New Mexico and extreme northwestern Mexico have the highest species diversity (Fig. 230), it is likely the degree of habitat and associated floral complexity largely due to the occurrence of desert and desert-grassland transitional habitats in these areas. Species diversity drops markedly beyond the area where these habitats occur.

Similarly, in *Cleonidius*, maximum species diversity is again noted in these same areas, but is also high throughout

the rest of Arizona and New Mexico, western Texas, central California, and southern Nevada, Utah, Colorado and Wyoming (Fig. 231). Species diversity thus appears again to be influenced by degree of arid habitat complexity. However, high species diversity is more widespread in *Cleonidius* because of the higher level of association of the species in the genus with more widespread grassland and woodland habitats (Fig. 240). Although evolution in this genus has likely also been influenced by increasing aridity in southwestern North America, these influences are not as clearly indicated as in *Apleurus*. In *Cleonidius*, adaptation to the most arid habitats is low, and to the least arid, relatively high. Furthermore, there is no clearly evident evolutionary trend for species of the genus to be found in increasingly arid habitats (Fig. 240), or at successively lower altitudes as are noted in *Apleurus*. High diversity in *Cleonidius* thus extends outside the area of the highest habitat complexity and especially into areas of grassland, woodland, and desert-grassland transitional habitats. High degree of association with less arid habitats also explains the broader geographic and altitudinal range of the species in the genus and perhaps also the occurrence of the genus in the Palearctic region as well. In addition to the effects of habitat on the evolution of the group, unlike *Apleurus*, which are all apparently associated with Compositae, species of *Cleonidius* are associated with a much broader taxonomic array of plants (Fig. 241). Clearly, host plant shifts are

an additional factor to consider in the evolution of this latter genus.

In the present study I address the hypothesis that evolution and speciation within *Apleurus* and *Cleonidius* closely parallel increasing aridity and associated changes in distribution and composition of habitat types in the southwestern United States and Mexico. History of distribution and composition of habitats in these areas has been dealt with extensively in paleobotanical studies by Axelrod (1950, 1975, 1979, 1985), Axelrod and Raven (1985) and Raven and Axelrod (1978) and thus interpretation of the historical biogeography of *Apleurus* and *Cleonidius* draws largely upon the results of these publications.

There appear to be three possible origins of arid land faunas in North America, only two of which are likely for Cleonini. First, as discussed by Axelrod (1975), a continuous North American-western Eurasian belt of Madrean-Tethyan sclerophyll woodland vegetation may have existed in the early Paleogene, with subsequent vicariance of this habitat, as a result of Atlantic Ocean floor spreading, proposed to account for any present disjunctions in distributions. Secondly, and perhaps the generally more favourable hypothesis, Webb (1977) notes the marked similarity between savanna-adapted mammal faunas in Asia and North America in the late Eocene (postdating similarities between the faunas of Europe and North America), and invokes subsequent vicariance of these habitats across Beringia as



responsible for present distributions. A third possibility, South American origins, as noted by Raven (1963) and Wells and Hunziker (1976) for various plants, does not appear likely, because no Cleonini are known from that continent (Wibmer and O'Brien 1986).

There are no reasons to doubt that *Apleurus* is North American in origin and likely evolved *in situ*, on Compositae, in isolation from its sister genus, the Palearctic *Chromoderus*, as a result of vicariance of sclerophyll woodland or savanna habitats in Holarctica either between western Europe and North America no later than early Eocene or between eastern Asia and North America no earlier than the late Eocene (Fig. 242) respectively. Accepting that *Apleurus* evolved in North America and on Compositae, investigation of the role of history proceeds by examination of distributions and aspects of natural history with respect to the reconstructed phylogeny.

First, aside from *A. hystrix*, the next two most plesiotypic species, *A. aztecus* and *A. lutulentus*, are exclusively and primarily mesquite-grassland and grassland inhabiting species, respectively (Fig. 238). This, and the increased association of more derived species with semi-desert and desert habitats, demonstrates a clear trend from mesquite-grassland and grassland to desert habitation within species of the genus (Fig. 238), and indicates that mesquite-grasslands, grassland, or a precursory habitat type not present today, or present but lacking association with

any *Apleurus* species, was likely the ancestral habitat for the genus. This habitat was likely a grassland or savanna habitat, because no *Apleurus* are known from woodland habitats, plesiotypic *Apleurus* are currently associated with this or a similar type of habitat (Fig. 238), no eastern North American *Apleurus* are known, and *Chromoderus* species judging from their distributions, are likely steppe or grassland inhabitants. In view of this, a trans-Beringian savanna or grassland vicariance origin of *Apleurus* (Fig. 242) is to be preferred over the Madrean-Tethyan vicariance of Axelrod (1975), which would predict southwestern North America-Mediterranean area relationships, presence of plesiotypic species in woodland habitats, presence of plesiotypic species in southeastern North America, and presence of closely related taxa in similar extralimital woodland habitats.

Grasses appear to be present in North America in the early Tertiary as evidenced by the first appearance of grass pollen in the Paleocene, and the first grass microfossils and first appearance of structural adaptations in mammals to open habitats (including especially, the initial albeit slight development of high crowned teeth resistant to the abrasive properties of grasses) in the Oligocene (Clayton 1981; Webb 1977).

Savanna or woodland precursors of mesquite-grassland and grassland and desert habitats likely first emerged as distinct habitats in the mid- to late-Eocene as a result of

a slowly expanding dry climate and a marked increase in seasonal aridity fostering *in situ* adaptation by already existing subhumid plants to these new and increasingly semi-arid conditions (Axelrod 1950, 1975, 1979; Axelrod and Raven 1985; Cronquist 1978; Webb 1977). Arboreal vertebrates are less frequent in fossil deposits of this age, a number of North American groups developed special adaptations to savanna habitation, and new, savanna-adapted groups emigrated from Asia where they had already developed their adaptations for this type of life (Webb 1977). Adaptation and radiation in animals and plants (especially dicotyledonous herbs) continued into the Oligocene as arid habitats proceeded to expand and diversify (Axelrod 1979; Cronquist 1978; Webb 1977). At this time semi-arid woodland or savanna habitats were apparently relatively extensive in North America (Axelrod 1975) and were likely interspersed with localized grasslands. Both faunas of late Eocene (Webb 1977), and floras of the Oligocene (Axelrod 1979), document relationships between taxa in southern California and those in the Rocky Mountain region (Axelrod 1975; Raven and Axelrod 1978). Axelrod (1979, Figure 6) and Raven and Axelrod (1978) further note that a number of congeneric "closely related" woodland taxa are presently disjunct in insular and maritime southern California and the highlands of central Mexico. A continuum may therefore have existed between these two areas and perhaps also the southern Rocky Mountain region from perhaps the late Eocene, at the

earliest, into the Miocene (Axelrod 1950; Raven and Axelrod 1978). The present woodland in the uplands of central Mexico is the habitat thought to be most like the ancestral Oligocene-Miocene woodland, hence the name Madrean for this vegetation type (Axelrod 1975, 1979). Development of increasingly arid habitats intermediate between these areas throughout the later Tertiary, likely resulted in at least some of these present disjunctions (Axelrod 1975).

This geographic pattern applies well to *Apleurus* because the most plesiotypic species of *Apleurus*, *A. hystrix*, is found in sand dunes in southern California, whereas the next most plesiotypic two species, *A. aztecus* and *A. lutulentus*, are found inland in central Mexico and in central Mexico but also in Arizona, New Mexico, Texas and adjacent states, respectively (Fig. 242). This pattern, although between a California sand dune species and Mexican or southern Rocky Mountain mesquite-grassland and grassland taxa, is geographically similar to the relict disjunctions noted above for semi-arid woodland plants and is suggestive that the same may also apply to taxa initially adapted to savanna or grasslands. If isolation and the relict nature of *A. hystrix* is a result of an early vicariance of California and inland semi-arid grassland or savanna habitats, we might expect similar phylogenetic and geographic relationships to those in *Apleurus* in other taxa in which a structurally markedly distinct species is restricted to one of the southern California coastal dune habitats. Unfortunately,

although endemics are known to occur in many California dune fields (Andrews *et al.* 1979), phylogenetic and geographic relationships of the endemics have not been studied in detail. Most extensive survey efforts have also concentrated principally on inland dune fields (Andrews *et al.* 1979; Hardy and Andrews 1976) Some of these endemics are also not of a relict nature and are likely a product of the later Tertiary climatic changes that fostered the rapid and numerous speciation events thought to have contributed so extensively to the marked levels of endemism in the California flora (Stebbins and Major 1965; Raven and Axelrod 1978). Perhaps relicts such as *A. hystrix* in California, are managing to survive in restricted or marginal habitats having changed habitat associations from the grassland or savanna habitat perhaps due to exclusion by newly evolved competitively superior species, or due to the changing nature and ephemeral status of habitats to which they were initially adapted. Raven and Axelrod (1978) note that among plants, a number of Tertiary relicts have survived in California owing to its sheltered and equable climate.

Grassland habitat only reached its present form and more extensive distribution during the Miocene and early Pliocene (but especially at the Miocene-Pliocene transition) (Axelrod 1950, 1985; Webb 1977). Because of this, it is unlikely that divergence between *A. aztecus* and the ancestor of the remainder of *Apleurus* took place before this time. Restriction of *A. aztecus* to mesquite-grasslands, and

primary association of *A. luteolentus* with mesquite-grasslands and grasslands, suggests that the ancestor of the subgenus *Apleurus*, like the ancestor of the genus *Apleurus* as noted, was also associated with grassland-like habitats. Present restriction of *A. aztecus* to central Mexico is unlikely to have been a result of competitive exclusion by *A. luteolentus* in the grasslands of the United States, for *A. luteolentus* coexists with *A. aztecus* in central Mexico (Figs. 207, 210). Rather, a late Miocene or early Pliocene north-south vicariance of Plains and Mexican Plateau grasslands, as a result of increasing aridity and development of more extensive semi-desert habitats (Webb 1977) can account for the distribution, with a subsequent dispersal of *A. luteolentus* south through desert-grassland transitional habitat into central Mexican mesquite-grasslands and the Hidalgo Desert region (term of Shreve 1940; Axelrod 1979) of a later Pleistocene origin. I suggest this later dispersal route was through a desert-grassland transitional habitat, not grassland or mesquite-grassland, because of the absence of concomitant dispersal north by *A. aztecus*. Webb (1977) has drawn attention to the role of central Mexico, noting that Miocene faunules from this area play a focal role in the history of the semi-arid biota of North America. Similarly, Cronquist (1978) notes that the tribe Heliantheae of the Asteraceae (Compositae herein) has its present center of diversity in the arid highlands of central Mexico, suggesting that this

area has been important in the evolutionary history of the group.

Increased aridity during the middle Pliocene resulted in increased extent of semi-desert habitats and likely the origin of the ancestor of the remainder of *Apleurus* in western desert-grassland transitional habitats, and *A. lutulentus* in more easterly, primarily grassland, habitats (Fig. 243).

Divergence of the *A. porosus*-*A. jacobinus*-*A. angularis* and *A. saginatus*-*A. albovestitus* lineages is more problematical, for extant species of the two groups, especially *A. angularis* and *A. albovestitus*, do not differ much in their habitat associations or distributions.

In the *A. saginatus*-*A. albovestitus* lineage, an east-west vicariance of desert-grassland and semi-desert habitats respectively (Axelrod 1979, figure 4) as a result of late Quaternary aridity following the increased precipitation of the late Pliocene and early Pleistocene (Axelrod 1979) likely resulted in the origin of *A. saginatus* and *A. albovestitus* respectively (Fig. 245). Although *A. albovestitus* is presently a widespread species, examination of intraspecific variation shows the most plesiotypic form to be that from coastal Californian semi-desert (possession of large glabrous abdominal patches, deeper and larger punctation; see "Geographic variation" of *A. albovestitus*). Presence of other forms of *A. albovestitus* to the east in other semi-desert and desert habitats is therefore

suggestive of recent dispersal followed by post-glacial local or regional selection pressures in various regional habitats which are hypothesized to be largely of a recent origin in the late Quaternary. (Spaulding *et al.* 1985). Unlike *A. albovestitus*, *A. saginatus* has not expanded its range and is still found only in the desert-grassland transition bordering the eastern portion of the Sonoran desert (Fig. 206).

In the *A. porosus*-*A. angularis*-*A. jacobinus* lineage, patterns of vicariance (Fig. 244) among the three species are clearly evident for they are allopatric (Figs. 208, 210-211). Because it has only been during the last four million years that Baja California reached its present position and acquired a notably drier climate and coastal foggy desert (Axelrod 1979), origin of *A. porosus* does not likely predate that time. Increased aridity in the area likely allowed entry of a desert-grassland-adapted ancestor from the north, as Baja California was reaching its present position. This form is presumed to have evolved *in situ* into *A. porosus*. An east-west vicariance of desert-grassland and semi-desert-adapted populations of the ancestor of *A. jacobinus*-*A. angularis*, as in the *A. albovestitus*-*A. saginatus* lineage, likely accounts for the origin and present distribution of *A. jacobinus* and *A. angularis*. Present distributions of *A. porosus*, *A. jacobinus*, and *A. angularis* may be maintained as allopatric by competitive exclusion or by selection pressures promoting reproductive



isolation.

For *Apleurus* then, increasing aridity and concomitant changes in habitat structure and availability appears to be the scenario most consistent with ideas and evidence about paleoenvironments and their distribution in space and time in southwestern North America.

In *Cleonidius*, the pattern is somewhat different. *Cleonidius* species, in general, appear to parallel some of the aspects of evolution of *Apleurus* species or lineages, but differ markedly in various others.

As mentioned, at least one species of *Cleonidius* is known from the Palearctic Region. However, its relationships are at present unclear. The manner in which the vittate elytral scale pattern is produced is apparently fundamentally different from that in other *Cleonidius*, and argues for exclusion from the *C. americanus* group, where it would otherwise be placed, and likely for sister-group status to the rest of *Cleonidius* (see Phylogeny section).

Unlike *Apleurus*, *Cleonidius* does not appear to be North American in origin because the Palearctic species component of *Cleonidius* appears to be sister to the Nearctic species component (Fig. 246). Thus, a Palearctic origin for the genus is to be preferred with subsequent entry into North America of this ancestral stock. It is most likely that a trans-Beringian grassland or savanna vicariance of this stock (Fig. 246), such as that proposed for the origin of *Apleurus*, resulted in the evolution of Nearctic *Cleonidius*.

The evidence for this is as follows: 1, Palearctic *Cleonidius* are eastern in their distribution and are apparently associated with grassland or steppe habitat; 2, association with grasslands is widespread in *Cleonidius* (although not restricted to primitive lineages or species) (Fig. 240); 3, association with woodland, although widespread, is only in more derived species (Fig. 240) (suggestive of secondary entry into that habitat type, a hypothesis supported by patterns of host plant association); and 4, the two eastern North American species are unrelated and also of a generally more derived nature. This trans-Beringian savanna or grassland event is likely to have occurred only post-late Eocene.

Within *Cleonidius* there are five of what I consider to be main, perhaps interrelated, differences from *Apleurus*: 1, species of *Cleonidius* are known from western xerophytic conifer woodland (Fig. 240); 2, species of *Cleonidius* are known from eastern North America (Figs. 215, 229); 3, species exhibit primary host plant associations with one of a various number of plant families (Fig. 241); 4, habitat association overlap (Fig. 240), altitudinal range overlap (Fig. 233), and association with less arid habitats are high; and 5, comparison of habitat associations or altitudinal ranges with the reconstructed phylogeny does not demonstrate a clear correlation with increasingly arid habitats (Fig. 240) or with lower altitudes, respectively. These differences are certainly suggestive of a different

evolutionary history, one influenced less so by shifts to increasingly arid habitats, but rather by host plant shifts, allopatric speciation within a habitat type, and habitat shifts to both more and less arid types at both lower and higher elevations over a more extensive geographic range.

There are, however, some similarities. Distributions of three of the four species groups can be geographically centered in certain areas. The *C. porcollis* group has five of six species distributed in whole or in part in California (Fig. 257); the *C. americanus* group has six of seven species distributed in whole or in part in the southern Rocky Mountain region (Fig. 258); and, the *C. boucardi* group has three of four species in the central to southern Rocky Mountain region or central Mexico (the most primitive being in central Mexico) (Fig. 259). With certain restrictions, namely the question of the origin of the *C. lobigerinus* group, this pattern brings to mind the California-central Mexico-southern Rocky Mountain pattern in primitive *Apleurus* (Fig. 242) because the common ancestor of the *C. boucardi* group (Mexico) and the *C. lobigerinus* group (questionable origin) is sister to the *C. americanus* group (southern Rocky Mountains), the common ancestor of which is sister to the *C. porcollis* group (California).

This pattern is suggestive of similar historical -- processes accounting for diversification in the basal lineages of the two genera, although it is possible that in *Cleonidius* the ancestral habitat association is with

sclerophyll woodland rather than grasslands or savanna for the two were undoubtedly intermixed (Axelrod 1979). As such, an Eocene to ~~Miocene~~ vicariance of California, southern Rocky Mountain region, and central Mexico was likely responsible for the initial diversification within the genus (Fig. 246).

Parsimony, however, dictates preference for an association with grasslands or savanna habitat because most *C. americanus* group members have a primary association with that habitat type; *C. lobigerinus* and *C. porlcollis* group members, although not primarily associated with grasslands, are not primarily associated with woodlands; and primitive *C. boucardi* members are associated with grasslands (Fig. 240). Furthermore, the two species associated with western xerophytic conifer woodland and the single species associated with eastern North American mesophytic evergreen forest, are more or less structurally derived and not relictual in distribution. Similarly, the other eastern North American species is also structurally derived, and although restricted in distribution, is the sister-species of a structurally similar western North American species (Fig. 236).

The best evidence, however, for association with xerophytic woodland being secondarily derived concerns host plant associations. *Cleonidius texanus* and *C. quadrilineatus* are both associated primarily with Rosaceae and Rhamnaceae and each have their two closest relatives associated

exclusively with Leguminosae (Fig. 241). This, and presence of infrequent association of *C. texanus* and *C. quadrilineatus* with Leguminosae (but not of relatives on Rosaceae or Rhamnaceae), suggests Leguminosae was the plesiotypic host plant for these lineages and that Rosaceae and concomitantly association with xerophytic woodland, derived.

Indeed, the widespread distribution and abundance of species associations with Leguminosae suggests that association with this family, or an ancestral form of this family, Rosaceae and Rhamnaceae (the families are phylogenetically closely related and placed in the order Rosales [Cronquist 1968]), may be ancestral for *Cleonidius* as a whole. Even more parsimonious, however, is the hypothesis that association with Compositae is ancestral for the genus. Association with this family is equally widespread and abundant in the genus, but unlike Leguminosae, is found particularly in plesiotypic species in the *C. poricollis*, *C. americanus*, and *C. lobigerinus* species groups, not in the more derived species, as is the association with Leguminosae. Association with Compositae is also known in numerous other Cleoninae (e.g. *Apleurus*, *Cleonis*, *Cyphocleonus*, some *Lixus*). Accordingly, preference must herein be given to a plesiotypic association with Compositae, but final resolution of these competing hypotheses rests largely on determination of host plant relationships in those *Cleonidius* species in which they are

as yet unknown, particularly, those of a primitive phylogenetic position and especially those of the Palearctic species component.

Further supportive of the more recent trans-Beringian vicariance of savanna or grassland are 1, non-relictual geographic or structural status of any *Cleonidius*; 2, lack of extensive structural divergence between species perhaps indicative of less available time for evolutionary change to have occurred; and 3, likelihood that *C. vibex* and relatives are the sister group to Nearctic *Cleonidius* (see Phylogeny section).

Axelrod's (1975) Madrean-Tethyan hypothesis has not been given much support from other studies. Floras of the eastern Gulf Coast and southeastern North America do not show evidence of arid-adapted taxa (Graham 1965). The hypothesis also does not receive any clear support from the biogeographic analyses presented herein. Clearly, I believe we must accept an ancestral association with grassland or savanna, as in *Apleurus*. Diversification in *Cleonidius*, however, has apparently been much more complex than in *Apleurus* as evidenced by: lack of a clear correlation between phylogeny and habitat association (Fig. 240); numerous different and variable habitat associations of the species (Fig. 240); different host plant associations (Fig. 241); and present relatively extensive species sympatry and altitudinal range overlap (Fig. 233).

Within species groups, caution must be exercised because degree of confidence in various of the hypothesized relationships is low (Fig. 237). Within the *C. poricollis* group, monophyly of *C. subcylindricus*-*C. texanus*-*C. longinasus* is strongly supported by three primary characters. Relationships of this lineage and the remaining three species in the species group are however much less clear and are left as an unresolved trichotomy (Fig. 237). Assuming, as noted, that the *C. poricollis* species group evolved in California (Fig. 246), it is likely that, as conditions became more arid, there was an entry into semi-desert habitat in the *C. eustictorrhinus*-*C. poricollis* lineage (Fig. 247). The hypothesized ancestral association with Compositae was retained in this lineage and *C. modestus*, and the ancestor of the *C. subcylindricus*-*C. texanus*-*C. longinasus* lineage likely shifted onto Leguminosae. *C. eustictorrhinus* and *C. poricollis* may then have diverged in part as a result of an east-west vicariance between California and the Great Basin Region with subsequent dispersal of *C. poricollis* back into California (Fig. 248). This event likely also involved, or may have even been exclusively due to, a host plant shift involving Compositae and the phylogenetically unrelated (Cronquist 1968) Cruciferae. Evidence for this is as follows. The sister-species *C. americanus* and *C. frontalis* (of the *C. americanus* group) have similar geographic distributions to *C. eustictorrhinus* and *C. poricollis*; *C. americanus* is

associated with Compositae in California; host plant associations of *C. frontalis* are not known. In *C. eustictorrhinus* and *C. porlicollis* the opposite is true; the widespread *C. porlicollis* is found on Cruciferae whereas plant associations of the California *C. eustictorrhinus* are not known. Furthermore, in the *C. lobigerinus* group, both *C. lobigerinus* and *C. calandroides* are associated with Cruciferae, but *C. lobigerinus* is also markedly associated with Compositae. Patterns of Cruciferae-Compositae association between closely related taxa therefore (and, as noted, Leguminosae-Rosaceae/Rhamnaceae associations of a similar nature), appear to repeat within *Cleonidius* (Fig. 241) and may thus represent more general patterns of host plant relationships. In view of this, association with Compositae can be hypothesized for *C. eustictorrhinus*, and association with Cruciferae for *C. frontalis*. If *C. modestus* ultimately proves sister-species of the *C. eustictorrhinus*-*C. porlicollis* lineage then evidence for *C. eustictorrhinus* being associated with Compositae is more marked, because then only one shift from Compositae in the evolution of the three species is required. Within *C. modestus* there has been a continued adaptation to more markedly arid lands (Fig. 240), perhaps interrelated with association with Compositae (note relatively high level of association with arid habitats in *Apleurus*, all of which are on Compositae; association of *C. lobigerinus*, the only other *Cleonidius* that is associated with desert habitats, is also



with Compositae).

In the *C. subcylindricus*-*C. texanus*-*C. longinasus* lineage there appears to be independent entry into xerophytic and mesophytic evergreen woodlands in the central-southern Rocky Mountain region and eastern North America by *C. texanus* and *C. subcylindricus* respectively as evidenced by the pattern of host plant associations. An initial vicariance in this lineage between *C. subcylindricus* in eastern North America and the *C. texanus*-*C. longinasus* lineage in western North America, both on Leguminosae, was likely followed by a host plant shift from Leguminosae to Rosaceae and Rhamnaceae and concurrent shift to xerophytic woodland habitat by *C. texanus* and to mesophytic woodland habitat by *C. subcylindricus* (Fig. 249).

Within the *C. americanus* group, most of the species are primarily associated with grasslands; *C. americanus* and *C. notolomus* are associated exclusively with Pacific semi-desert and desert-grassland transition respectively (Fig. 240). Vicariance patterns are clearly evident in the present distributions of species in this group. A California-Great Basin or southern Great Plains vicariance can be proposed to account for the origin of the *C. americanus*-*C. frontalis* lineage and the ancestor of the remainder of the *C. americanus* group (Fig. 250). Within the former lineage, a California-inland vicariance again very likely explains the evolution of *C. americanus* and *C. frontalis* (Fig. 251). This was likely concomitant with a

shift to more arid habitats, especially by *C. americanus*.

Origins of *C. canescens* and of the *C. notolomus*-*C. collaris* and *C. puberulus*-*C. infrequens* lineages are not clear. However, a north-south vicariance of both *C. collaris* and *C. notolomus* (Fig. 253), and of *C. puberulus* and *C. infrequens* (Fig. 252), accounts for the present distribution of these species. Of the four, only *C. notolomus* has changed from a grassland association to desert-grassland transitional habitat (Fig. 240). Few structural differences between these sister-species, and the allopatric distributions favour a relatively recent occurrence of these speciation events. Association with Compositae is hypothesized as plesiotypic for the species group.

Origin of the *C. lobigerinus* group is unclear principally because of the widespread distribution and eurytopic nature of *C. lobigerinus* (Fig. 228). It was undoubtedly concurrent with the hypothesized initial radiation of species group lineages and while an east coast isolation of *C. calandroides* likely accounts for the origin of that species, it is not clear where the ancestor of *C. lobigerinus* and *C. calandroides* evolved. Association with Compositae is proposed as ancestral for this species group as well.

In the *C. boucardi* group, there was likely a Mexican highlands isolation of *C. boucardi* in mesquite-grassland and a southern Rocky Mountain isolation of the *C. trivittatus*-*C. quadrilineatus*-*C. placidus* lineage in grassland habitat, an

event brought about by increasing aridity and the formation of intermediate desert-grassland transitional or semi-desert habitats (Fig. 254). In the *C. trivittatus*-*C. quadrilineatus*-*C. placidus* lineage, east-west vicariance of *C. trivittatus* and *C. quadrilineatus*-*C. placidus* in grassland and semi-desert habitats respectively, appears to account for the origin of *C. trivittatus* (Fig. 255). Subsequent entry into xerophytic evergreen woodland habitat by the semi-desert adapted ancestor, which, as in the *C. poricollis* group was associated with Leguminosae, and a host plant shift to Rosaceae and Rhamnaceae, resulted in *C. quadrilineatus* in the eastern xerophytic woodlands and *C. placidus* in the semi-desert habitat of California (Fig. 256). Present occurrence of *C. quadrilineatus* in California is likely a recent dispersal event. Ancestral host plant relationships of this group are also likely with Compositae.

Summarizing, both *Apleurus* and *Cleonidius* are hypothesized to have entered North America in savanna or grassland habitats (or their precursors) across Beringia during late Eocene time. In both genera there appears to be an initial association with Compositae and vicariance of lineages in grassland or mesquite-grassland between California, central Mexico, and the southern Rocky Mountains. Further speciation events in *Apleurus* appear due primarily to shifts to atypical habitat types brought about by increased aridity during the later Tertiary; there is little within habitat-type allopatric speciation. host plant

shifts are not a factor for all *Apleurus* species remain associated with Compositae. Host plant shifts, however, primarily between Leguminosae-Rosaceae and Compositae-Cruciferae, and, to a lesser degree, within-habitat-type allopatric speciation, appear to predominate as the causes of further speciation in *Cleonidius*, most species of which remain associated with plesiotypic less-arid habitat types such as grasslands and desert-grasslands.

#### 5.4 Evolutionary trends and adaptations

##### 5.4.1 Adaptive trends to arid conditions

Numerous apparent adaptations to arid habitats are known in Coleoptera, including Curculionidae (Crowson 1981). Some of these are exhibited in Cleoninae.

1. Wing length and flightlessness.-- Adults of various species of North American Cleoninae are apterous or brachypterous to varying degrees and in varying frequency. Some, such as *Stephanocleonus confusus* or *Apleurus aztecus*, have the elytra more or less solidly fused together along the median suture. The cuticle can also be thick and hard in many of these same individuals. Flightlessness appears more frequently in traditional Cleonini than in traditional Lixini.

In *Apleurus* there is no clear phylogenetic trend in this character although adults of the closely related *A. angularis*, *A. jacobinus* and *A. porosus* are all more or less apterous. Lack of a trend in *Apleurus* is perhaps surprising in view of the correlation of phylogenetic position with occupation of increasingly arid habitats (Fig. 238). Similarly in *Cleonidius*, there is no clear phylogenetic trend in this character. Adults of species of the *C. americanus* group are predominantly brachypterous but they are mostly associated with grassland habitats and not more arid habitats. Undoubtedly, degree of aridity is not the sole factor influencing the frequency of brachyptery and although the aforementioned adaptations may serve to prevent desiccation in arid environments, the spatial distribution of the habitat-type or of host plants, likely also influences the tendency towards or away from brachyptery. Compromise between these differing selective forces likely results in the wing-length polymorphisms noted in some species and the lack of a clear trend.

Species in the southwestern United States with populations on both sides of the Continental Divide, in which individuals are flightless, are structurally more distinct in the north, where the divide is high in elevation, as compared to the south, where the divide is low in elevation and where gene flow appears to occur. Species exhibiting similar distributions but which are composed of macropterous individuals do not show such patterns of

variation.

2. Diel activity.-- There is no evidence to indicate distinct peaks of diel activity in North American Cleoninae which might indicate avoidance of diurnal stressful conditions in an arid habitat. Such diel activity is known in other arid land Curculionidae (C.W. O'Brien, E.L. Sleeper, pers. comms.).

3. Burrowing adaptations.-- Modifications of legs for burrowing are not known in North American Cleoninae. This is not surprising in view of 2, above. There are, however, adults of some species that possess long erect surface vestiture which may prove correlated with life on sand dunes and which may facilitate movement through sandy substrates. Such long vestiture is known in various obligate sand dune Scarabaeidae and other Curculionidae.

4. Ventral tarsal pilose vestiture.-- Within *Apleurus* (Figs. 234-235), and to a lesser extent also in *Cleonidius* (Figs. 236-237), there is a clear trend towards reduction in ventral tarsal pilose vestiture. This reduction has been reasonably suggested to be due to living predominantly on the ground rather than on plants (Kissinger 1964; Crowson 1981) and is widespread in many arid-adapted Curculionidae. In *Apleurus* the trend is as expected, because those more derived species in more arid environments, where plants are more widely spaced, and where it is less desirable to spend time exposed on the plant, lack ventral tarsal pilose vestiture or have pilose pads that are markedly reduced. In

*Cleonidius* however, this is not clearly so, for although more arid habitat species (e.g. *C. poricollis*) possess reduced pilose pads, it is the members of the *C. americanus* group which exhibit the greatest frequency of lack of ventral tarsal pilose vestiture. This is perhaps surprising, for the species in this group are associated largely with plesiotypic grassland habitats; predicted adaptive correlates are thus not always evident in *Cleonidius*.

#### 5.4.2 Other evolutionary trends

5. Host plant associations.-- No trends in host plant relationships are evident in *Apleurus*, because all species appear associated with Compositae (Fig. 239). In *Cleonidius* however, two repeated patterns warrant mention (Fig. 241). Leguminosae-Rosaceae/Rhamnaceae associations are found in the sister-species *C. longinasus*-*C. texanus* and *C. placidus*-*C. quadrilineatus*, and Compositae-Cruciferae associations are found in *C. lobigerinus*-*C. calandroides* and are hypothesized in *C. americanus*-*C. frontalis* and *C. eustictorrhifus*-*C. poricollis*. It is unlikely that the Compositae-Cruciferae association is of a phylogenetic basis, but rather reflects the co-occurrence (perhaps recent) of these two plant taxa in the same, more arid habitats. The Leguminosae-Rosaceae/Rhamnaceae association, however, is likely phylogenetically based for the taxa involved are closely related and the host plant shift is concomitant

with a shift in habitat association.

Groups in which sympatry is low, altitude range overlap low, and habitat association overlap low, tend to have species which differ little in host plant relationships. Increasing levels of these factors appear to be correlated with more varied host associations. Host plant associations, largely unknown in the *C. americanus* group, can thus be predicted to vary little in terms of their taxonomic range and thus largely Compositae or Cruciferae.

There is no trend from generalist to specialist feeding in either *Apleurus* or *Cleonidius*.

6. Habitat associations.-- As noted in detail in the biogeography section, there is a clear trend in *Apleurus* for phylogenetically more derived species to be associated with increasingly arid habitats (Fig. 238) and lower elevations. This is not so in *Cleonidius* wherein most species are associated with grassland or grassland-desert habitats (Fig. 240).

7. Size.-- Within *Apleurus* there is a tendency towards increased size (Figs. 234-235); such is not clearly evident in *Cleonidius*. Increased size in an arid environment may serve to aid in prevention of water loss for surface area to volume ratios would be lower in larger individuals which are, notably, found in more arid environments. In *Cleonidius* however, the largest species are those in the *C. boucardi* group. These are not found in notably arid habitats but rather grasslands, semi-desert and xerophytic evergreen



woodlands.

7. Genitalia.-- Form of male genitalia including the internal sac is very conservative in species of both *Apleurus* (Figs. 99-104) and *Cleonidius* (Figs. 177-196) and form of female genitalia is very conservative in *Cleonidius* (Figs. 157-176). Such is not so in females of *Apleurus* wherein there is a clear trend towards reduction in size of the stylus on gonocoxite II and an increasingly subapical position of attachment of the same (Figs. 89-96). This trend is likely related to the oviposition habits of the species but such are not known in detail for any North American species of Cleoninae.

Form of the aedeagus (Figs. 31-36) and of the internal sac (Figs. 43-48) differs markedly between species of *Stephanocleonus*, in contrast to the lack of differences in *Cleonidius* and *Apleurus*. Reasons for this are unknown.

8. Elytral scale patterns.-- Elytral scale patterns are generally uniform within and between *Apleurus* species except for *A. angularis* (Figs. 72-74) and *A. albovestitus* (Figs. 75-79) wherein various forms are recognizable. Correlation of particular scale patterns with geographic distribution suggests a local adaptive significance, one I believe associated with crypsis on the ground substrate or in debris under host plants. A thermoregulatory role however, may apply to those *A. albovestitus* in the lower Colorado River drainage. In *Apleurus* there are no species which possess a distinctly vittate scale pattern; most are more or less

mottled in general appearance.

In *Cleonidius* elytral patterns vary between species but not to a significant extent within a species (with the exception of *C. canescens*). Adaptive significance of the vittate pattern (e.g. Fig. 131) found in most *Cleonidius* is likely one of concealment on host plants or in debris under the host plant. Crypsis on the ground substrate is not likely, except perhaps in the *C. americanus* group, for otherwise, most species have moderately developed ventral tarsal pilose vestiture indicating life predominantly on plants. Only in the *C. americanus* group is there predominance of species in which individuals have reduced ventral tarsal pilose vestiture. Also in this group are found the only species with mottled elytral scale patterns (e.g. Figs. 124-125). These patterns are similar to those of *Apleurus* species, and likely serve as an adaptation to life on the ground substrate.

Vittate (and margined [Figs. 118-123]) patterns predominate in species of *Cleonidius* with extensive ventral tarsal pilosity; non-vittate patterns predominate in species with reduced tarsal vestiture. Wide ranging species with reduced ventral pilose tarsal vestiture in *Apleurus* exhibit geographic variation in scale pattern, whereas restricted species or those wide-ranging species with extensive ventral pilose vestiture do not show variation in elytral scale pattern. In *Cleonidius* nearly all species exhibit little or no variation in elytral scale pattern (except for *C.*

*canescens* of the *C. americanus* group, which like *A. angularis*, has more or less vittate and non-vittate forms occurring on the west and east sides of the Continental Divide). Within the *C. americanus* group, which contains mostly allopatric species with reduced ventral vestiture, there is the most extensive within-species-group variation in scale pattern. Other species groups do not show within group variation in scale pattern. It thus appears that there is an inverse correlation in both *Apleurus* and *Cleonidius* between degree of variation in elytral pattern and extent of ventral tarsal pilose vestiture, but only seen in species or species groups distributed over large geographic areas.

### 5.5 Patterns and predictions

From results of this study of Cleoninae, some tentative generalizations can be made as regards arid adapted faunas in North America.

1. Preference must go to a post-late Eocene vicariance of savanna or grassland habitat, or precursory habitat, across Beringia as the original mode of isolation of Nearctic arid land taxa which possess Palearctic relatives also inhabiting arid lands (Figs. 242, 246). There is no evidence for direct Madrean-Tethyan connections as proposed by Axelrod (1975).

2. The most plesiotypic species or lineages in the Nearctic region will be found on Compositae in various

combinations of the mesquite-grasslands of the highlands of central Mexico, the southern Rocky Mountain grasslands, and in one of various habitats in California which may be relictual or predominantly grassland in nature (Figs. 242, 246).

3. Arid-adapted species in eastern North America are more recent in origin, not relictual in nature, and are related to a species or a lineage of few species in western North America.

4. California is the geographical area recognizable as possessing the most arid land endemics. All have their closest relatives outside of California.

5. Evolution in arid-adapted taxa may involve a trend towards occupation of increasingly arid habit by successively more derived species (Fig. 238). This is accompanied by few or no host plant shifts (Fig. 239), and may be limited to non-speciose groups. Sympatry, altitudinal range overlap, and habitat association overlap is low in these taxa.

6. Alternatively to 5 above, host plant shifts (Fig. 241) permit sympatry, altitudinal range overlap and habitat association overlap and are perhaps associated only with more speciose groups. No clear correlation between phylogenetic position of the species and degree of aridity of habitat occupied is evident (Fig. 240).

7. Species inhabiting North American desert habitats are of recent origin (but not necessarily of derived

phylogenetic position), and are related to taxa associated with precursory North American non-desert habitats although not in a generally predictable manner. That regional deserts are composed of taxa from non-desert arid habitats and are of recent origin is supported. No regional desert endemics are known in *Apleurus* or *Cleonidius*.

8. Contrary to statements of other authors (Axelrod 1985), grassland endemics are known, are primitive or derived in their phylogenetic position, and should thus be expected in other arid land adapted taxa.

9. Compositae-Cruciferae and Leguminosae-Rosaceae/Rhamnaceae host plant shifts may represent general patterns to be found in other phytophagous taxa. Shifts from Leguminosae to Rosaceae and Rhamnaceae may prove generally associated with entry into western upland xerophytic evergreen woodland habitat by some phytophagous taxa. These shifts likely have a phylogenetic basis as indicated by the fact that the families are placed in the order Rosales (Cronquist 1968).

In contrast, Compositae-Cruciferae host plant shifts are likely due to occurrence of both the two plant and two weevil taxa in the same habitat types. No phylogenetic basis can be inferred because the two families are not phylogenetically closely related but are placed in separate, unrelated subclasses (Cronquist 1968).

## 6. Conclusions and future research

Scientific studies should not only attempt to answer specific questions but should also strive to pose questions which might prove the basis for further inquiry. More than anything else, I believe this study draws attention to the need for examination of world faunas or at the least, regional faunas but placed in a world perspective. Traditionally, North American Cleonini have all been treated as *Cleonis*, but as now should be clearly evident, the relationships of many taxa in North America are actually quite phylogenetically distant.

Relationships of *Apleurus* are largely resolved. Unfortunately, this cannot be said with the same confidence for *Cleonidius*. World study of the speciose and structurally diverse genus *Lixus* should help resolve the still problematical relationships and even species constituency of *Cleonidius*. What other Old World *Lixus* may require placement in *Cleonidius* and subsequent study of large numbers of individuals of these species and determination of what their relationships are to North American *Cleonidius*, should help to resolve the phylogenetic placement of the Old World *Cleonidius*, further resolve the interrelationships of North American species, and answer some still remaining questions of biogeographic importance. This will be a large task, for within *Lixus* are placed a great number of species. More complete knowledge of structural diversity of *Lixus* species and the phylogenetic relationships of component

species-groups or subgenera should ultimately lead to a clearer understanding of suprageneric relationships in Cleoninae and a well substantiated tribal classification.

Not to be forgotten are the interrelationships among genera of traditional Cleonini. A start at a study of these relationships leading, I hope, to a generic reclassification, has been made as part of this present thesis. Results are not presented in detail here; only characters of relevance to our understanding of the North American fauna have been included. Study of representatives of generic-group names not yet examined is required, as is study of more species in more speciose genera to assure proper placement and uniformity of features that appear to prove phylogenetically valuable at the generic level.

Even within the North American fauna itself, work remains to be done. For *Apleurus* and *Cleonidius*, I feel confident that no new species will be discovered; for *Stephanocleonus*, largely because the genus is northern and Holarctic, I suspect that one or more new species remain to be discovered, perhaps in the Yukon Territory and Alaska. Whether any of the North American species are Holarctic in distribution (and perhaps conspecific with nominal Old World species) also needs to be determined for it has not been possible to examine types or representatives of even a small number of the many Palearctic species. Only when this has been done can phylogenetic relationships of the species in this genus be reconstructed.

For all Cleoninae, more host plant information needs to be compiled. Hosts, even plant associations, are not at all known for *Stephanocleonus* species; the relationship with Compositae needs to be further substantiated in *Apleurus* species; and, various of the species of *Cleonidius* do not have hosts or even plant associations established, and those that are established need variable degrees of further supportive data. With host plant shifts appearing to play an integral role in *Cleonidius*, more such information can only increase understanding of the evolution of species in that genus and determine whether predictions about specific host plant relationships made herein, and whether predictions that these patterns are repeated and general are borne out.

Immature stages of Cleoninae are largely unknown and unstudied. Collection and study of eggs, larva and pupae, particularly in North America, may provide characters of phylogenetic use in substantiating relationships as based on adult characters, or, in resolving problematical relationships or those in which confidence is low.



## 7. Literature cited

- Anderson, D.M. 1956. Notes on *Cleonus piger* (Scop.) in the United States (Coleoptera; Curculionidae). *The Coleopterists' Bulletin* 10:81-85.
- Anderson, E.M. 1914. Insects collected in the Okanagan Valley, 1913. *Proceedings of the Entomological of British Columbia* 4:54-58.
- Anderson, R.S. 1984a. Taxonomic studies of *Cleonis piger* (Scop.) and *Cyphocleonus achates* (Fahrs.) (Coleoptera; Curculionidae; Cleoninae) from *Centaurea* as biological agents for control of *Centaurea diffusa* (Compositae) in Canada. Unpublished Report for Agriculture Canada, Regina, Saskatchewan. 49 pp + 8 figures.
- Anderson, R.S. 1984b. *Connatichela artemisiae*, a new genus and species of weevil from the Yukon Territory (Coleoptera: Curculionidae: Leptopiinae): Taxonomy; paleontology, and biogeography. *The Canadian Entomologist* 116:1571-1580.
- Anderson, R.S. in press. Cleoninae Schoenherr, 1826 (Insecta; Coleoptera; Curculionidae). Proposed conservation. *Bulletin of the International Commission of Zoological Nomenclature*, in press.
- Anderson, R.S. and S.B. Peck. 1986. Geographic patterns of colour variation in North American *Nicrophorus* burying beetles (Coleoptera; Silphidae). *Journal of Natural History* 20:283-297.
- Andrews, F.G., A.R. Hardy, and D. Giuliani. 1979. The coleopterous fauna of selected California sand dunes. California Department of Food and Agriculture, Insect Taxonomy Laboratory. i + 142 pp. Sacramento, California.
- Arnaud, P.H. 1978. A host-parasite catalogue of North American Tachinidae (Diptera). *United States Department of Agriculture, Miscellaneous Publications* 1319. 860 pp.
- Arnett, R.H., Jr. 1960-1962. The beetles of the United States (a manual for identification). Catholic University Press. xi + 1112 pp. Washington D.C..
- Arnett, R.H., Jr., N.M. Downie, and H.E. Jaques. 1980. How to know the beetles (2nd edition). The Pictured Key Nature Series. Wm. C. Brown Co.. viii + 416 pp. Dubuque, Iowa.

- Aslam, N.A. 1961. An assessment of some internal characters in the higher classification of the Curculionidae S.I. (Coleoptera). *The Transactions of the Royal Entomological Society of London* 113:417-489.
- Aslam, N.A. 1963. On the genera of Indo-Pakistan Cleoninae and Hylobiinae (Coleoptera: Curculionidae). *Bulletin of the British Museum (Natural History), Entomology*. 13:47-66.
- Axelrod, D.I. 1950. Evolution of desert vegetation in western North America. pp. 215-306. In: Axelrod, D.I. Contributions to Paleontology. Studies in Late Tertiary Paleobotany. Carnegie Institute of Washington Publication 590, Washington D.C. 1-323 pp.
- Axelrod, D.I. 1975. Evolution and Biogeography of Madrean-Tethyan Sclerophyll Vegetation. *Annals of the Missouri Botanical Garden* 62:280-334.
- Axelrod, D.I. 1979. Age and Origin of Sonoran Desert Vegetation. *Occasional Papers of the California Academy of Sciences* 132:i-vi + 74 pp.
- Axelrod, D.I. 1985. Rise of the Grassland Biome, Central North America. *The Botanical Review* 51:163-201.
- Axelrod, D.I. and P.H. Raven. 1985. Origins of the Cordilleran flora. *Journal of Biogeography* 12:21-47.
- Ball, I.R. 1975. Nature and Formulation of Biogeographical Hypotheses. *Systematic Zoology* 24:407-430.
- Bargagli, P. 1884. Rassegna biologica di Rincofori Europei. *Bollettino della Società Entomologica Italiana* 16:149-258.
- Batra, S.W.T., J.R. Coulson, P.H. Dunn, and P.E. Boldt. 1981. Insects and fungi associated with *Carduus* thistles (Compositae). United States Department of Agriculture. Technical Bulletin 1616. 100 pp.
- Bedel, L. 1882-1888. Faune des Coléptères du bassin de la Seine. Rhynchophora. *Annales de la Société Entomologique de France* 6:1-442.
- Berg, F.G.C. 1884. (Communication). *Bulletin de la Société Entomologique de France* 1884:XCVIII-C.
- Billberg, G.J. 1820. Enumeratio insectorum in museo Gust. Joh. Billberg. Stockholm. 138 pp.
- Blackwelder, R.E. 1939. Fourth supplement 1933 to 1938 (inclusive) to the Leng Catalogue of Coleoptera of

- America, north of Mexico. Mount Vernon, New York. 146 pp.
- Blackwelder, R.E. 1947. Checklist of the Coleopterous insects of Mexico, Central America, the West Indies and South America. Part 5. *Bulletin. United States National Museum* (185)I-IV, 765-925.
- Blanchard, C.E. 1851. Fauna Chilena. Insectos. Coleópteros. pp. 285-563. *In: Gay, C. Historia física y política de Chile. Volume 5. Zoología. Paris.*
- Blatchley, W.S. 1914. Notes on the winter and early spring Coleoptera of Florida with descriptions of new species. *The Canadian Entomologist* 46:247-251.
- Blatchley, W.S. 1930. Blatchleyana. A list of the published writings of W.S. Blatchley, A.B., A.M., LL.D. of Indianapolis, Indiana and Dunedin, Florida. Together with a chronology of his life: the fixation of types of new genera and species described by him, etc. etc. 77 pp. Indianapolis, Indiana.
- Blatchley, W.S. and C.W. Leng. 1916. Rhynchophora or weevils of North Eastern America. The Nature Publishing Company. Indianapolis, Indiana. 682 pp.
- Bleasdel, G.G. 1937. The Rhynchophora of Iowa. *Iowa State College Journal of Science* 11:405-445.
- Boheman, C.H. 1836. *In: Schoenherr, C.J. Genera et species Curculionidum, cum synonymia hujus familiae. Species novae aut hactenus minus cognitae, descriptionibus a Dom. Leonardo Gyllenhal, C.H. Boheman, et entomologis aliis illustratae. Volume 3, Part 1, 505 pp. Paris.*
- Böving, A.G. 1927. Immature stages of *Eumycterus* (?) *saccharides* Barber, with comments on the classification of the tribe Barini (Coleoptera: Curculionidae). *Proceedings of the Entomological Society of Washington* 29:151-159.
- Bradley, J.C. 1930. A manual of the genera of beetles of America, north of Mexico. x + 360 pp. Ithaca, New York. Mimeographed.
- Brèthes, J. 1910. Coleópteros argentinos y bolivianos. *Anales de la Sociedad Científica Argentina* 69:205-227.
- Brimley, C.S. 1938. The insects of North Carolina. Being a list of the insects of North Carolina and their close relatives. North Carolina Department of Agriculture. 560 pp.

- Brown, D.E. (ed.). 1982. Biotic Communities of the American Southwest-United States and Mexico. *Desert Plants* 4:1-342.
- Brown, J.H. and A.C. Gibson. 1983. Biogeography. The C.V. Mobby Company, St. Louis. i-xii + 643 pp.
- Brown, W.J. 1940. Notes on the American distribution of some species of Coleoptera common to the European and North American continents. *The Canadian Entomologist* 72:65-78.
- Bruhn, A.F. 1947. The external male genitalia of some Rhynchophora. *The Great Basin Naturalist* 8:1-35.
- Buchner, P. 1933. Studien an Intracellularen Symbionten. VII. Die Symbiontischen Einrichtungen der Rüsselkäfer. *Zeitschrift für Morphologie und Ökologie der Tiere*. 26:709-777.
- Buchner, P. 1953. Endosymbiosis of Animals with Plant Microorganisms. Revised English Version. Interscience Publishers, New York.
- Burke, H.R. and D.M. Anderson. 1976. Systematics of larvae and pupae of American Curculionoidea: Status report, historical review and bibliography. *The Southwestern Entomologist* 1:56-73.
- Capiomont, G. and M.C.-E. Leprieur. 1873. Monographie des Lixus. Deuxième partie. *Annales de la Société Entomologique de France* 5:41-64.
- Casey, T.L. 1891. Coleopterological Notices. III. *Annals of the New York Academy of Sciences* 6:9-214.
- Casey, T.L. 1904. On some new Coleoptera, including five new genera. *The Canadian Entomologist* 36:312-324.
- Champion, G.C. 1902-1906. Biologia Centrali-Americana. Insecta. Coleoptera. Rhynchophora. Curculionidae. Curculionidae (part). Volume 4, Part 4, viii + 750 pp.
- Chevrolat, L.A.A. 1860. Description de Coléoptères nouveaux d'Algérie. *Revue et Magasin de Zoologie, Series 2* 12:75-82.
- Chevrolat, L.A.A. 1873. Mémoire sur les Cléonides. *Mémoires de la Société Royale des Sciences de Liège, Series 2* 5(6):viii-118.
- Chittenden, F.H. 1911. Some miscellaneous results of the work of the Bureau of Entomology - IX. Insects injurious to the loco weeds. *United States Department*

- of Agriculture. Bureau of Entomology. Bulletin. 64:33-42.
- Clayton, W.D. 1981. Evolution and Distribution of Grasses. *Annals of the Missouri Botanical Garden*. 68:5-14.
- Cody, M.L. 1975. Towards a Theory of Continental Species Diversities. pp. 214-257. In: Cody, M.L. and J.M. Diamond, Eds. *Ecology and Evolution of Communities*. Bellknapp Press of Harvard University, Cambridge. i-xiv + 545 pp.
- Cody, M.L. and J.M. Diamond, Eds. 1975. *Ecology and Evolution of Communities*. Bellknapp Press of Harvard University, Cambridge. i-xiv + 545 pp.
- Coope, G.R. 1979. Late Cenozoic Fossil Coleoptera: Evolution, Biogeography, and Ecology. *Annual Review of Ecology and Systematics*. 10:247-267.
- Correll, D.S. and M.C. Johnston, 1970. A manual of the vascular plants of Texas. Texas Research Foundation. Renner, Texas. xvi + 1881 pp.
- Croizat, L. 1982. Vicariance/Vicariism, Panbiogeography, "Vicariance Biogeography," etc. : A Clarification. *Systematic Zoology* 31:291-304.
- Cronquist, A. 1968. The evolution and classification of flowering plants. Thomas Nelson and Sons, London. xii + 396 pp.
- Cronquist, A. 1978. The Biota of the Intermountain Region in Geohistorical Context. *Great Basin Naturalist Memoirs* 2:3-15.
- Crosby, C.R. and W.E. Blauvelt. 1930. A European beetle found in New York (Coleop.: Curculionidae). *Entomological News* 41:164.
- Crowson, R.A. 1955. The natural classification of the families of Coleoptera. Nathaniel Lloyd, London. 187 pp.
- Crowson, R.A. 1981. The biology of the Coleoptera. Academic Press. London. xii + 802 pp.
- Csiki, E. 1934. Coleopterorum Catalogus, pars 134, Curculionidae: Subfam. Cleoninae, 152 pp.
- Danks, H.V. 1981. Arctic arthropods. A review of systematics and ecology with particular reference to the North American fauna. Entomological Society of Canada. viii + 608 pp. Ottawa, Ontario.

- Dejean, P.F.M.A. 1821. Catalogue de la collection de Coléoptères de M. le Baron Dejean. viii + 136 pp. Paris.
- Eldredge, N. 1979. Cladism and Common Sense. pp.165-198. In: Cracraft, J. and N. Eldredge, Eds. Phylogenetic Analysis and Paleontology. Columbia University Press, New York. 1-233 pp.
- Eldredge, N. and J. Cracraft. 1980. Phylogenetic Patterns and the Evolutionary Process. Columbia University Press, New York. i-ix + 1-349 pp.
- Ely, C.R. 1913. The food plant of *Cleonus calandroides* Rand. *Proceedings of the Entomological Society of Washington* 15:104-105.
- Endler, J.A. 1977. Geographic variation, speciation, and clines. Monographs in Population Biology 10. Princeton University Press, Princeton. i-ix + 1-246 pp.
- Endler, J.A. 1982. Problems in Distinguishing Historical from Ecological Factors in Biogeography. *American Zoologist* 22:441-452.
- Essig, E.O. 1958. Insects and mites of Western North America (revised edition). The MacMillan Co.. xiv + 1050 pp. New York, New York.
- Everts, J.E. 1903. Coleoptera Neerlandica. De Schildvleugelige Insecten van Nederland en het Aangrenzend Gebied. Tweede Deel Martinus Nijhoff. iv + 798 pp. + 8 pl. 'Sgravenhage.
- Fabricius, J.C. 1792. Entomologia systematica emendata et aucta. Secundum classes, ordines, genera, species adjectis synonymis, locis, observationibus, descriptionibus. Volume 1. xx + 538 pp. Hafniae.
- Fabricius, J.C. 1801. Systema eleutheratorum secundum ordines, genera, species: adiectis synonymis, locis, observationibus, descriptionibus. Volume 2. 687 pp. Kiliae.
- Fähræus, O.I. von. 1842. In: Schoenherr, C.J. Genera et species Curculionidum, cum synonymia hujus familiae. Species novae aut hactenus minus cognitae, descriptionibus a Dom. Leonardo Gyllenhal, C.H. Boheman, et entomologis aliis illustratae. Volume 6, Part 2, 495 pp. Paris.
- Fall, H.C. 1897. A list of the Coleoptera of the Southern California islands, with notes and descriptions of new

- species. *The Canadian Entomologist* 29:233-244.
- Fall, H.C. 1901. List of the Coleoptera of southern California, with notes on habits and distribution and description of new species. *Occasional Papers of the California Academy of Sciences* 8:1-282.
- Fall, H.C. 1913. A brief review of our species of *Magdalis*, with notes and descriptions of other North American Rhynchophora. *Transactions of the American Entomological Society* 39:23-72.
- Fall, H.C. and T.D.A. Cockerell. 1907. The Coleoptera of New Mexico. *Transactions of the American Entomological Society* 33:145-272.
- Faust, J. 1890. Notizen über Rüsselkäfer. *Stettiner Entomologische Zeitung* 51:109-116.
- Faust, J. 1904. Revision der Gruppe Cléonides vrais. *Deutsche Entomologische Zeitschrift* 1904:177-284.
- Fletcher, J. 1906. The entomological record, 1905. 36th. *Annual Report of the Entomological Society of Ontario* 1905:90-105.
- Gaffney, E.S. 1979. An Introduction to the Logic of Phylogeny Reconstruction. pp. 79-111. In: Cracraft, J. and N. Eldredge, Eds. *Phylogenetic Analysis and Paleontology*. Columbia University Press, New York. 1-233 pp.
- Gebler, F.A. von. 1830. Bemerkungen über die Insecten Sibiriens, vorzüglich des Altai. *Ledebours Reise* 2(3):1-228.
- Gemminger, M. and E. von Harold. 1871. *Catalogus Coleopterorum hucusque descriptorum synonymicus et systematicus*. Volume 8. Curculionidae. pp. 2181-2668. Monachii.
- Gibson, A. 1914. The entomological record, 1913. 44th. *Annual Report of the Entomological Society of Ontario* 1913:116-139.
- Gmelin, J.F. 1790. *Caroli a Linné Systema Naturae per regna tria naturae, secundum classes, ordines, genera, species, cum characteribus, differentiis, synonymis, locis*. Editio decima tertia, aucta, reformata. Beer, Lipsiae. Volume 1, Part 4 (Coleoptera, Hemiptera). pp. 1517-2224.
- Goeze, J.A.E. 1777. *Entomologische Beiträge zu des Ritter Linné swelfth Ausgabe des Natursystems*. Volume 1. 736

pp. Leipzig.

- Goudge, T.A. 1961. *The Ascent of Life. A Philosophical Study of the Theory of Evolution.* University of Toronto Press, Toronto. 236 pp.
- Gould, S.J. 1985. *The Flamingo's Smile. Reflections in Natural History.* W.W. Norton and Company. 476 pp. New York.
- Graham, A. 1965. Origin and evolution of the biota of southeastern North America: evidence from the fossil plant record. *Evolution* 18:571-585.
- Gyllenhal, L. 1834. *In: Schoenherr, C.J. Genera et species Curculionidum, cum synonymia hujus familiae. Species novae aut hactenus minus cognitae, descriptionibus a Dom. Leonardo Gyllenhal, C.H. Boheman, et entomologis aliis illustratae.* Volume 2, Part 1, 326 pp., Part 2, 329-669 pp. Paris.
- Hardy, A.R. and F.G. Andrews. 1976. A final report to the office of endangered species in contract 14-16-0008-966. California Department of Food and Agriculture, Insect Taxonomy Laboratory. 53 pp. Sacramento, California.
- Hatch, M.H. 1971. *The beetles of the Pacific Northwest. Part V: Rhipicerioidea, Sternoxi, Phytophaga, Rhynchophora, and Lamellicornia.* University of Washington Publications in Biology 16. xiv + 662 pp.
- Hecht, M.K. and J.L. Edwards. 1976. The determination of parallel or monophyletic relationships; The proteid salamanders - a test case. *American Naturalist* 100:653-677.
- Hecht, M.K. and J.L. Edwards. 1977. The methodology of phylogenetic inference above the species level. pp. 3-51, *In: Hecht, M.K., P.C. Goody and B.M. Hecht, Eds., Major patterns in vertebrate evolution. Volume 14. NATO Advanced Study Institute Series A. Plenum Press, New York.* 908 pp.
- Hennig, W. 1966. *Phylogenetic systematics.* University of Illinois Press, Urbana. 263 pp.
- Henshaw, S. 1881-1882. Index to the Coleoptera described by J.L. LeConte, M.D. *Transactions of the American Entomological Society* 9:197-272.
- Henshaw, S. 1885. *List of the Coleoptera of America, north of Mexico.* American Entomological Society. 161 pp. Philadelphia, Pennsylvania.



- Herbst, J.F.W. 1795. Natursystem aller bekannten in- und ausländischen Insekten, als eine Fortsetzung der von Buffonschen Naturgeschichte. Der Käfer. VI. Theil, xxiv + 520 pp. Berlin.
- Hicks, S.D. 1947. Additional notes on Coleoptera taken in Essex County and southern Ontario. *The Canadian Entomologist* 79:117-119.
- Hicks, S.D. 1949. A note on the occurrence of *Cleonus plger*, a European weevil near Ottawa, Ontario. *The Coleopterists' Bulletin* 3:7.
- Horn, G.H. 1894. The Coleoptera of Baja California. *Proceedings of the California Academy of Sciences, Series 2* 4(1):302-449.
- Hughes, O.L., C.R. Harrington, J.A. Janssens, J.V. Matthews, Jr., R.E. Morlan, N.W. Rutter and C.E. Schweger. 1981. Upper Pleistocene stratigraphy, paleoecology, and archaeology of the northern Yukon interior, eastern Beringia 1. Bonnet Plume Basin. *Arctic* 34:329-365.
- Hull, D.L. 1974. Philosophy of Biological Science. Prentice-Hall Inc., Englewood Cliffs. i-xii + 1-148 pp.
- Kearney, T.H. and R.H. Peebles. 1960. Arizona Flora. University of California Press, Berkeley. i-viii + 1085 pp.
- Kingsolver, J.M. 1961. Evolutionary studies in the weevil genus *Lixus* (Coleoptera, Curculionidae). Unpublished Ph.D. Thesis. University of Illinois, Urbana. 108 pp.
- Kingsolver, J.M. 1972. A catalogue of the genus *Lixus* F. for America north of Mexico (Coleoptera: Curculionidae). *The Coleopterists' Bulletin* 26:151-154.
- Kirby, W. 1837. Part the fourth and last. The insects. In: Richardson, J. Fauna Boreali-Americana; or the zoology of the northern parts of British America: containing descriptions of the objects of natural history collected on the late Northern Land Expeditions, under command of Captain Sir John Franklin, R.N. xxxix + 325 pp.
- Kirkland, R.L. and R.D. Goeden. 1977. Descriptions of the immature stages of imported puncturevine weevils, *Microlarinus tareynii* and *M. lypriformis*. *Annals of the Entomological Society of America* 70:583-587.
- Kirkland, R.L. and R.D. Goeden. 1978a. Biology of *Microlarinus tareynii* (Col.: Curculionidae) on

- puncturevine in Southern California. *Annals of the Entomological Society of America* 71:13-18.
- Kirkland, R.L. and R.D. Goeden. 1978b. Biology of *Microlarinus lypriformis* (Col.: Curculionidae) on puncturevine in Southern California. f 2>Annals of the Entomological Society of America 71:65-69.
- Kissinger, D.G. 1964. Curculionidae of America, north of Mexico. A key to the genera. Taxonomic Publications. v + 143 pp. South Lancaster, Massachusetts.
- Kissinger, D.G. 1970. Curculionidae tribe Ophyrastini of North America (Coleoptera). Taxonomic Publications. vi + 238 pp. South Lancaster, Massachusetts.
- Knoch, A.W. 1781. Beiträge zur Insectengeschichte. Leipzig, Schwickert. Stück 1. 10, 98u 3p. 6 col. Taf.
- Krombein, K.V. 1979. Superfamily Sphecoidea. pp. 1573-1740. In: Krombein, K.V. et al. Catalog of Hymenoptera in America North of Mexico. Volume 2: Apocrita (Aculeata). Smithsonian Institution Press. xvi + 2209 pp. Washington D.C.
- Krombein, K.V. et al.. 1979. Catalog of Hymenoptera in America North of Mexico. Volumes 1, 2, 3. Smithsonian Institution Press. Washington D.C.
- Kumar, R., R.J. Lavigne, J.E. Lloyd and R.E. Pfadt. 1976. Insects of the Central Plains Experiment Range, Pawnee National Grassland. Agricultural Experiment Station, University of Wyoming, Science Monograph No. 32. iv + 74 pp.
- Kuschel, G. 1949. Los "Curculionidae" del extremo norte de Chile (Coleoptera, Curcul. ap.6°). *Acta Zoológica Lilloana* 8:5-54.
- Kuschel, G. 1950. I. Nuevos Curculionidae de Bolivia y Perú. II. Notas a algunas especies de Brèthes. *Revista del Museo de La Plata* 6(Zool. 38):69-116.
- Lacordaire, J.T. 1863. Histoire naturelle des insectes. Genera des Coléoptères ou exposé méthodique et critique de tous les genres proposés jusqu'ici dans cet ordre d'insectes. Volume 6. pp. 1-608, 615-637. Paris.
- La Ferla, A. 1939. Contributo alla conoscenza del Cleono del Carciofo (*Cleonus piger* Scop.). *Bollettino del Laboratorio di Entomologia della Facoltà agraria in Portici* 3:3-33.
- Latreille, P.A. 1810. Considérations générale sur l'ordre

- natural des animaux composant les classes des crustacés, des arachnides, et des insectes; avec un tableau méthodique de leurs genres, disposés en familles. Schoell, Paris. 444 pp.
- LeConte, J.L. 1850. General remarks upon the Coleoptera of Lake Superior. Article IV, pp. 201-242 + 8 pl. In: Agassiz, J.L.R. and J.E. Cabot. Lake Superior: Its physical character, vegetation and animals. 461 pp. Boston, Massachusetts.
- LeConte, J.L. 1858. Descriptions of new species of Coleoptera, chiefly collected by the United States and Mexican Boundary Commission, under Major W.H. Emory, U.S.A. *Proceedings of the Academy of Natural Sciences of Philadelphia*. 10:59-89.
- LeConte, J.L. 1859a. The Coleoptera of Kansas and eastern New Mexico. *Smithsonian Contributions to Knowledge* 2(6):i-vi + 58 pp.
- LeConte, J.L. 1859b. The complete writings of Thomas Say on the entomology of North America. Volume 1. xxiv + 412 pp.
- LeConte, J.L. 1876a. Tribe IV. Cleonini, pp. 144-160. In: LeConte, J.L. and G.H. Horn. The Rhynchophora of America, north of Mexico. *Proceedings of the American Philosophical Society* 15(96):i-xvi + 1-455.
- LeConte, J.L. 1876b. Appendix. pp. 412-442. In: LeConte, J.L. and G.H. Horn. The Rhynchophora of America, north of Mexico. *Proceedings of the American Philosophical Society* 15(96):i-xvi + 1-455.
- LeConte, J.L. 1878. The Coleoptera of the Alpine Regions of the Rocky Mountains. *Bulletin of the United States Geological and Geographical Survey of the Territories* 4(2) 1878:447-480.
- LeConte, J.L. and G.H. Horn. 1883. Classification of the Coleoptera of North America. Smithsonian Institution. 567 pp. Washington D.C.
- Leng, C.W. 1919. The Rhynchophora (except Ipidae). pp. 19E-21E In: Report of the Canadian Arctic Expedition. Volume 3: Insects, part E: Coleoptera. pp. 3E-27E.
- Leng, C.W. 1920. Catalog of the Coleoptera of America, north of Mexico. x + 470 pp. The Cosmos Press. Cambridge, Massachusetts.
- Leng, C.W. and A.J. Mutchler. 1933. Second and third supplements 1925 to 1932 (inclusive) to Catalogue of

- the Coleoptera of America, North of Mexico. 112 pp. The Cambridge Press. Cambridge, Massachusetts.
- Leonard, M.D. 1926. A list of the insects of New York with a list of the spiders and certain other allied groups. *Cornell University, Agricultural Experiment Station. Memoir # 101.* 1121 pp.
- Linnaeus, C. 1758. *Systema naturae, per regna tria naturae, secundum classes, ordines, genera, species cum characteribus, differentiis, synonymis, locis, ed. 10, reformata, Volume 1,* 823 pp. Holmiae.
- Linnaeus, C. 1767. *Systema naturae, per regna tria naturae, secundum classes, ordines, genera, species cum characteribus, differentiis, synonymis, locis, Vol. 1, pars 2, editio duodecima reformata,* pp. 533-1327. Holmiae.
- Livingston, B.E. and F. Shreve. 1921. The distribution of vegetation in the United States, as related to climatic conditions. xiv + 590 pp. Carnegie Institute of Washington, Washington D.C..
- Lohse, G.A. 1983. Unterfamilie: Cleoninae. pp. 7-29. In: Freude, H., K.W. Harde, and G.A. Lohse. *Die Käfer Mitteleuropas. Band 11.* Goecke and Evers, Krefeld. 342 pp.
- Lukjanovitsh, T. 1926. Bemerkungen über Systematik und geographische Verbreitung der Curculioniden (Coleoptera). *Entomologicheskoe Obozrenie* 20: 140-143.
- Maddison, W.P., M.J. Donoghue, and D.R. Maddison. 1984. Out-group analysis and parsimony. *Systematic Zoology* 33:83-103.
- Maddox, D.M. 1976. History of weevils on puncturevine in and near the United States. *Weed Science* 24:414-419.
- Mann, J.S. and R.A. Crowson. 1983. On the occurrence of mid-gut caeca, and organs of symbiont transmission, in leaf beetles. *The Coleopterists Bulletin* 37:1-15.
- Mannerheim, G.C.G. 1843. Beitrag zur Käfer-Fauna der Aleutischen Inseln der Insel Sitkha und Neu-Californiens. *Bulletin de la Société Impériale des Naturalistes de Moscou* 16(2):175-314.
- Marcovitch, S. 1923. A root weevil (*Stephanocleonus plumbeus* Lec.) reared from strawberry. *The Canadian Entomologist* 55:218.
- Marshall, G.A.K. 1932. Notes on the Hylobiinae (Col.,

- Curc.). *Annals and Magazine of Natural History, Series 10* 9:341-355.
- Matthews, J.V. Jr. 1974. Quaternary environments at Cape Deceit (Seward Peninsula, Alaska): Evolution of a tundra ecosystem. *Geological Society of America, Bulletin* 85:1353-1384.
- Matthews, J.V. Jr. 1975. Insects and plant macrofossils from two Quaternary exposures in the Old Crow-Porcupine region, Yukon Territory, Canada. *Arctic and Alpine Research* 7:249-259.
- Matthews, J.V. Jr. 1977. Tertiary Coleoptera fossils from the North American Arctic. *The Coleopterists Bulletin* 31:297-308.
- Mayr, E. 1963. Animal species and evolution. Belknap Press, Harvard University. Cambridge, Massachusetts. 797 pp.
- Melsheimer, F.E. 1853. Catalogue of the described Coleoptera of the United States. xvi + 174 pp.
- Morgan, A.V., A. Morgan, A.C. Ashworth and J.V. Matthews, Jr. 1984. Late Wisconsin fossil beetles in North America. pp. 354-363. In: Porter, S.C., Ed. Late-Quaternary Environments of the United States. Volume 1. The Late Pleistocene. University of Minnesota Press, Minneapolis.
- Morimoto, K. 1962. Comparative morphology and phylogeny of the superfamily Curculionoidea of Japan. *Journal of the Faculty of Agriculture, Kyushu University* 11:331-373.
- Morlan, R.E. and J.V. Matthews, Jr. 1983. Taphonomy and paleoecology of fossil insect assemblages from Old Crow River (CRH-15) Northern Yukon Territory, Canada. *Géographie physique et Quaternaire* 37:147-157.
- Motschulsky, T.V. von. 1845. Observations sur la musée entomologique de l'Université Impériale de Moscou. *Bulletin de la Société Impériale des Naturalistes de Moscou* 18:332-388.
- Motschulsky, T.V. von. 1860. Coléoptères rapportés par M. Sévertsef des Steppes méridionales des Kirghises. *Bulletin de l'Académie Impériale des Sciences de St.-Petersbourg* 2:513-544.
- Mott, R.J., T.W. Anderson and J.V. Matthews, Jr. 1981. Late-glacial paleoenvironments of sites bordering the Champlain Sea based on pollen and macrofossil evidence. pp. 129-171. In: W.C. Mahaney, Ed. Quaternary Paleoclimate. Geabstracts, Norwich, England.

- Müller, O.F. 1776. Zoologiae Danicae prodromus, seu animalium Daniae et Norwegiae indigenarum characteres, nomina, et synonyma imprimis popularium. xxxii + 282 pp. Hafniae.
- Munz, P.A. and D.D. Keck. 1968. A California Flora. University of California Press. Berkeley, California. 1681 pp.
- O'Brien, C.W. and G.J. Wibmer. 1978. Numbers of genera and species of Curculionidae (Coleoptera). *Entomological News* 89:89-92.
- O'Brien, C.W. and G.J. Wibmer. 1982. Annotated checklist of the weevils (Curculionidae *sensu lato*) of North America, Central America, and the West Indies (Coleoptera: Curculionoidea). *Memoirs of the American Entomological Society* 34:x + 382 pp.
- O'Brien, C.W. and G.J. Wibmer. 1984. Annotated checklist of the weevils (Curculionidae *sensu lato*) of North America, Central America, and the West Indies. Supplement 1. *The Southwestern Entomologist* 9:286-307.
- Olivier, A.G. 1807. Entomologie, ou histoire naturelle des insectes, avec leurs caractères génériques et spécifiques, leur description, leur synonymie, et leur figure enluminée. Coléoptères. Volume 5. 612 pp. Paris.
- Pallas, P.S. 1781. Icones insectorum praesertim Rossiae Sibiriaeque peculiarium quae collegit et descriptionibus illustravit. Walther, Erlangae. 56 pp.
- Patterson, C. 1981. Methods of Paleobiogeography. pp. 446-489. In: Nelson, G. and D.E. Rosen, Eds. Vicariance Biogeography. A Critique. Columbia University Press, New York. i-xviii + 593 pp.
- Peschken, D.P. 1984. Host range of *Lema cyaneella* (Coleoptera: Chrysomelidae), a candidate for biocontrol of Canada Thistle, and of four stenophagous, foreign thistle insects in North America. *The Canadian Entomologist* 116:1377-1384.
- Petri, K. 1905. Bestimmungs-Tabelle der mir bekannt gewordenen Arten der Gattung *Lixus* Fab. aus Europa und der angrenzenden Gebieten. *Wiener Entomologische Zeitung* 24:33-48, 101-117, 155-167.
- Petri, K. 1912. Ein neuer *Lixus* aus Turkestan und Bemerkungen zu meiner Bestimmungstabelle des Genus *Lixus* Fabr.. *Annales Historico-naturales Musei Nationalis Hungarici, Budapest* 10:277-280.

- Pianka, E.R. 1975. Niche relations of desert lizards. pp. 292-314. In: Cody, M.L. and J.M. Diamond, Eds. *Ecology and Evolution of Communities*. Bellknap Press of Harvard University, Cambridge. i-xiv + 545 pp.
- Pierce, W.D. 1907. On the biologies of the Rhynchophora of North America. *Studies from the Zoological Laboratory, The University of Nebraska* 78:249-319.
- Platnick, N.I. and G. Nelson. 1978. A method of analysis for historical biogeography. *Systematic Zoology* 27:1-16.
- Randall, J. 1838. Descriptions of new species of coleopterous insects inhabiting the state of Massachusetts. *Boston Journal of Natural History* 2:34-52.
- Raven, P.H. 1963. Amphitropical relationships in the floras of North and South America. *Quarterly Review of Biology* 38:151-177.
- Raven, P.H. and D.I. Axelrod. 1978. Origin and relationships of the California flora. *University of California Publications in Botany* 72:i-viii + 1-134.
- Reitter, E. 1916. *Fauna Germanica. Die Käfer des Deutschen Reiches. Nach der analytischen Methode bearbeitet. Band V. K.G. Lutz' Verlag. Stuttgart. 343 pp.*
- Rzedowski, J. 1978. *Vegetacion de Mexico. Editorial Limusa. 432 pp. Mexico City.*
- Sanders, H.O. 1960. The female genitalia and spermathecae of some of the Rhynchophora. *The Great Basin Naturalist* 20:1-22.
- Say, T. 1831. Descriptions of new species of Curculionites of North America, with observations of some of the species already known. 30 pp. New Harmony, Indiana.
- Scheinert, W. 1933. Symbiose und Embryonalentwicklung bei Rüsselkäfern. *Zeitschrift für Morphologie und Ökologie der Tiere* 27:76-128.
- Schenkling, S. and G.A.K. Marshall. 1931. *Coleopterorum Catalogus. Pars 174. Curculionidae: Cylindrorrhinae. pp. 1-23.*
- Schoenherr, C.J. 1823. *Tabula synoptica familiae curculionidum. Isis Oken, heft X, columns 1132-1146.*
- Schoenherr, C.J. 1826. *Curculionidum disposito methodica cum generum characteribus, descriptionibus atque*

observationibus variis, seu prodromus ad synonymiae insectorum, partena 4, x + 338 pp. Lipsiae.

- Schoenherr, C.J. 1834. Genera et species Curculionidum, cum synonymia hujus familiae. Species novae aut hactenus minus cognitae, descriptionibus a Dom. Leonardo Gyllenhal, C.H. Boheman, et entomologis aliis illustratae. Volume 2, Part 1. 1-326 pp. Part 2. 329-669 pp. Paris.
- Scopoli, J.A. 1763. Entomologia Carniolica exhibens insecta Carnioliae indigena distributa in ordines, genera, species, varietates. Methodo Linnaeana. xxxiv + 420 pp. Vindobonae.
- Scudder, S.H. 1893. Tertiary Rhynchophorous Coleoptera of the United States. *Monographs of the United States Geological Survey* 21:1-206.
- Shreve, F. 1942. The desert vegetation of North America. *Botanical Review* 195-246.
- Silfverberg, H. (Ed.) 1979. Enumeratio Coleopterorum Fennoscandiae et Daniae. vi + 79 pp. Helsinki.
- Sleeper, E.L. 1969. Two new Cleoninae from the Rita Blanca Lake deposits (Coleoptera: Curculionidae). *Geological Society of America, Memoirs* 113:131-133.
- Smith, P.W. 1957. An analysis of post-Wisconsin biogeography of the Prairie Peninsula region based on distributional phenomena among terrestrial vertebrate populations. *Ecology* 38:205-218.
- Solari, F. 1941. Revisione dei *Neoplinthus* Italiani ed alcune note di sistematica generale dei Curculionidi (Coleoptera). *Memoire Societa Entomologica Italiana* 20:43-90.
- Spaulding, W.G., E.B. Leopold, and T.R. Van Devender. 1985. Late Wisconsin Paleoeecology of the American Southwest. pp 259-293. In: Porter, S.C., Ed. Late-Quaternary Environments of the United States. Volume 1. The Late Pleistocene. University of Minnesota Press, Minneapolis.
- Sprague, P.S. and E.P. Austin. 1875. On the species of Coleoptera described by Mr. J.W. Randall. *Proceedings of the Boston Society of Natural History* 17:373-385.
- Stebbins, G.L. and J. Major. 1965. Endemism and speciation in the California flora. *Ecological Monographs* 35:1-35.
- Tanner, V.M. 1934. Coleoptera of Zion National Park No. II.



*Annals of the Entomological Society of America*  
27:43-49.

- Tanner, V.M. 1966. Rhynchophora beetles of the Nevada Test Site. *Brigham Young University Science Bulletin, Biological Series* 8(2):1-35.
- Ter-Minasyan, M.E. 1963. Phylogeny of beetles of the subfamily Cleoninae (Coleoptera, Curculionidae). *Entomological Review* 42:420-423.
- Ter-Minasyan, M.E. 1978. Weevils of the subfamily Cleoninae in the fauna of the USSR. Tribe Lixini. Zoological Institute, Academy of Sciences of the USSR. vii + 166 pp. Amerind Publishing Co., New Delhi. (Translated from Russian; originally published in 1967).
- Ter-Minasyan, M.E. 1979. Review of the weevil genus *Stephanocleonus* Motsch. (Coleoptera; Curculionidae). *Insects of Mongolia*. Number 6. pp. 184-342.
- Ting, P.C. 1936. The mouthparts of the coleopterous group Rhynchophora. *Microentomology* 1:93-114.
- Van Dyke, E.C. 1953. New Coleoptera from western North America (Carabidae, Throscidae, Curculionidae). *The Pan-Pacific Entomologist* 29:98-101.
- Villers, C.J. 1789. Caroli Linnaei entomologia. Band 1. Lugduni, Piestre et Delamolliere. 24u 765p. 3 Taf.
- Vuilleumier, F. and D. Simberloff. 1980. Ecology versus History as Determinants of Patchy and Insular Distributions in High Andean Birds. *Evolutionary Biology* 12:235-379.
- Watrous, L.E. and Q.D. Wheeler. 1981. The out-group comparison method of character analysis. *Systematic Zoology* 30:1-11.
- Webb, S.D. 1977. A history of savanna vertebrates in the New World. Part I: North America. *Annual Review of Ecology and Systematics* 8:355-380.
- Wells, P.V. and J.H. Hunziker. 1976. Origin of the creosote bush (*Larrea*) deserts of southwestern North America. *Annals of the Missouri Botanical Garden* 63:843-861.
- Wheeler, A.G., Jr. and D.R. Whitehead. 1985. *Larinus planus* (F.) in North America (Coleoptera: Curculionidae: Cleoninae) and comments on biological control of Canada thistle. *Proceedings of the Entomological Society of Washington* 87:751-758.

- Whitehead, D.R. 1972. Classification, phylogeny, and zoogeography of *Schizogenius* Putzeys (Coleoptera: Carabidae: Scaratini). *Quaestiones Entomologicae* 8:131-348.
- Wibmer, G.J. and C.W. O'Brien. 1986. Annotated checklist of the weevils (Curculionidae *sensu lato*) of South America (Coleoptera: Curculionidae). *Memoirs of the American Entomological Institute* 39:i-xvi + 1-563 pp.
- Wickham, H.F. 1889. Collecting notes. *Entomologica Americana* 5:77-78.
- Wickham, H.F. 1896. A list of some Coleoptera from the southern portions of New Mexico and Arizona. *Bulletin from the Laboratories of Natural History, State University of Iowa* 3:153-171.
- Wickham, H.F. 1902. A catalog of the Coleoptera of Colorado. *Bulletin of the State of Illinois Laboratory of Natural History* 5:217-310.
- Wickham, H.F. 1909. A list of the Coleoptera of Iowa. *Bulletin from the Laboratories of Natural History of the State University of Iowa* 6:1-40.
- Wickham, H.F. 1911. Fossil Coleoptera from Florissant, with descriptions of several new species. *Bulletin of the American Museum of Natural History* 30:53-69.
- Wickham, H.F. 1912. On some fossil Rhynchophorous Coleoptera from Florissant, Colorado. *Bulletin of the American Museum of Natural History* 31:41-55 + 4 pl.
- Wilcox, J., D.C. Mote, and L. Childs. 1934. The root weevils injurious to Strawberries in Oregon. Oregon State Agricultural Experiment Station, Station Bulletin 330. 109 pp.
- Wiley, E.O. 1981. Phylogenetics. The Theory and Practice of Phylogenetic Systematics. Wiley-Interscience, John Wiley and Sons, New York. i-xvi + 1-439 pp.
- Yothers, W. 1916. Bud weevils and other bud-eating insects of Washington. State College of Washington Agricultural Experiment Station Bulletin 124. 43 pp. + 6 plates.
- Zoubkoff, B. 1829. Notice sur un nouveau genre et quelques nouvelles espèces de Coléoptères. *Bulletin de la Société Impériale des Naturalistes de Moscou* 1:147-168, tab. 2.
- Zoubkoff, B. 1833. Nouveaux Coléoptères recueillis en Turcménie. *Bulletin de la Société Impériale des*

- Naturalistes de Moscou* 6:310-340.
- Zwölfer, H. and P. Harris. 1984. Biology and host specificity of *Rhinocyllus conicus* (Froel.) (Col. Curculionidae), a successful agent for biocontrol of the thistle, *Carduus nutans* L. *Zeitschrift für Angewandte Entomologie* 97:36-62.
- Zwölfer, H. and M. Preiss. 1983. Host selection and oviposition behaviour in West-European ecotypes of *Rhinocyllus conicus* Froel. (Col. Curculionidae). *Zeitschrift für Angewandte Entomologie* 95:113-122.

8. Appendix 1

Table 87. Chronological relationships of Cleonidius species

<i>C. porricollis</i>	S	<i>C. eustictognathus</i>	S
<i>C. eubictorhinus</i>	BS P	<i>C. modestus</i>	A A A
<i>C. modestus</i>	A A A	<i>C. subcylindricus</i>	BS S A A
<i>C. subcylindricus</i>	BS S A A	<i>C. longinasus</i>	BS S BS A BS
<i>C. longinasus</i>	BS S BS A BS	<i>C. porticollis</i>	S S P A S S
<i>C. texanus</i>	S S P A S S	<i>C. americanus</i>	BS P BS A BS BS P
<i>C. americanus</i>	BS P BS A BS BS P	<i>C. frontalis</i>	BS A BS A A NS A BS
<i>C. frontalis</i>	BS A BS A A NS A BS	<i>C. canescens</i>	P A A A A S A S S
<i>C. canescens</i>	P A A A A S A S S	<i>C. infrequens</i>	S A A A A A A S S
<i>C. infrequens</i>	S A A A A A A S S	<i>C. puberulus</i>	S A A A A A A S S
<i>C. puberulus</i>	S A A A A A A S S	<i>C. collaris</i>	P A S A A S A S S
<i>C. collaris</i>	P A S A A S A S S	<i>C. notolomus</i>	BS S BS A BS BS S BS BS S
<i>C. notolomus</i>	BS S BS A BS BS S BS BS S	<i>C. lobigerinus</i>	A A A NS A A A A A A A
<i>C. lobigerinus</i>	A A A NS A A A A A A A	<i>C. calandroides</i>	P A P A NS NS A A A NS
<i>C. calandroides</i>	P A P A NS NS A A A NS	<i>C. boucardi</i>	BS P BS A BS BS P BS BS S
<i>C. boucardi</i>	BS P BS A BS BS P BS BS S	<i>C. trivittatus</i>	S S P A S S S P A A A A A
<i>C. trivittatus</i>	S S P A S S S P A A A A A	<i>C. placidus</i>	BS S BS A BS BS S BS BS S
<i>C. placidus</i>	BS S BS A BS BS S BS BS S	<i>C. quadrilineatus</i>	BS S BS A BS BS S BS BS S
<i>C. quadrilineatus</i>	BS S BS A BS BS S BS BS S		

S, sympatric; BS, broadly sympatric; NS, narrowly sympatric; P, parapatric; A, allopatric.



Table 69. Character states in *Cleonidius* species and related taxa

Character	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	
<i>C. poricolis</i>	1	1	1	1	1	2	1	1	1	1	2	1	3	r	1	1	3	2	2	2	2	2	1
<i>C. eustictorrhinus</i>	1	1	1	1	2	2	1	1	1	1	2	1	3	1	1	1	2	2	2	2	2	2	1
<i>C. modestus</i>	1	1	1	3	1	2	1	1	1	1	2	2	1	1	1	1	2	1	2	2	2	2	1
<i>C. subcylindricus</i>	1	1	1	1	1	2	2	1	2	2	2	1	2	1	2	1	1	1	2	2	2	2	2
<i>C. longinasus</i>	1	-1	1	1	1	2	2	1	2	2	2	2	1	1	1	1	1	3	1	2	2	2	2
<i>C. texanus</i>	1	1	1	1	1	2	2	1	2	2	2	2	1	1	1	1	3	1	2	2	2	2	2
<i>C. americanus</i>	2	1	2	1	2/3	2	1	1	1	1	2	2	2	1	1	1	3	2	2	2	2	2	1
<i>C. frontalis</i>	2	1	-2	1	2/3	2	1	1	1	1	2	1	2	1	1	1	3	2	1	2	2	2	1
<i>C. canescens</i>	1	2	1	1	2/3	2	1	1	1	1	2	1	3	1	3	1	2	2	2	2	1	1	1
<i>C. infrequens</i>	1	3	1	1	2	2	1	1	1	1	2	2	1	2	2	1	2	2	1	2	2	1	1
<i>C. puberulus</i>	1	3	1	1	2	2	1	1	1	1	2	1	2	1	1	1	2	3	2	1	1	1	1
<i>C. collaris</i>	1	1	1	1	3	2	1	1	1	1	2	2	2	1	2	2	2	2	2	1	1	1	1
<i>C. notolomus</i>	1	1	1	1	2	2	1	1	1	1	2	1	2	2	1	2	2	2	2	1	1	1	1
<i>C. lobigerinus</i>	1	1	1	1/2	2	2	1	1	1	1	2	1	3	1	1	1	3	1	2	2	2	1	1
<i>C. calandroides</i>	1	1	1	1/2	2	2	1	1	1	1	2	1	3	1	1	1	2	1	2	2	2	1	1
<i>C. boucardi</i>	1	1	1	1	2	2	1	2	1	1	1	1	1	1	1	2	1	1	1	1/2	2	1	1
<i>C. trivittatus</i>	1	1	1	1	2	2	1	2	1	1	1	1	3	1	2/3	1	2	2	1/2	2	1	1	1
<i>C. piacidus</i>	1	1	1	1	1	2	1	2	1	1	1	1	3	1	2/3	1	3	2	2	1	1	1	1
<i>C. quadrilineatus</i>	1	1	1	1	2	2	1	2	1	1	2	1	3	1	3	1	3	2	2	2	2	1	1
<i>C. vibex</i>	1	2	1	1	2	2	1	1	1	1	2	1	3	2	1	1	1	3	2	1	2	1	1
<i>Lixoglyptus</i>	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	2
<i>Cylindropterus</i>	1	1	1	1	/	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1

Table 70. Plesiotypic and apotypic states of characters used in phylogenetic analysis of *Apieurus* species and related taxa

Character	Plesiotypic	Apotypic
1. Epistoma, apical margin	-emarginate	-rounded and produced
2. Prementum, setae	-setae lacking or only one large seta per side	-2 or more large setae per side
3. Pronotum, tubercles, development and shape of anterior angles	-tubercles absent	-tubercles present, anterior angles obtuse (2) -tubercles present, anterior angles acute (3)
4. Tarsus, ventral pilose vestiture	-present, extensive	-present, moderately reduced in extent (2) -present, markedly reduced in extent or absent (3)
5. Female, stylus, size	-present, large	-present small (2) -absent (3)
6. Female, stylus, position	-present, apical	-present; antepical (2) -absent, antepical sclerotized ridge present (3)
7. Male, abdominal sternum VII, dorsally directed tooth at apical margin	-absent	-present
8. Male, abdominal sternum VIII, anterior angles of each sternite	-basal projection absent	-basal projection present, slightly to moderately developed (2) -basal projection present, markedly developed (3)
9. Size, length of elytra	-X < 7.0 mm	-7.0 mm < X < 9.0 mm (2) -X > 9.0 mm (3)
10. Eye, prominence and convexity	-markedly convex and prominent	-slightly to moderately prominent and convex (2) -flat (3)
11. Suberect or erect vestiture, length	-absent or indistinct	-present, short (2) -present, long (3)
12. Rostrum, carina	-present	-absent
13. Rostrum, declivity at apex	-flat to slightly declivous	-moderately to steeply declivous

14. Maxillary palpus, setae  
 -one large seta on palpifer, one on stipes  
 -one large seta on palpifer
15. Pronotum, width  
 -greatest at base to subequal at base and tubercles  
 -greatest at tubercles
16. Pronotum, punctation  
 -small and shallow  
 -large and deep
17. Pronotum, median carina  
 -absent  
 -present, various in extent
18. Prosternum, impressions and swellings  
 -impression anterolaterad of fore-coxal cavity; no swelling present  
 -impression anterolaterad of fore-coxal cavity; low rounded swelling present immediately anterior to impression (2)  
 -impression anterolaterad of fore-coxal cavity; high sharp swelling present immediately anterior to impression (3)  
 -impression anterolaterad of fore-coxal cavity; low rounded swelling present immediately anterior to impression and low rounded swelling present immediately in front of fore-coxal cavity (4)
19. Elytra, elevation of intervals  
 -all intervals equally flat or slightly elevated and convex  
 -sutural and intervals 3, 5, 7, and 9 slightly to markedly elevated and convex
20. Fore-tibia of female, apex  
 -uncus and large second spur present  
 -uncus and at most small second spur present
21. Fore-tibia, inner margins  
 -large denticles present  
 -small denticles present
22. Female abdominal sternum VIII, lateral arms  
 -more or less straight throughout length  
 -arcuate
23. Female abdominal tergum VII, median longitudinal carina  
 -absent  
 -present, slightly developed (2)  
 -present, markedly developed (3)
24. Female abdominal tergum VIII, median longitudinal carina  
 -absent  
 -present, slightly developed (2)  
 -present, markedly developed (3)
25. Male, aedeagus, curvature  
 -markedly arcuate at base  
 -more or less evenly arcuate throughout length
26. Male, aedeagus, apex  
 -spatulate  
 -not spatulate



27. Male, aedeagus, internal sac, paired lateral sclerite at midlength -present  
 -absent
28. Male, aedeagus, internal sac, shape -dorsal median pocket low, two pairs of lobes  
 -dorsal median pocket high, one pair of lobes  
 -absent
29. Male, aedeagus, internal sac, lobe A -present  
 -absent
30. Male, aedeagus, internal sac, apical sclerite complex, median projection -present, long  
 -present, short and indistinct or absent
31. Male, aedeagus, internal sac, apical sclerite complex, sclerotization of base and sides or apical pocket -well-sclerotized, dark brown in color  
 -lightly-sclerotized, pale brown in color

Table 71. Pleistotypic and apotypic states of characters used in phylogenetic analysis of *Cleonidius* species and related taxa

Character	Pleistotypic	Apotypic
1. Submentum, degree of ventral swelling	-flat	-swollen
2. Rostrum, ratio of length to width	-moderately elongate, $0.45 < X < 0.65$	-very elongate, $X < 0.45$ (-1) -short, robust, $0.65 < X < 0.77$ (2) -very short, robust, $X > 0.77$ (3)
3. Rostrum, degree of lateral swelling	-not swollen at midlength	-swollen at midlength
4. Frons/vertex, surface sculpture	-area behind eye with wrinkles of cuticle	-area behind eye with curved sulcus (2) -area behind eye with deep, straight sulcus (3)
5. Elytra, scale pattern	-marginal (see text)	-vittate (see text) (2) -mottled (see text) (3)
6. Abdomen of female, ornamentation of abdominal sternum VII	-basal margin uniformly flat	-basal margin at middle internally with variously developed glabrous shiny tubercles
7. Male, aedeagus, internal sac, height of dorsal median pocket	-low	-high
8. Male, aedeagus, internal sac, apex of dorsal median pocket	-rounded or truncate in dorsal view	-emarginate or sinuate in dorsal view
9. Male, aedeagus, internal sac, lobe B	-present	-absent
10. Male, aedeagus, internal sac, lobe D	-present	-absent
11. Size, length of elytra	- $X > 9$ mm	- $X < 9$ mm
12. Rostrum, punctuation	-apical portion with punctures moderately large, dense and moderately deep	-apical portion with punctures sparse, moderately small and shallow
13. Rostrum, median tumescence	-absent	-present, slightly developed (2) -present, markedly developed (3)
14. Rostrum, lateral margins	-rounded	-sharp

- 15. Pronotum, anterolateral margins
  - postocular lobe rounded, slightly well-developed (2)
  - postocular lobe absent, margin straight (3)
- 16. Pronotum, impressions
  - basal median area shallowly to moderately deeply impressed
  - basal median area deeply impressed, lateral margins variously longitudinally impressed
- 17. Suberect or erect vestiture, length
  - absent
  - present, short to moderate (2)
  - present, long (3)
- 18. Tarsus, ventral pilose vestiture
  - present, extensive (3)
- 19. Fore-tibia of female, inner margin-large denticles present
  - small denticles present
- 20. Female abdominal sternum VIII, length of basal arm
  - long
  - short
- 21. Female abdominal sternum VIII, shape of lateral arms
  - arcuate
  - more or less straight, divergent
- 22. Male, aedeagus, internal sac, lobe
  - present
  - absent

**Figures 1 to 4.** Lateral view of head of various Cleoninae.

**Figure 1.** *Microlarinus* species (Scale bar = 0.8 mm).

**Figure 2.** *Rhinocyllus conicus* (Scale bar = 1.6 mm).

**Figure 3.** *Larinus planus* (Scale bar = 1.6 mm).

**Figure 4.** *Lixus scrobicollis* (Scale bar = 1.6 mm).

**Figures 5 to 6.** Apex of tibia of various Cleoninae (Scale bar = 0.5 mm).

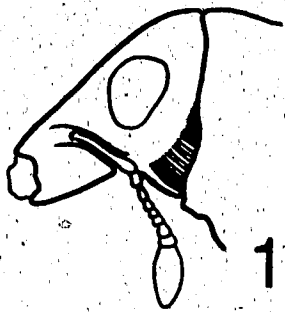
**Figure 5.** *Cleonidius infrequens*.

**Figure 6.** *Apleurus lutulentus*.

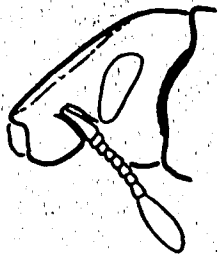
**Figures 7 to 8.** Ventral view of prosternum of various Cleoninae (Scale bar = 1.6 mm).

**Figure 7.** *Apleurus* species.

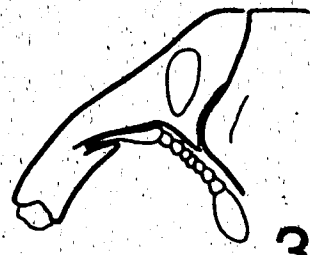
**Figure 8.** *Cleonidius* species.



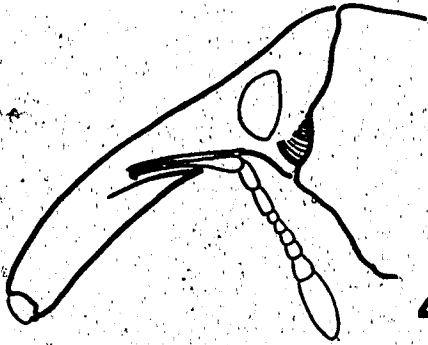
1



2



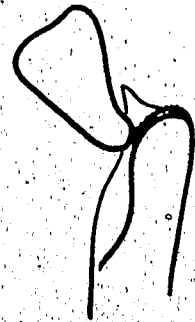
3



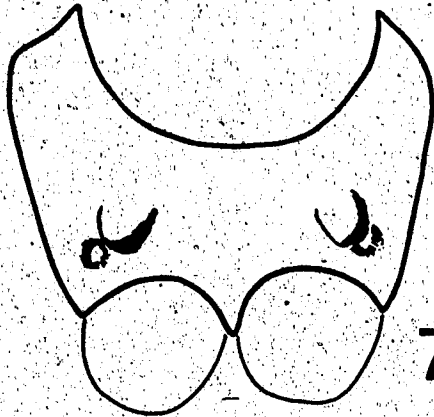
4



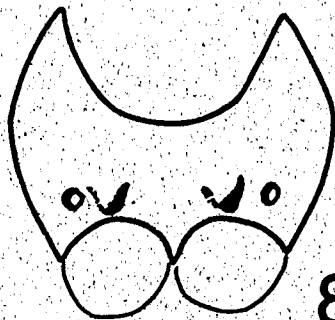
5



6



7



8

**Figures 9 to 18. *Cleonis plgra*.**

**Figure 9.** Lateral view of head (Scale bar = 1.1 mm).

**Figure 10.** Dorsal view of head (Scale bar = 1.1 mm).

**Figure 11.** Lateral view of internal sac of male (Scale bar = 0.7 mm).

**Figure 12.** Dorsal view of internal sac of male (Scale bar = 0.7 mm).

**Figure 13.** Lateral view of aedeagus of male (Scale bar = 1.3 mm).

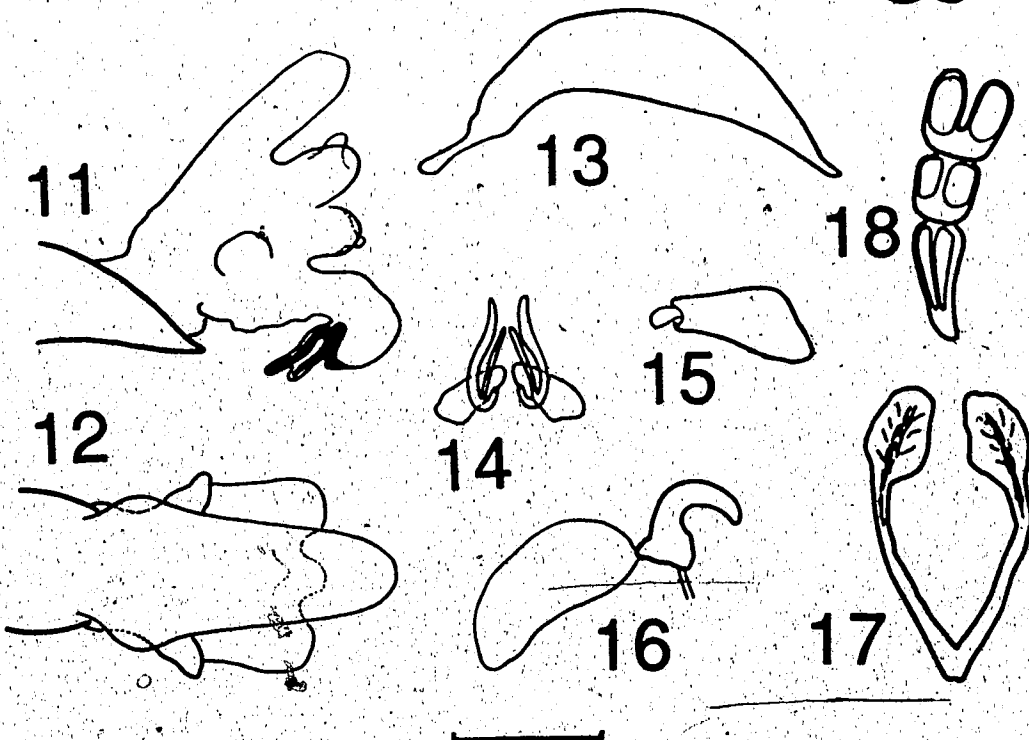
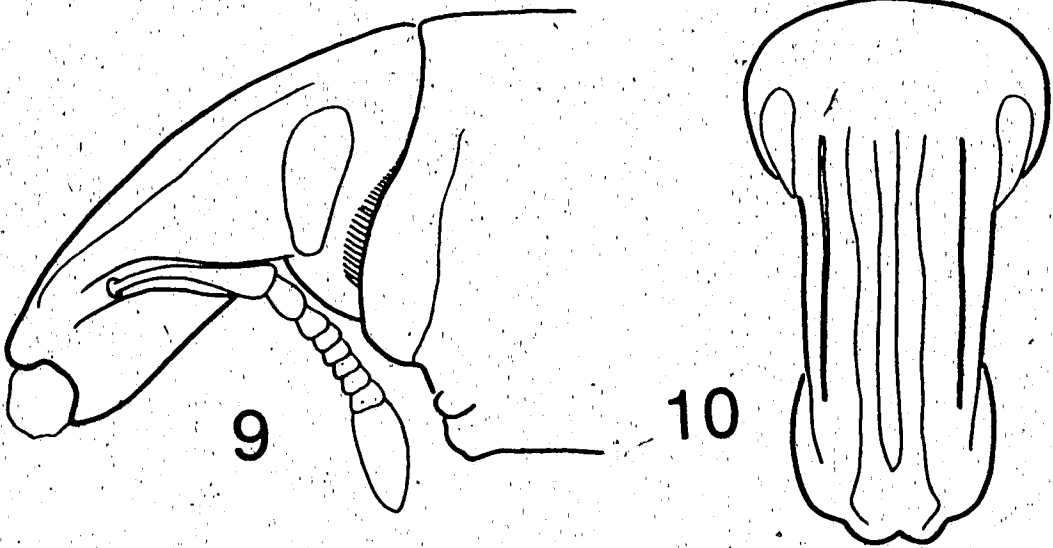
**Figure 14.** Ventral view of apical sclerite complex of internal sac of male (Scale bar = 0.6 mm).

**Figure 15.** Lateral view of gonocoxite II and stylus of female (Scale bar = 1.3 mm).

**Figure 16.** Spermatheca of female (Scale bar = 0.7 mm).

**Figure 17.** Ventral view of sternum VIII of female (Scale bar = 1.3 mm).

**Figure 18.** Ventral view of metatarsus (Scale bar = 1.3 mm).



Figures 19 to 24.

Figure 19. Dorsal habitus of *Cleonis pigra*. Scale bar = 5.0 mm.

Figure 20. Dorsal habitus of *Stephanocleonus confusus*. Scale bar = 5.0 mm.

Figure 21. Dorsal habitus of *Stephanocleonus immaculatus*. Scale bar = 5.0 mm.

Figure 22. Dorsal habitus of *Stephanocleonus plumbeus*. Scale bar = 5.0 mm.

Figure 23. Dorsal habitus of *Stephanocleonus stenothorax*. Scale bar = 5.0 mm.

Figure 24. Dorsal habitus of *Apleurus (Gibbostethus) hystrix*. Scale bar = 2.5 mm.

---





19



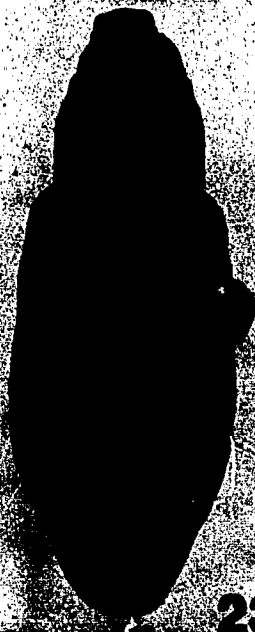
20



21



22



23



24



Figures 25 to 30. Head of *Stephanocleonus* species (Scale bar = 2.4 mm).

Figure 25. *Stephanocleonus confusus*.

a. Lateral view.

b. Dorsal view.

Figure 26. *Stephanocleonus cristaticollis*.

a. Lateral view.

b. Dorsal view.

Figure 27. *Stephanocleonus immaculatus*.

a. Lateral view.

b. Dorsal view.

Figure 28. *Stephanocleonus parshus*.

a. Lateral view.

b. Dorsal view.

Figure 29. *Stephanocleonus plumbeus*.

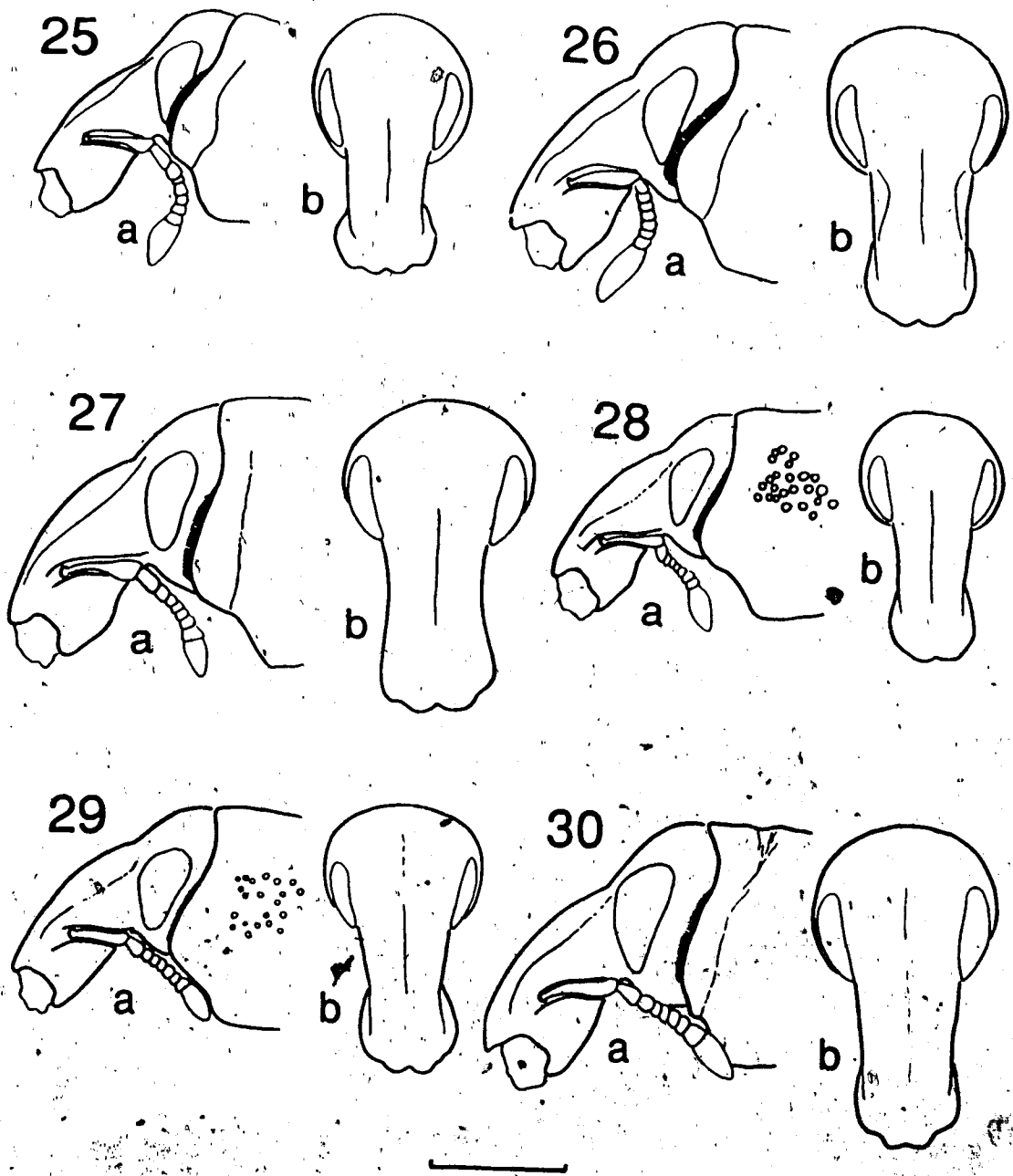
a. Lateral view.

b. Dorsal view.

Figure 30. *Stephanocleonus stenothorax*.

a. Lateral view.

b. Dorsal view.



Figures 31 to 36. Male genitalia of *Stephanocleonus* species  
(Scale bar = 1.3 mm).

Figure 31. *Stephanocleonus confusus*.

- a. Lateral view of aedeagus.
- b. Ventral view of apex of aedeagus.
- c. Ventral view of sternite of sternum VII.

Figure 32. *Stephanocleonus cristaticollis*.

- a. Lateral view of aedeagus.
- b. Ventral view of apex of aedeagus.
- c. Ventral view of sternite of sternum VII.

Figure 33. *Stephanocleonus immaculatus*.

- a. Lateral view of aedeagus.
- b. Ventral view of apex of aedeagus.
- c. Ventral view of sternite of sternum VII.

Figure 34. *Stephanocleonus parshus*.

- a. Lateral view of aedeagus.
- b. Ventral view of apex of aedeagus.
- c. Ventral view of sternite of sternum VII.

Figure 35. *Stephanocleonus plumbeus*.

- a. Lateral view of aedeagus.
- b. Ventral view of apex of aedeagus.
- c. Ventral view of sternite of sternum VII.

Figure 36. *Stephanocleonus stenothorax*.

- a. Lateral view of aedeagus.
- b. Ventral view of apex of aedeagus.
- c. Ventral view of sternite of sternum VII.

Figures 37 to 42. Ventral view of sternum VIII of female  
*Stephanocleonus* species (Scale bar = 1.3 mm).

Figure 37. *Stephanocleonus confusus*.

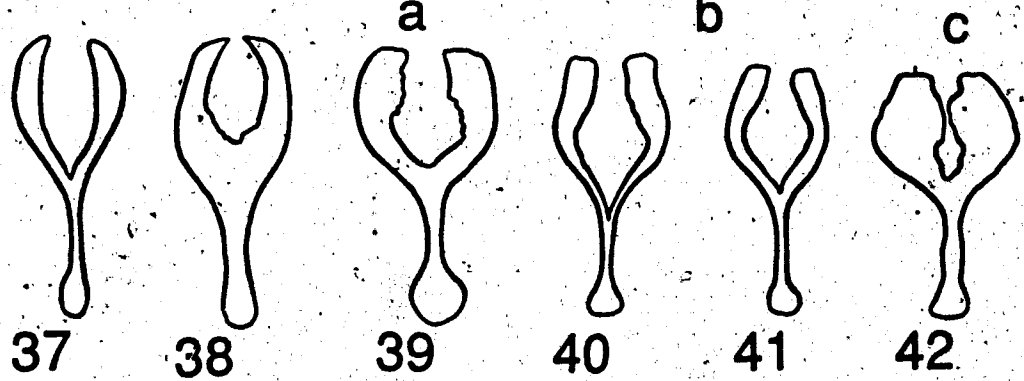
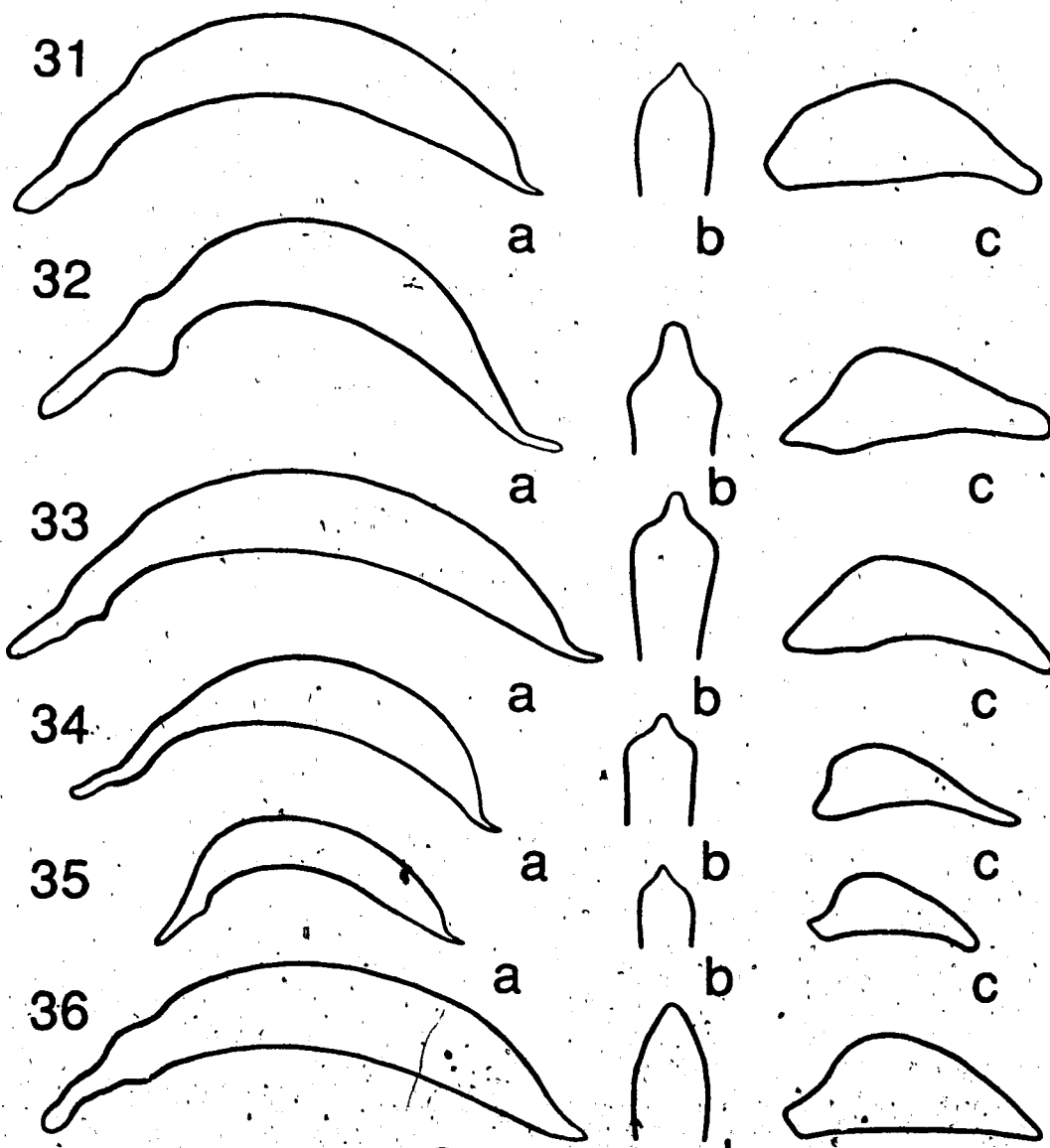
Figure 38. *Stephanocleonus cristaticollis*.

Figure 39. *Stephanocleonus immaculatus*.

Figure 40. *Stephanocleonus parshus*.

Figure 41. *Stephanocleonus plumbeus*.

Figure 42. *Stephanocleonus stenothorax*.



Figures 43 to 48. Internal sac of male *Stephanocleonus* species (Scale bar = 0.7 mm).

Figure 43. *Stephanocleonus confusus*.

- a. Lateral view.
- b. Dorsal view.

Figure 44. *Stephanocleonus cristicollis*.

- a. Lateral view.
- b. Dorsal view.

Figure 45. *Stephanocleonus immaculatus*.

- a. Lateral view.
- b. Dorsal view.

Figure 46. *Stephanocleonus parshus*.

- a. Lateral view.
- b. Dorsal view.

Figure 47. *Stephanocleonus plumbeus*.

- a. Lateral view.
- b. Dorsal view.

Figure 48. *Stephanocleonus stenothorax*.

- a. Lateral view.
- b. Dorsal view.

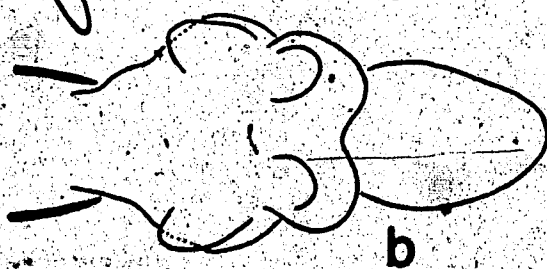
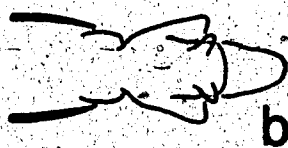
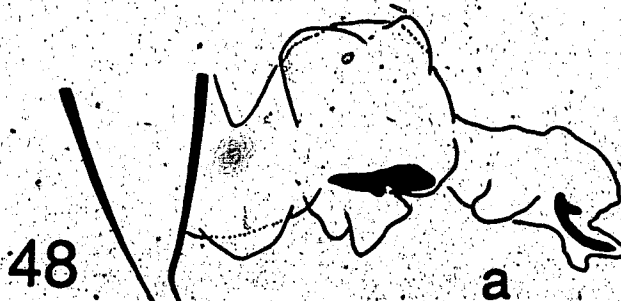
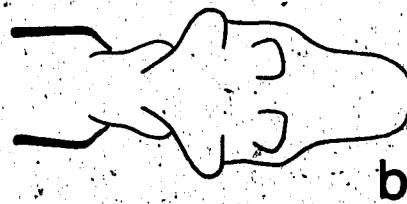
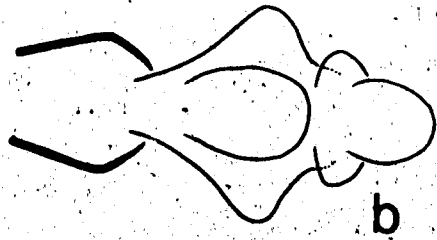
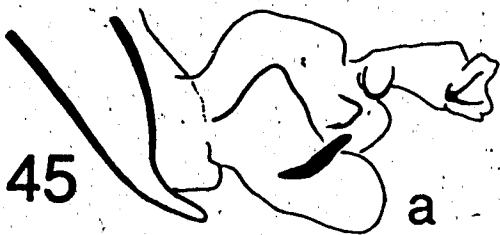
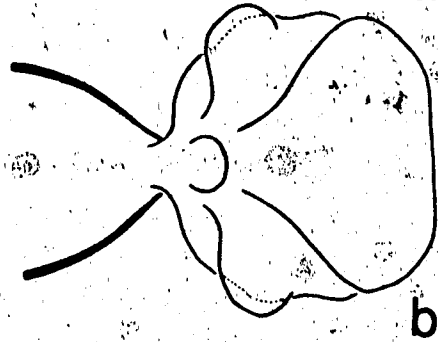
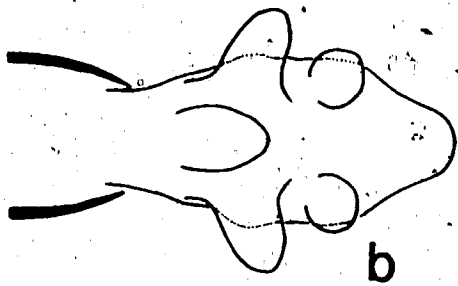
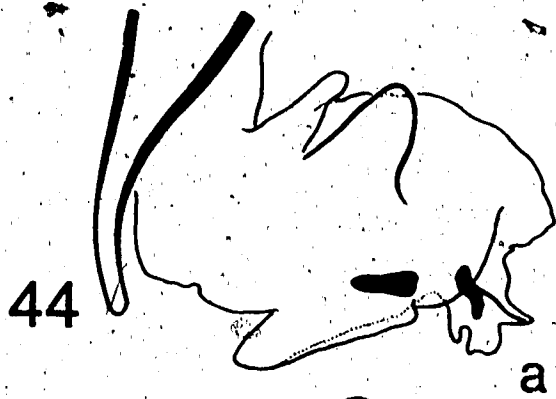
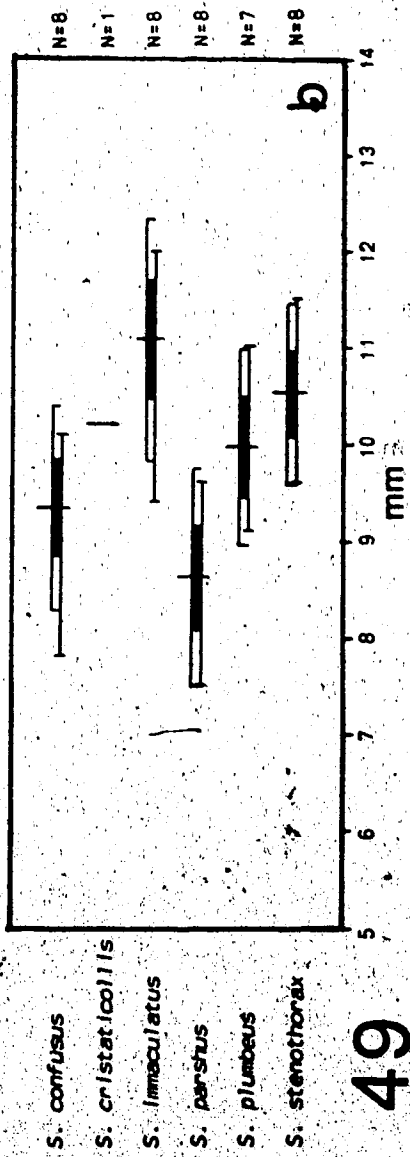
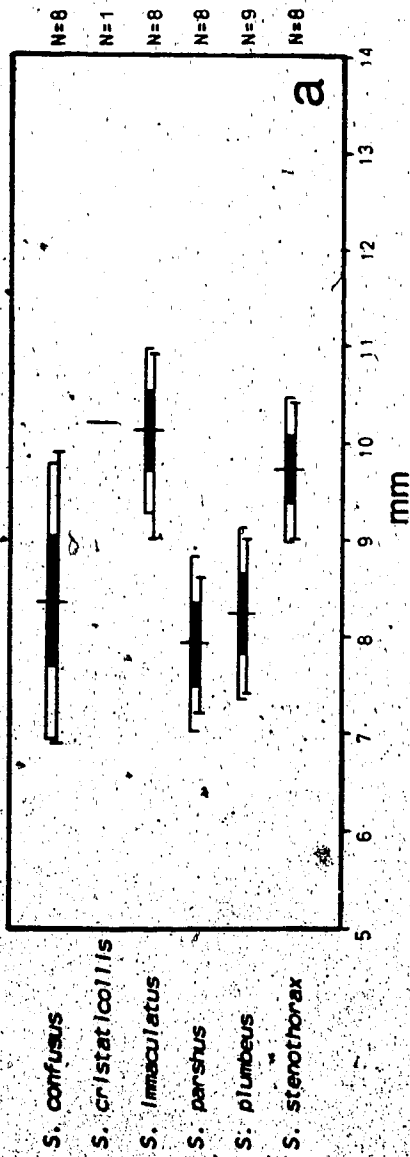


Figure 49. Hubbs-Hubbs diagram illustrating variation among specimens of *Stephanocleonus* species; length of elytra (LEl): a, males; b, females. The range of each species is indicated by a horizontal line and the mean by a long vertical line. A solid rectangle represents two standard errors on either side of the mean and a hollow rectangle represents 1.5 standard deviations on either side of the mean. Only mean and range are shown for samples with less than five specimens.

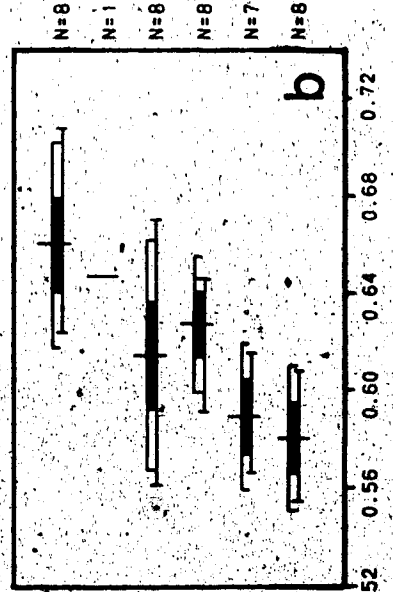
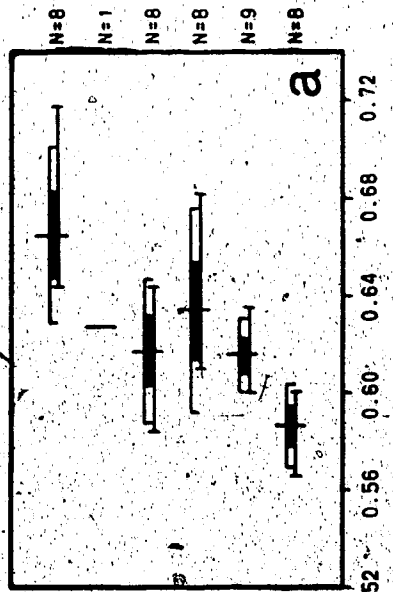
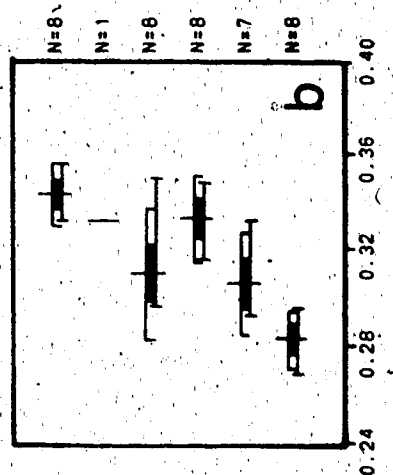
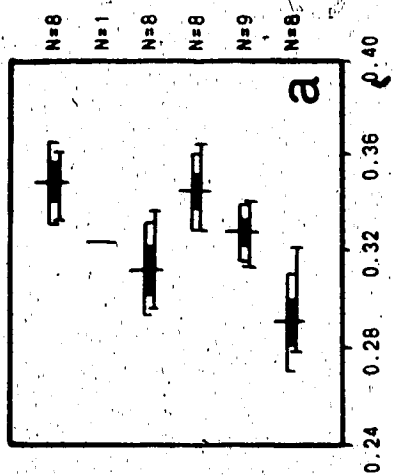




**Figures 50 to 51.**

**Figure 50.** Hubbs-Hubbs diagram illustrating variation among specimens of *Stephanocleonus* species; form of elytra (WE1M/LE1): a, males; b, females. See caption for Figure 49 for explanation.

**Figure 51.** Hubbs-Hubbs diagram illustrating variation among specimens of *Stephanocleonus* species; relative length of pronotum compared to elytra (LP/LE1): a, males; b, females. See caption for Figure 49 for explanation.



51

50

Figure 52. Hubbs-Hubbs diagram illustrating variation among specimens of *Stephanocleonus* species; form of rostrum (WRA/LR): a, males; b, females. See caption for Figure 49 for explanation.

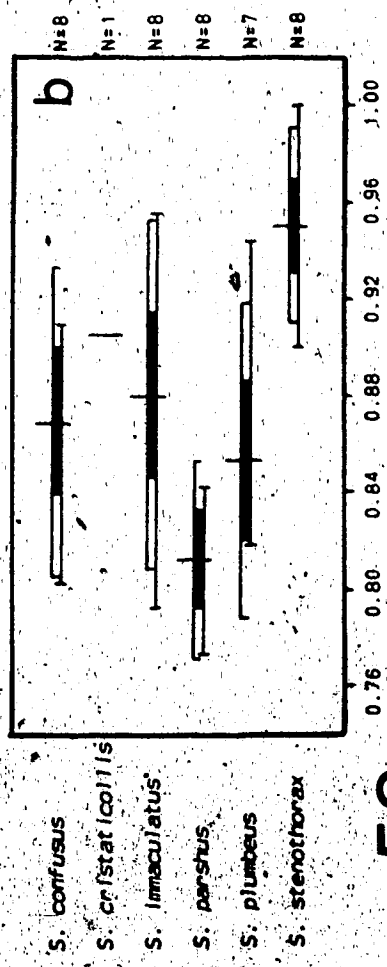
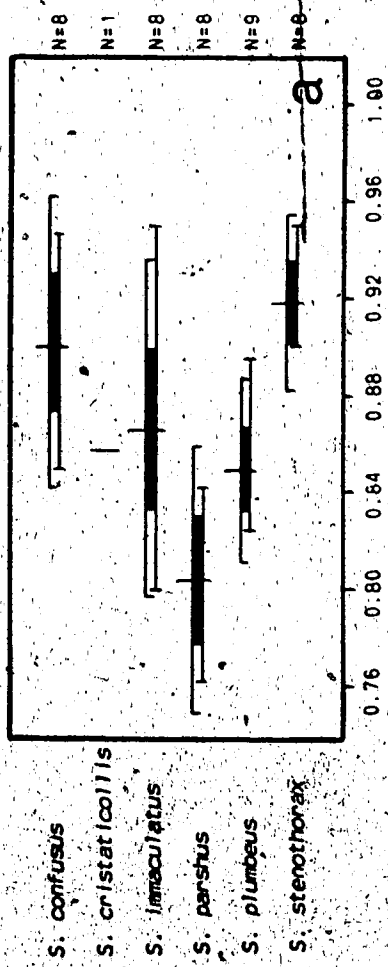
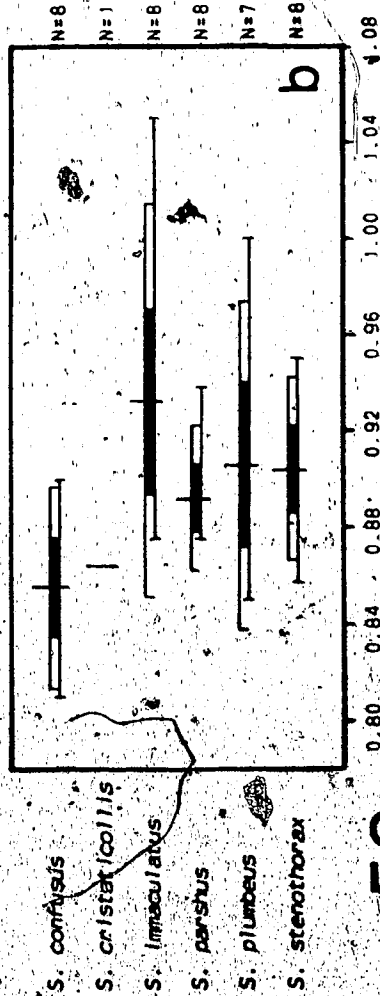
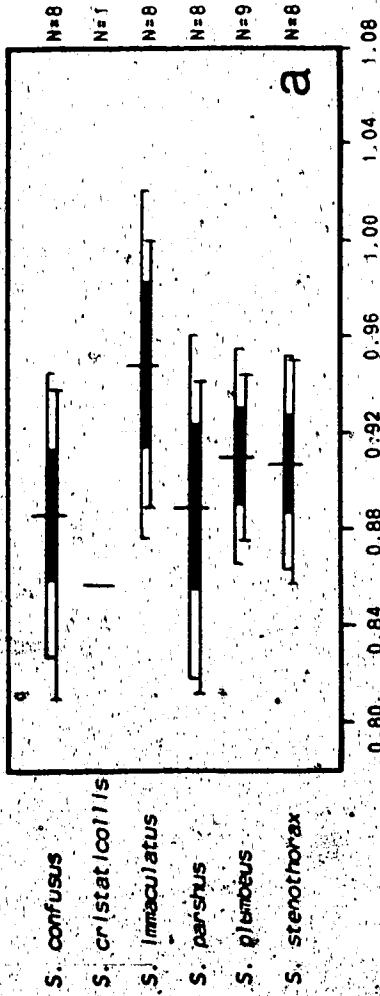


Figure 53. Hubbs-Hubbs diagram illustrating variation among specimens of *Stephanocleonus* species; relative width of frons compared to rostrum (WRA/WF): a, males; b, females. See caption for Figure 49 for explanation.



Figures 54 to 67.

Figure 54. Lateral view of gonocoxite II and stylus of female *Stephanocleonus*. Scale bar = 0.7 mm.

Figure 55. Spermatheca of female *Stephanocleonus*. Scale bar = 0.7 mm.

Figure 56. Genitalia of *Chromoderus fasciatus*.

a. Lateral view of aedeagus of male. Scale bar = 1.3 mm.

b. Lateral view of internal sac of male. Scale bar = 0.7 mm.

c. Dorsal view of internal sac of male. Scale bar = 0.7 mm.

d. Ventral view of apical sclerite complex of internal sac of male. Scale bar = 0.6 mm.

e. Ventral view of sternum VIII of female. Scale bar = 1.3 mm.

Figure 57. Genitalia of *Cnemodontus limpidus*.

a. Lateral view of aedeagus of male. Scale bar = 1.3 mm.

b. Lateral view of internal sac of male. Scale bar = 0.7 mm.

c. Dorsal view of internal sac of male. Scale bar = 0.7 mm.

d. Ventral view of apical sclerite complex of internal sac of male. Scale bar = 0.6 mm.

e. Ventral view of sternum VIII of female. Scale bar = 1.3 mm.

Figure 58. Lateral view of mesosternum of *Apleurus* (*Gibbostethus*) *hystrix*. Scale bar = 1.3 mm.

Figure 59. Lateral view of mesosternum of *Apleurus* (*Apleurus*) *lutulentus*. Scale bar = 1.3 mm.

Figure 60. Spermatheca of female *Apleurus*. Scale bar = 0.7 mm.

Figure 61. Ventral view of ligula of *Apleurus* (*Gibbostethus*) *hystrix*. Scale bar = 0.2 mm.

Figure 62. Ventral view of ligula of *Apleurus* (*Apleurus*) *lutulentus*. Scale bar = 0.4 mm.

Figure 63. Ventral view of ligula of *Apleurus* (*Apleurus*) *sagrinatus*. Scale bar = 0.4 mm.

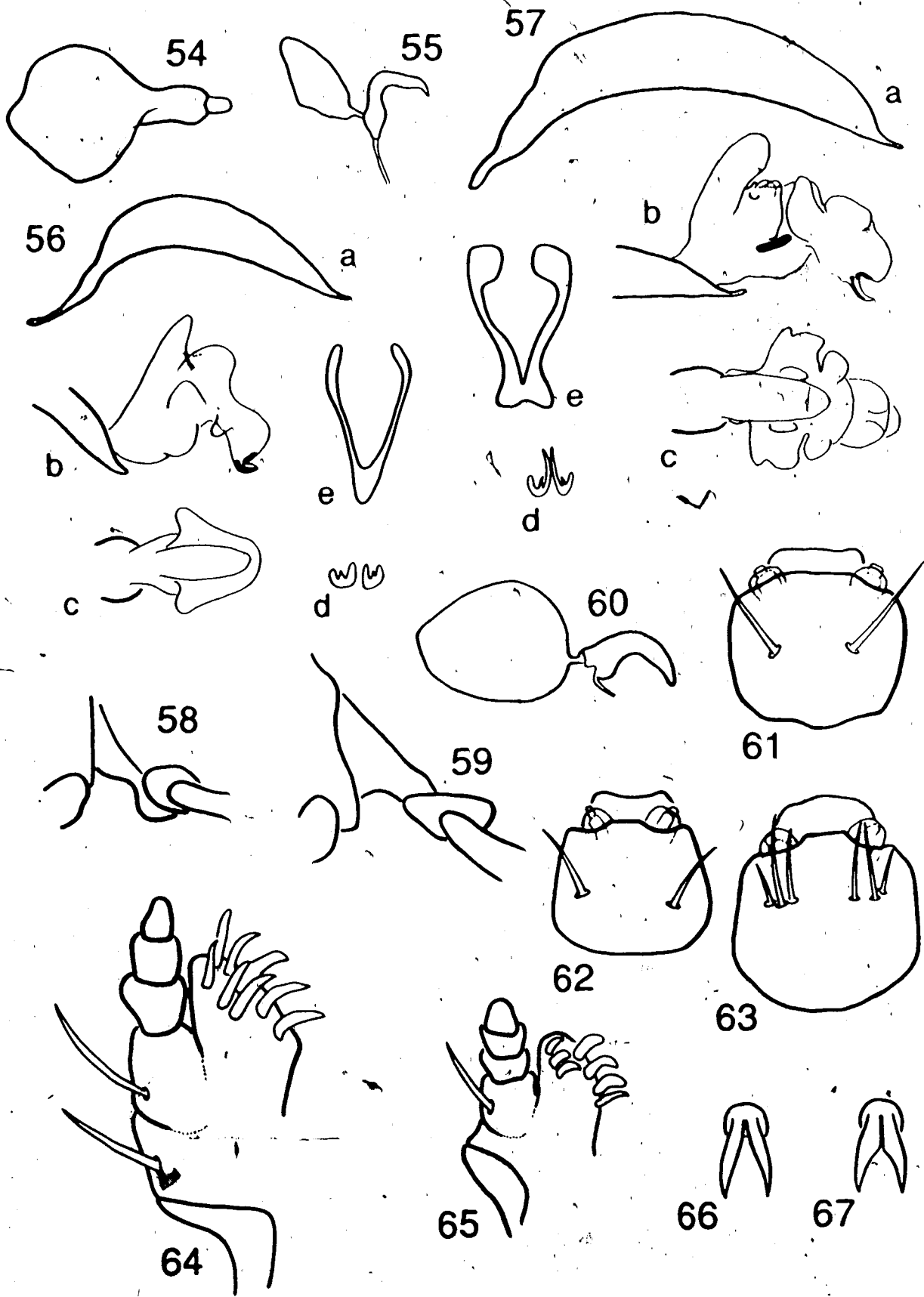
Figure 64. Ventral view of maxilla of *Apleurus* (*Apleurus*) *sagrinatus*. Scale bar = 0.4 mm.

Figure 65. Ventral view of maxilla of *Apleurus* (*Gibbostethus*) *hystrix*. Scale bar = 0.4 mm.

Figure 66. Tarsal claws of *Apleurus angularis*. Scale bar = 0.7 mm.

Figure 67. Tarsal claws of *Apleurus jacobinus*. Scale bar = 0.7 mm.





Figures 68 to 74.

Figure 68. Dorsal habitus of *Apleurus aztecus*. Scale bar = 2.0 mm.

Figure 69. Dorsal habitus of *Apleurus lutulentus*. Scale bar = 4.0 mm.

Figure 70. Dorsal habitus of *Apleurus porosus*. Scale bar = 4.0 mm.

Figure 71. Dorsal habitus of *Apleurus jacobinus*. Scale bar = 4.0 mm.

Figure 72. Dorsal habitus of *Apleurus angularis* (3 mi. N Baker, Nevada). Scale bar = 4.0 mm.

Figure 73. Dorsal habitus of *Apleurus angularis* (Colorado Springs, Colorado). Scale bar = 4.0 mm.

Figure 74. Dorsal habitus of *Apleurus angularis* (Baboquivaria Mountains, Arizona). Scale bar = 3.0 mm.



68



69



70



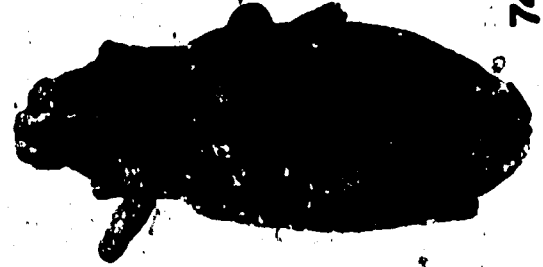
71



72



73



74

Figures 75 to 80:

Figure 75. Dorsal habitus of *Apleurus albovestitus* (18 mi. SW Mendota, California). Scale bar = 5.0 mm.

Figure 76. Dorsal habitus of *Apleurus albovestitus* (16 mi. S. Vidal, California). Scale bar = 5.0 mm.

Figure 77. Dorsal habitus of *Apleurus albovestitus* (Virgin, Utah). Scale bar = 5.0 mm.

Figure 78. Dorsal habitus of *Apleurus albovestitus* (Phoenix, Arizona). Scale bar = 5.0 mm.

Figure 79. Dorsal habitus of *Apleurus albovestitus* (Yuma, Arizona). Scale bar = 5.0 mm.

Figure 80. Dorsal habitus of *Apleurus saginatus*. Scale bar = 5.0 mm.



77



78



79



80



Figures 81 to 88. Head of *Apleurus* species (Scale bar = 2.4 mm).

Figure 81. *Apleurus hystrix*.

a. Lateral view.

b. Dorsal view.

Figure 82. *Apleurus aztecus*.

a. Lateral view.

b. Dorsal view.

Figure 83. *Apleurus lutulentus*.

a. Lateral view.

b. Dorsal view.

Figure 84. *Apleurus porosus*.

a. Lateral view.

b. Dorsal view.

Figure 85. *Apleurus angularis*.

a. Lateral view.

b. Dorsal view.

Figure 86. *Apleurus jacobinus*.

a. Lateral view.

b. Dorsal view.

Figure 87. *Apleurus saginatus*.

a. Lateral view.

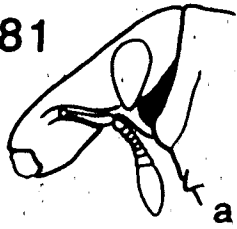
b. Dorsal view.

Figure 88. *Apleurus albovestitus*.

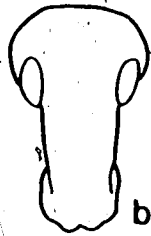
a. Lateral view.

b. Dorsal view.

81

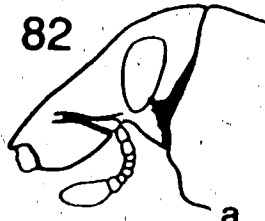


a

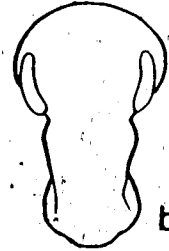


b

82

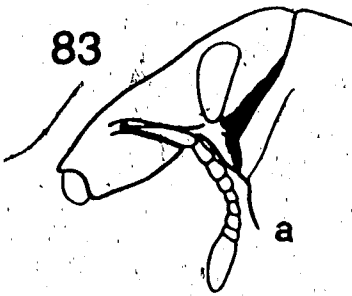


a

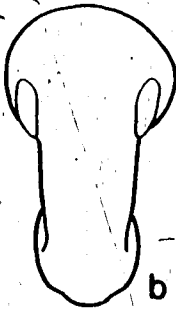


b

83

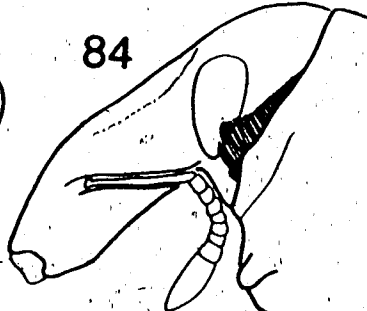


a

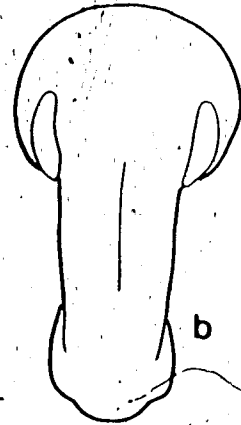


b

84

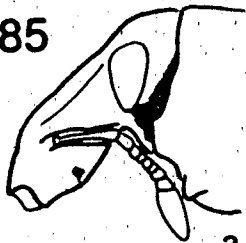


a



b

85

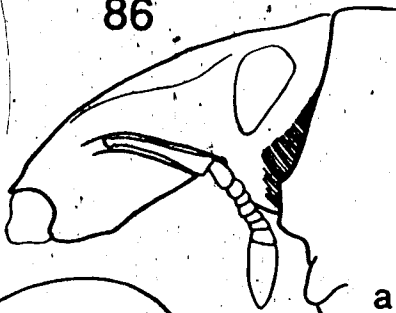


a

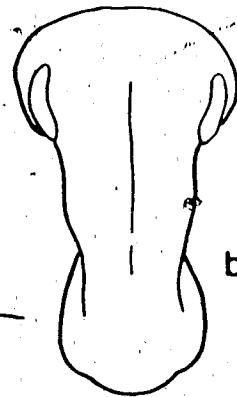


b

86

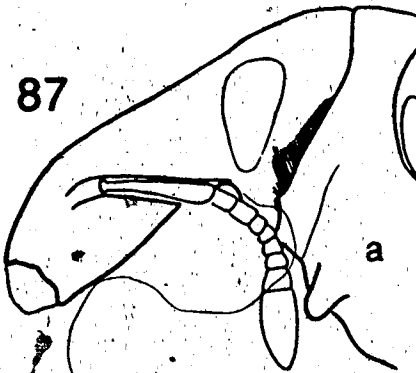


a

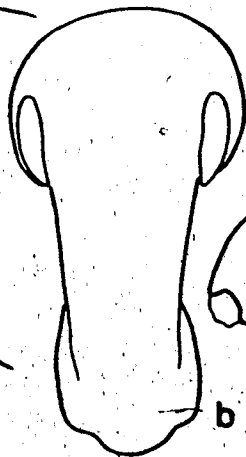


b

87

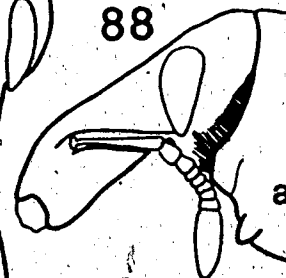


a

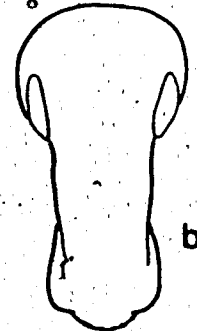


b

88



a



b

Figures 89 to 96. Female genitalia of *Apleurus* species.

Figure 89. *Apleurus hystrix* (Scale bar = 1.0 mm).

a. Ventral view of sternum VIII.

b. Lateral view of gonocoxite II and stylus.

Figure 90. *Apleurus aztecus*.

a. Ventral view of sternum VIII. Scale bar = 2.0 mm.

b. Lateral view of gonocoxite II and stylus.  
Scale bar = 1.0 mm.

Figure 91. *Apleurus lutulentus*.

a. Ventral view of sternum VIII. Scale bar = 2.0 mm.

b. Lateral view of gonocoxite II and stylus.  
Scale bar = 1.0 mm.

Figure 92. *Apleurus porosus*.

a. Ventral view of sternum VIII. Scale bar = 2.0 mm.

b. Lateral view of gonocoxite II and stylus.  
Scale bar = 1.0 mm.

Figure 93. *Apleurus angularis*.

a. Ventral view of sternum VIII. Scale bar = 2.0 mm.

b. Lateral view of gonocoxite II and stylus.  
Scale bar = 1.0 mm.

Figure 94. *Apleurus jacobinus*.

a. Ventral view of sternum VIII. Scale bar = 2.0 mm.

b. Lateral view of gonocoxite II and stylus.  
Scale bar = 1.0 mm.

Figure 95. *Apleurus aldovestitus*.

a. Ventral view of sternum VIII. Scale bar = 2.0 mm.

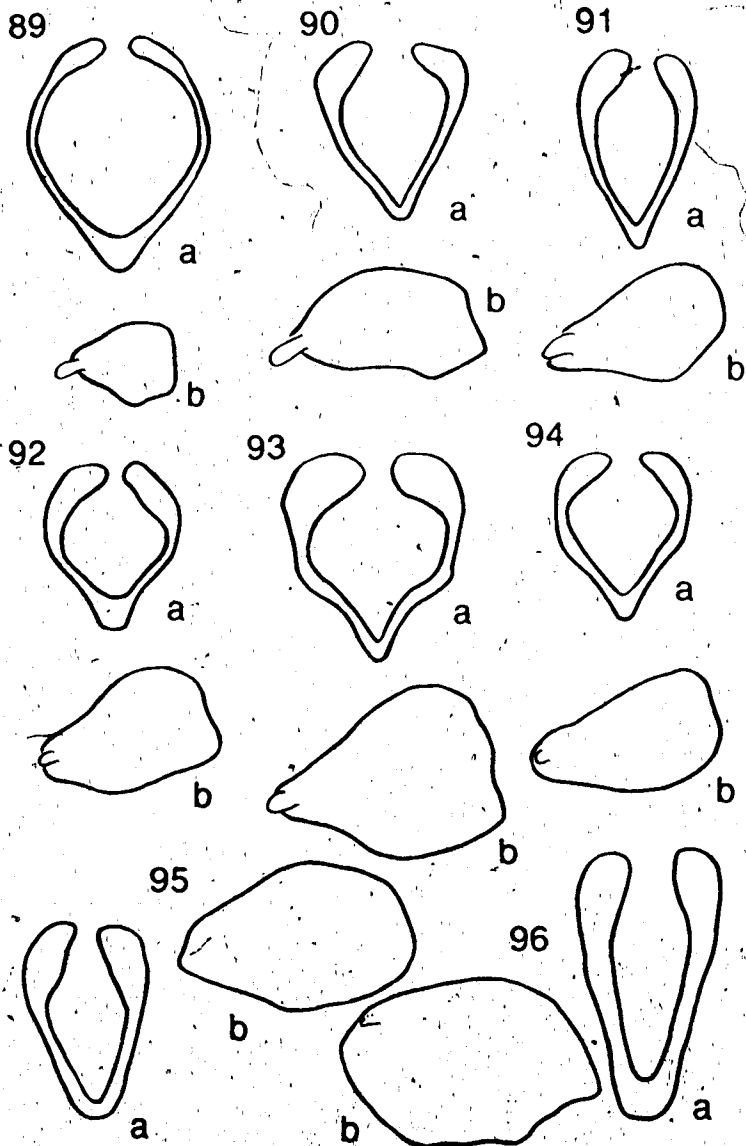
b. Lateral view of gonocoxite II and stylus.  
Scale bar = 1.0 mm.

Figure 96. *Apleurus saginatus*.

a. Ventral view of sternum VIII. Scale bar = 2.0 mm.

b. Lateral view of gonocoxite II and stylus.  
Scale bar = 1.0 mm.





Figures 97 to 99. Male genitalia of *Apleurus* species.

Figure 97. *Apleurus hystrix*.

- a. Lateral view of internal sac. Scale bar = 0.4 mm.
- b. Dorsal view of internal sac. Scale bar = 0.4 mm.
- c. Ventral view of apical sclerite complex of internal sac. Scale bar = 0.3 mm.
- d. Lateral view of aedeagus. Scale bar = 1.3 mm.
- e. Ventral view of aedeagus. Scale bar = 1.3 mm.
- f. Ventral view of sternite of sternum VII. Scale bar = 1.3 mm.

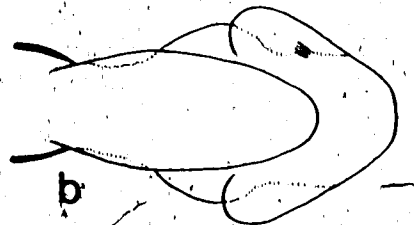
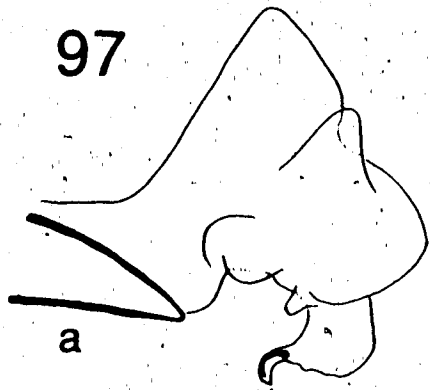
Figure 98. *Apleurus aztecus*.

- a. Lateral view of internal sac. Scale bar = 0.7 mm.
- b. Dorsal view of internal sac. Scale bar = 0.7 mm.
- c. Lateral view of aedeagus. Scale bar = 1.3 mm.
- d. Ventral view of apical sclerite complex of internal sac. Scale bar = 0.6 mm.

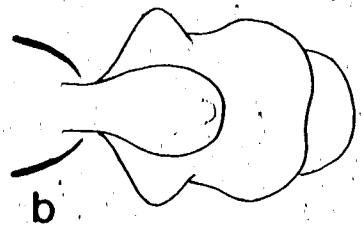
Figure 99. *Apleurus lutulentus* (Scale bars as in Fig. 98).

- a. Lateral view of internal sac.
- b. Dorsal view of internal sac.
- c. Lateral view of aedeagus.
- d. Ventral view of apical sclerite complex of internal sac.

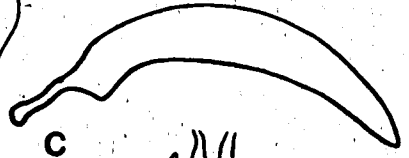
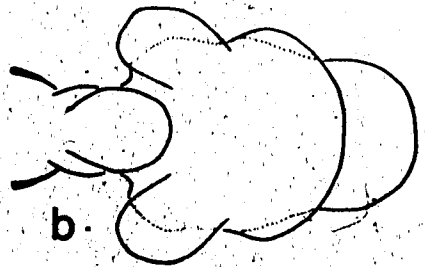
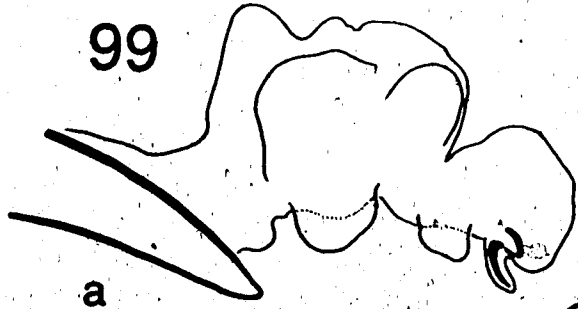
97



98



99



Figures 100 to 102. Male genitalia of *Apleurus* species.

Figure 100. *Apleurus angularis*.

- a. Lateral view of internal sac. Scale bar = 0.7 mm.
- b. Dorsal view of internal sac. Scale bar = 0.7 mm.
- c. Lateral view of aedeagus. Scale bar = 1.3 mm.
- d. Ventral view of apical sclerite complex of internal sac. Scale bar = 0.6 mm.

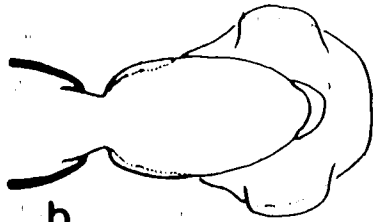
Figure 101. *Apleurus jacobinus* (Scale bars as in Fig. 100).

- a. Lateral view of internal sac.
- b. Dorsal view of internal sac.
- c. Lateral view of aedeagus.
- d. Ventral view of apical sclerite complex of internal sac.
- e. Ventral view of sternite of sternum VII.

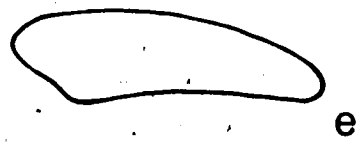
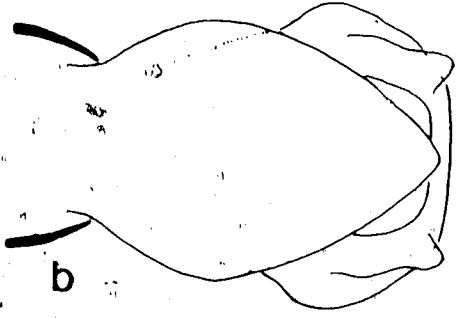
Figure 102. *Apleurus porosus* (Scale bars as in Fig. 100).

- a. Lateral view of internal sac.
- b. Dorsal view of internal sac.
- c. Lateral view of aedeagus.

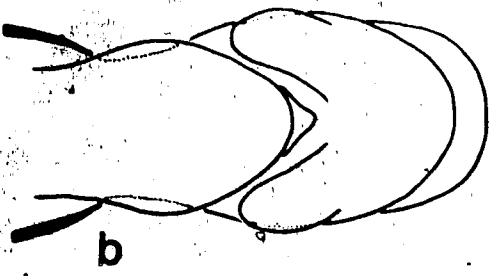
100



101



102



Figures 103 to 104. Male genitalia of *Apleurus* species.

Figure 103. *Apleurus albovestitus*.

- a. Lateral view of internal sac. Scale bar = 0.7 mm.
- b. Dorsal view of internal sac. Scale bar = 0.7 mm.
- c. Lateral view of aedeagus. Scale bar = 1.3 mm.
- d. Ventral view of apical sclerite complex of internal sac. Scale bar = 0.6 mm.
- e. Ventral view of sternite of sternum VII. Scale bar = 1.3 mm.

Figure 104. *Apleurus saginatus* (Scale bars as in Fig. 103).

- a. Lateral view of internal sac.
- b. Dorsal view of internal sac.
- c. Lateral view of aedeagus.
- d. Ventral view of apical sclerite complex of internal sac.

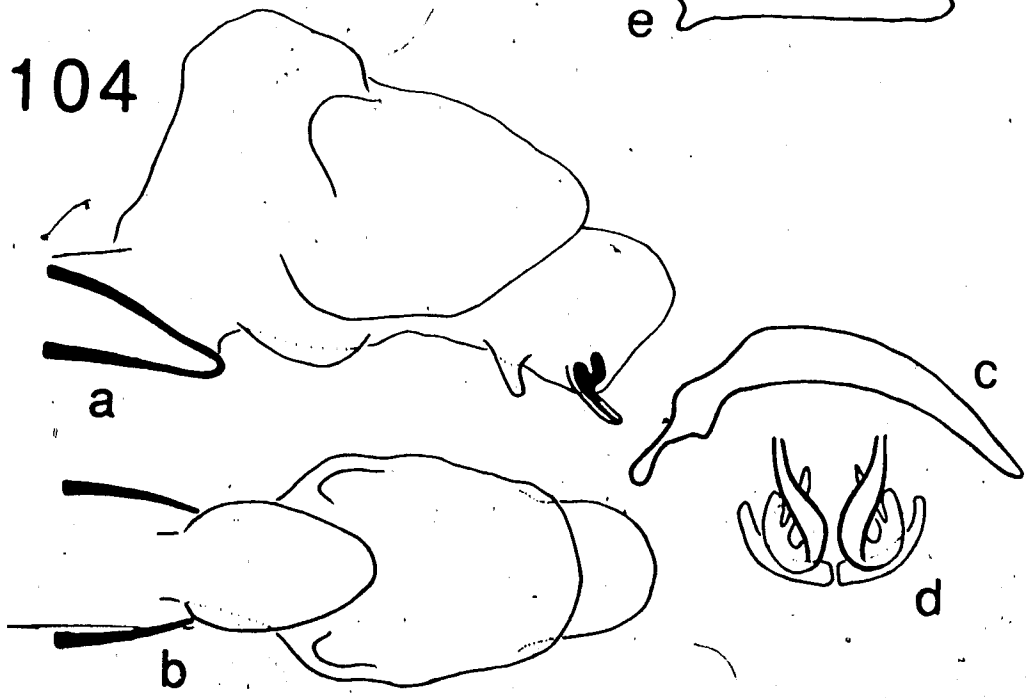
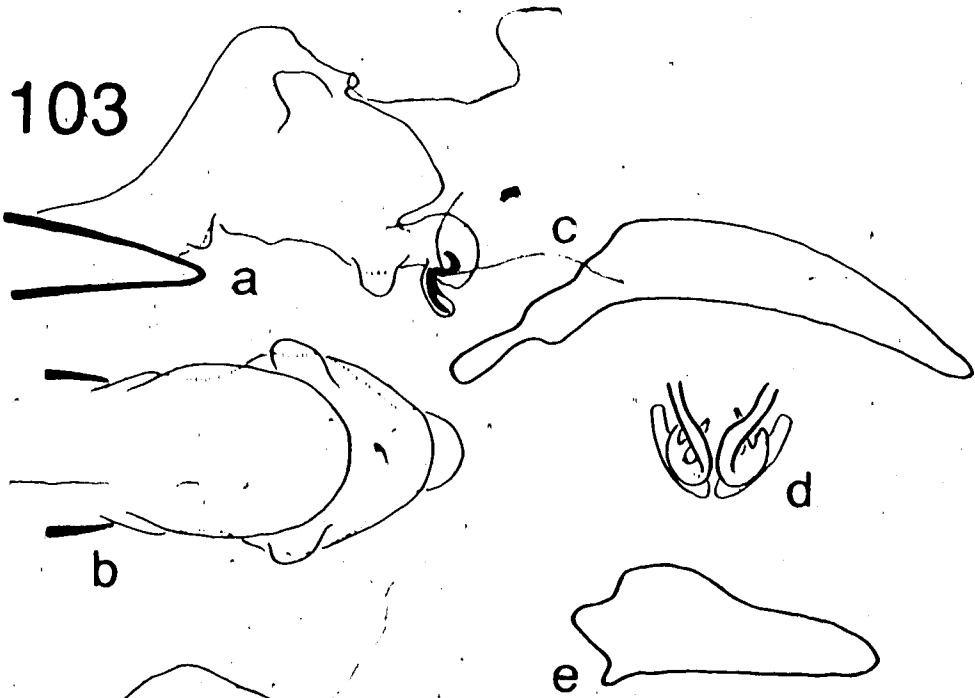
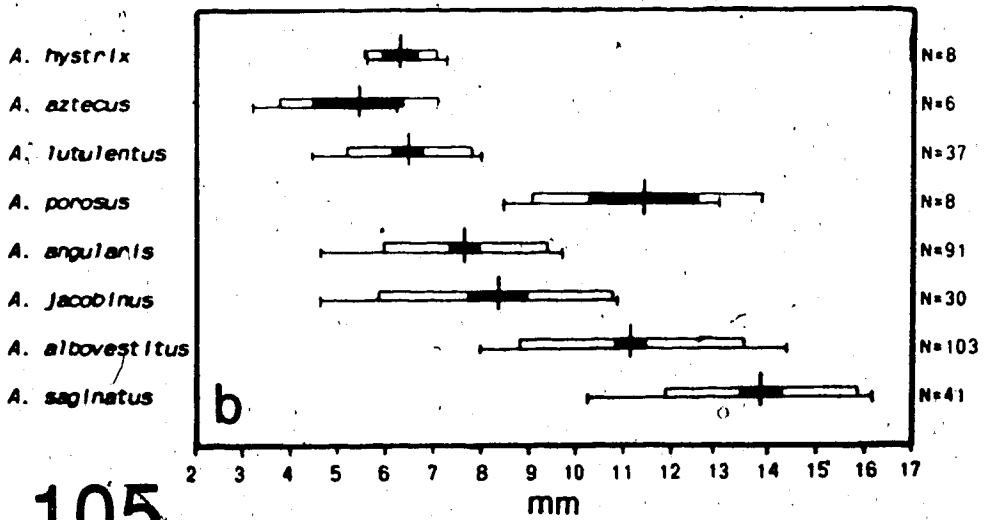
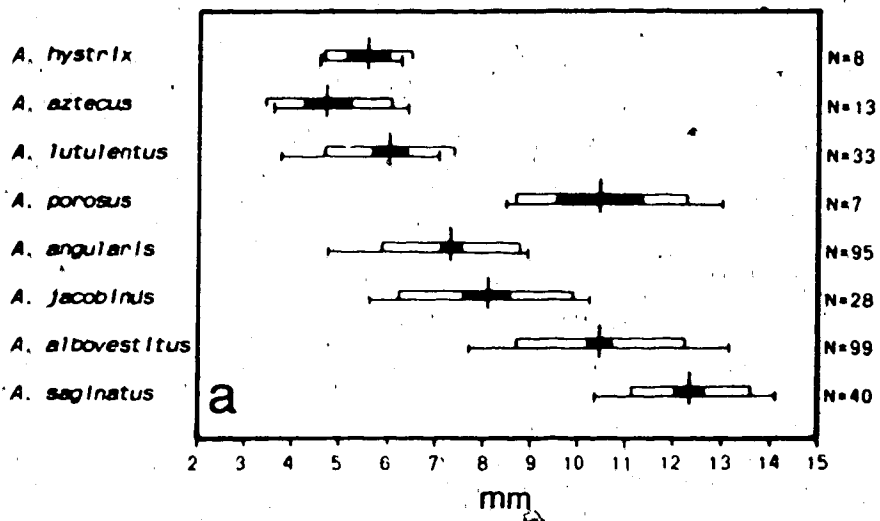


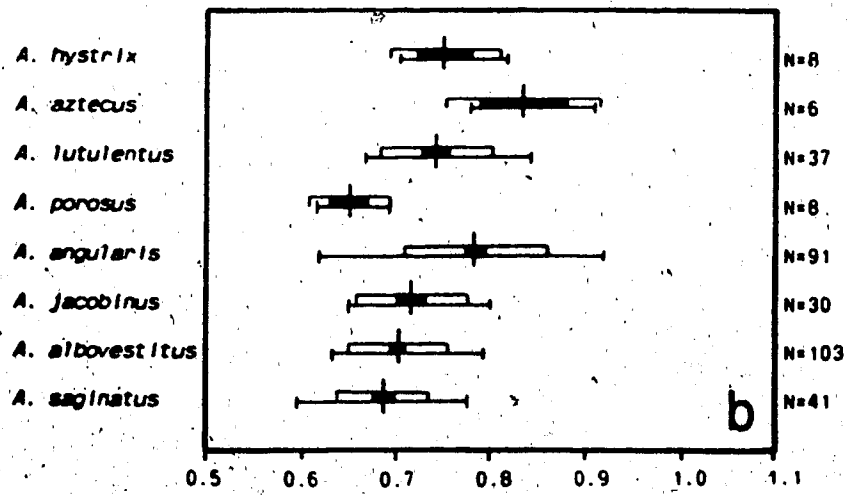
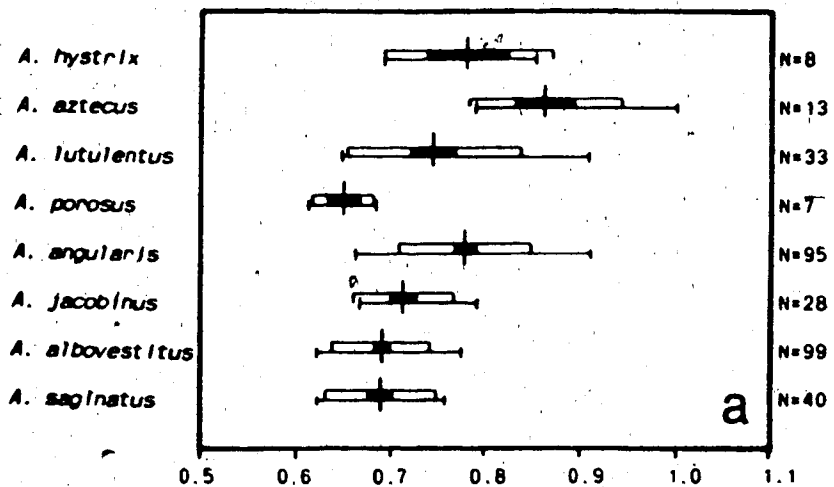
Figure 105. Hubbs-Hubbs diagram illustrating variation among specimens of *Apleurus* species; length of elytra (LE1): a, males; b, females. See caption for Figure 49 for explanation.





105

Figure 106. Hubbs-Hubbs diagram illustrating variation among specimens of *Apleurus* species; form of rostrum (WRA/LR): a, males; b, females. See caption for Figure 49 for explanation.

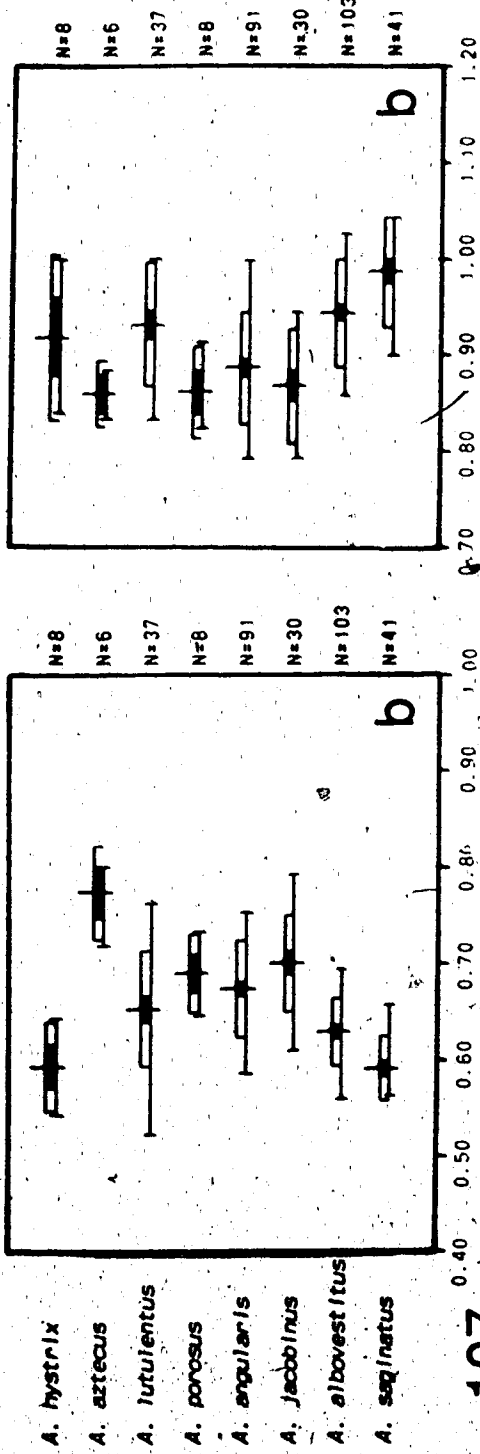
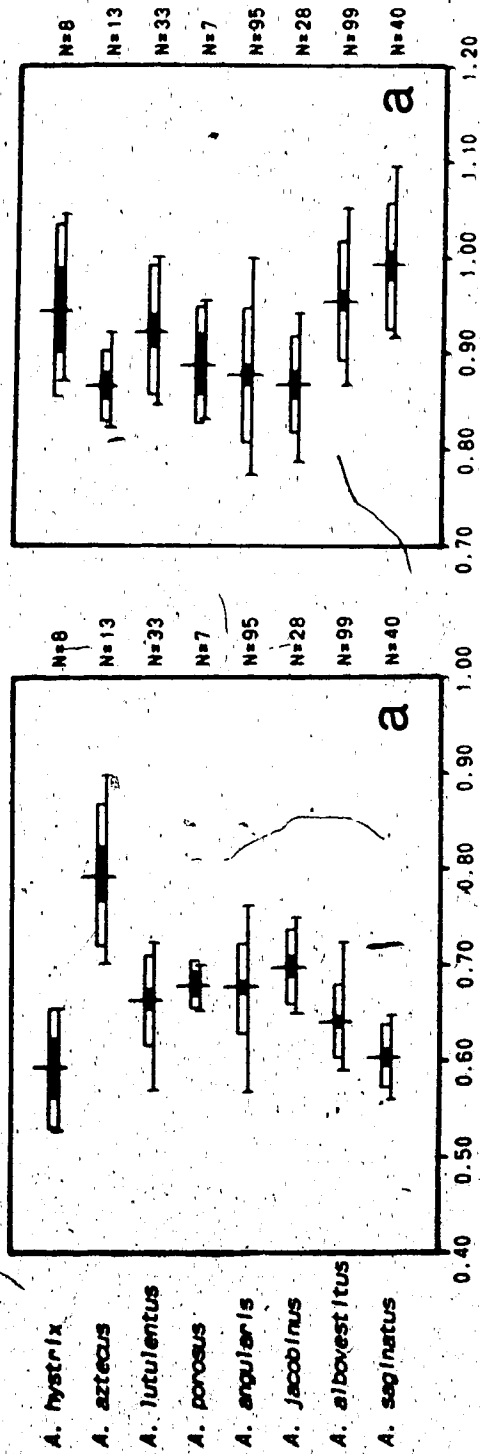


Figures 107 to 108.

Figure 107. Hubbs-Hubbs diagram illustrating variation among specimens of *Apleurus* species; form of elytra (WELM/LE1): a, males; b, females. See caption for Figure 49 for explanation.

Figure 108. Hubbs-Hubbs diagram illustrating variation among specimens of *Apleurus* species; width of frons compared to width at apex of rostrum (WRA/WF): a, males; b, females. See caption for Figure 49 for explanation.

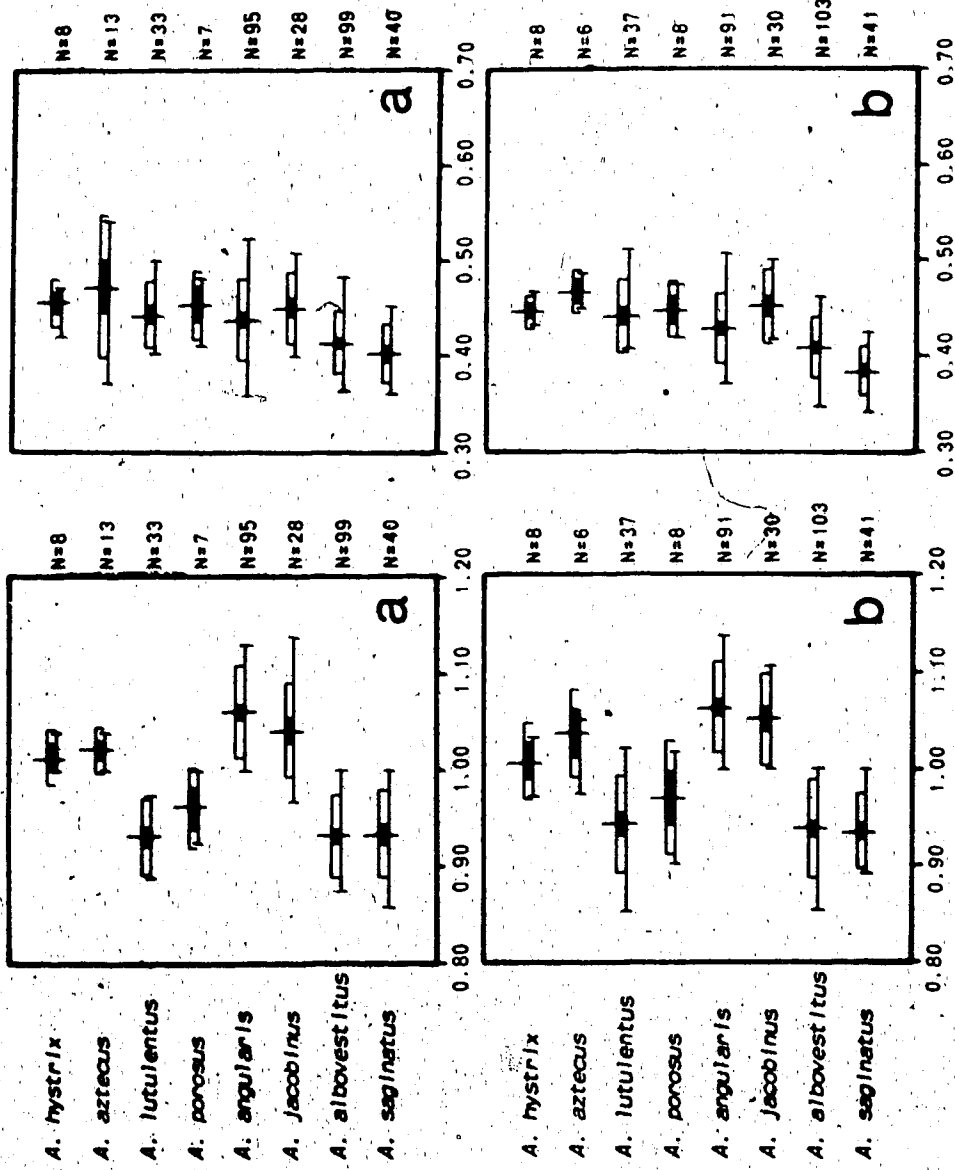
---



Figures 109 to 110.

Figure 109. Hubbs-Hubbs diagram illustrating variation among specimens of *Apleurus* species; form of pronotum (WPT/WPB): a, males; b, females. See caption for Figure 49 for explanation.

Figure 110. Hubbs-Hubbs diagram illustrating variation among specimens of *Apleurus* species; length of pronotum compared to length elytra (LP/LE1): a, males; b, females. See caption for Figure 49 for explanation.



109 110

Figures 111 to 117.

Figure 111. Male genitalia of *Cylindropterus luxerj*.  
Scale bar = 0.7 mm.

- a. Lateral view of internal sac.
- b. Dorsal view of internal sac.

Figure 112. Male genitalia of *Lixus (Lixoglyptus) spartii*. Scale bar = 0.7 mm.

- a. Lateral view of internal sac.
- b. Dorsal view of internal sac.

Figure 113. Ventral view of sternum VIII of female *Cylindropterus luxerj*. Scale bar = 1.3 mm.

Figure 114. Ventral view of sternum VIII of female *Lixus (Lixoglyptus) spartii*. Scale bar = 1.3 mm.

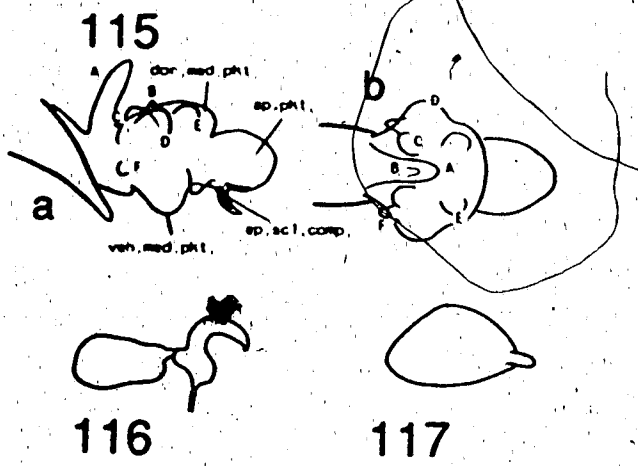
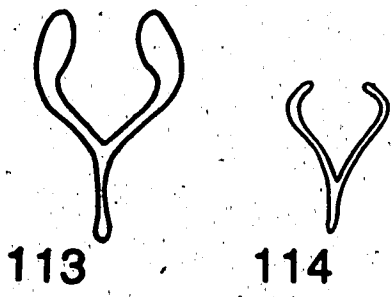
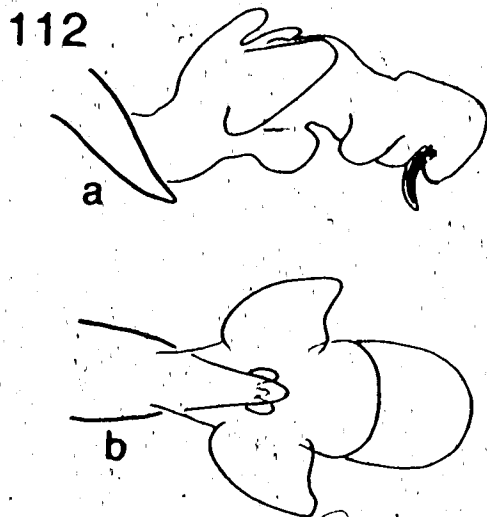
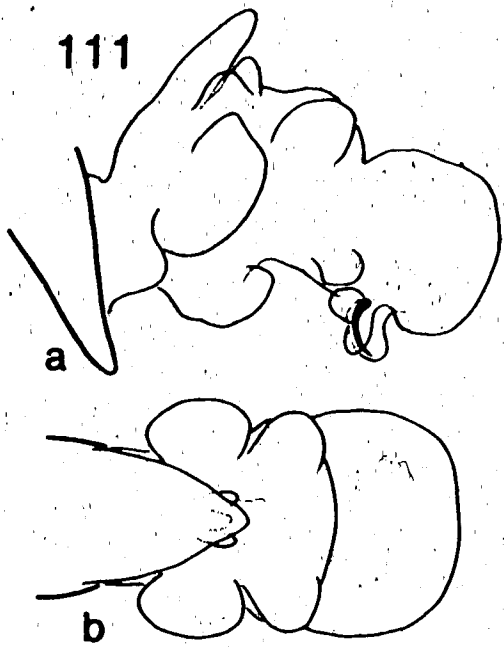
Figure 115. Internal sac of male *Cleonidius*. Lobes and pockets of internal sac labelled as noted (Scale bar = 0.7 mm).

- a. Lateral view.
- b. Dorsal view.

Figure 116. Spermatheca of female *Cleonidius*. Scale bar = 0.3 mm.

Figure 117. Lateral view of gonocoxite II and stylus of *Cleonidius*. Scale bar = 0.7 mm.





Figures 118 to 123.

Figure 118. Dorsal habitus of *Cleonidius poricollis*.

Scale bar = 4.0 mm.

Figure 119. Dorsal habitus of *Cleonidius*

*eustictorrhinus*. Scale bar = 4.0 mm.

Figure 120. Dorsal habitus of *Cleonidius modestus*.

Scale bar = 4.0 mm.

Figure 121. Dorsal habitus of *Cleonidius*

*subcylindricus*. Scale bar = 4.0 mm.

Figure 122. Dorsal habitus of *Cleonidius longinasus*.

Scale bar = 4.0 mm.

Figure 123. Dorsal habitus of *Cleonidius texanus*. Scale

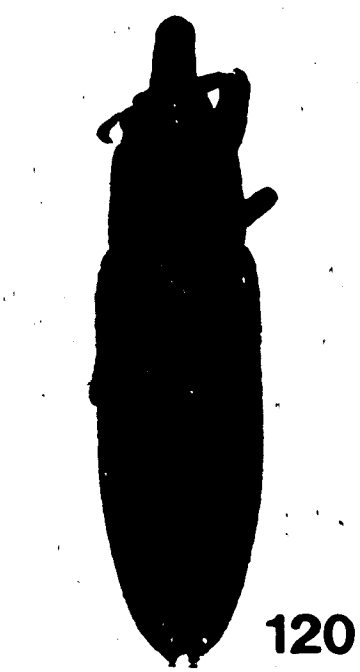
bar = 4.0 mm.



118



119



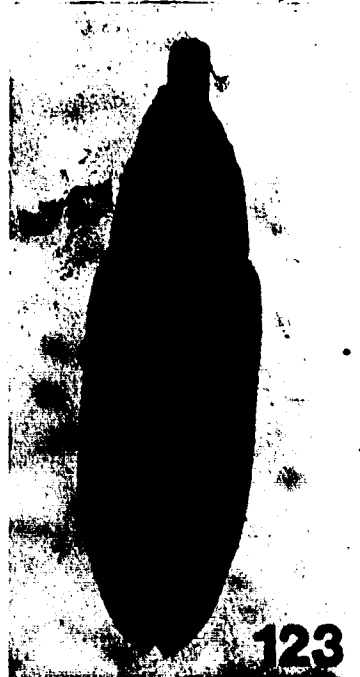
120



121



122



123

Figures 124 to 130.

Figure 124. Dorsal habitus of *Cleonidius americanus*.

Scale bar = 4.0 mm.

Figure 125. Dorsal habitus of *Cleonidius frontalis*.

Scale bar = 4.0 mm.

Figure 126. Dorsal habitus of *Cleonidius canescens*.

Scale bar = 4.0 mm.

Figure 127. Dorsal habitus of *Cleonidius infrequens*.

Scale bar = 4.0 mm.

Figure 128. Dorsal habitus of *Cleonidius puberulus*.

Scale bar = 4.0 mm.

Figure 129. Dorsal habitus of *Cleonidius collaris*.

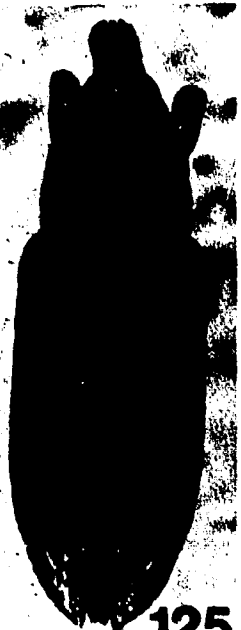
Scale bar = 4.0 mm.

Figure 130. Dorsal habitus of *Cleonidius notolomus*.

Scale bar = 4.0 mm.



124



125



126



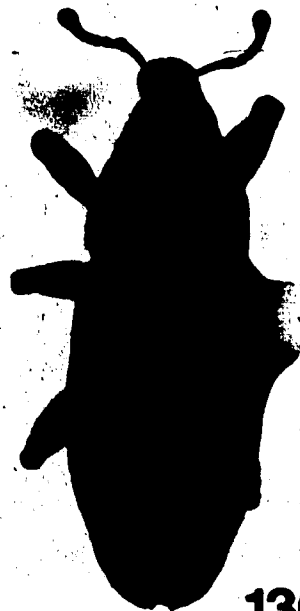
127



128



129



130

Figures 131 to 136.

Figure 131. Dorsal habitus of *Cleonidius lobigerinus*.

Scale bar = 4.0 mm.

Figure 132. Dorsal habitus of *Cleonidius calandroides*.

Scale bar = 4.0 mm.

Figure 133. Dorsal habitus of *Cleonidius boucardi*.

Scale bar = 5.0 mm.

Figure 134. Dorsal habitus of *Cleonidius trivittatus*.

Scale bar = 5.0 mm.

Figure 135. Dorsal habitus of *Cleonidius placidus*.

Scale bar = 5.0 mm.

Figure 136. Dorsal habitus of *Cleonidius*

*quadrilineatus*. Scale bar = 4.0 mm.



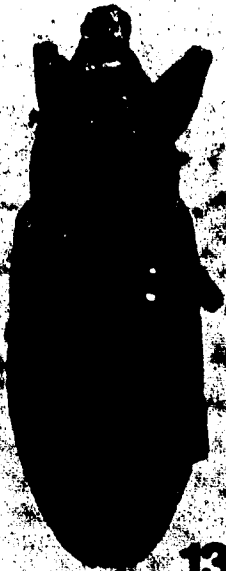
131



132



133



134



135



136

Figures 137 to 142. Head of *Cleonidius* species (Scale bar = 2.3 mm).

Figure 137. *Cleonidius poricollis*.

- a. Lateral view of head.
- b. Dorsal view of head.

Figure 138. *Cleonidius eustictorrhinus*.

- a. Lateral view of head.
- b. Dorsal view of head.

Figure 139. *Cleonidius modestus*.

- a. Lateral view of head.
- b. Dorsal view of head.

Figure 140. *Cleonidius subcylindricus*.

- a. Lateral view of head.
- b. Dorsal view of head.

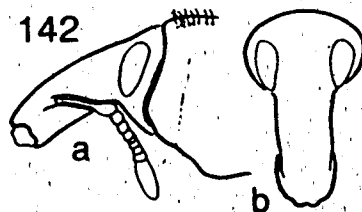
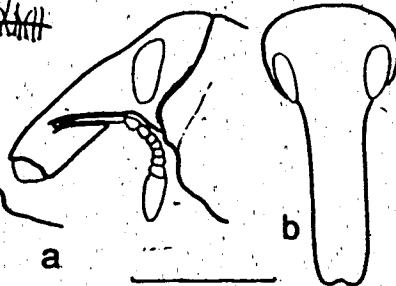
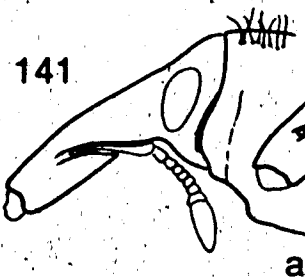
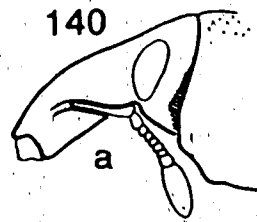
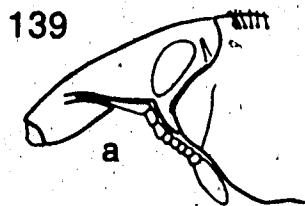
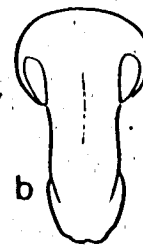
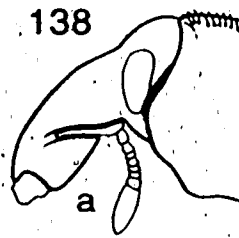
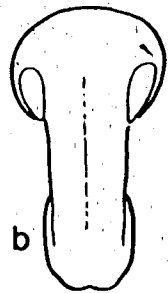
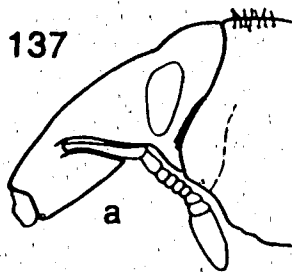
Figure 141. *Cleonidius longinatus*.

- a. Lateral view of heads of male and female.
- b. Dorsal view of head of female.

Figure 142. *Cleonidius texanus*.

- a. Lateral view of head.
- b. Dorsal view of head.





Figures 143 to 149. Head and femur of *Cleonidius* species  
(Scale bar = 2.3 mm).

Figure 143. *Cleonidius americanus*.

- a. Lateral view of head.
- b. Dorsal view of head.
- c. Lateral view of pro-tibia of female.

Figure 144. *Cleonidius frontalis*.

- a. Lateral view of head.
- b. Dorsal view of head.
- c. Lateral view of pro-tibia of female.

Figure 145. *Cleonidius canescens*.

- a. Lateral view of head.
- b. Dorsal view of head.

Figure 146. *Cleonidius infrequens*.

- a. Lateral view of head.
- b. Dorsal view of head.

Figure 147. *Cleonidius puberulus*.

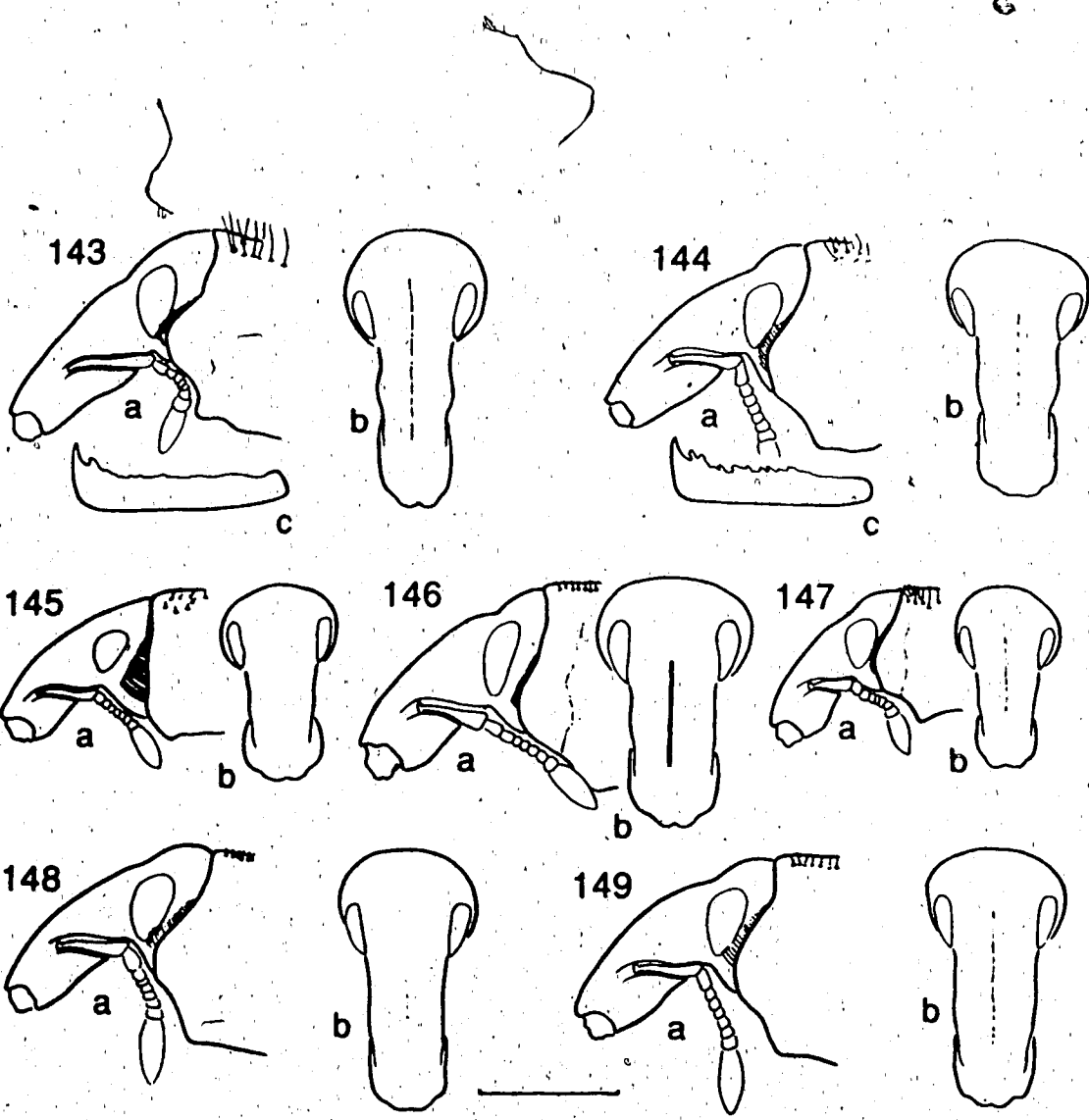
- a. Lateral view of head.
- b. Dorsal view of head.

Figure 148. *Cleonidius collaris*.

- a. Lateral view of head.
- b. Dorsal view of head.

Figure 149. *Cleonidius notolomus*.

- a. Lateral view of head.
- b. Dorsal view of head.



Figures 150 to 156. Head of *Cleonidius* species (Scale bar = 2.3 mm).

Figure 150. *Cleonidius lobigerinus*.

- a. Lateral view of head.
- b. Dorsal view of head.

Figure 151. *Cleonidius calandroides*.

- a. Lateral view of head.
- b. Dorsal view of head.

Figure 152. *Cleonidius boucardi*.

- a. Lateral view of head.
- b. Dorsal view of head.

Figure 153. *Cleonidius trivittatus*.

- a. Lateral view of head.
- b. Dorsal view of head.

Figure 154. *Cleonidius placidus*.

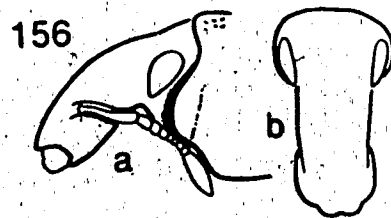
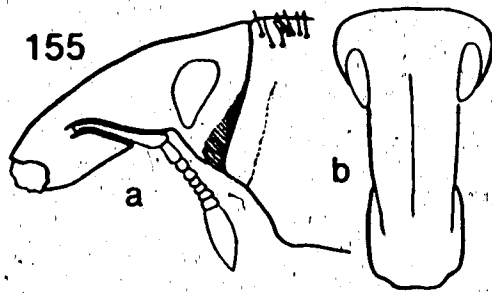
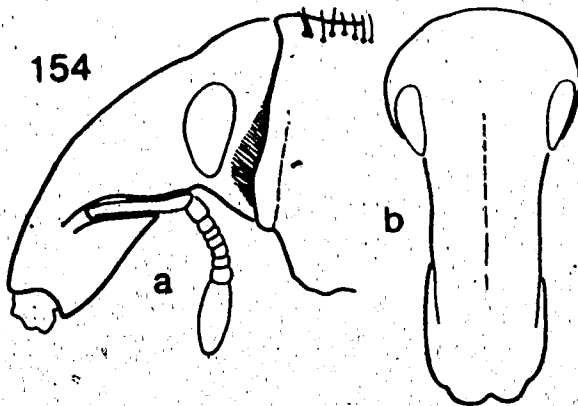
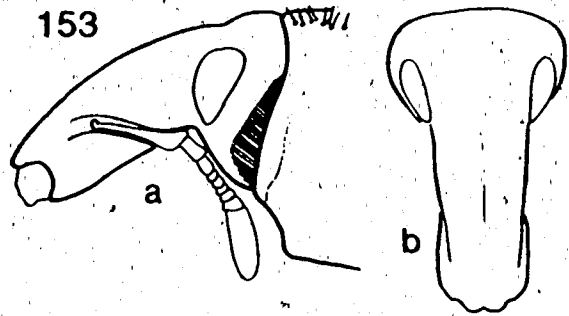
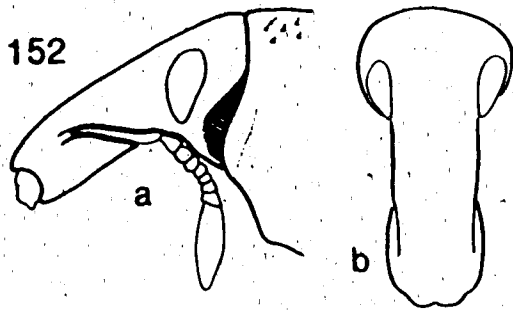
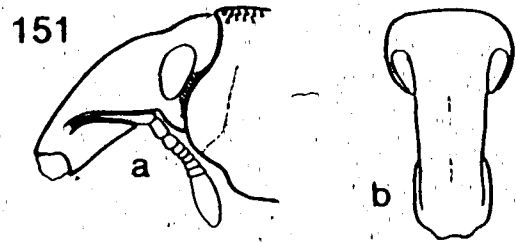
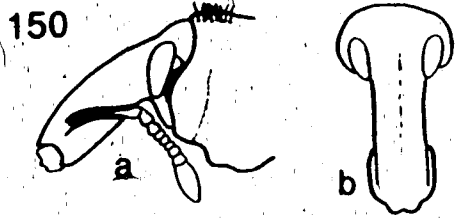
- a. Lateral view of head.
- b. Dorsal view of head.

Figure 155. *Cleonidius quadrilineatus*.

- a. Lateral view of head.
- b. Dorsal view of head.

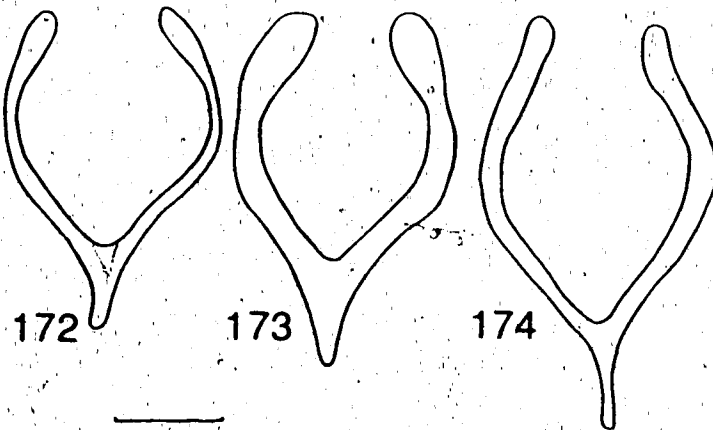
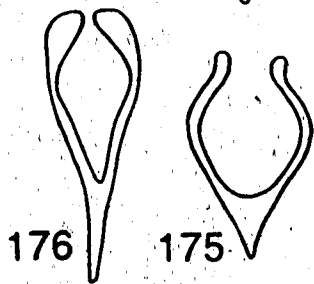
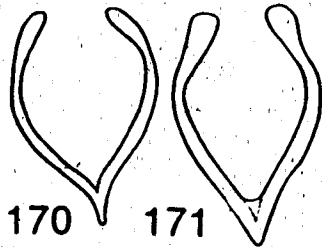
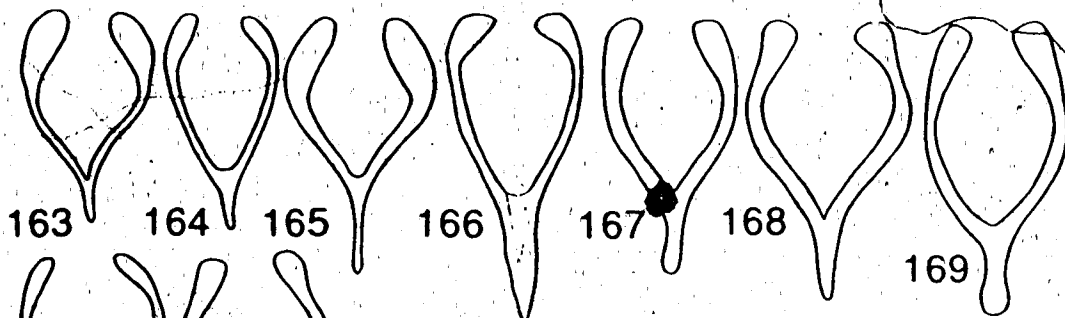
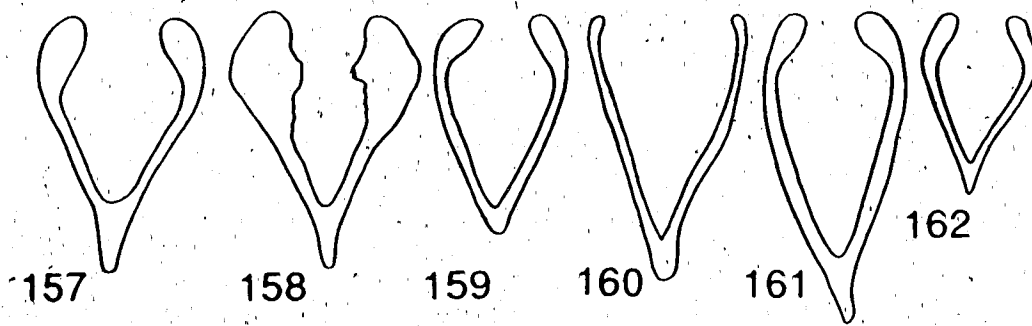
Figure 156. *Cleonidius vibex*.

- a. Lateral view of head.
- b. Dorsal view of head.



Figures 157 to 176. Ventral view of sternum VIII of female  
*Cleonidius* species (Scale bar = 0.8 mm).

- Figure 157. *Cleonidius porlcollis*.
- Figure 158. *Cleonidius eustictorrhinus*.
- Figure 159. *Cleonidius modestus*.
- Figure 160. *Cleonidius subcylindricus*.
- Figure 161. *Cleonidius longinasus*.
- Figure 162. *Cleonidius texanus*.
- Figure 163. *Cleonidius americanus*.
- Figure 164. *Cleonidius frontalis*.
- Figure 165. *Cleonidius canescens*.
- Figure 166. *Cleonidius infrequens*.
- Figure 167. *Cleonidius puberulus*.
- Figure 168. *Cleonidius notolomus*.
- Figure 169. *Cleonidius collaris*.
- Figure 170. *Cleonidius lobigerinus*.
- Figure 171. *Cleonidius calandroides*.
- Figure 172. *Cleonidius boucardi*.
- Figure 173. *Cleonidius trivittatus*.
- Figure 174. *Cleonidius placidus*.
- Figure 175. *Cleonidius quadrilineatus*.
- Figure 176. *Cleonidius vibex*.



Figures 177 to 182. Male genitalia of *Cleonidius* species.

Figure 177. *Cleonidius poricollis*.

- a. Lateral view of internal sac of male. Scale bar = 0.7 mm.
- b. Dorsal view of internal sac of male. Scale bar = 0.7 mm.
- c. Lateral view of aedeagus of male. Scale bar = 1.3 mm.

Figure 178. *Cleonidius eustictorrhinus*. Scale bars as in Fig. 177.

- a. Lateral view of internal sac of male.
- b. Dorsal view of internal sac of male.
- c. Lateral view of aedeagus of male.

Figure 179. *Cleonidius modestus*. Scale bars as in Fig. 177.

- a. Lateral view of internal sac of male.
- b. Dorsal view of internal sac of male.
- c. Lateral view of aedeagus of male.

Figure 180. *Cleonidius subcylindricus*. Scale bars as in Fig. 177.

- a. Lateral view of internal sac of male.
- b. Dorsal view of internal sac of male.
- c. Lateral view of aedeagus of male.

Figure 181. *Cleonidius longinasus*. Scale bars as in Fig. 177.

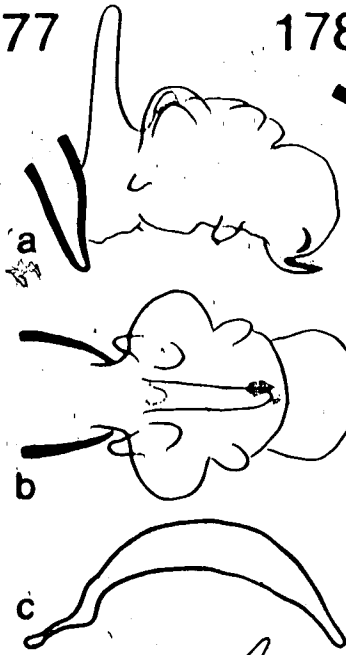
- a. Lateral view of internal sac of male.
- b. Dorsal view of internal sac of male.
- c. Lateral view of aedeagus of male.

Figure 182. *Cleonidius texanus*. Scale bars as in Fig. 177.

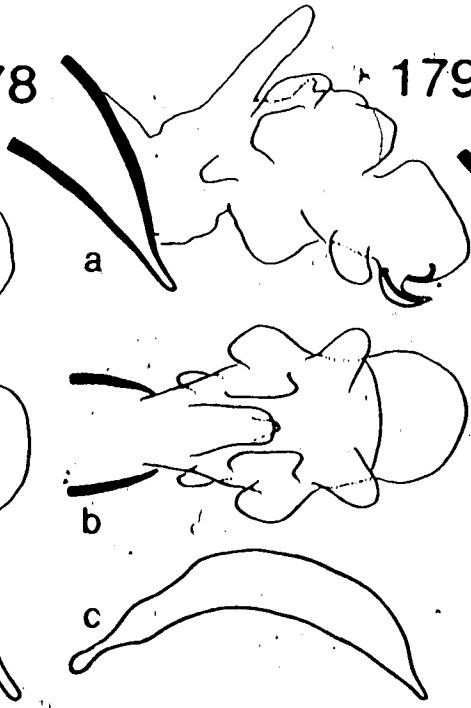
- a. Lateral view of internal sac of male.
- b. Dorsal view of internal sac of male.
- c. Lateral view of aedeagus of male.



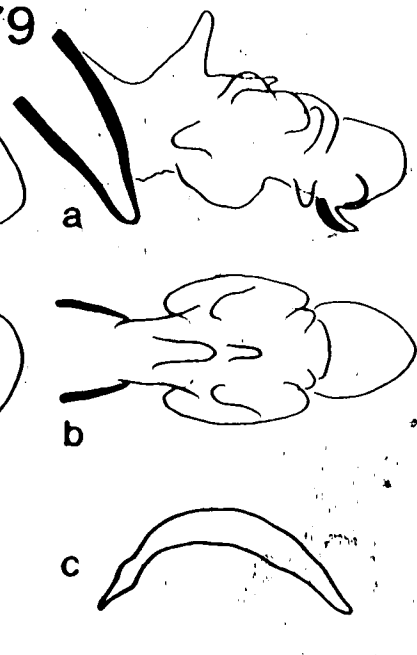
177



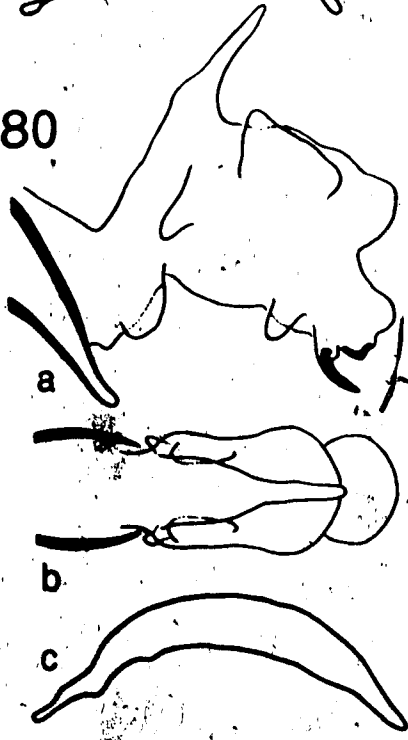
178



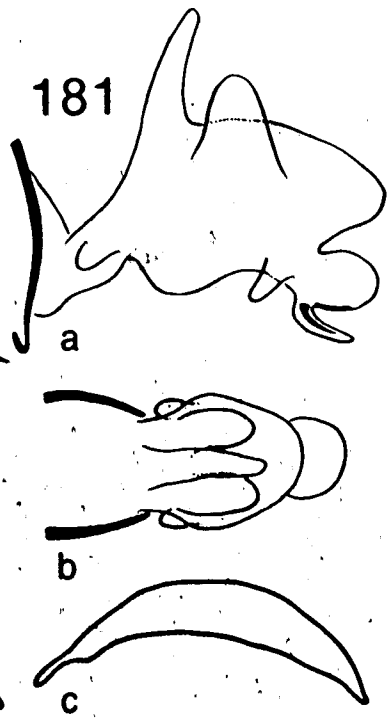
179



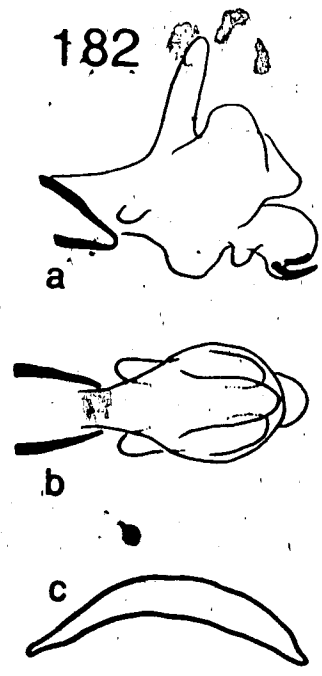
180



181



182



Figures 183 to 189. Male genitalia of *Cleonidius* species.

Figure 183. *Cleonidius americanus*.

- a. Lateral view of internal sac of male. Scale bar = 0.7 mm.
- b. Dorsal view of internal sac of male. Scale bar = 0.7 mm.
- c. Lateral view of aedeagus of male. Scale bar = 1.3 mm.

Figure 184. *Cleonidius frontalis*. Scale bars as in Fig. 183.

- a. Lateral view of internal sac of male.
- b. Dorsal view of internal sac of male.
- c. Lateral view of aedeagus of male.

Figure 185. *Cleonidius canescens*. Scale bars as in Fig. 183.

- a. Lateral view of internal sac of male.
- b. Dorsal view of internal sac of male.
- c. Lateral view of aedeagus of male.

Figure 186. *Cleonidius infrequens*. Scale bars as in Fig. 183.

- a. Lateral view of internal sac of male.
- b. Dorsal view of internal sac of male.
- c. Lateral view of aedeagus of male.

Figure 187. *Cleonidius puberulus*. Scale bars as in Fig. 183.

- a. Lateral view of internal sac of male.
- b. Dorsal view of internal sac of male.
- c. Lateral view of aedeagus of male.

Figure 188. *Cleonidius notolomus*. Scale bars as in Fig. 183.

- a. Lateral view of internal sac of male.
- b. Dorsal view of internal sac of male.
- c. Lateral view of aedeagus of male.

Figure 189. *Cleonidius collaris*. Scale bars as in Fig. 183.

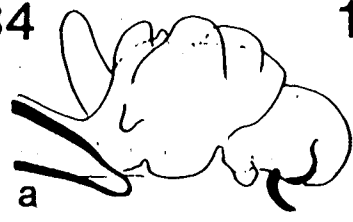
- a. Lateral view of internal sac of male.
- b. Dorsal view of internal sac of male.
- c. Lateral view of aedeagus of male.

183



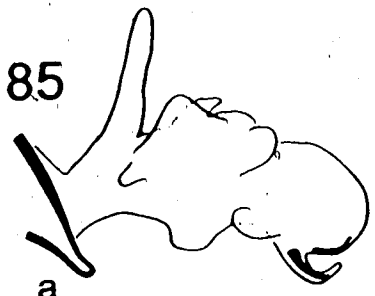
a

184

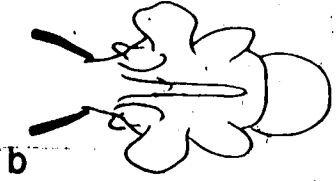


a

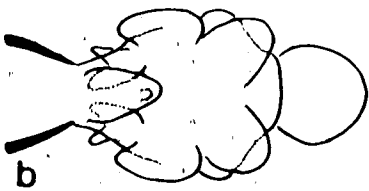
185



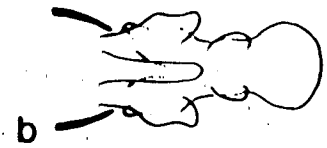
a



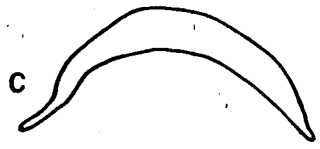
b



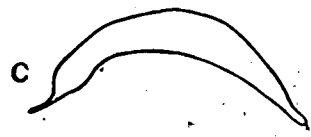
b



b



c

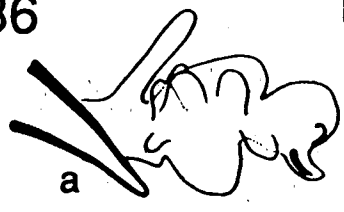


c



c

186



a

187



a

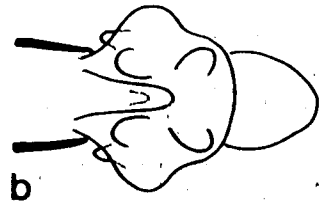
188



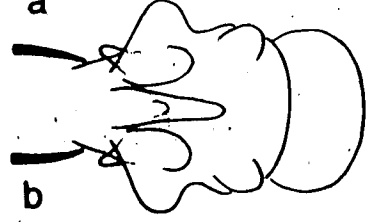
a



b



b



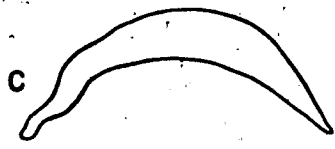
b



c

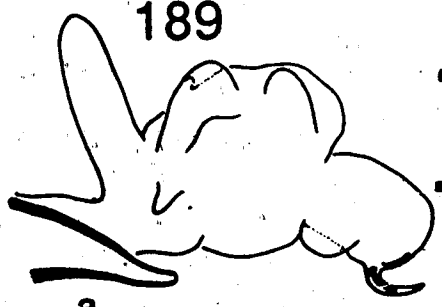


c

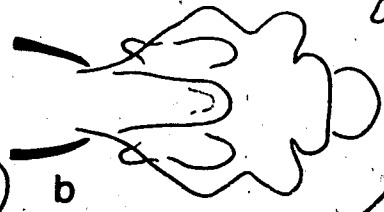


c

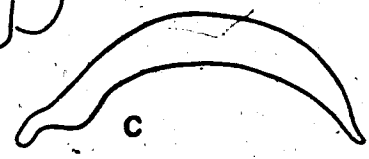
189



a



b



c

Figures 190 to 196. Male genitalia of *Cleonidius* species.

Figure 190. *Cleonidius lobigerinus*.

- a. Lateral view of internal sac of male. Scale bar = 0.7 mm.
- b. Dorsal view of internal sac of male. Scale bar = 0.7 mm.
- c. Lateral view of aedeagus of male. Scale bar = 1.3 mm.

Figure 191. *Cleonidius calandroides*. Scale bars as in Fig. 190.

- a. Lateral view of internal sac of male.
- b. Dorsal view of internal sac of male.
- c. Lateral view of aedeagus of male.

Figure 192. *Cleonidius boucardi*. Scale bars as in Fig. 190.

- a. Lateral view of internal sac of male.
- b. Dorsal view of internal sac of male.
- c. Lateral view of aedeagus of male.

Figure 193. *Cleonidius trivittatus*. Scale bars as in Fig. 190.

- a. Lateral view of internal sac of male.
- b. Dorsal view of internal sac of male.
- c. Lateral view of aedeagus of male.

Figure 194. *Cleonidius placidus*. Scale bars as in Fig. 190.

- a. Lateral view of internal sac of male.
- b. Dorsal view of internal sac of male.

Figure 195. *Cleonidius quadrilineatus*. Scale bars as in Fig. 190.

- a. Lateral view of internal sac of male.
- b. Dorsal view of internal sac of male.
- c. Lateral view of aedeagus of male.

Figure 196. *Cleonidius vibex*. Scale bars as in Fig. 190.

- a. Lateral view of internal sac of male.
- b. Dorsal view of internal sac of male.
- c. Lateral view of aedeagus of male.

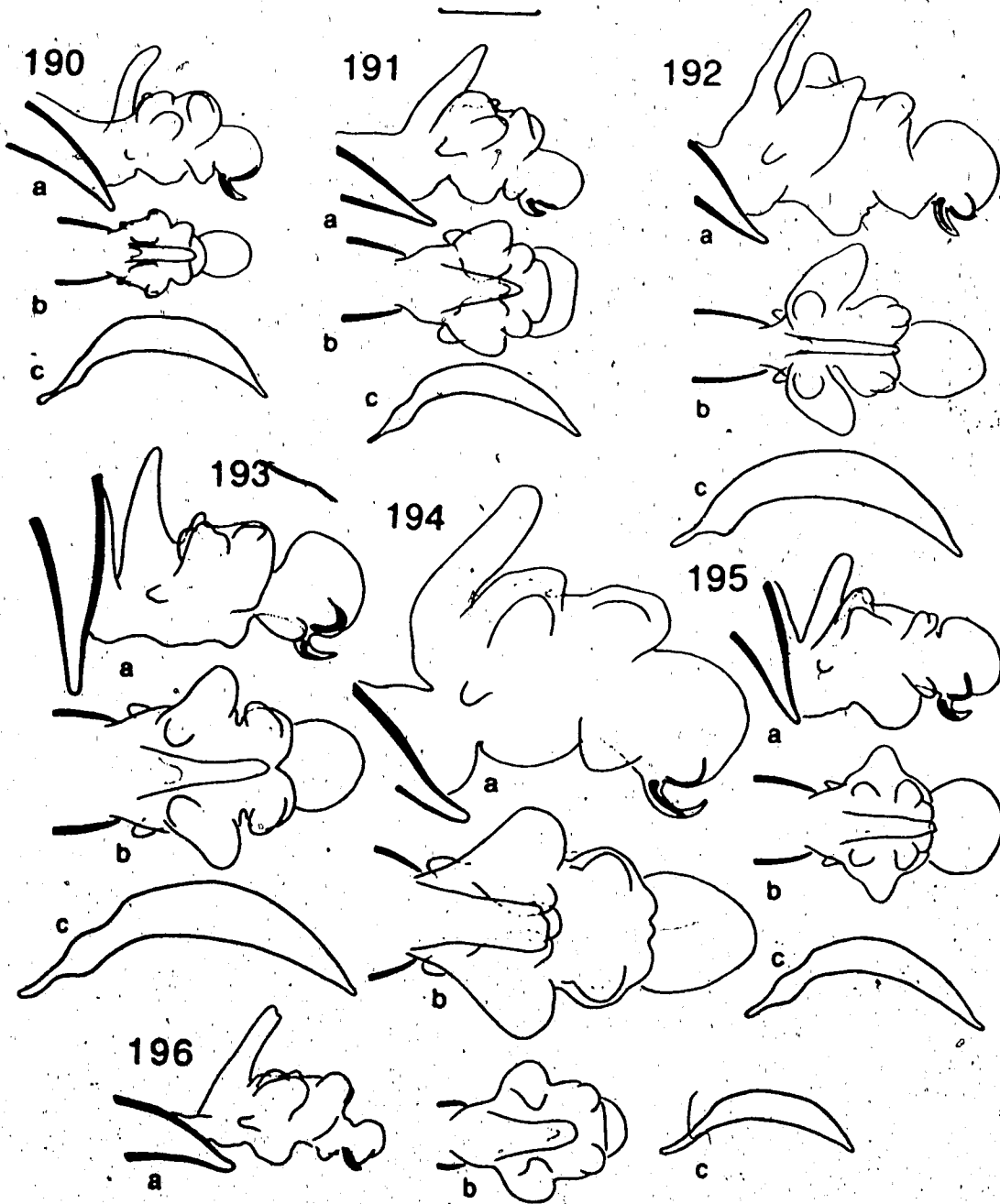


Figure 197. Hubbs-Hubbs diagram illustrating variation among specimens of *Cleonidius* species; length of elytra (LE1): a, males; b, females. See caption for Figure 49 for explanation.

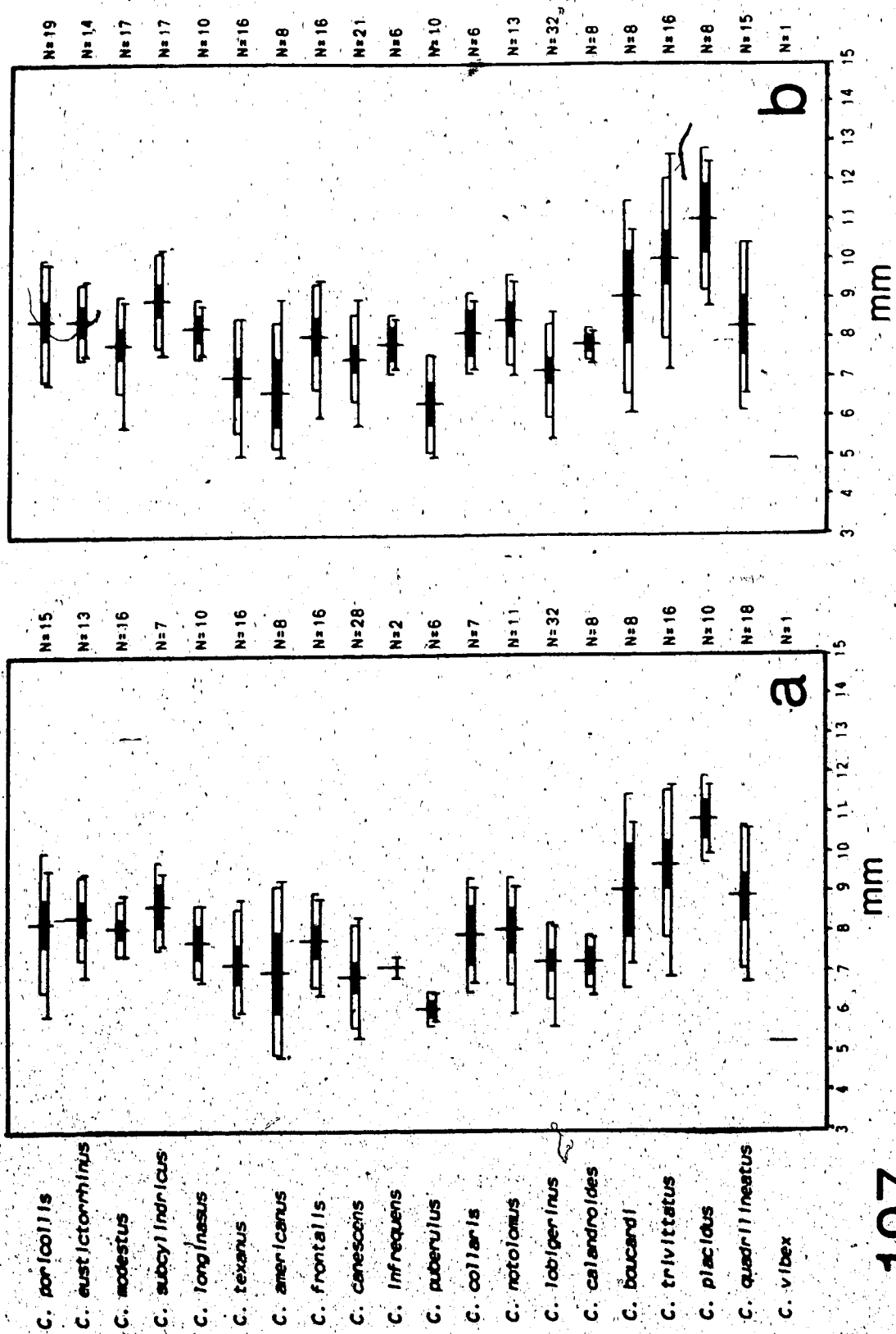


Figure 198. Hubbs-Hubbs diagram illustrating variation among specimens of *Cleonidius* species; form of rostrum (WRA/LR): a, males; b, females. See caption for Figure 49 for explanation.



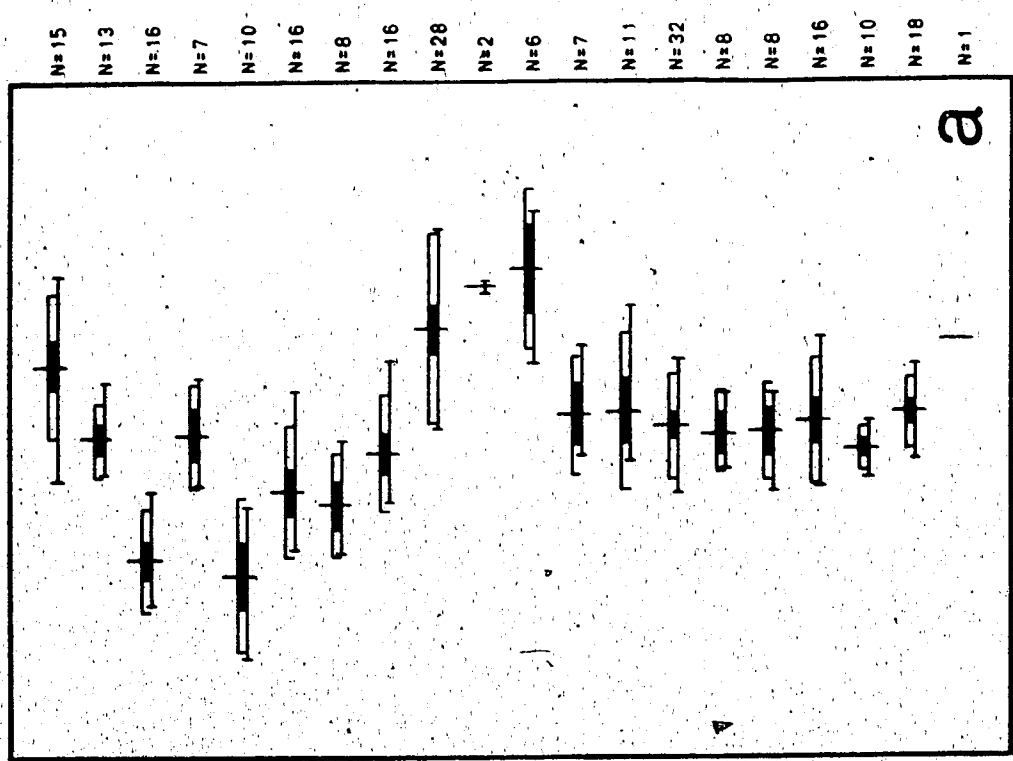
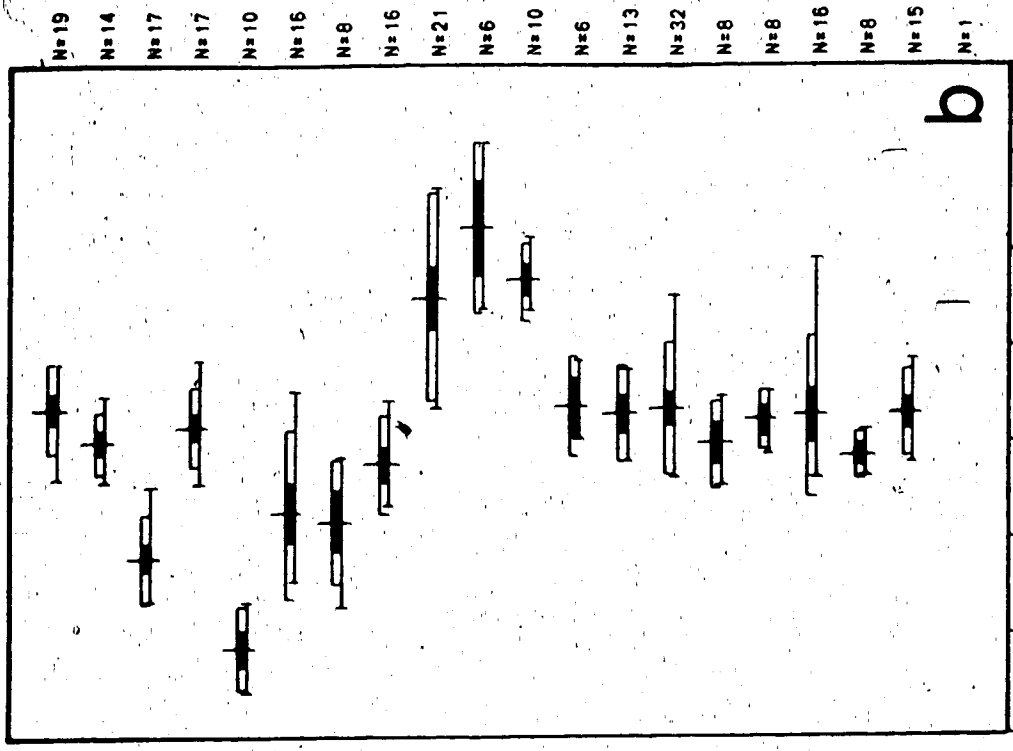
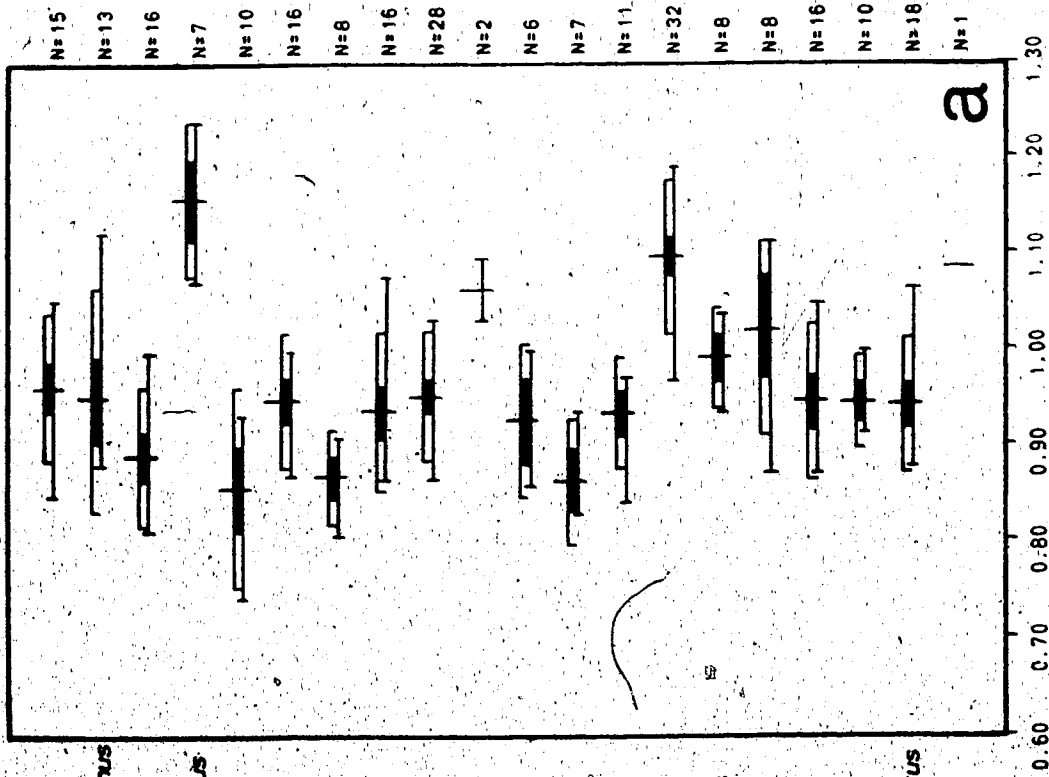
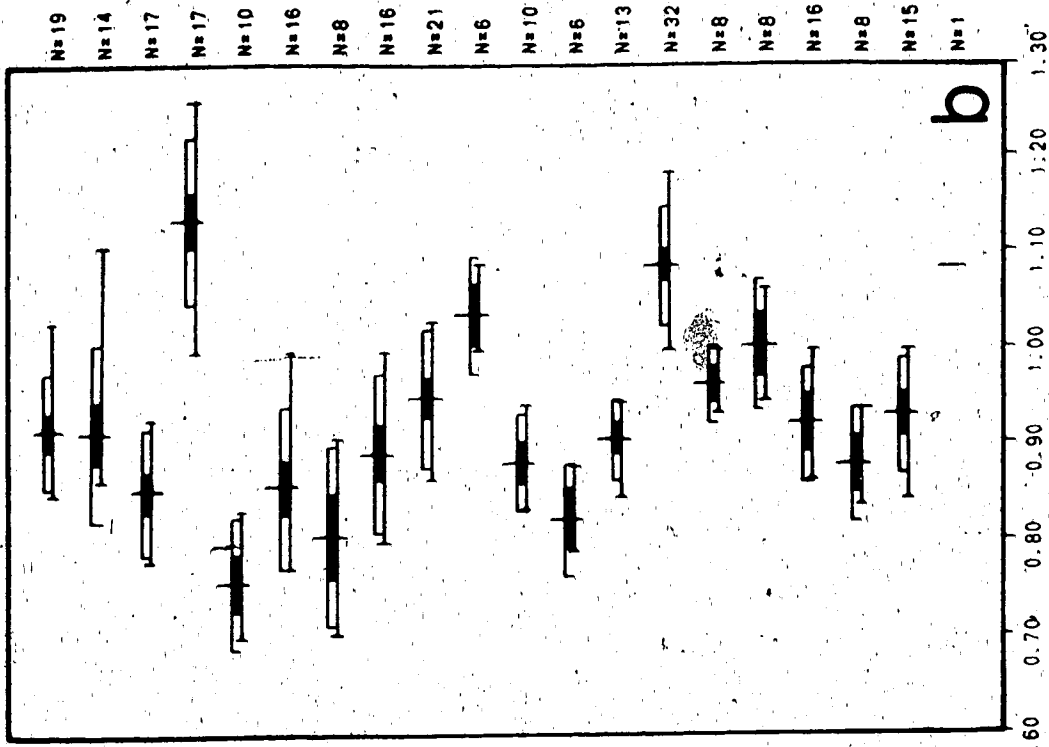


Figure 199. Hubbs-Hubbs diagram illustrating variation among specimens of *Cleonidius* species; width frons compared to width at apex of rostrum (WRA/WF): a, males; b, females. See caption for Figure 49 for explanation.

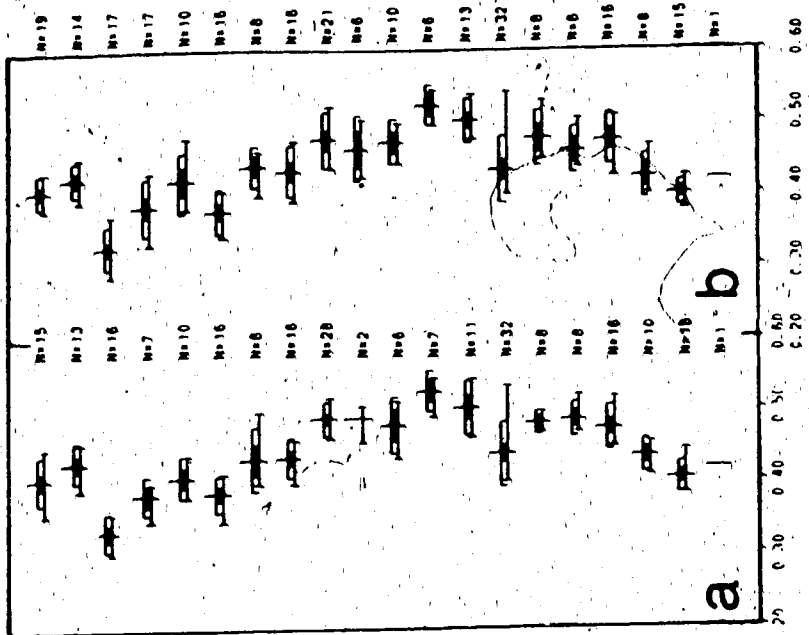


- C. porcolliis*
- C. eustictorhinus*
- C. modestus*
- C. subcylindricus*
- C. longinasus*
- C. texanus*
- C. americanus*
- C. frontalis*
- C. canescens*
- C. infrequens*
- C. puberulus*
- C. collaris*
- C. notolarius*
- C. lobigerinus*
- C. calandroides*
- C. boucardi*
- C. trivittatus*
- C. placidus*
- C. quadrilineatus*
- C. vibex*

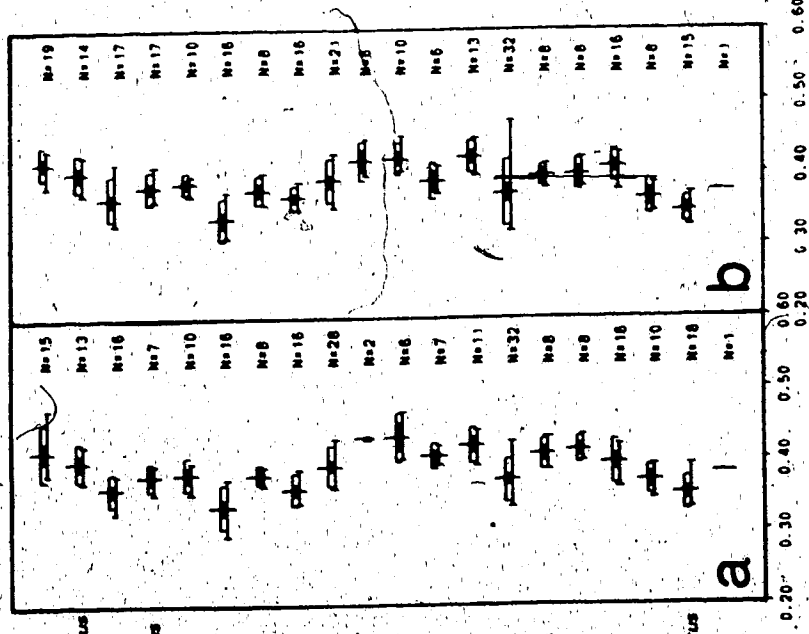
Figures 200 to 201.

Figure 200. Hubbs-Hubbs diagram illustrating variation among specimens of *Cleonidius* species; length pronotum compared to length of elytra (LP/LE1): a, males; b, females. See caption for Figure 49 for explanation.

Figure 201. Hubbs-Hubbs diagram illustrating variation among specimens of *Cleonidius* species; form of elytra (WE1M/LE1): a, males; b, females. See caption for Figure 49 for explanation.



201



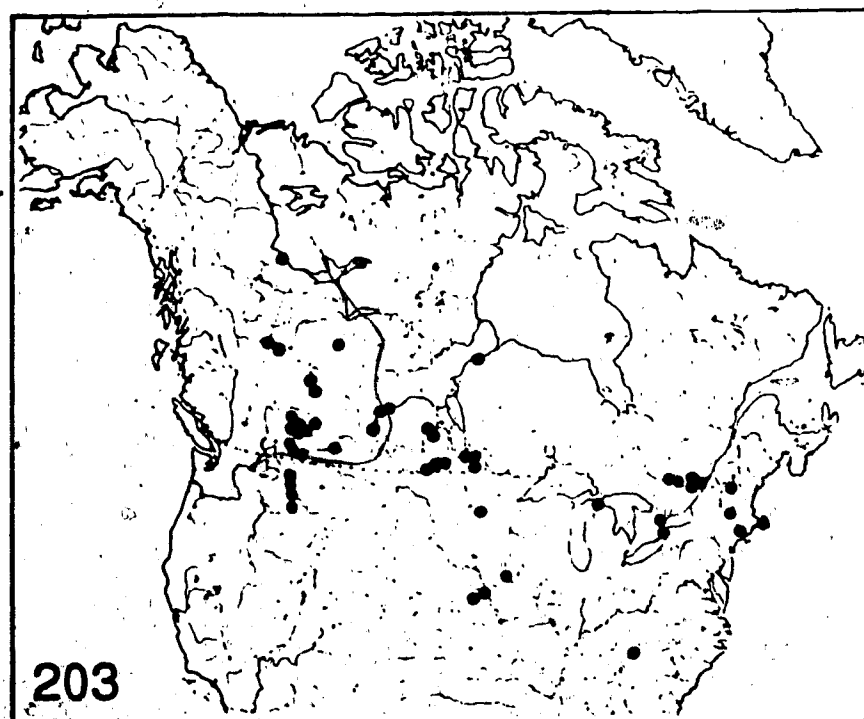
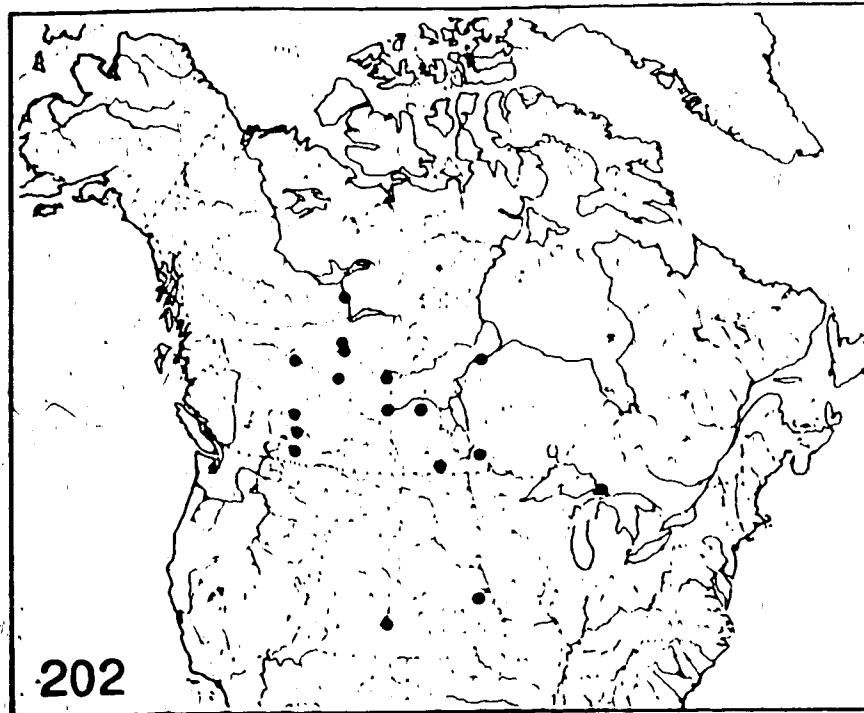
200

- C. parvicolis*
- C. eustictomus*
- C. modestus*
- C. subcylindricus*
- C. longinervis*
- C. texanus*
- C. americanus*
- C. frontalis*
- C. canescens*
- C. infrequens*
- C. subannulus*
- C. collaris*
- C. nototomus*
- C. jobigerinus*
- C. callanovoides*
- C. boucardi*
- C. trivittatus*
- C. placidus*
- C. quadrilineatus*
- C. vibax*

Figures 202 to 203.

Figure 202. Collection localities for *Stephanocleonus*  
*immaculatus*.

Figure 203. Collection localities for *Stephanocleonus*  
*parshus*.

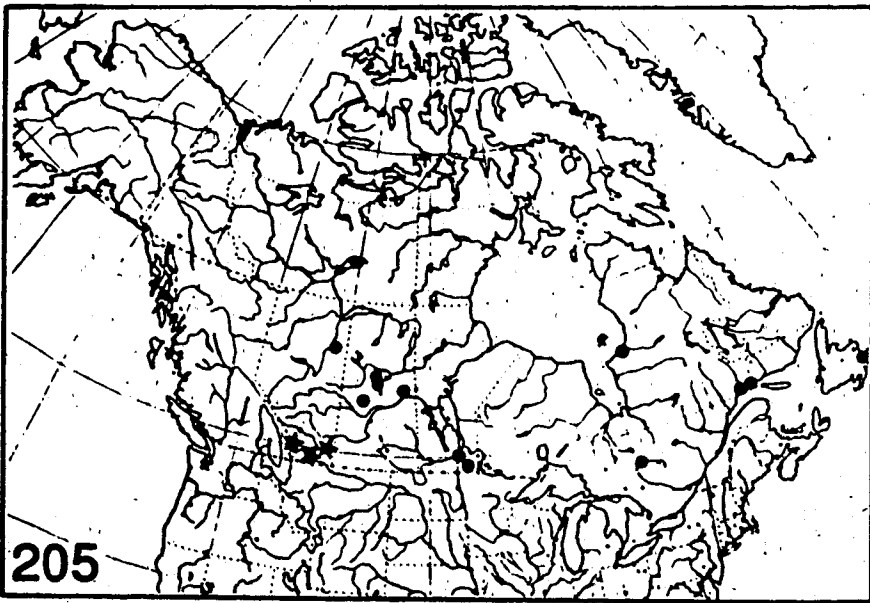
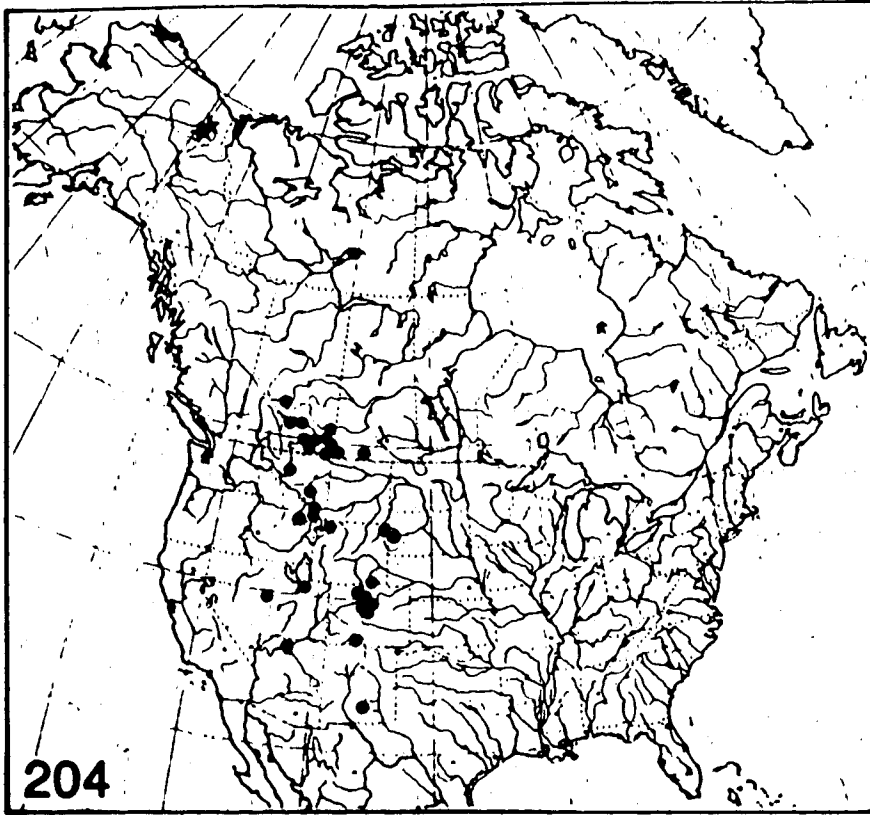


Figures 204 to 205.

Figure 204. Collection localities for *Stephanocleonus confusus* (dots) and *Stephanocleonus stenothorax* (stars).

Figure 205. Collection localities for *Stephanocleonus plumbeus* (dots) and *Stephanocleonus cristaticollis* (stars).





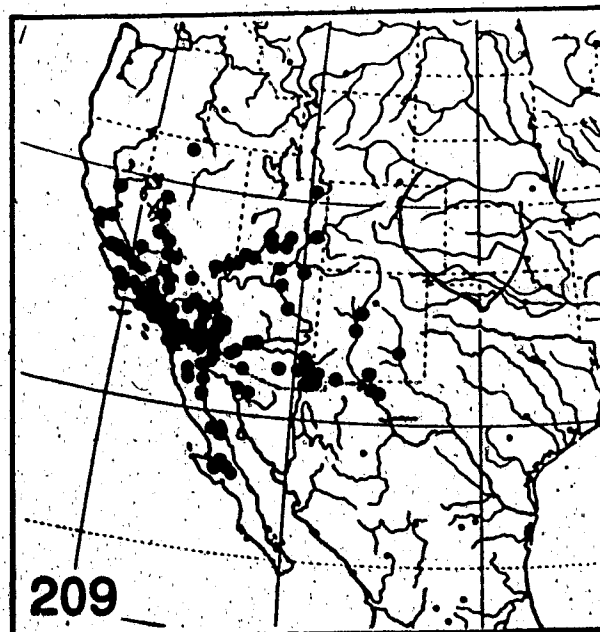
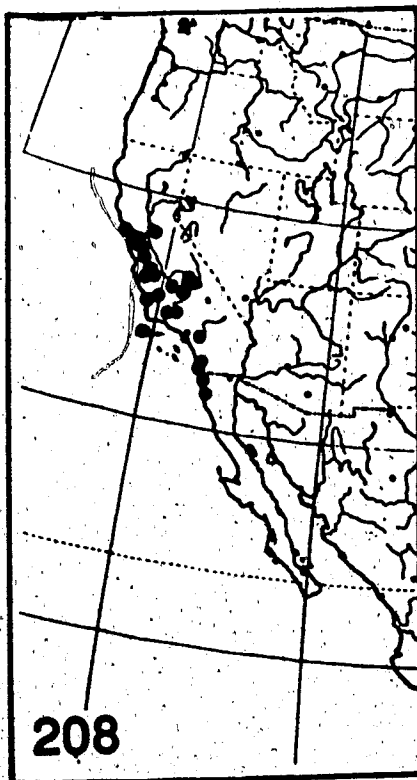
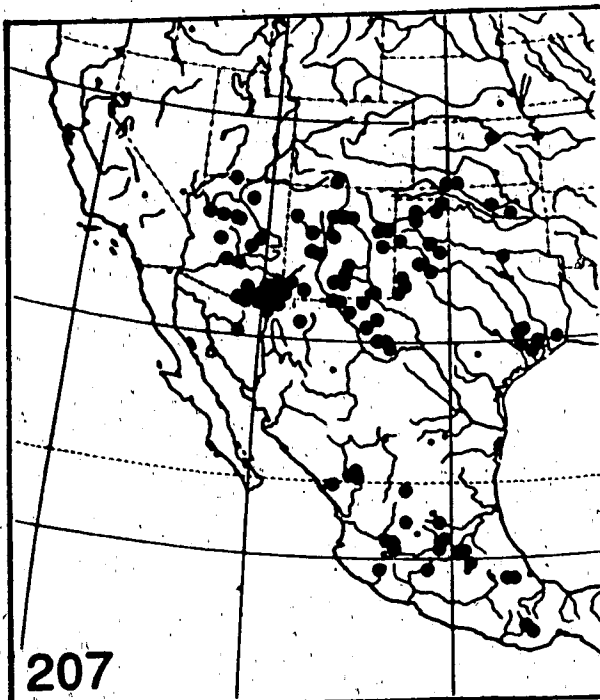
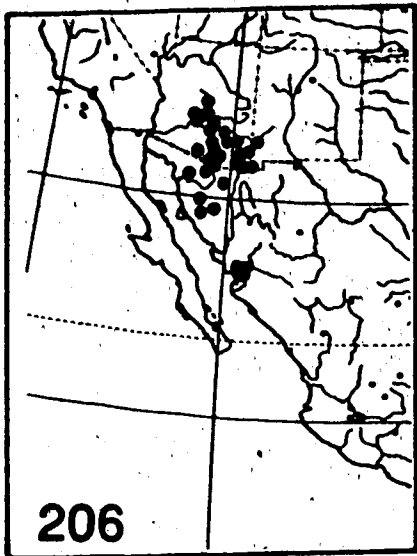
Figures 206 to 209.

Figure 206. Collection localities for *Apleurus saginatus*.

Figure 207. Collection localities for *Apleurus lutulentus*.

Figure 208. Collection localities for *Apleurus jacobinus*.

Figure 209. Collection localities for *Apleurus albovestitus*.



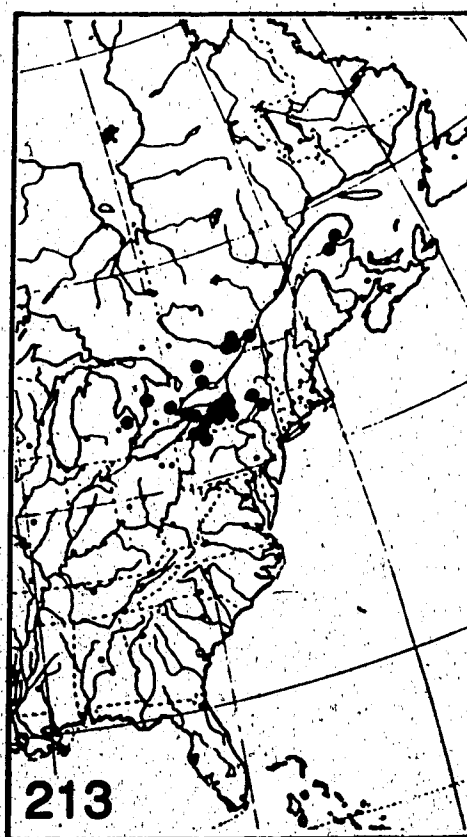
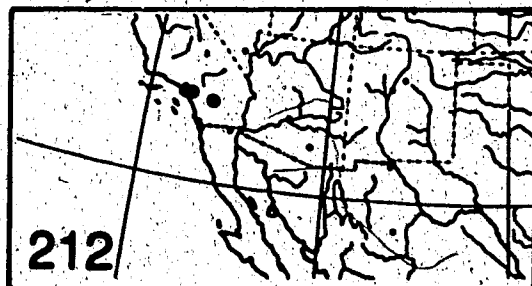
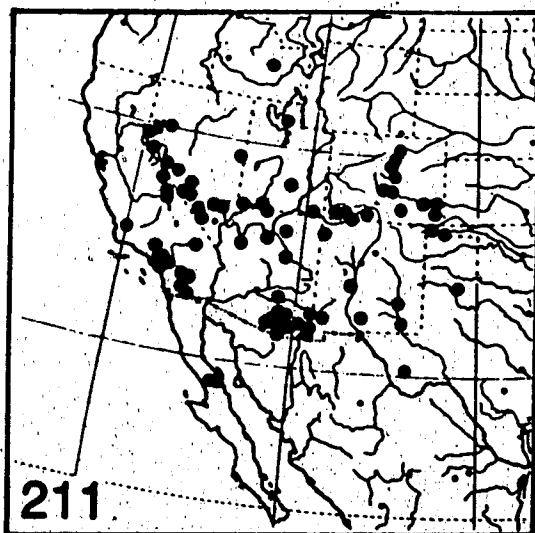
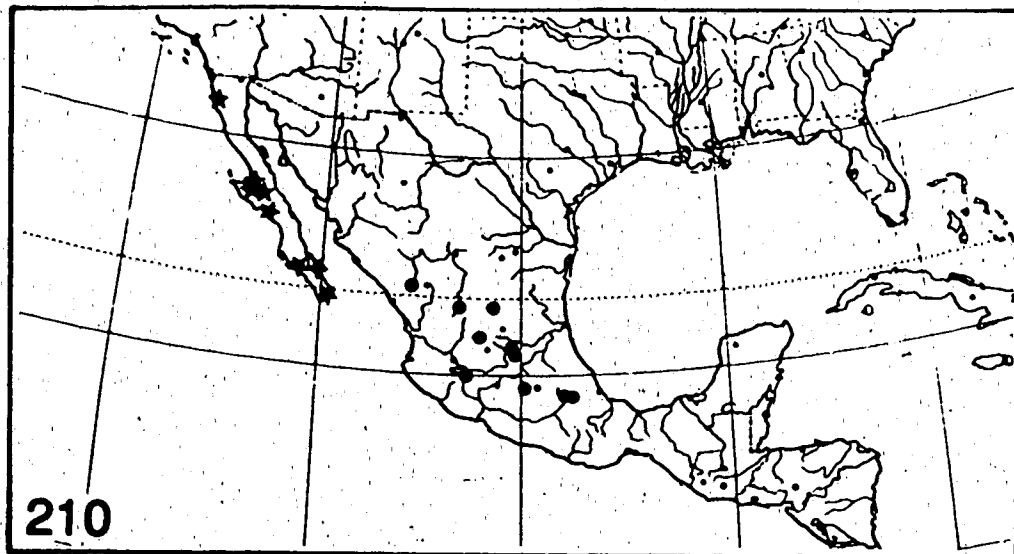
Figures 210 to 213.

Figure 210. Collection localities for *Apleurus aztecus* (dots) and *Apleurus porosus* (stars).

Figure 211. Collection localities for *Apleurus angularis*.

Figure 212. Collection localities for *Apleurus hystrix*.

Figure 213. Collection localities for *Cleonis plura*.



Figures 214 to 219.

Figure 214. Collection localities for *Cleonidius longinasus*.

Figure 215. Collection localities for *Cleonidius subcylindricus*.

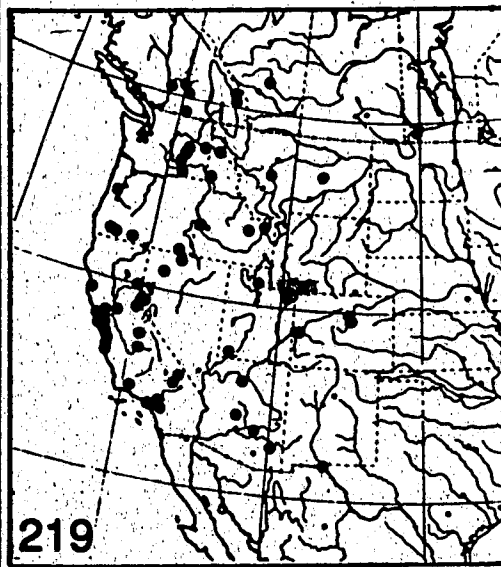
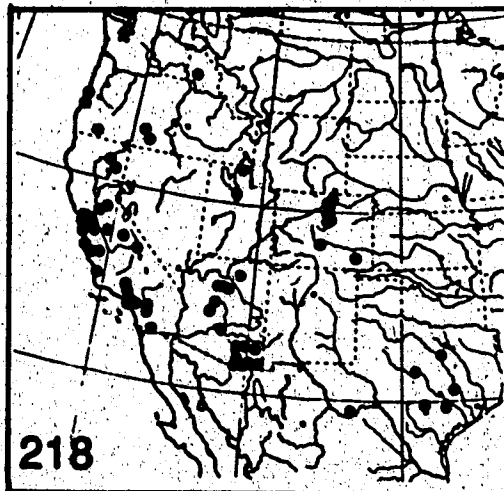
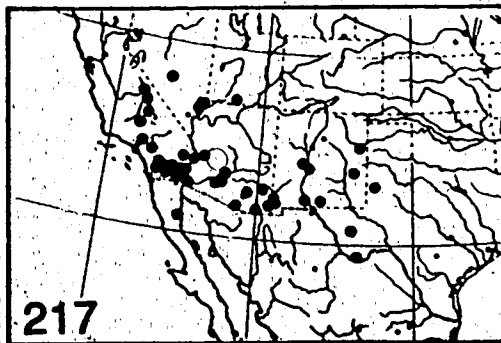
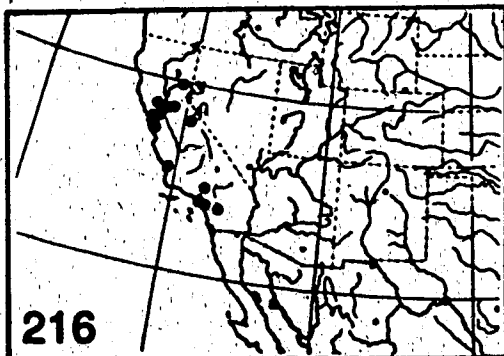
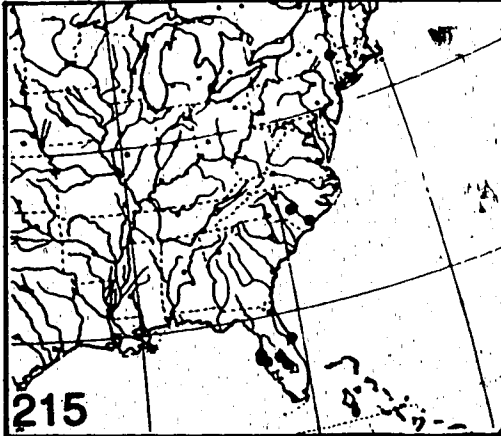
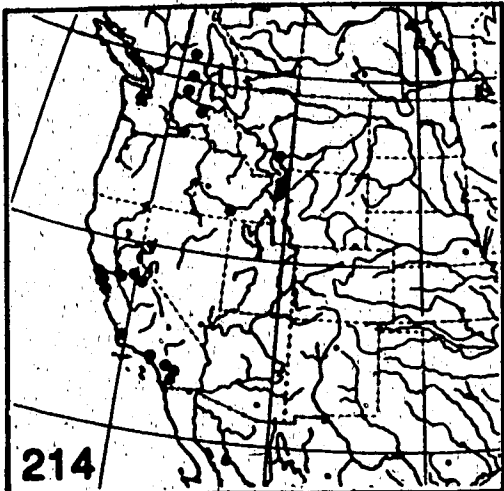
Figure 216. Collection localities for *Cleonidius eustictorrhinus*.

Figure 217. Collection localities for *Cleonidius modestus*.

---

Figure 218. Collection localities for *Cleonidius texanus*.

Figure 219. Collection localities for *Cleonidius poricollis*.



Figures 220 to 225.

Figure 220. Collection localities for *Cleonidius trivittatus*.

Figure 221. Collection localities for *Cleonidius boucardi*.

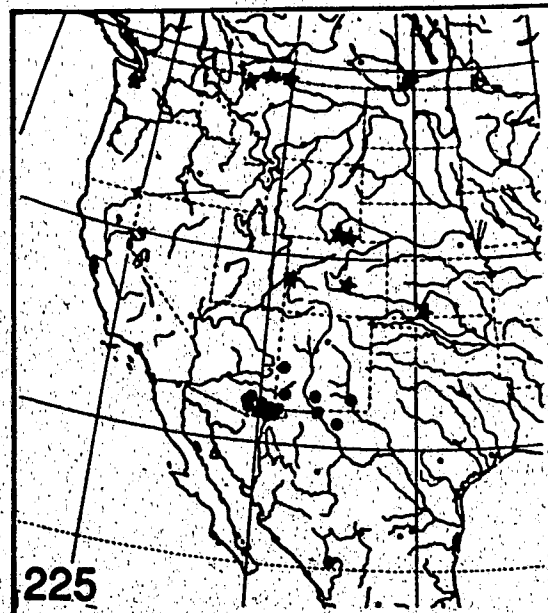
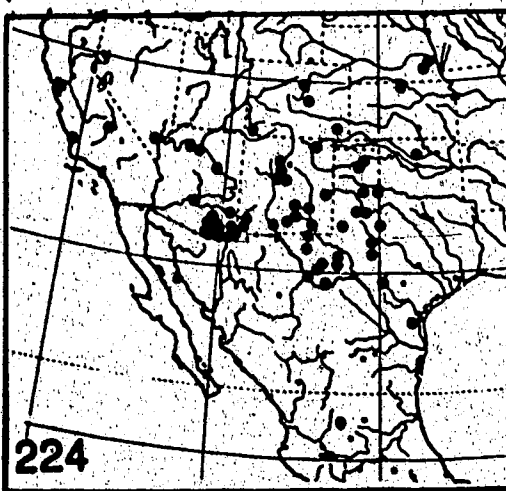
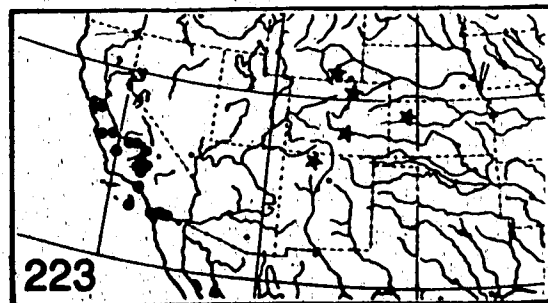
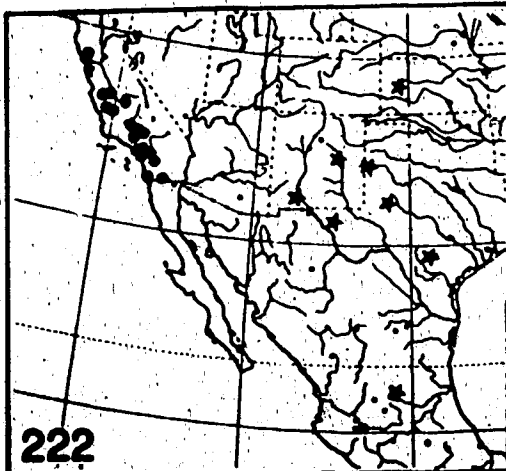
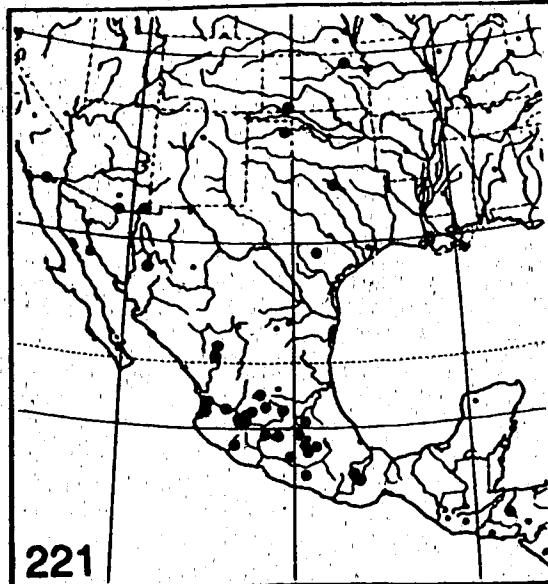
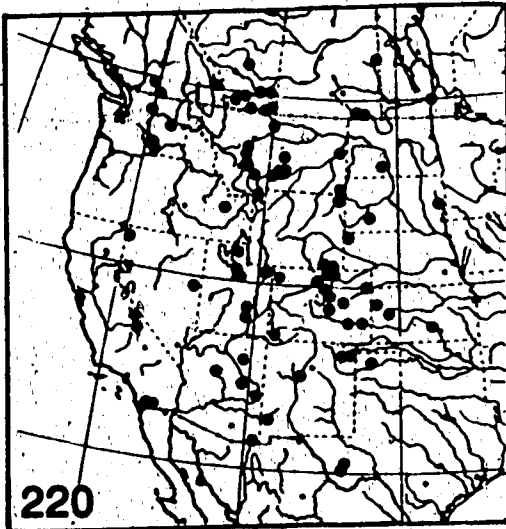
Figure 222. Collection localities for *Cleonidius placidus* (dots) and *Cleonidius infrequens* (stars).

Figure 223. Collection localities for *Cleonidius americanus* (dots) and *Cleonidius collaris* (stars).

Figure 224. Collection localities for *Cleonidius quadrilineatus*.

Figure 225. Collection localities for *Cleonidius notolomus* (dots) and *Cleonidius puberulus* (stars).





Figures 226 to 229.

Figure 226. Collection localities for *Cleonidius frontalis*.

Figure 227. Collection localities for *Cleonidius canescens*.

Figure 228. Collection localities for *Cleonidius lobigerinus*.

Figure 229. Collection localities for *Cleonidius calandroides*.

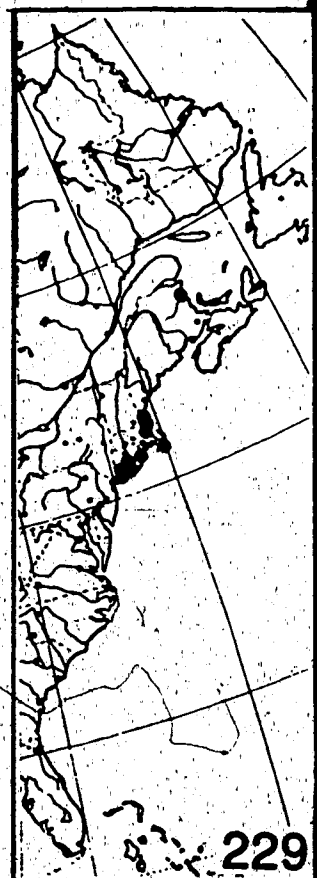
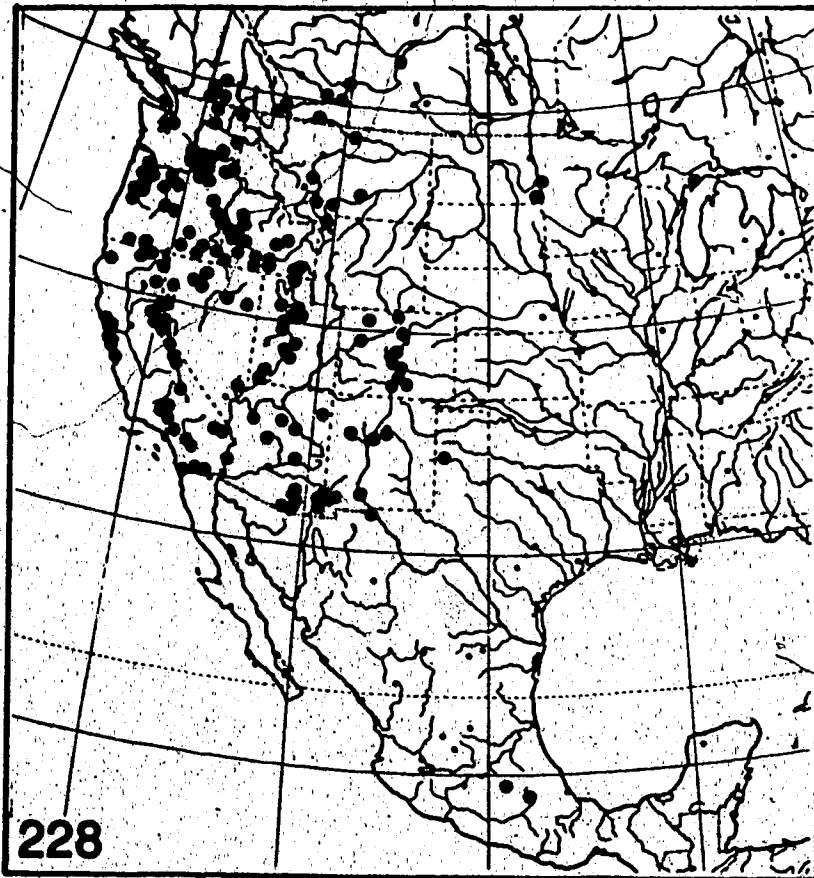
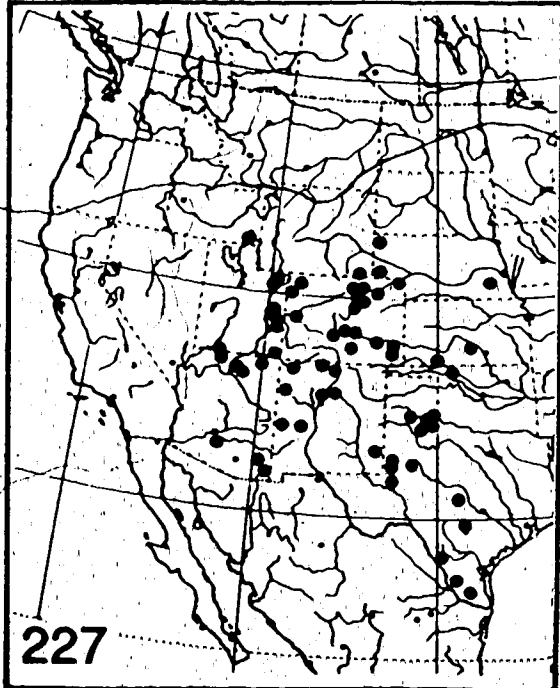
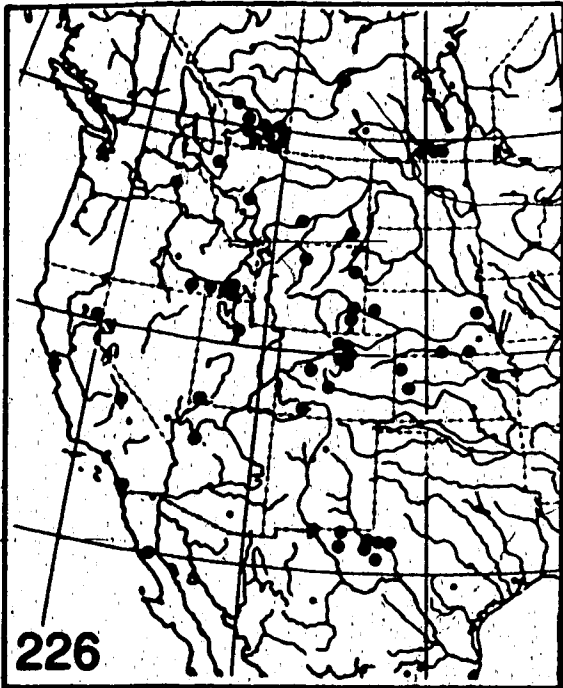


Figure 230. Numbers of species of *Apleurus* in 5° intervals.

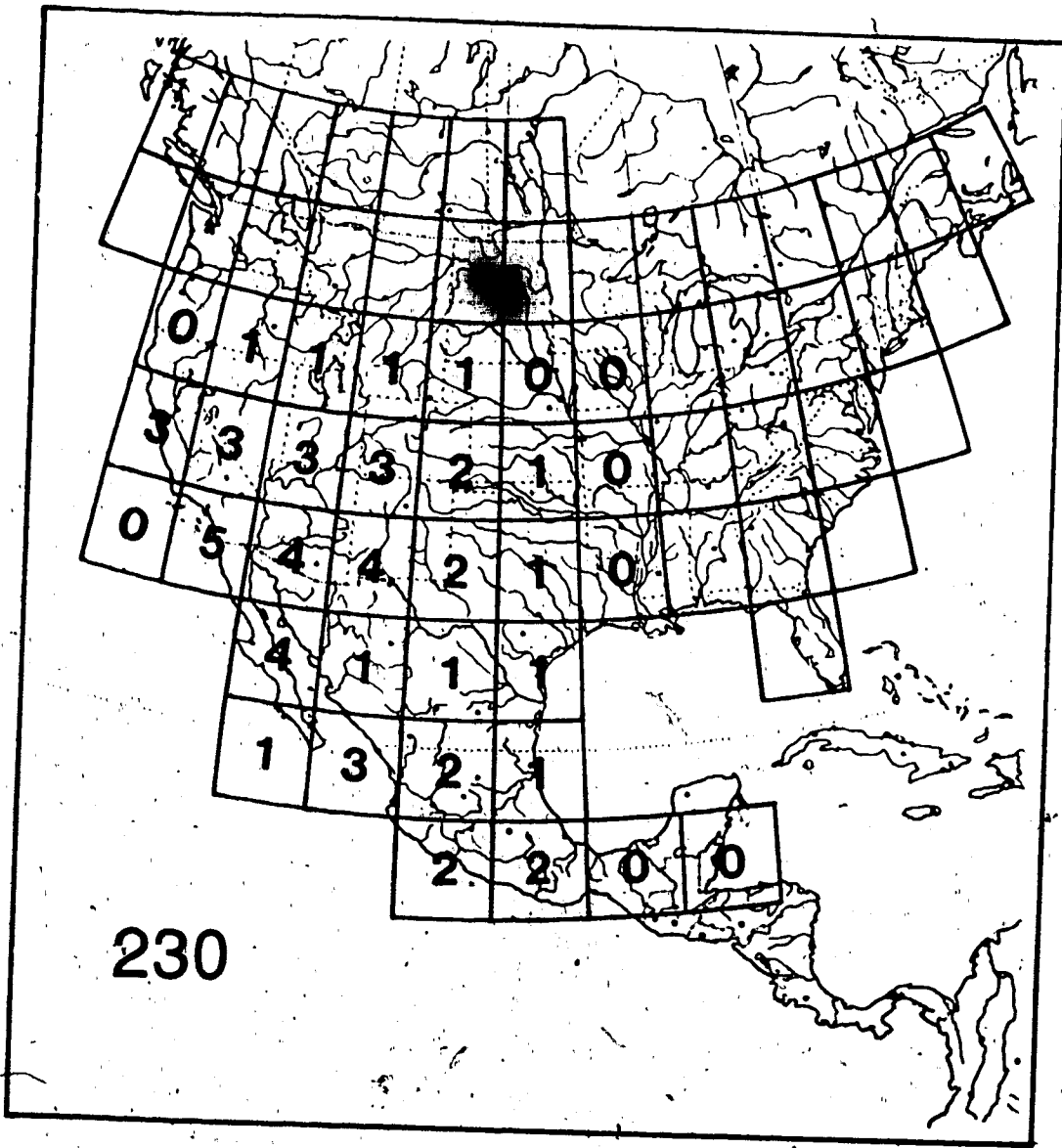


Figure 231. Numbers of species of *Cleonidius* in 5° intervals.

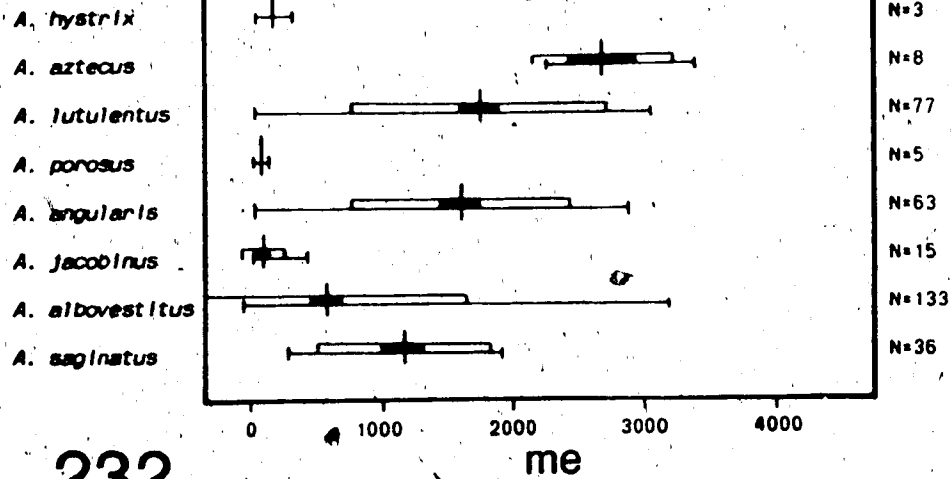


Figures 232 to 233.

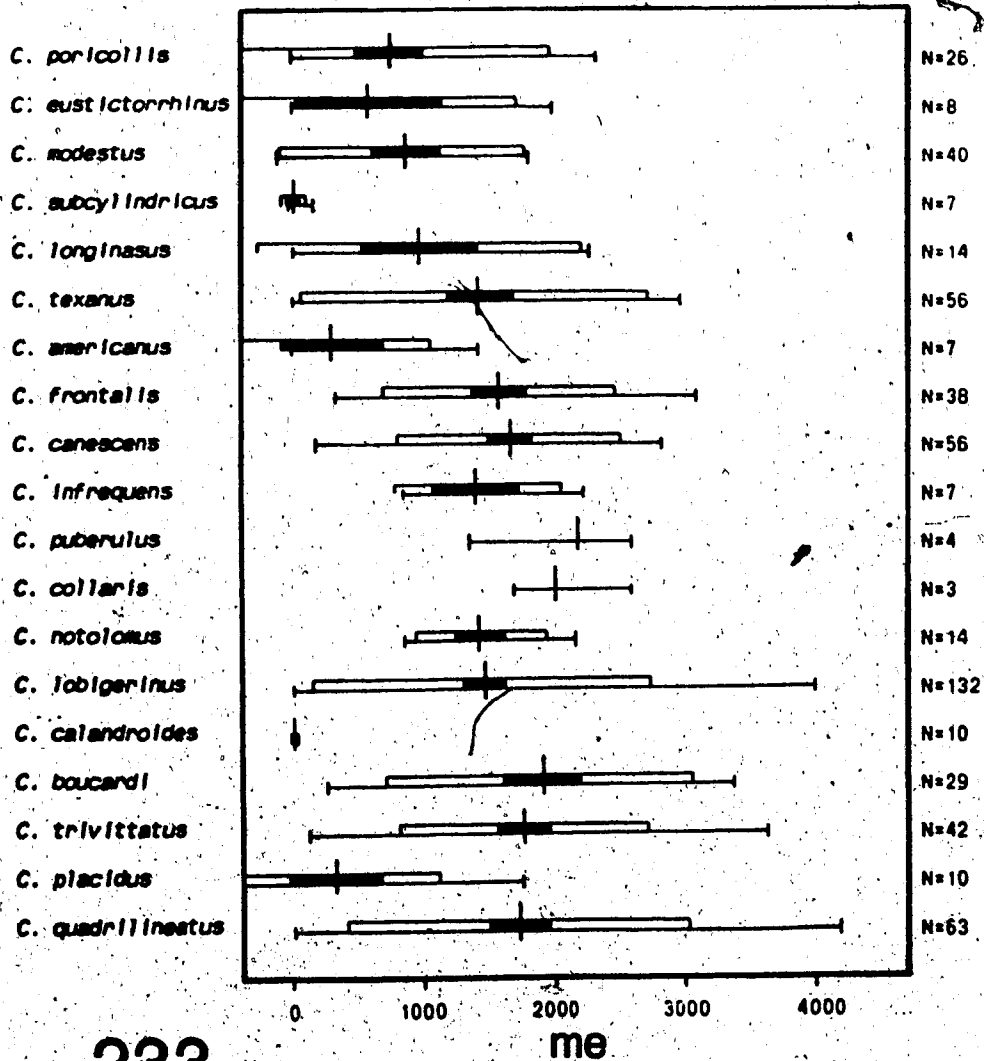
Figure 232. Hubbs-Hubbs diagram illustrating variation among specimens of *Apleurus* species; elevation. See caption for Figure 49 for explanation.

Figure 233. Hubbs-Hubbs diagram illustrating variation among specimens of *Cleonidius* species; elevation. See caption for Figure 49 for explanation.



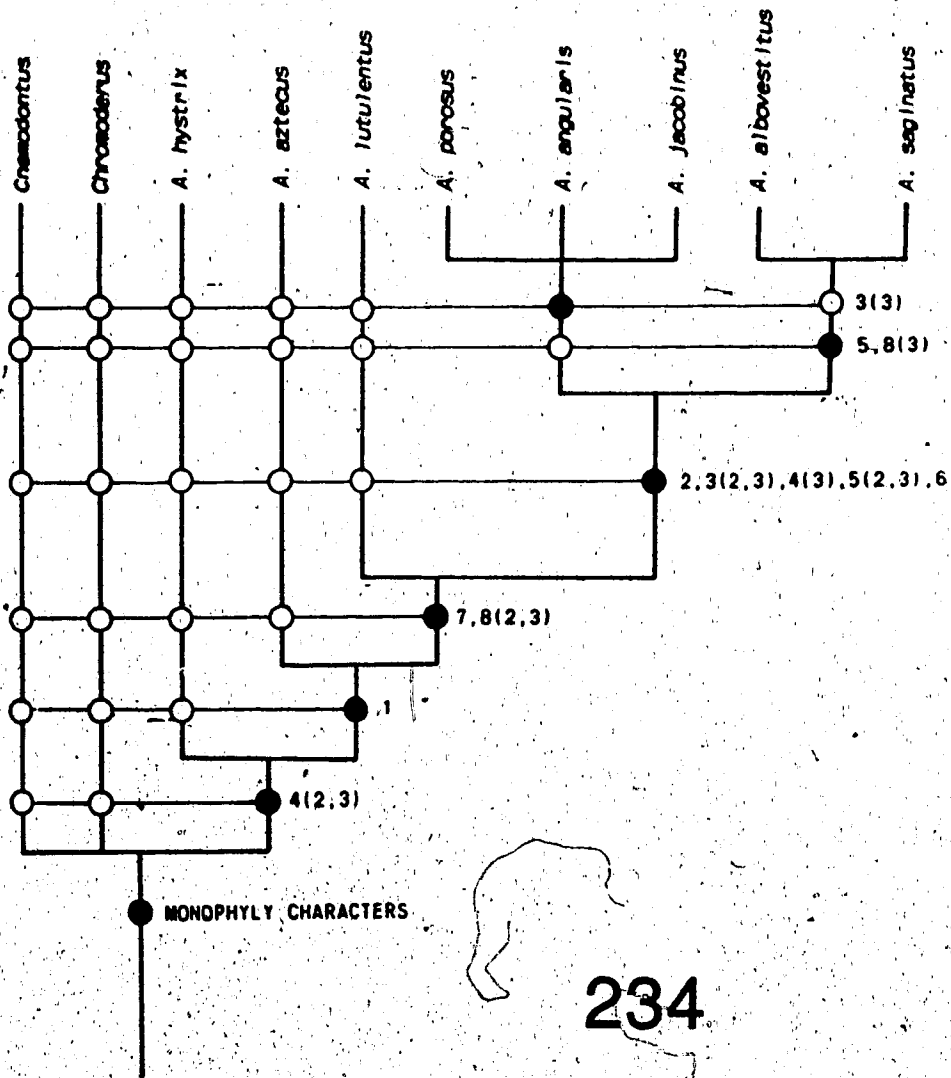


232



233

Figure 234. Reconstructed phylogeny based on primary characters illustrating hypothesized phylogenetic relationships among *Apleurus* species and related out-groups.. Dots denote apotypic character states, open circles plesiotypic states. See "Phylogeny" section and Tables 68 and 70 for discussion of characters and states (numbers in parentheses).



8

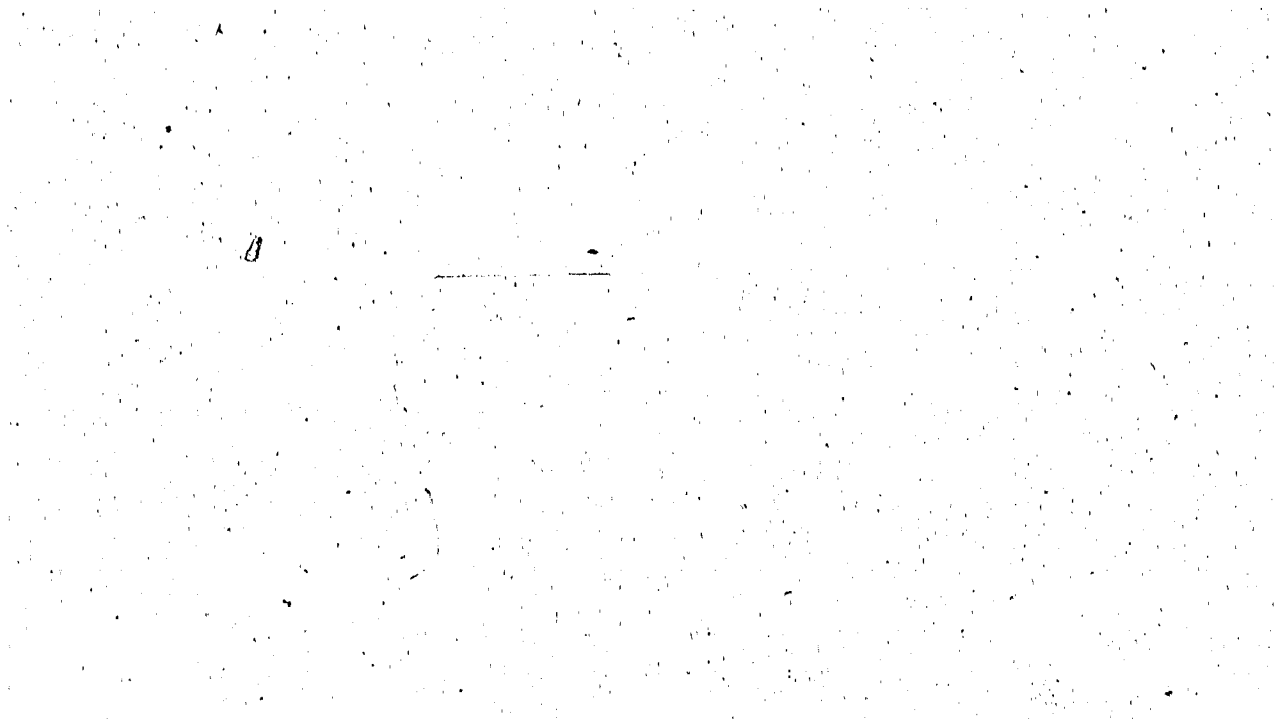
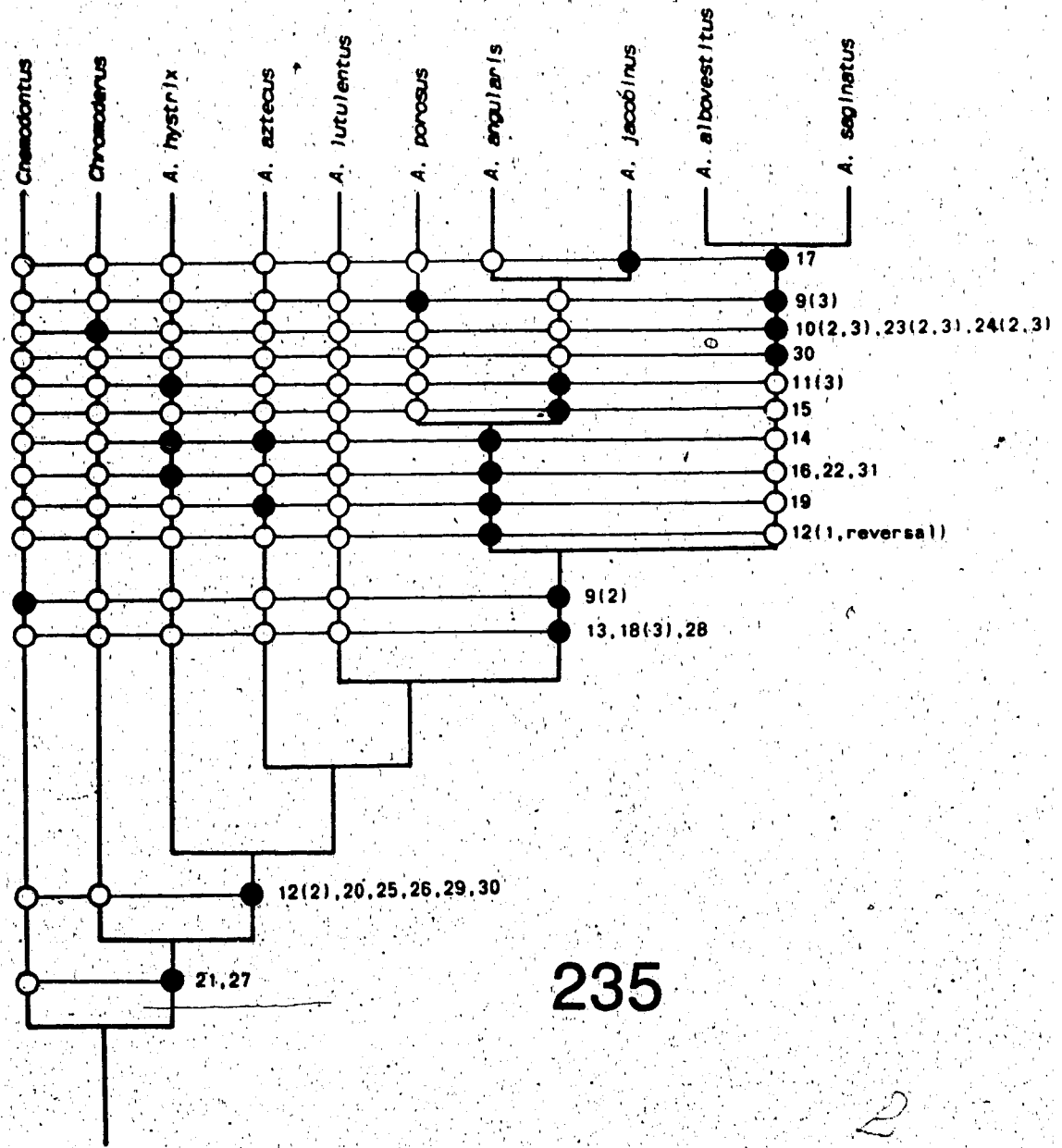
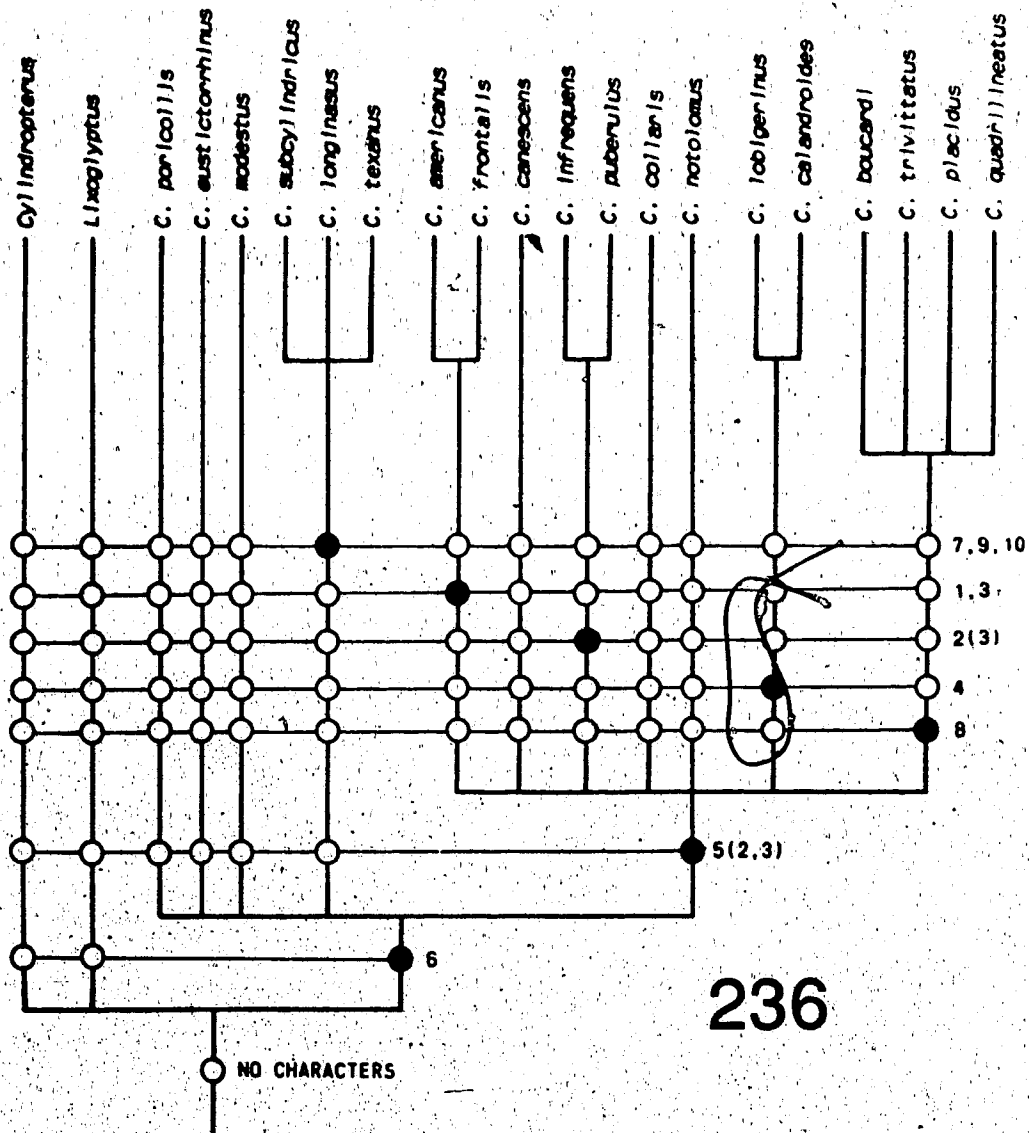


Figure 235. Reconstructed phylogeny based on primary and secondary characters illustrating hypothesized phylogenetic relationships among *Apleurus* species and related out-groups. Dots denote apotypic character states, open circles plesiotypic states. See "Phylogeny" section and Tables 68 and 70 for discussion of characters and states (numbers in parentheses).



J

Figure 236. Reconstructed phylogeny based on primary characters illustrating hypothesized phylogenetic relationships among *Cleonidius* species and related out-groups. Dots denote apotypic character states, open circles plesiotypic states. See "Phylogeny" section and Tables 69 and 71 for discussion of characters and states (numbers in parentheses).



236

Figure 237. Reconstructed phylogeny based on primary and secondary characters illustrating hypothesized phylogenetic relationships among *Cleonidius* species and related out-groups. Dots denote apotypic character states, open circles plesiotypic states. See "Phylogeny" section and Tables 69 and 71 for discussion of characters and states (numbers in parentheses).



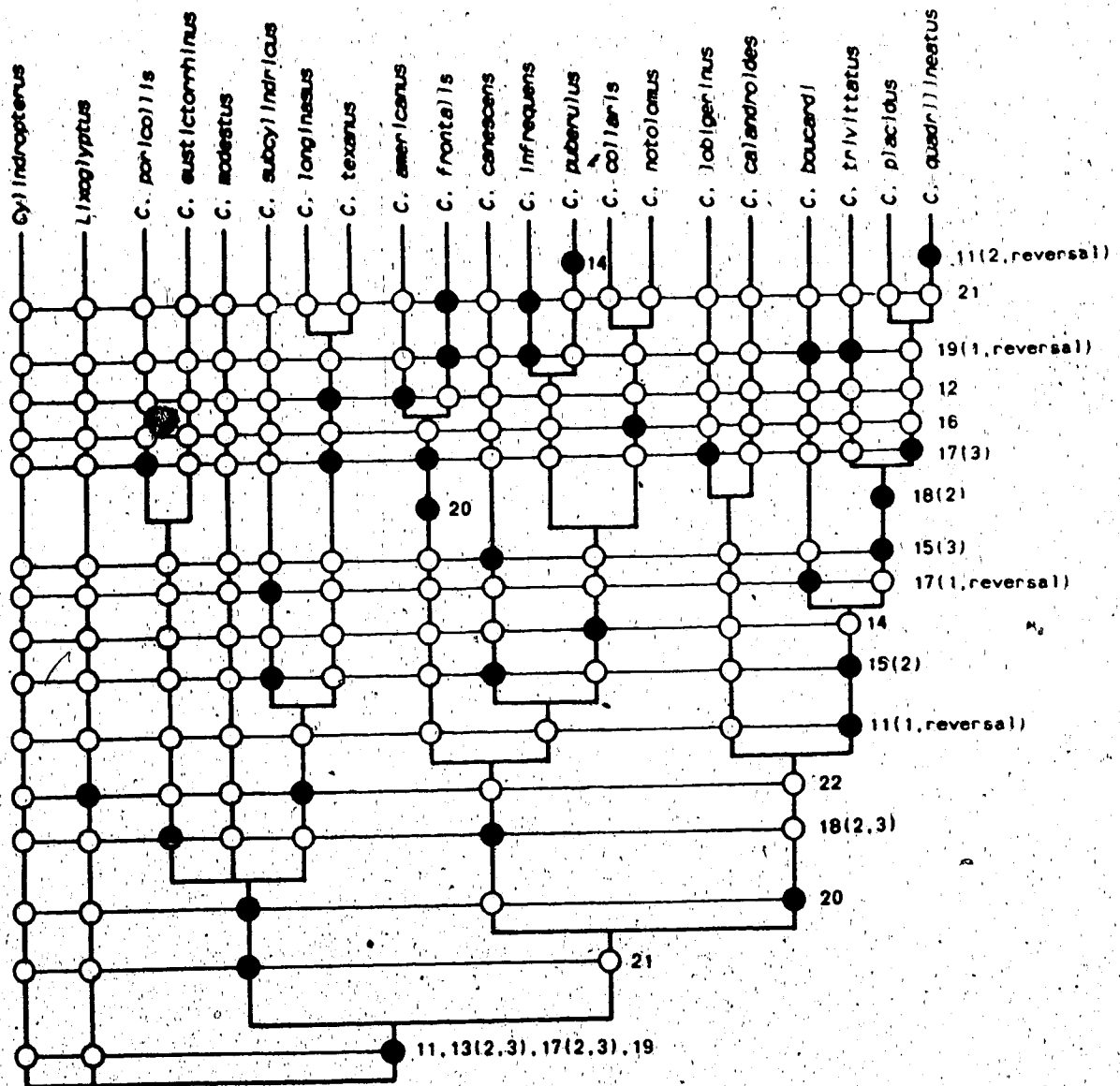


Figure 238. Habitat associations of *Apleurus* species. Solid box denotes marked association, half-box slight association. Phylogeny following Figure 235.

Mojave Desert  
 Sonoran Desert  
 Chihuahuan Desert  
 Great Basin Desert  
 Texas Semi-desert  
 Pacific Semi-desert  
 Desert-Grassland  
 Grassland  
 Grassland-Decid For.  
 Mesquite Grassland

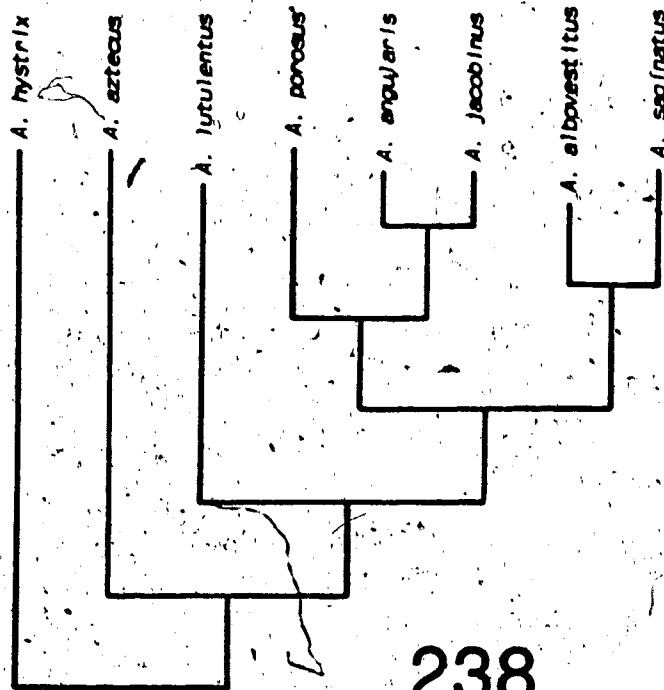
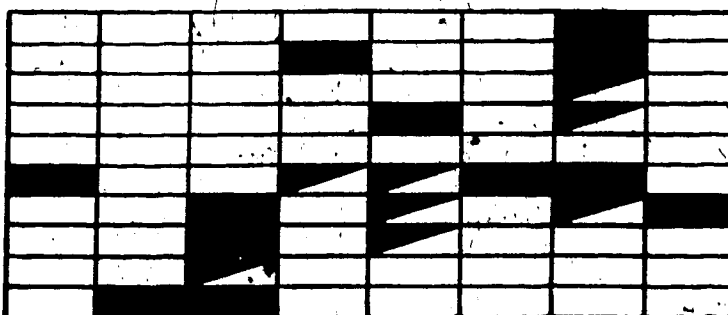


Figure 239. Plant associations of *Apleurus* species. Solid box denotes marked association, half-box slight association, check mark few collection records. Plant taxa are listed alphabetically. Phylogeny following Figure 235.

Asclepiadaceae			✓✓		✓		✓	
Chenopodiaceae			✓✓		✓		✓	
Compositae								
Cruciferae							✓✓	
Ephedraceae							✓✓✓	
Graminae							✓✓	
Leguminosae	✓		✓✓			✓✓	✓✓	
Malvaceae			✓✓			✓✓	✓✓	
Polygonaceae							✓✓	
Rosaceae							✓✓	
Solenaceae							✓✓	
Umbelliferae						✓	✓	
Zygophyllaceae							✓	

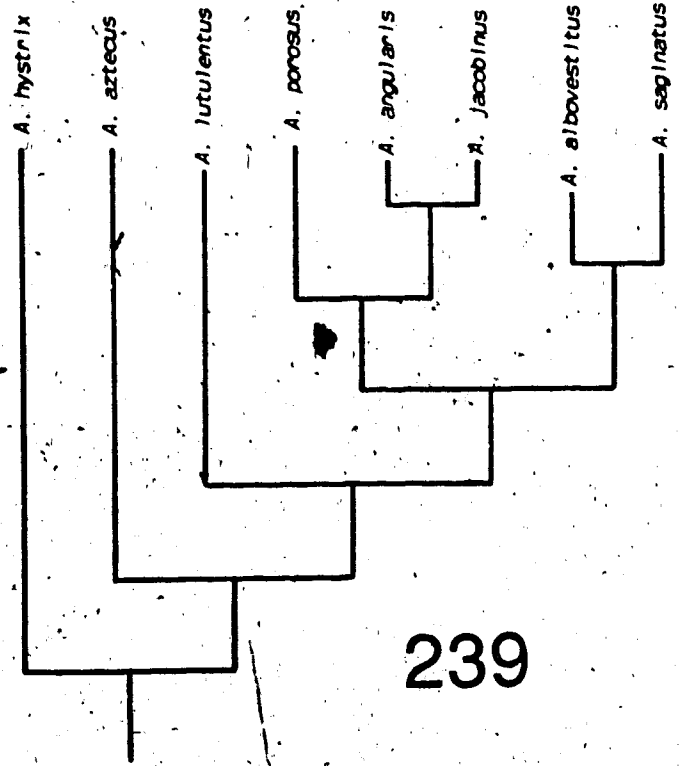
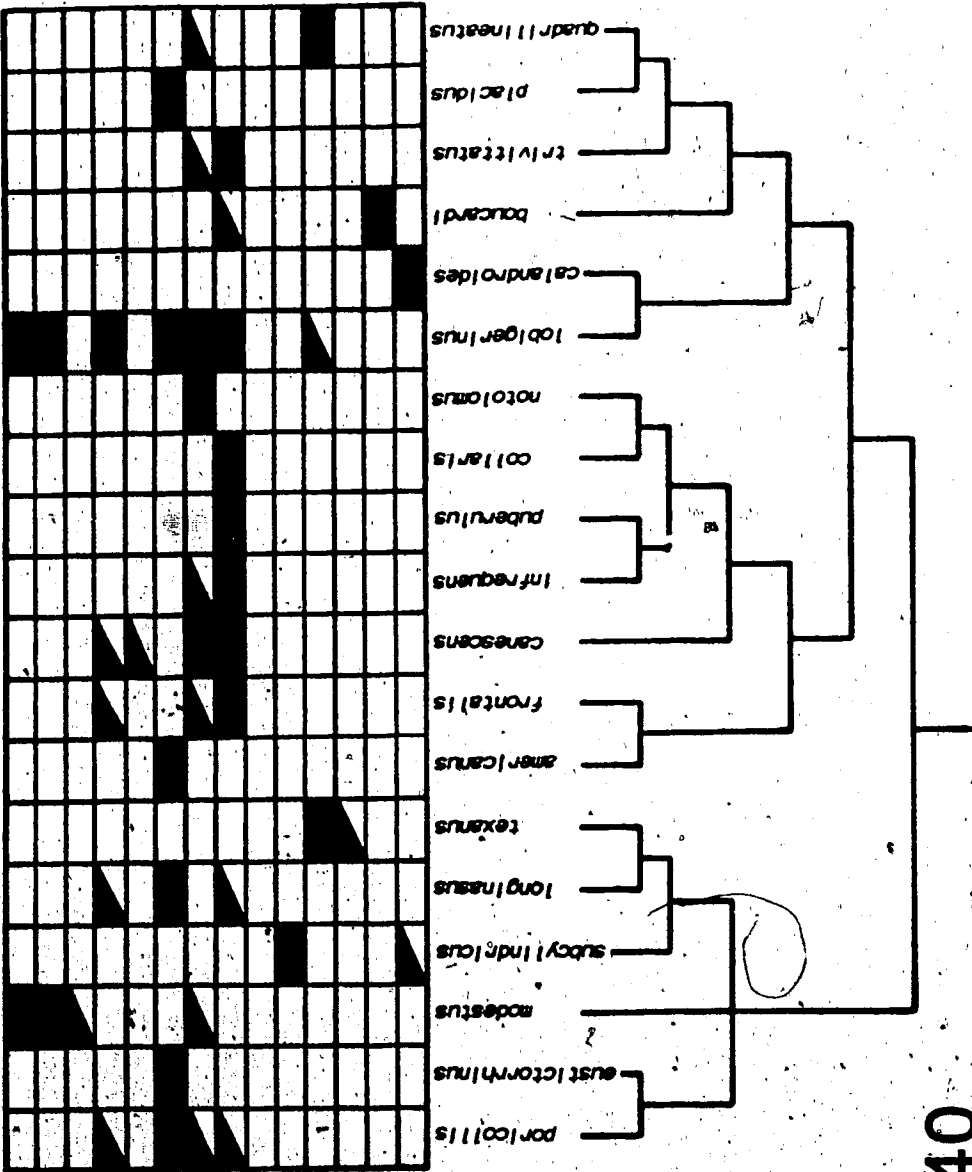


Figure 240. Habitat associations of *Cleonidius* species.

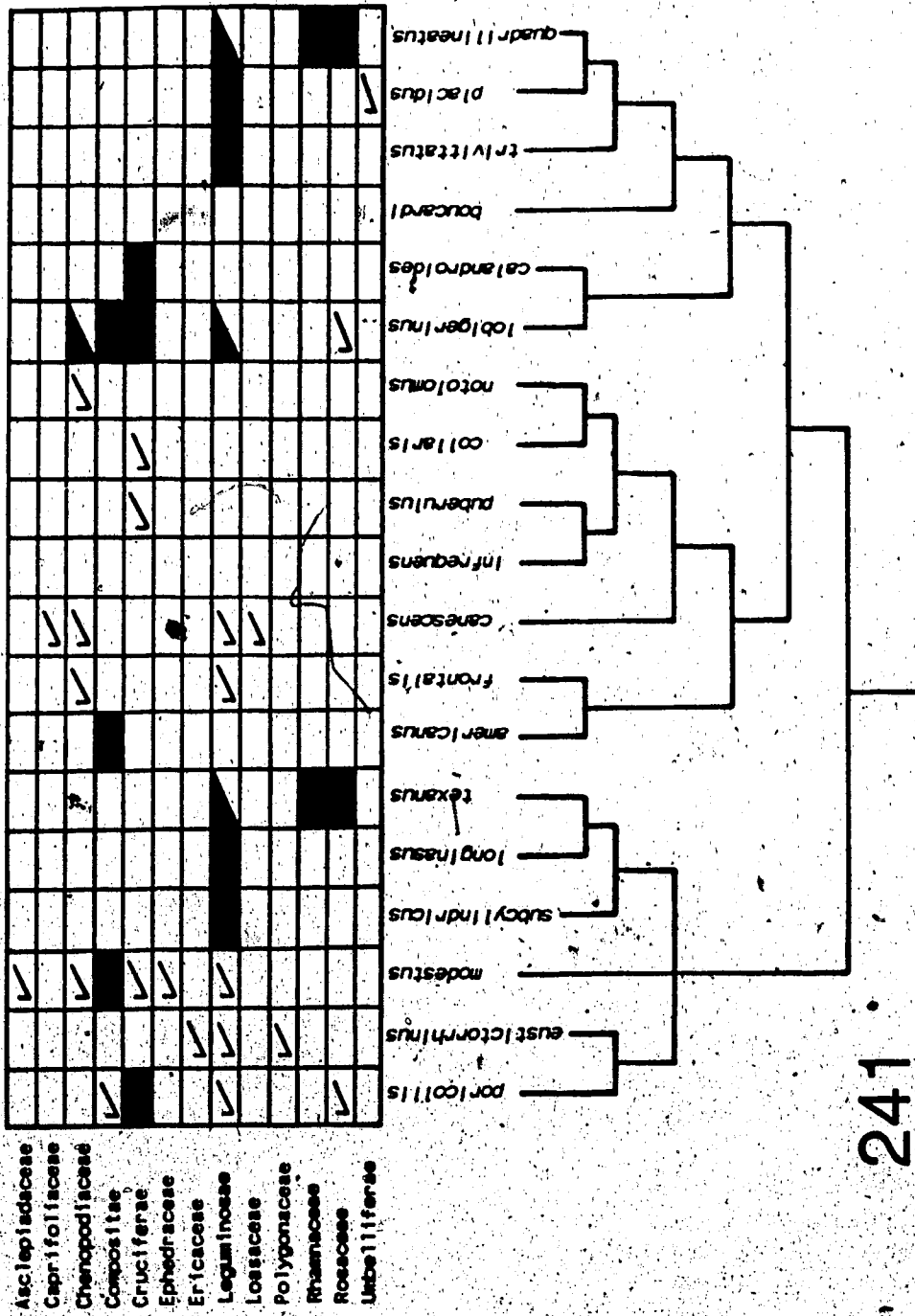
Solid box denotes marked association, half-box slight association. Phylogeny following Figure 237.



240

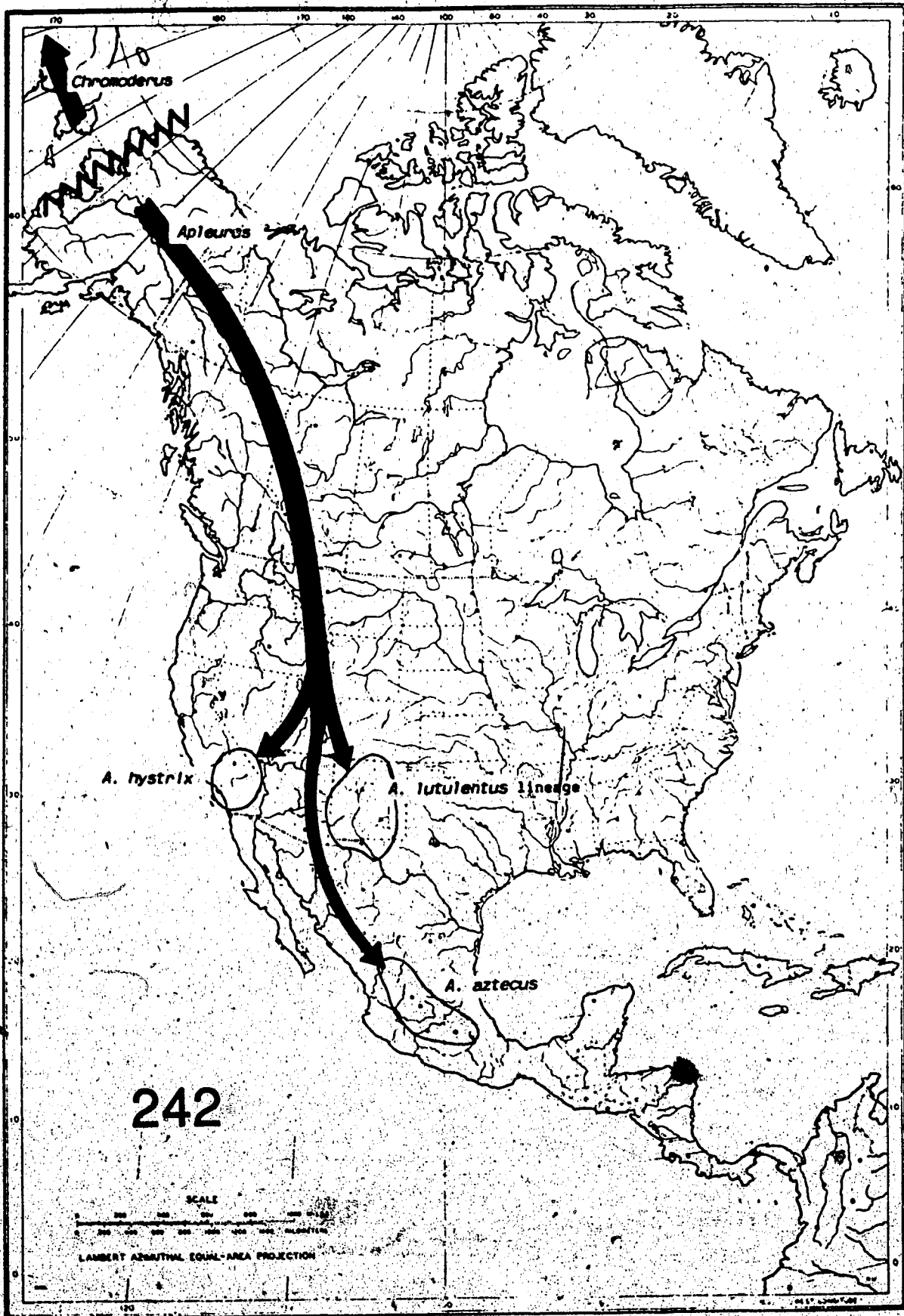
Figure 241. Plant associations of *Cleonidius* species. Solid box denotes marked association, half-box slight association, check mark few collection records. Plant associations are unknown for some species. Plant taxa are listed alphabetically. Phylogeny following Figure 237.





241

Figure 242. Origin and initial diversification of *Apleurus* in North America: *Apleurus hystrix*, *A. aztecus*, and *A. lutulentus*-*A. porosus*-*A. angularis*-*A. jacobinus*-*A. albovestitus*-*A. saginatus* lineage. See text for detailed explanation. Zig-zagged line indicates barrier formation.



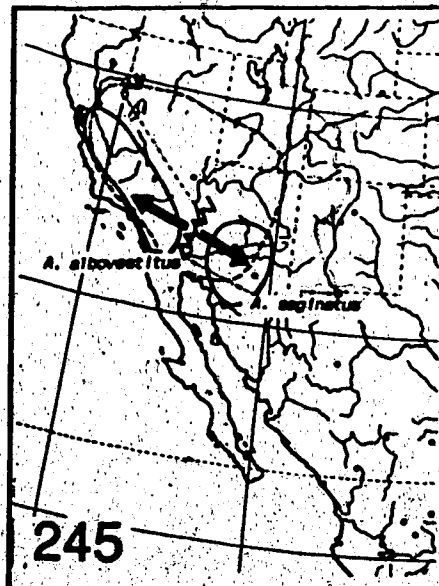
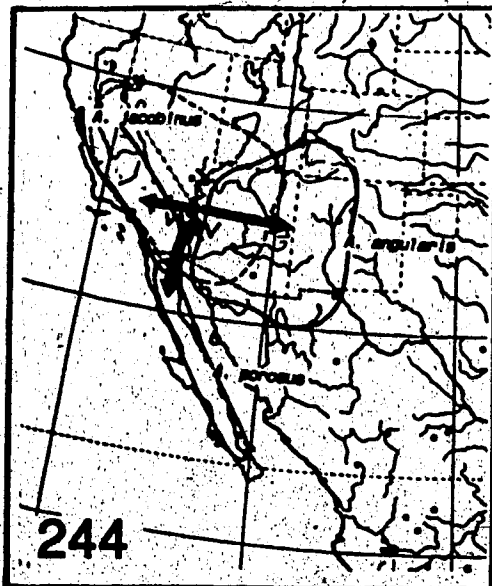
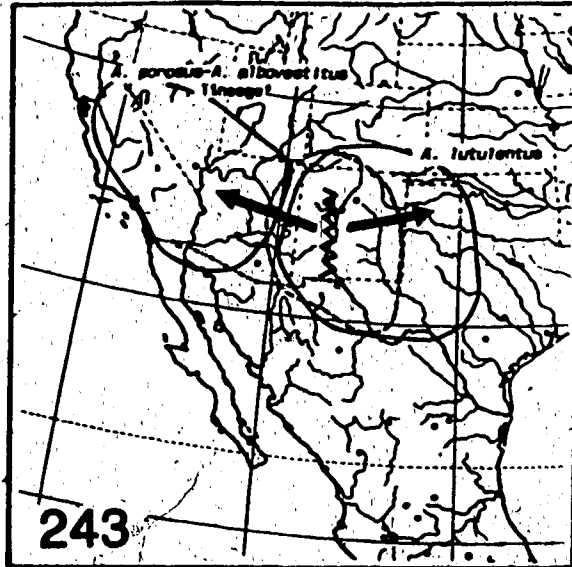
242

Figures 243 to 245.

Figure 243. Diversification of *Apleurus* in North America: *Apleurus lutulentus* and *A. porosus*-*A. angularis*-*A. jacobinus*-*A. albovestitus*-*A. saginatus* lineage. See text for detailed explanation. Zig-zagged line indicates barrier formation. Dashed line indicates hypothesized ancestral distribution. Solid lines indicate present distributions.

Figure 244. Diversification of *Apleurus* in North America: *Apleurus angularis*, *A. jacobinus* and *A. porosus*. See text and caption for Figure 243 for detailed explanation.

Figure 245. Diversification of *Apleurus* in North America: *Apleurus albovestitus* and *A. saginatus*. See text and caption for Figure 243 for detailed explanation.



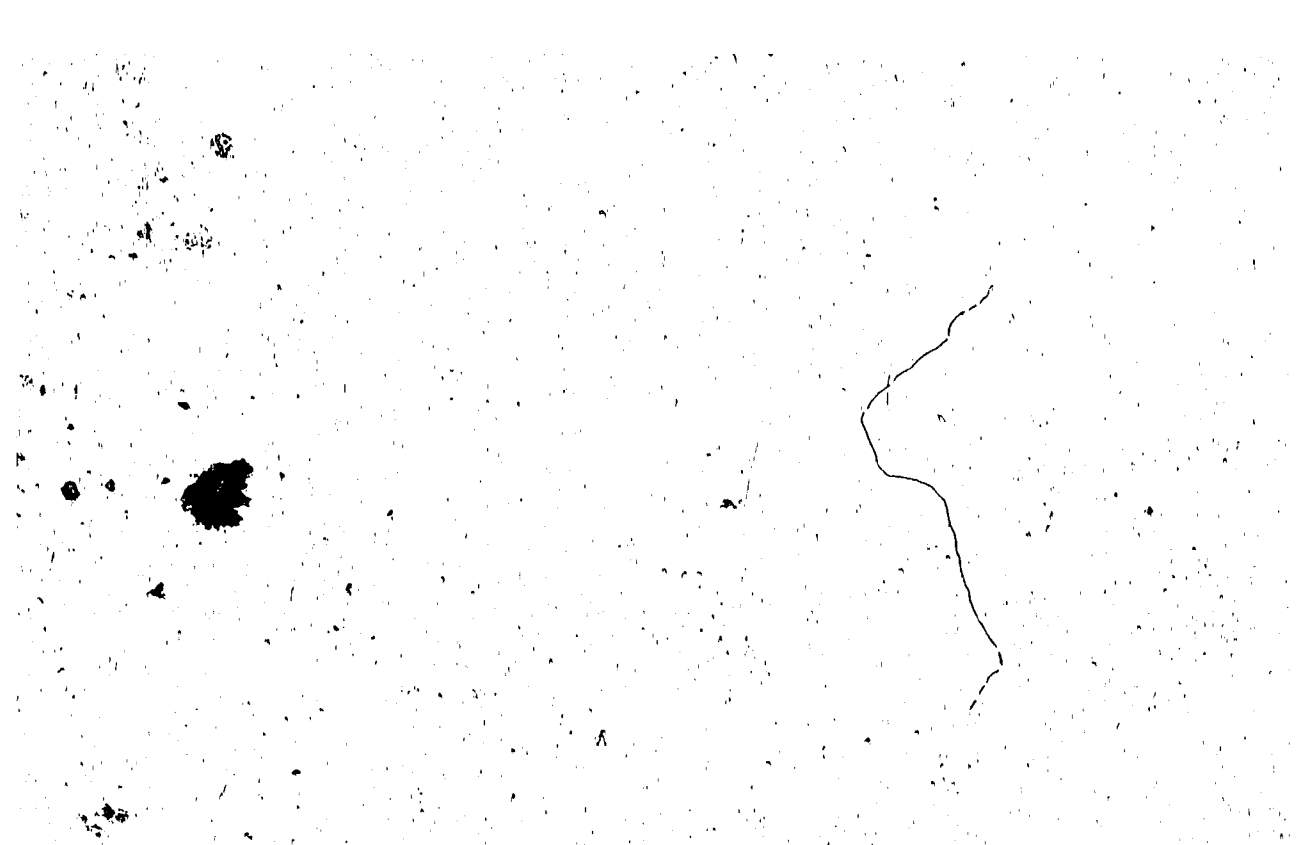
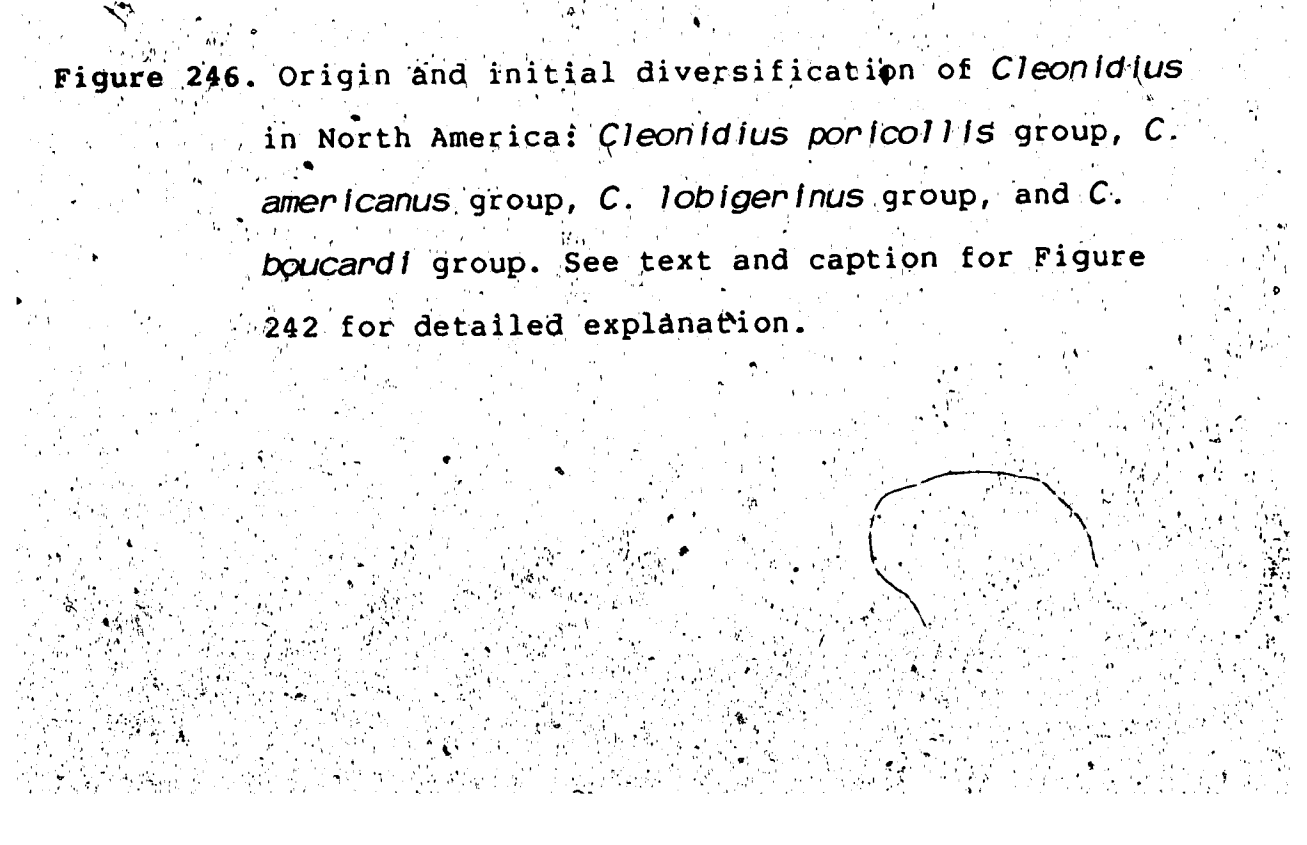
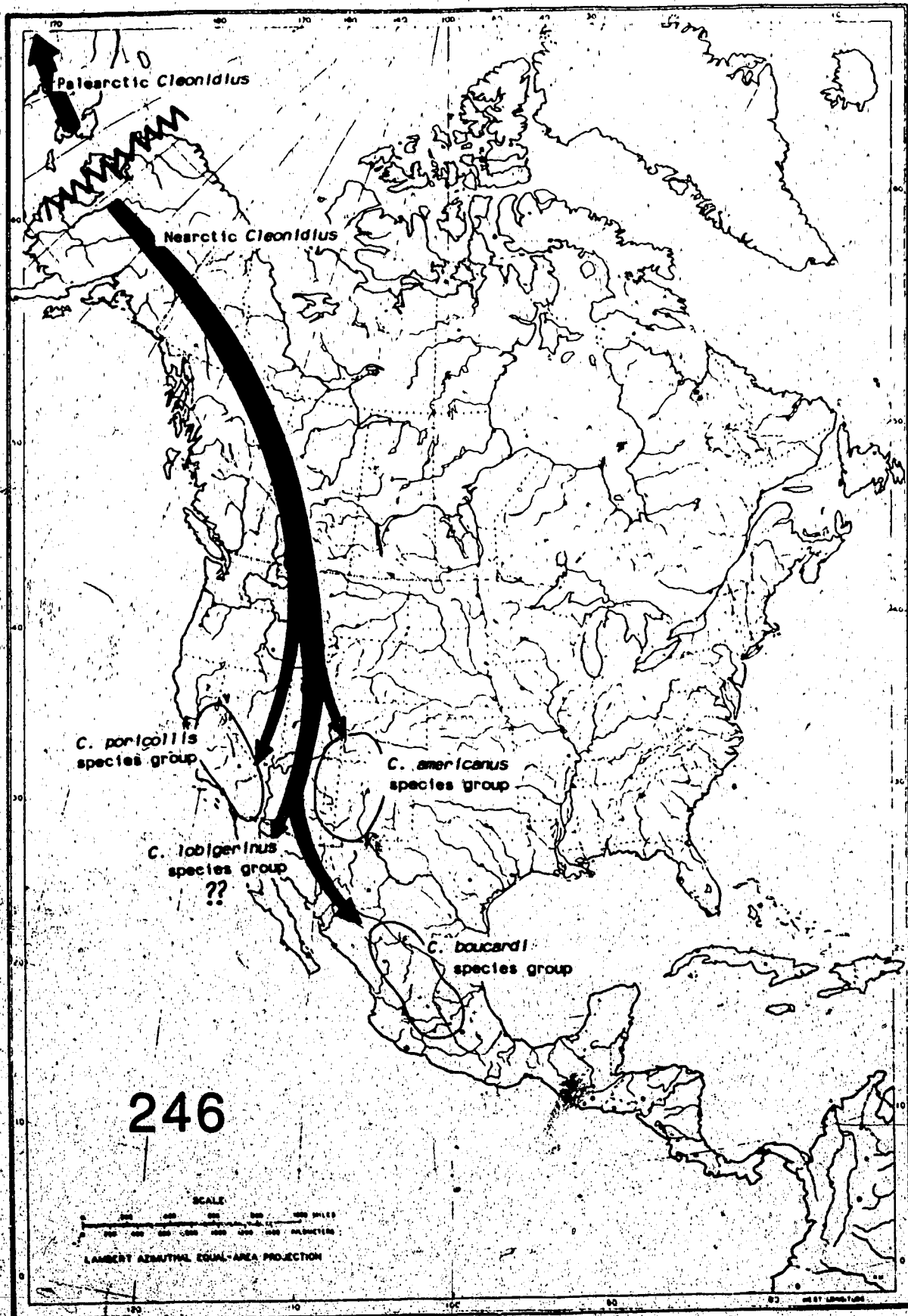


Figure 246. Origin and initial diversification of *Cleonidius* in North America: *Cleonidius porlicollis* group, *C. americanus* group, *C. lobigerinus* group, and *C. boucardi* group. See text and caption for Figure 242 for detailed explanation.





246

SCALE  
0 200 400 600 800 1000 MILES  
0 200 400 600 800 1000 KILOMETERS  
LAMBERT AZIMUTHAL EQUAL-AREA PROJECTION

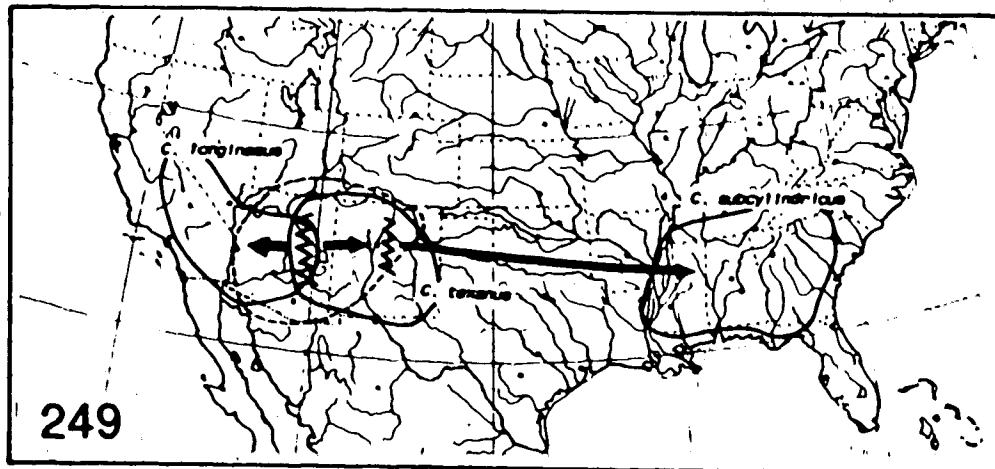
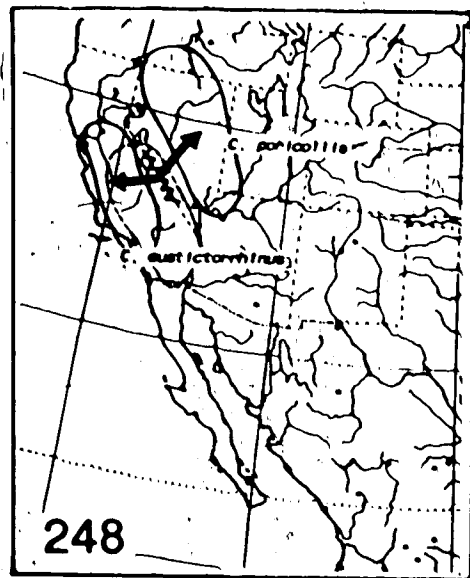
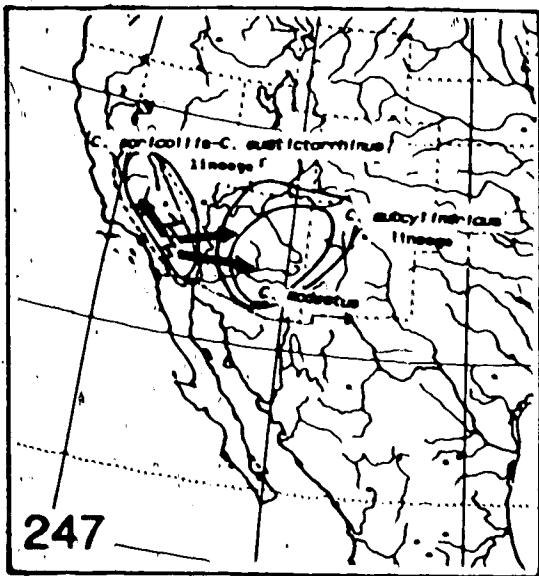
Figures 247 to 249.

Figure 247. Diversification of *Cleonidius* in North America: *Cleonidius poricollis*-*C. eustictorrhinus*, *C. modestus*, and *C. subcylindricus*-*C. longinasus*-*C. texanus* lineages. See text and caption for Figure 243 for detailed explanation.

Figure 248. Diversification of *Cleonidius* in North America: *Cleonidius poricollis* and *C. eustictorrhinus*. See text and caption for Figure 243 for detailed explanation.

Figure 249. Diversification of *Cleonidius* in North America: *Cleonidius subcylindricus*, *C. longinasus* and *C. texanus*. See text and caption for Figure 243 for detailed explanation.





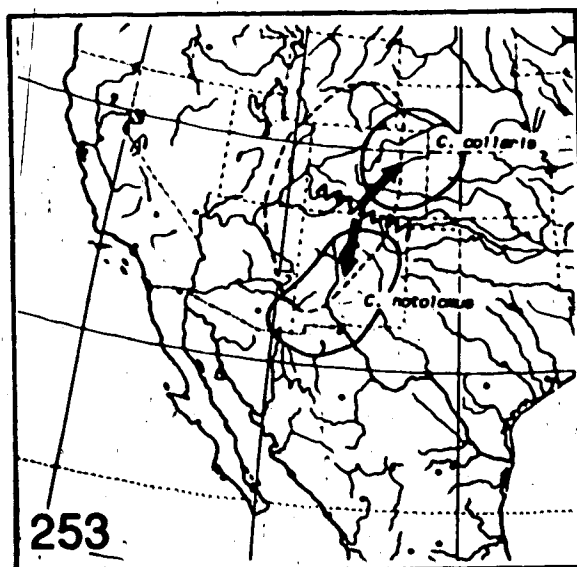
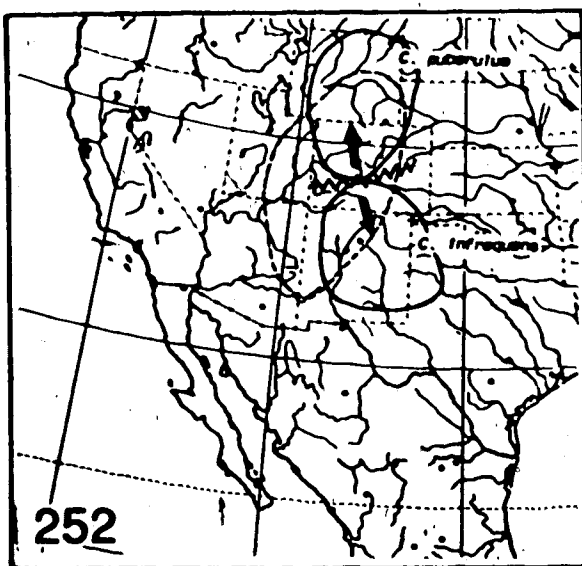
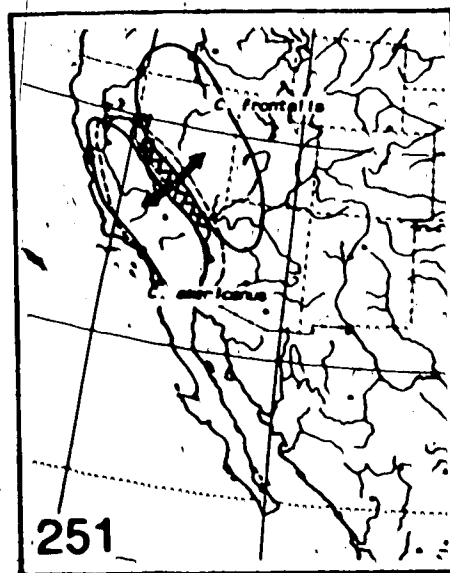
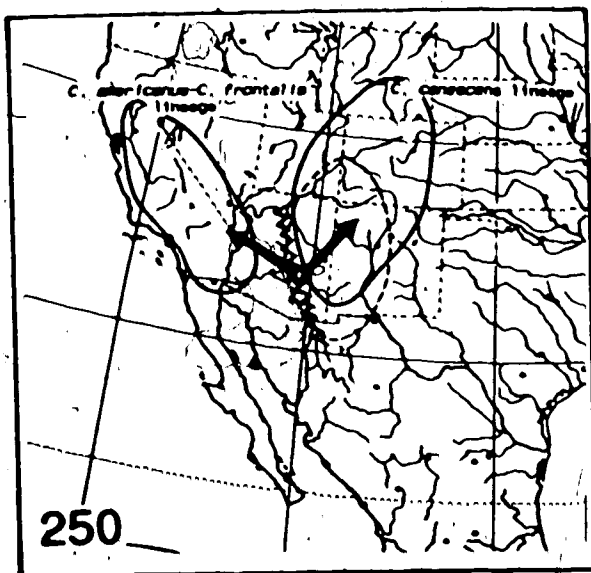
Figures 250 to 253.

Figure 250. Diversification of *Cleonidius* in North America: *Cleonidius americanus*-*C. frontalis*, *C. canescens*-*C. infrequens*-*C. puberulus*-*C. collaris*-*C. notolomus* lineages. See text and caption for Figure 243 for detailed explanation.

Figure 251. Diversification of *Cleonidius* in North America: *Cleonidius americanus* and *C. frontalis*. See text and caption for Figure 243 for detailed explanation.

Figure 252. Diversification of *Cleonidius* in North America: *Cleonidius puberulus* and *C. infrequens*. See text and caption for Figure 243 for detailed explanation.

Figure 253. Diversification of *Cleonidius* in North America: *Cleonidius collaris* and *C. notolomus*. See text and caption for Figure 243 for detailed explanation.



Figures 254 to 256.

Figure 254. Diversification of *Cleonidius* in North

America: *Cleonidius boucardi* and *C.*

*trivittatus*-*C. placidus*-*C. quadrilineatus*

lineage. See text and caption for Figure 243 for detailed explanation.

Figure 255. Diversification of *Cleonidius* in North

America: *Cleonidius trivittatus* and *C.*

*placidus*-*C. quadrilineatus* lineage. See text and

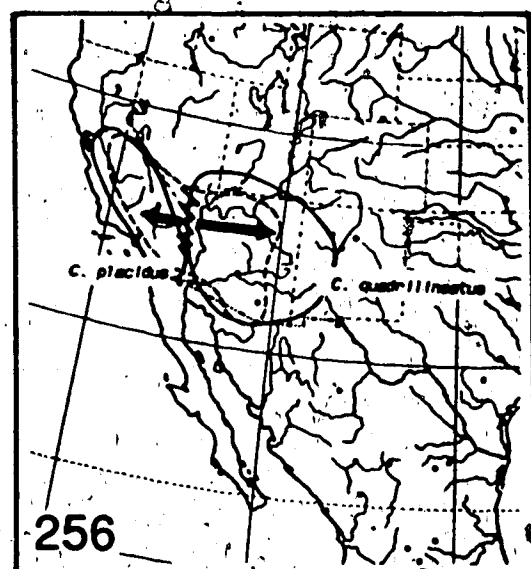
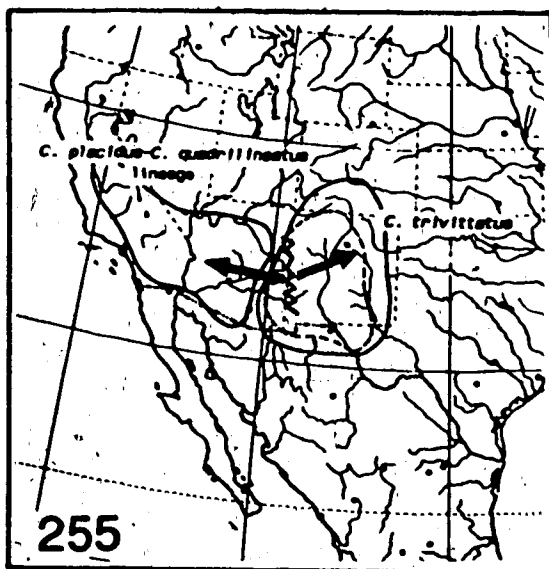
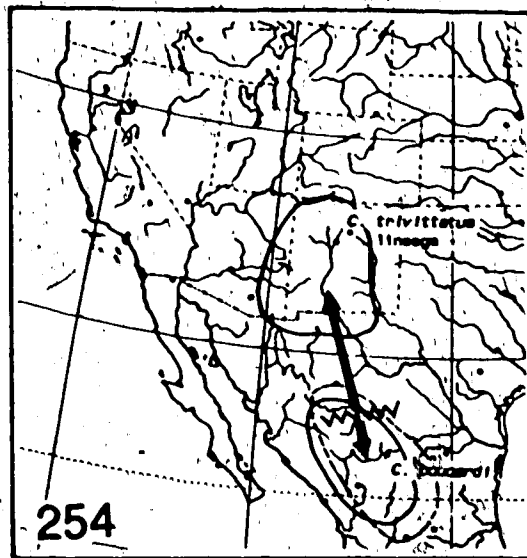
caption for Figure 243 for detailed explanation.

Figure 256. Diversification of *Cleonidius* in North

America: *Cleonidius placidus* and *C.*

*quadrilineatus*. See text and caption for Figure

243 for detailed explanation.



Figures 257 to 258.

Figure 257. Geographic ranges of species in the  
*Cleonidius poricollis* species group.

Figure 258. Geographic ranges of species in the  
*Cleonidius americanus* species group.

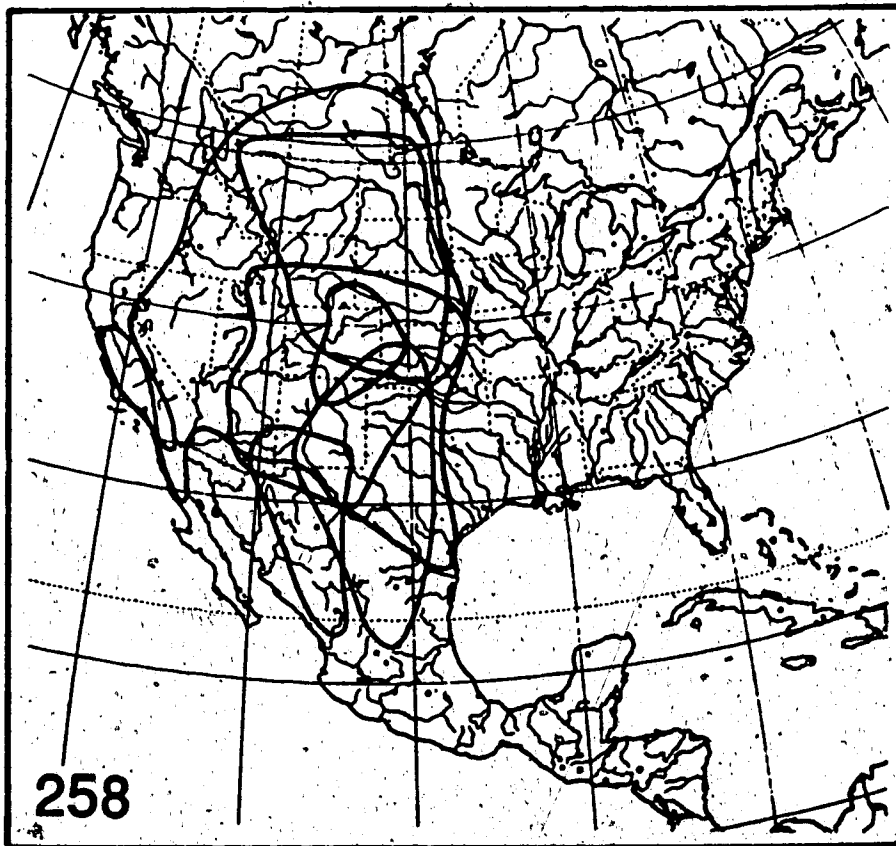
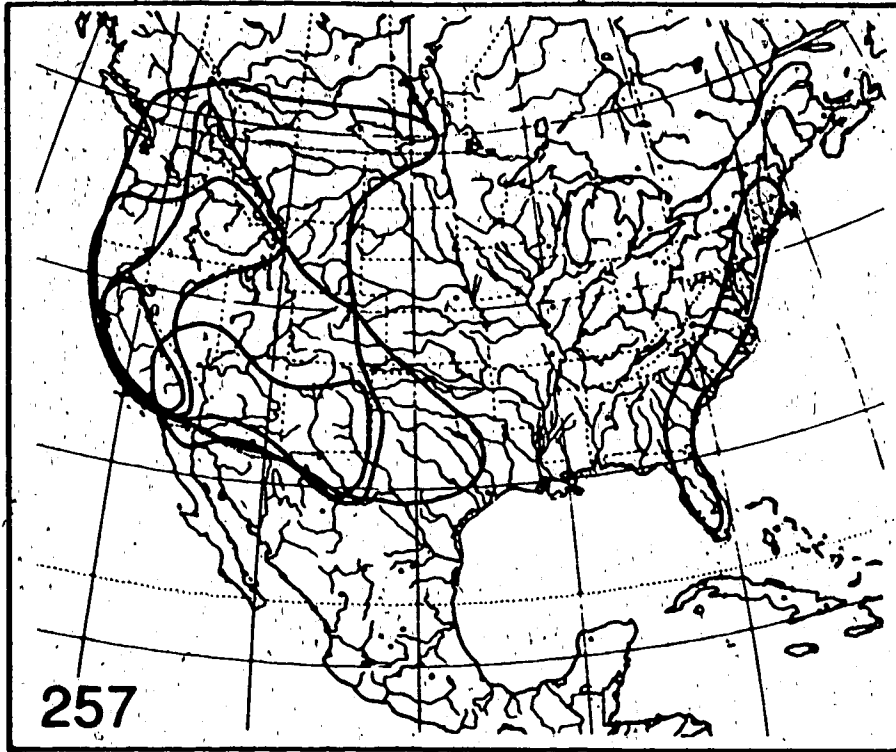
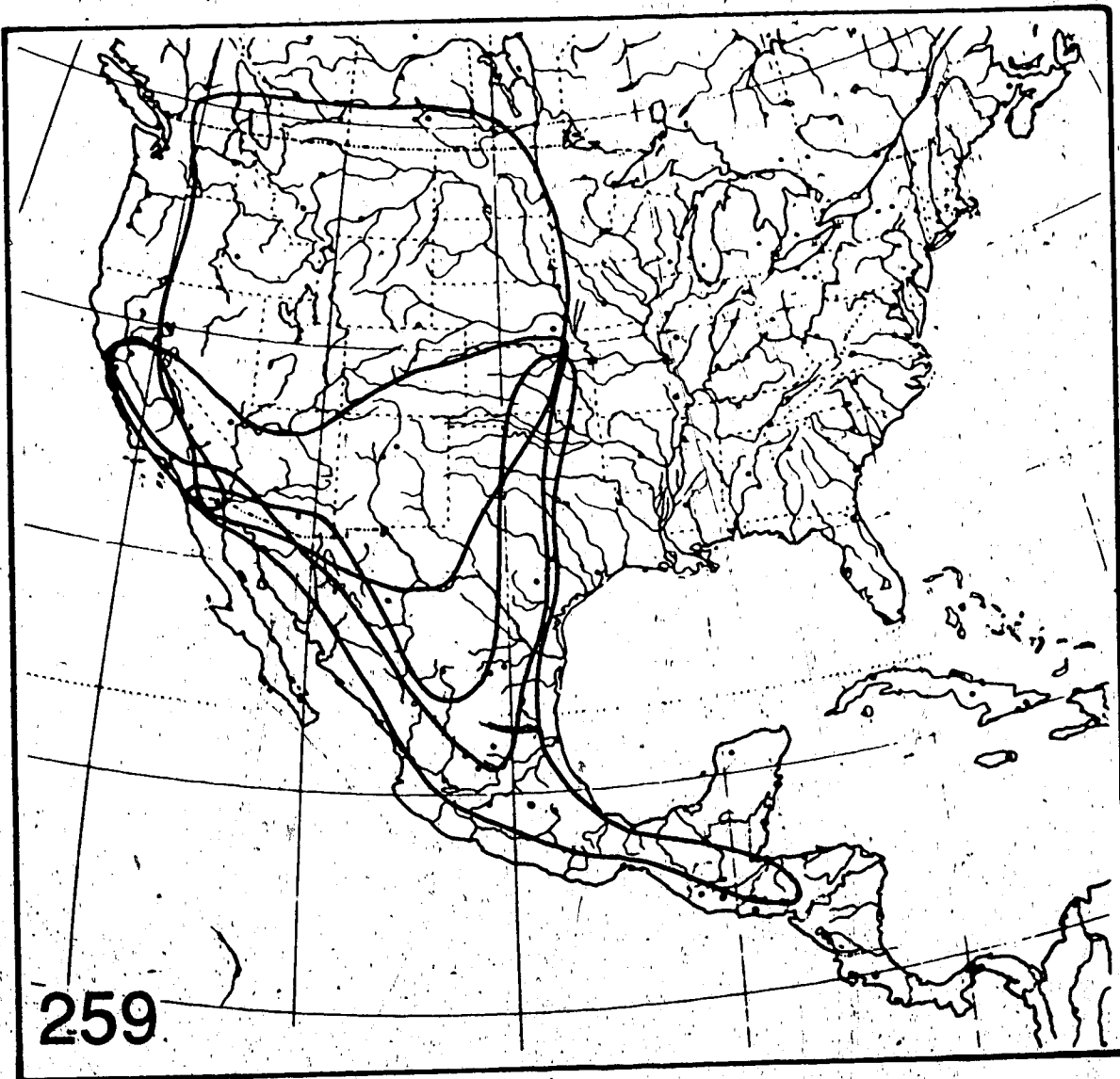


Figure 259. Geographic ranges of species in the *Cleonidius*  
*boucardi* species group.





259

10. **Appendix 3**: Species of genus-group names of Cleoninae examined for purposes of determining relationships of New World taxa. Genus-group name is listed first, followed by species examined on loan from BMNH. An asterisk indicates that the species examined is the type-species of that generic-group name. No asterisk indicates that the type species was not examined because, no type-species has yet been designated for that genus-group name; or, type species was not available for study in the BMNH.

<i>Adosomus</i> Faust	<i>A. granulatus</i> (Mannerheim)
<i>Amblysomus</i> Faust	<i>A. brevis</i> (Fähræus) *
<i>Ammocleonus</i> Bedel	<i>A. hieroglyphicus</i> (Olivier) *
<i>Atactogaster</i> Faust	<i>A. orientalis</i> (Chevrolat) *
<i>Bothynoderes</i> Schoenherr	<i>B. punctiventris</i> (Germar) *
<i>Brachyoleonus</i> Faust	<i>B. fronto</i> (Fischer von Waldheim) *
<i>Calodemus</i> Faust	<i>C. vetustus</i> Faust
<i>Centrocleonus</i> Chevrolat	<i>C. fallax</i> (Fähræus) *
<i>Chromoderus</i> Motschulsky	<i>C. albidus</i> (Fabricius) * [ = <i>C. fasciatus</i> (Vill.) ]
<i>Chromonotus</i> Motschulsky	<i>C. vittatus</i> (Hochhuth) *
<i>Chromosomus</i> Motschulsky	<i>C. fischeri</i> (Fähræus) *
<i>Cnemodontus</i> Chevrolat	<i>C. limpidus</i> (Gyllenhal)
<i>Coniocleonus</i> Motschulsky	<i>C. excoriatus</i> (Gyllenhal) [type species, <i>C. carinirostris</i> (Gyllenhal)]
<i>Conorhynchus</i> Motschulsky	<i>C. conirostris</i> (Gebler) *
<i>Cosmogaster</i> Faust	<i>C. lateralis</i> (Gyllenhal) *
<i>Cylindropterus</i> Chevrolat	<i>C. luxeri</i> Chevrolat *
<i>Cyphocleonus</i> Motschulsky	<i>C. cenchrus</i> (Pallas) *

<i>Epirhynchus</i> Schoenherr	<i>E. humerosus</i> Faust [type species, <i>E. argus</i> (Sparrm.)]
<i>Eumecops</i> Hochhuth	<i>Eumecops</i> sp. [type species, <i>E. kittaryi</i> Hochhuth]
<i>Eurycleonus</i> Bedel	<i>E. baluchicus</i> Marshall [type species, <i>E. gigas</i> Mars.]
<i>Gonocleonus</i> Chevrolat	<i>G. helferi</i> Chevrolat *
<i>Isomerus</i> Motschulsky	<i>I. granosus</i> (Zoubkoff) *
<i>Koenigius</i> Heyd.	<i>K. palaestinus</i> Heyd. *
<i>Leucochromus</i> Motschulsky	<i>L. imperialis</i> (Zoubkoff) *
<i>Leucomigus</i> Motschulsky	<i>L. candidatus</i> (Pallas) *
<i>Leucosomus</i> Motschulsky	<i>L. ophthalmicus</i> (Rossi) *
<i>Liocleonus</i> Motschulsky	<i>L. clathratus</i> (Olivier) *
<i>Lixocleonus</i> Marshall	<i>L. incanus</i> Marshall *
<i>Lixomorphus</i> Faust	<i>L. ocularis</i> (Fabricius) *
<i>Mecaspis</i> Schoenherr	<i>M. palmatus</i> (Olivier)
<i>Menocleonus</i> Faust	<i>M. implicatus</i> Faust
<i>Microcleonus</i> Faust	<i>M. panderi</i> (Fischer von Waldheim) *
<i>Nemoxenus</i> Faust	<i>N. zebra</i> (Chevrolat)
<i>Neocleonus</i> Chevrolat	<i>N. sannio</i> (Herbst) *
<i>Nemimonyx</i> Faust	<i>N. perturbans</i> Faust *
<i>Pachycerus</i> Schoenherr	<i>P. cordiger</i> (Germar) [type species, <i>P. scabrosus</i> Brullé]
<i>Pentatropis</i> Faust	<i>P. sparsus</i> (Fåhraeus) [type species, <i>P. formosus</i> (Fåhraeus)]
<i>Phaulosomus</i> Faust	<i>P. kilimanus</i> Faust
<i>Pleurocleonus</i> Motschulsky	<i>P. quadrivittatus</i> (Zoubkoff) *

<i>Porocleonus</i> Motschulsky	<i>P. candidus</i> (Olivier) *
<i>Prionorhinus</i> Chevrolat	<i>P. sordidatus</i> (Gyllenhal)
<i>Pseudocleonus</i> Chevrolat	<i>P. cinereus</i> (Schrank)
<i>Rhabdorhynchus</i> Motschulsky	<i>R. anchusae</i> Chevrolat [type species, <i>R. menetriesi</i> (Gyllenhal)]
<i>Stephanocleonus</i> Motschulsky	<i>S. flaviceps</i> (Pallas) *
<i>Temnorhinus</i> Chevrolat	<i>T. brevirostris</i> (Gyllenhal) *
<i>Tetragonothorax</i> Chevrolat	<i>T. retusus</i> (Fabricius) *
<i>Trichocleonus</i> Motschulsky	<i>T. leucophyllus</i> (Fischer von Waldheim) *
<i>Xanthochelus</i> Chevrolat	<i>X. longus</i> Chevrolat *
<i>Xenomacrus</i> Faust	<i>X. glacialis</i> (Herbst)