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Title of Thesis - Titre de la thèse

Ion Turbulence in a CO₂ Laser/Plasma Interaction

Degree for which thesis was presented / Grade pour lequel cette thèse fut présentée

Ph.D.

Year this degree conferred / Année d'obtention de ce grade

1985

University - Université

University of Alberta

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by

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YARUB S. AL-SHIRAIDA

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 ELECTRICAL ENGINEERING

EDMONTON, ALBERTA

FALL 1985

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DEDICATION

To my parents for their love and support.

ABSTRACT

A comprehensive investigation of the ion turbulence generated in a laser/plasma interaction experiment has been carried out using the high power laser facility available in the Department of Electrical Engineering at the University of Alberta. A CO_2 laser beam with focused intensity of $5 \times 10^{12} \text{ w/cm}^2$ was used to ionize and heat an oxygen gas target, thereby producing plasma with an electron temperature, T_e , of $\sim 100 \text{ eV}$ and density $n_s \lesssim 10^{19} \text{ cm}^{-3}$.

Ruby laser Thomson scattering was utilized to study the non-equilibrium ion fluctuations generated in the plasma by the presence of a strong electromagnetic wave. Considerable enhancement of these ion fluctuations over the thermal level was observed for two plasma regimes (short and long density scale length regimes). In the long scale length regime the magnitude of density fluctuations along with temporal and spectral features of the scattered ruby light indicate that ion turbulence can account for the anomalous absorption of CO_2 laser energy observed in earlier experiments. In the long scale length regime a general measurement of the ion fluctuation wavenumber spectrum induced in the plane of the CO_2 laser electric field and high speed streak measurements of the Thomson scattered light were performed for the first time and are presented in this thesis.

The source of ion turbulence in laser produced plasmas is discussed, placing particular emphasis on ion waves generated by parametric instabilities. The general plasma dispersion equation is

solved numerically for the ion response in the presence of a strong pump to establish instability growth rates. Such analyses indicate that the Brillouin instability, filamentation and off-resonance parametric decay (PD) and oscillating two stream (OTS) instabilities can be easily excited and are believed to be the main source for the observed enhanced ion fluctuations in our experiment. In particular for fluctuations induced parallel to the electric field of the incident CO_2 laser pump wave, PD and OTS are potential sources of turbulence leading to anomalous absorption effects observed experimentally.

ACKNOWLEDGEMENT

I would like to express my sincere gratitude to my supervisor, Dr. A.A. Offenberger, for his excellent guidance, ideas, encouragement and financial support which made this work possible. I feel very fortunate to have been able to work with Dr. Offenberger.

I would like to extend particular thanks to Dr. W. Rozmus who provided many important discussions and comments on the whole thesis in general and the second chapter in particular. Without his invaluable ideas this chapter would not have been accomplished.

Ideas and advices of Dr. A. Ng during the early stage of this work as well as his comments on the first draft of the thesis are greatly appreciated. Discussion and ideas of Dr. R. Fedosejevs and Dr. D. McKen are also acknowledged.

I am very grateful to my sincere friend Dr. F. Soliman for spending much of his valuable time in reading through the thesis. His friendship and encouragement are greatly acknowledged. In addition, I am extremely grateful to Mrs. Soliman for offering her tremendous effort in the preparing and typing of the thesis.

This work would not have been possible without the technical assistance of the machine shop staff, in particular H. Gans, K. Doerrbecker and G. Soligo, and the optics technologist, B. Arnold. I am also grateful to D. Way-Nee and M. Billing for their technical assistance and advice.

I must also thank my fellow graduate students, R. Giles, W. Tighe, S. Au, M. Abdel-Halim, D. Simeoni and G. Philip for their friendship and co-operation.

Finally, I would like to extend my sincere thanks and appreciation to my wife, children and parents for their patience, understanding and prayer.

TABLE OF CONTENTS

	PAGE
CHAPTER 1 INTRODUCTION	1
CHAPTER 2 ION TURBULENCE IN LASER PLASMA INTERACTIONS	7
2.1 Ion Wave Instability	7
2.2 Ion Waves Induced by Parametric Processes	11
2.2.1 Introduction	11
2.2.2 Numerical solutions of the Drake dispersion relation.	14
2.2.3 Off-resonance oscillating two stream and parametric decay instabilities.	20
2.2.4 Saturation of ion waves.	28
2.3 Absorption Due to Ion Turbulence	34
CHAPTER 3 EXPERIMENTAL METHODS AND TECHNIQUES	40
3.1 Introduction	40
3.2 Laser Plasma Experiment	40
3.2.1 Carbon dioxide laser.	40
3.2.2 Gas jet and target chamber.	44
3.3 Measurements of Plasma Parameters	50
3.3.1 Infrared (IR) radiation detection.	50
3.3.2 Electron temperature (T_e).	52
3.3.3 Electron density (n) measurement.	54
3.4 Thomson Scattering Experiment	62
3.4.1 Outline of the experiment.	62
3.4.2 Scattering arrangements.	70

	PAGE
CHAPTER 4 EXPERIMENTAL RESULTS AND DISCUSSION	79
4.1 Introduction	79
4.2 Ion Turbulence and Absorption	88
4.2.1 Laser light absorption in CO_2 laser plasma interaction.	88
4.2.2 Scattering parameters.	92
4.2.3 Temporal measurements of ion fluctuations.	93
4.2.4 Spectral measurements.	95
4.2.5 Ion fluctuation level.	104
4.3 Results of Case II	108
4.3.1 Introductory remarks.	108
4.3.2 $S(\omega)$ measurements.	109
4.3.3 Streak measurements.	112
4.3.4 Sources of ion turbulence.	114
CHAPTER 5 SUMMARY AND CONCLUSION	128
APPENDIX A SOLUTION OF THE DISPERSION EQUATION	136
APPENDIX B MODELING OF THE CO_2 LASER	147
B.1 Gas Kinetics of the CO_2 - N_2 -He Mixture	147
B.2 Numerical Solution of the Rate Equations	151
APPENDIX C LABEL INVERSION	155
APPENDIX D NUMERICAL SOLUTION OF THE $S(\omega, \vec{r})$ FUNCTION	160
APPENDIX E DECONVOLUTION OF SPECTRA	163
REFERENCES	131

LIST OF TABLES

TABLE		PAGE
3.1	Features of the CO ₂ TEA Laser	42
3.2	Features of the Q-switched Ruby Laser	60

LIST OF FIGURES

FIGURE		PAGE
2.1	Electron and ion distribution functions in the presence of an electron heat flux \bar{Q} .	8
2.2	Growth rate and real frequency of ion wave for Brillouin scattering and filamentation versus k/k_D for oxygen plasma ($z=6$) with $T_i = T_e = 100$ eV, $n/n_c = 0.5$ and $v_0/v_e = 2$.	15
2.3	Growth rate and real frequency of ion wave for Brillouin scattering and filamentation versus k/k_D for oxygen plasma ($z=6$) with $T_i = T_e = 100$ eV, $n/n_c = 0.75$ and $v_0/v_e = 2$.	16
2.4	Growth rate and real frequency of ion wave for Brillouin scattering and filamentation versus k/k_D for oxygen plasma ($z=6$) with $T_i = T_e = 100$ eV, $n/n_c = 0.25$ and $v_0/v_e = 2$.	17
2.5	Growth rate and real frequency of ion wave for Brillouin scattering and filamentation versus k/k_D for oxygen plasma ($z=6$) with $T_i = T_e = 100$ eV, $n/n_c = 0.5$ and $v_0/v_e = 1$.	18
2.6	Growth rate of ion wave instability (PD off-resonance) versus k/k_D for oxygen plasma ($z=6$) with $T_i = T_e = 100$ eV, $n/n_c = 0.5$ and $v_0/v_e = 1$.	22
2.7	Growth rate of ion wave instability (PD off-resonance) versus k/k_D for oxygen plasma ($z=6$) with $T_i = T_e = 100$ eV, $n/n_c = 0.6$ and $v_0/v_e = 1$.	23
2.8	Growth rate of ion wave instability (PD and OTS off-resonance) versus k/k_D for oxygen plasma ($z=6$) with $T_i = T_e = 100$ eV, $n/n_c = 0.7$ and $v_0/v_e = 1$.	24

FIGURE	Y	PAGE
2.9	Growth rate of ion wave instability (PD and OTS off-resonance) versus k/k_D for oxygen plasma ($z=6$) with $T_i = T_e = 100$ eV, $n/n_c = 0.8$ and $v_0/v_e = 1$.	25
2.10	Growth rate of ion wave instability (PD and OTS off-resonance) versus k/k_D for oxygen plasma ($z=6$) with $T_i = T_e = 100$ eV, $n/n_c = 0.9$ and $v_0/v_e = 1$.	26
2.11	Thresholds of the most easily excited modes of PD and OTS off-resonance instabilities as a function of density for oxygen plasma ($z=6$) with $T_i = T_e = 100$ eV.	27
2.12	Growth rate of PD and OTS off-resonance instabilities versus k/k_D for oxygen plasma ($z=6$) with $T_i = T_e = 100$ eV, $n/n_c = 0.8$ and $v_0/v_e = 1, 2, 3$ and 4 .	29
2.13	The growth rate (of PD and OTS off-resonance) and the real frequency (of PD off-resonance) of the most easily excited ion modes in an oxygen plasma ($z=6$) with $T_i = T_e = 100$ eV, $n/n_c = 0.7$ and $v_0/v_e = 1$.	30
2.14	Real frequency of the ion waves of PD off-resonance instability as a function of k/k_D for an oxygen plasma ($z=6$) with $T_i = T_e = 100$ eV, $v_0/v_e = 1$ and $n/n_c = 0.5, 0.6, 0.7, 0.8$ and 0.9 .	31
2.15	The growth rate (of PD and OTS off-resonance) and the ratio γ/ω (of PD off-resonance) of the most easily excited ion modes versus T_i/T_e for an oxygen plasma ($z=6$) with $v_0/v_e = 1$ and $n/n_c = 0.7$.	33
2.16	$\text{Im} \frac{1}{\epsilon(\bar{k}_z, \omega_0)}$ averaged over a uniform spectrum of ion fluctuations as a function of density for a Maxwellian oxygen plasma with $T_i = T_e = 100$ eV.	38
3.1	Gain switched CO_2 laser output.	43
3.2	Output of the CO_2 laser model.	45

FIGURE		PAGE
3.3	Schematic diagram of the CO ₂ laser plasma interaction experiment and various diagnostics.	46
3.4	Gas target design.	48
3.5	Firing sequences of the CO ₂ laser.	49
3.6	Normalized x-ray intensity as a function of aluminum foil thickness. The circles are experimental data from the CO ₂ laser plasma experiment.	53
3.7	a) X-ray detector b) Typical x-ray signal for $T_e \ll 1$ keV.	55
3.8	Optical system for Fresnel's bimirror interferometer.	58
3.9	Subnanosecond switching of the ruby laser.	61
3.10	Typical fringe pattern with 25 micron fringe spacing.	63
3.11	Schematic diagram of the scattering geometry.	65
3.12	Thomson scattering experiment for $S(\omega, \vec{k})$ measurements.	73
3.13	Thomson scattering experiment for streak measurements.	75
3.14	Thomson scattering experiment for $S(\vec{k})$ measurements.	77
4.1	General behaviour of the CO ₂ laser plasma interaction: a) Transmitted CO ₂ laser signal, b) SBS signal, c) Thomson scattering signal.	80
4.2	Schematic diagram of the Thomson scattering experiment; a) Wave vectors at the incident, scattered and ion waves, b) Intersection of the CO ₂ laser beam, ruby laser beam and the collected light.	83
4.3	Unfolded interferogram of the plasma density of Case I at ~ 3 ns after the start of SBS.	85
4.4	Variation of the peak density ($r=0$) with the axial distance Z at ~ 3 ns after the start of SBS for Case I.	86
4.5	Unfolded interferogram of the plasma density of Case I at ~ 10 ns after the start of SBS.	87

FIGURE	PAGE
4.6 Unfolded interferogram of the plasma density of Case II at ~ 3 ns after the start of SBS.	89
4.7 Variation of the peak density ($r=0$) with the axial distance Z at ~ 3 ns after the start of SBS for Case II.	90
4.8 Calculated ruby Thomson scattering spectrum for a thermal oxygen plasma with $T_i = T_e = 100$ eV, $n = 0.8 n_c$, $z = 6$ and $\theta = 60^\circ$ showing the ion component (solid) and the electron component (dashed).	96
4.9 Calculated ruby Thomson scattering spectrum for a thermal oxygen plasma with $T_i = T_e = 100$ eV, $n = 0.4 n_c$, $z = 6$ and $\theta = 60^\circ$ showing the ion component (solid) and the electron component (dashed).	97
4.10 Variation of the ion component (of the ruby Thomson scattering spectrum) with ion temperature and degree of ionization for a thermal oxygen plasma with $T_e = 100$ eV, $n = 0.8 n_c$ and $\theta = 60^\circ$.	99
4.11 Spectra of scattered ruby laser light for $\theta = 60^\circ$ showing a) and b) unshifted typical spectra (for scattering in the X-Y plane), c) shifted spectrum for scattering in the Z-Y plane.	100
4.12 Spectrum of the scattered ruby laser light for $\theta = 22^\circ$ in the X-Y plane.	102
4.13 Ion turbulence spectrum $S(k)$ as a function of $k\lambda_D$ assuming an average density $n = 0.8 n_c$ and $T_e = 100$ eV (Case I). The Kadomtsev type spectrum is also shown (dashed).	107
4.14 Ion turbulence spectrum $S(k)$ as a function of $k\lambda_D$ assuming an average density $n = 0.7 n_c$ at $T_e = 100$ eV (Case II). The error bars indicate the standard deviations for thirty shots at average focused CO_2 laser intensity of $5 \times 10^{12} \text{w/cm}^2$.	111

FIGURE		PAGE
4.15	Schematic diagram of the Thomson scattering experiment to probe ion fluctuations induced by stimulated Brillouin backscattering.	113
4.16	Temporal behaviour of the Thomson scattered light (bottom) and the incident ruby light (top) in each plate showing different features; a) long lived fluctuations (~ 3 ns), b) short lived fluctuations (250 ps) with no structure, c) ~ 0.7 ns long fluctuations with fine structure.	115
4.17	Streak photograph of ~ 1.2 ns duration enhanced scattering event from ion turbulence occupying ~ 170 μm region.	117
4.18	Time evolution of the spatial distribution of turbulence for a 3 ns duration ($\theta = 16^\circ$ in the X-Y plane).	119
4.19	Time evolution of the spatial distribution of turbulence for a 0.8 ns duration ($\theta = 16^\circ$ in the X-Y plane).	120
4.20	Time evolution of the spatial distribution of turbulence showing a motion of the fluctuation region in the direction of the propagation of the CO_2 laser beam ($\theta = 16^\circ$ in the X-Y plane).	121
4.21	Numerical values of $S(k)$ versus k/k_D for oxygen plasma with $z = 6$ and $T_i = T_e = 100$ eV. Dashed curves correspond to $v_0/v_e = 0$. a) $v_0/v_e = 0.6$, $n/n_c = 0.6$, b) $v_0/v_e = 0.05$, $n/n_c = 0.8$.	125
A.1	Roots found using the bisection method.	137
B.2	Some of the low lying levels of the CO_2 and N_2 molecules.	148
C.1	Cross sectional view of a cylindrically symmetric plasma.	156
E.1	Effect of a finite slit on a spectrum. a) The spectrum as observed through a Gaussian slit of FWHM = 13 channels. b) The true spectrum. c) The recovery of the true spectrum from the observed one using the deconvolution technique.	166

CHAPTER 1

INTRODUCTION

A problem of crucial importance to laser fusion programs is how a laser light is absorbed by the plasma. It is believed that radiation can be absorbed in laser produced plasmas by different mechanisms depending on the laser parameters (intensity, wavelength and pulse duration) as well as the target material and configuration. These mechanisms may be classified into two types: (a) collisional absorption or inverse bremsstrahlung, (b) absorption due to collective effects in the plasma. In the inverse bremsstrahlung absorption (sometimes called classical absorption) an electron absorbs a photon as it moves from one free energy state to a more energetic state in the field of an ion. Absorption of the laser beam by a plasma will take place only in those regions where the density is lower than the critical density (at which the plasma frequency and the frequency of the incident radiation are the same). The absorbed fraction, A , calculated for the case of normal incidence of light on a plasma, the density of which varies linearly between $n = 0$ to a critical density n_c within a distance L , is given by [1]

$$A = 1 - \exp\left(-\frac{32}{15} \frac{L}{c} \nu_{ei}^c\right),$$

in which the electron-ion collisional frequency at the critical density may be written as

$$\nu_{ei}^c = 3 \times 10^{-6} z n_c \ln NT_e^{3/2},$$

where $\ln \Lambda$ is the usual coulomb logarithm ($\ln \Lambda = 5 - 10$), z is the average ionic charge and T_e is the electron temperature in eV. According to the above equations classical absorption is more significant for high values of z , low temperature plasma and short wavelength laser light (large n_c). For long wavelength lasers, such as CO_2 , and plasmas at temperatures of interest to laser-fusion, inverse bremsstrahlung mechanism is of less significance. It should also be pointed out that the absorption due to this mechanism decreases with profile steepening of the plasma near the critical density [2]. In addition, nonlinear effects and the disturbance in the plasma distribution function induced at higher laser intensities greatly reduce its effect [2,3].

Collective absorption, on the other hand, takes place as a result of various physical processes in the plasma. The most important of these processes are resonance absorption and parametric instabilities of the plasma. Resonance absorption occurs when the laser light is obliquely incident on a nonuniform plasma with the electric field of the incident wave being polarized in the plane of incidence (P-polarization). In this way the electric field perpendicular to the plane of the target can tunnel through from the point of reflection to the critical density, thus exciting an electron plasma wave. This wave, being electrostatic, cannot propagate outward and the energy is ultimately dissipated as heat in the plasma. The actual amount of resonance absorption of P-polarized light will depend on the angle of incidence θ and the density scale length. Contrary to inverse bremsstrahlung, resonance absorption is more significant at high plasma temperatures (high laser intensities), longer laser wavelengths and shorter plasma scale lengths.

The second type of collective absorption is that associated with parametric instabilities. The most important of these instabilities are: the oscillating two-stream instability, OTS; the parametric decay instability, PD (both occurring near the critical density); and the two-plasmon decay occurring near $n_c/4$. In these instabilities, called absorptive instabilities (to be distinguished from the scattering instabilities like stimulated Brillouin and Raman instabilities), the pump wave decays into two electrostatic waves. The electron plasma wave, which is one of the products, continues to grow at the expense of the pump wave until it saturates by pump depletion or any other non-linear mechanism. These high frequency oscillations, in turn, accelerate and heat the electrons. The effective collision frequency ν^* or the so-called anomalous collisionality (defined as the rate of transferring wave energy to particles) can be estimated quantitatively according to the following energy balance scheme:

Pump wave \rightarrow Electron plasma waves \rightarrow Plasma particles .

At saturation the effective collision frequency for OTS and PD instabilities is given by |4|

$$\nu^* \approx 4\gamma \left(\frac{\omega_{pe}}{\omega_0} \right)^2 = 4\gamma \frac{n}{n_c} \approx 4\gamma$$

where γ is the growth rate of the instability, ω_0 is the laser light frequency and ω_{pe} and n are the electron plasma frequency and density respectively. The parametric instabilities are intrinsically non-linear processes and will not be induced below a threshold light intensity (unlike resonance absorption which may occur at any laser light intensity). The threshold values depend on density scale length, plasma temperature, laser wavelength and substantially increase with the

inhomogeneity in the plasma. Although this mechanism is less important than resonance absorption it can still compete with inverse bremsstrahlung at higher intensities and longer laser wavelength. One of the disadvantages of this mechanism, as well as the resonance absorption mechanism, is the production of very energetic electrons which carry a portion of the absorbed energy, resulting in a very weak coupling with the background thermal plasma and thus an energy loss. In addition, these fast electrons may penetrate into and preheat the pellet core in laser-fusion pellets and raise its entropy which makes high compression difficult to achieve. For this reason, inverse bremsstrahlung is still considered the most favourable absorption mechanism in laser fusion.

It was pointed out by Dawson and Oberman [5] that absorption of radiation by a plasma can be enhanced in the presence of coherent short wavelength ion fluctuations. Recently [6-8] the interest in this subject has tremendously increased. Faehl and Kruer [6] derived an expression relating the damping rate of laser energy to the ion fluctuation level in a plasma (Chapter 2). According to this relation the damping rate could be of the same order or even higher than the classical collision frequency, ν_{ei} , in the presence of high level coherent ion fluctuations. Manheimer and Colombant [9] continued their investigations in an attempt to correlate experimental results on absorption of high intensity laser light by plasma with previous theories. These investigations clearly indicated that there is a discrepancy between theory and experiment. This led them to believe that ion turbulence is a possible candidate as an absorption mechanism that causes such a discrepancy.

The actual source and magnitude of ion fluctuations in laser plasma interactions remains somewhat hypothetical. Manheimer and his

colleagues [7-10] have postulated that the turbulence is produced by heat flow driven ion acoustic instabilities. Faehl and Kruer [6] and Forslund [11] have suggested that ion-ion streaming instabilities could create a high level of ion fluctuations. Moreover, there is only limited theoretical knowledge of the ion fluctuation spectra which may be generated by instabilities other than the current driven ion wave instability.

On the experimental side, ion turbulence has been measured directly in several CO₂ laser plasma experiments. Gray and Kilkenny [12] investigated the possible influence of ion turbulence in reducing thermal conductivity in a Z-pinch plasma heated by moderate intensity CO₂ laser radiation. Walsh and Baldis [13] and Giles and Offenberger [14] studied ion fluctuations associated with Brillouin scattering in high intensity CO₂ laser plasma interactions. Although substantial enhancements in the fluctuation level over thermal noise have been seen in these experiments no general measurement of the fluctuation level and spectrum has been made for high intensity laser produced plasmas and, in particular, for wave vector \vec{k} parallel to the direction of the incident electric field which is of importance for absorption (Chapter 2).

The previous survey clearly indicated that ion turbulence, if understood may provide an explanation for some of the physical processes that occur during laser/plasma interactions. This being the case, it appeared to us that ion turbulence in general, and its role in absorption of laser radiation in laser heated plasma experiments in particular, are two interesting topics for research. The main objective of this research is two fold; one is to study the cause as well as the

characteristics of ion fluctuations in high intensity laser produced plasmas; the other is to confirm the existence of ion turbulence as an absorption mechanism using temporal and spectral analysis.

In Chapter 2 a summary of ion wave instabilities, placing particular emphasis on parametric instabilities is first given. Numerical solutions of the plasma dispersion equation in the presence of a high intensity laser beam is carried out and the results are given. Curves illustrating the dependence of the growth rate of ion wave related parametric instabilities on different plasma parameters are presented. Finally the theory of light absorption by an ion turbulence mechanism is also highlighted.

Chapter 3 provides a detailed description of the laser plasma experimental setup and the various diagnostic techniques utilized for studying ion fluctuations.

In Chapter 4 the results of our experimental investigations as well as interpretation of these results are given. The measurements of the plasma parameters and basic characteristics are first presented. Results of temporal and spectral measurements and turbulence levels of the ion fluctuations (from Thomson scattering measurements) are provided with an estimation of the laser light being absorbed via ion turbulence. Measurements of the spectral form factor $S(k)$ and streak camera measurements of the ion fluctuations are also presented. Finally a discussion of the possible source of ion fluctuations in the present experiment is given.

Conclusions and suggestions for further investigations are given in Chapter 5.

CHAPTER 2

ION TURBULENCE IN LASER PLASMA INTERACTIONS

2.1 Ion Wave Instability

It has been shown theoretically that ion turbulence may explain the nonclassical plasma behaviour in absorption of laser radiation [1-3], heat transport of absorbed energy [4,5] and generation of energetic particles in laser plasma interactions [6]. The magnitude of these effects depends significantly on both turbulence level and spectrum. Although high fluctuation levels ($\delta n/n \sim 0.1$) have been measured in laser plasma experiments [7-12], the actual source of ion turbulence is not well understood. Manheimer and his colleagues [1] suggested that ion turbulence may be produced by heat flow driven acoustic instabilities; a mechanism that is similar to the well known current driven ion acoustic instability mechanism [13]. For high electron to ion temperature ratio ($zT_e/T_i \gg 1$), ion acoustic waves become weakly Landau damped and their phase velocity becomes much greater than the ion thermal velocity v_{ti} . This means that any distortion in the electron distribution function will easily drive these ion waves unstable. Such a distortion may occur in the presence of a strong electromagnetic wave. Figure 2.1 shows the electron and ion distribution functions for $T_e/T_i \gg 1$ in which the laser beam is propagating in the negative x direction. The laser energy flux absorbed by electrons will be converted into thermal energy flux (heat flux) \bar{Q} . As can be seen in the figure the electron distribution is distorted for $v_x < 0$ because

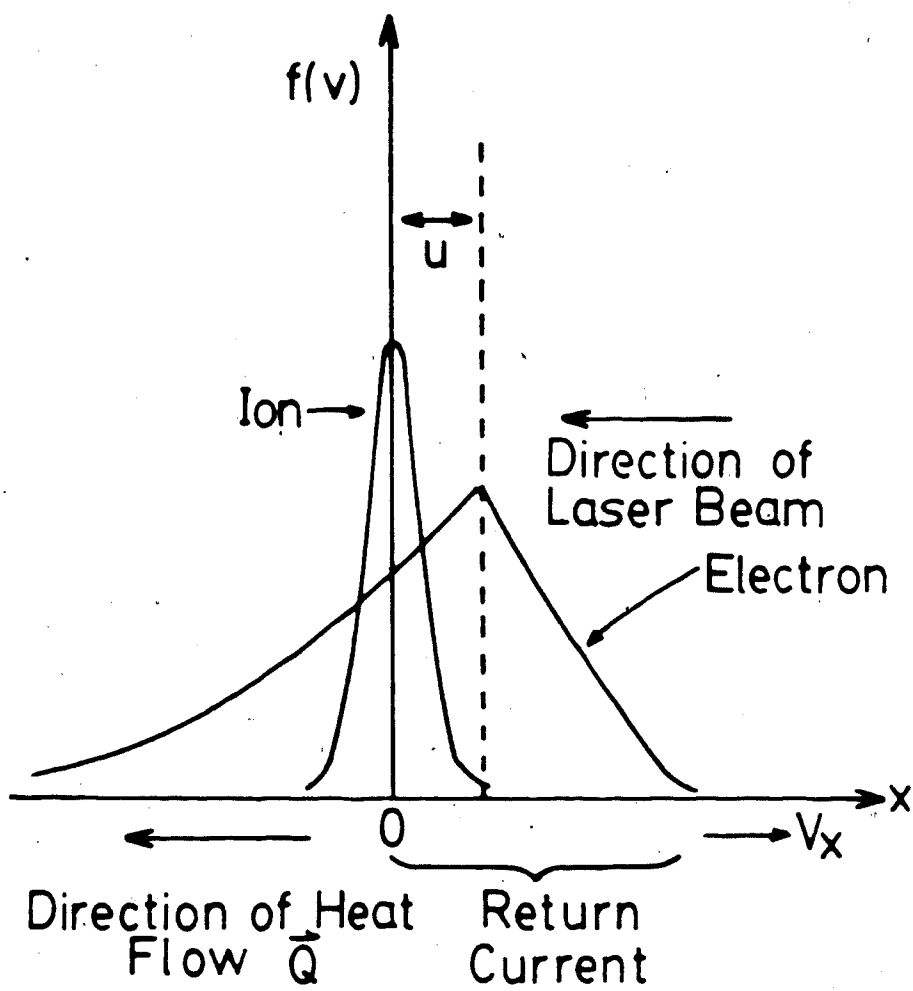


Fig. 2.1 Electron and ion distribution functions in the presence of an electron heat flux \vec{Q} .

heat is being carried in the $-X$ direction by electrons with speed $|u_x| > v_e$. In order to maintain charge neutrality a return current of colder electrons must flow in a direction opposite to the direction of the laser light propagation (i.e., $+X$ direction). Thus the electron distribution function for $v_x > 0$ is skewed to the right. In addition, the peak of the electron distribution function occurs at $v_x = +u$ rather than at $v_x = 0$. The ion waves will therefore be driven unstable if $u > c_s$ and $c_s > v_i$ (i.e., $zT_e > T_i$), where c_s is the sound speed and v_i is the ion thermal speed. The wave vector \vec{k} of the ion acoustic waves is antiparallel to the heat flux \vec{Q} . It is worth mentioning that the fluctuation level associated with this mechanism was found to saturate by ion trapping in the range $0.1 < \delta n/n < 0.2$ within a time of order $t \sim 10^3/\omega_{pe}$ (~ 5 p sec for CO_2 laser), where ω_{pe} is the electron plasma frequency [6]. Moreover, these fluctuations seem to be nearly uniformly distributed within a wide cone of an angle $\theta \sim 50 - 60^\circ$ around the direction of the return current. It is therefore expected that no significant fluctuations will be generated in the directions perpendicular to the direction of the laser beam.

Another mechanism that could be responsible for generating ion turbulence in laser produced plasmas is the ion-ion streaming instability proposed by Faehl and Kruer [3]. Such an instability occurs when different ion species are available (ions of different masses or different ionization degrees). The physics of this mechanism is easy to understand. When a plasma is heated a large ambipolar potential develops to prevent the hot electrons from leaving the target. Ions are then accelerated by this potential and also by the large ponderomotive potentials near the critical density. At low densities, where

ion-ion collisions are weak, separation of the different ion species takes place (due to the different expansion velocities). A simple estimate shows that this separation is obtained when $\nu_{ii} < v_i/L$, where ν_{ii} is the ion-ion collision frequency, v_i is the ion speed which the ions obtain by falling through the large potential, and L is the distance over which it occurs. Results of a simple simulation code [3] and others of a two-dimensional simulation code [14] show that a fluctuation level of $\sim 10\%$ could be reached in a plasma of two ion species such as H and D_2 . In real plasmas this instability may be severely restricted, especially in the hot region where almost all the ions are ionized to the highest possible level and ions of identical masses are used.

The third possible mechanism for generating ion turbulence in laser produced plasmas is the ion wave instability induced by different parametric processes. It is well established that ion waves can be excited via stimulated Brillouin instability in the underdense plasma and the $e-i$ decay and oscillating two stream instabilities near the critical density. Because of the small value of k for ion fluctuations induced by the Brillouin instability, the enhancement in classical absorption is of no importance. This will be explained in a later section. On the other hand, very little attention has been paid to other parametric instabilities which can excite ion waves in underdense plasmas and, particularly, for $k\lambda_D \sim 0(1)$ (where λ_D is the Debye length). In the next section the importance of parametric instabilities in generating ion turbulence will be studied in more detail.

2.2 Ion Waves Induced by Parametric Processes

2.2.1 Introduction.

Parametric processes refer to the nonlinear couplings of the electromagnetic wave with the natural wave modes in the plasma. These modes subsequently grow at the expense of the driving electromagnetic field which is called the pump. Such processes can only occur when the pump intensity exceeds a threshold value. To simply explain how the parametric instability takes place, assume the plasma is driven by a field (of a finite amplitude) which oscillates at a frequency ω_0 . If the plasma has an oscillating mode at a frequency ω (the initial amplitude of which may be as low as the noise level) which can couple to the pump at ω_0 then these two oscillations will beat together to produce sideband modes at frequencies $\omega_{\pm} = \omega_0 \pm \omega$. If the difference frequency ω_- happens to be another lightly damped mode of the plasma, it will beat with ω_0 at $\omega = \omega_0 - \omega_-$ and $\omega' = \omega_0 + \omega_-$. Thus the mode at ω is enhanced and will beat more strongly with ω_0 to give a stronger ω_- mode. It should be born in mind that for conservation of both energy and momentum (assuming the pump as a quantum being converted into quanta of other modes) it is required that the following conditions must be satisfied

$$\omega_0 = \omega + \omega_-, \quad \bar{k}_0 = \bar{k} + \bar{k}_-. \quad (2.1)$$

In laser plasma interactions the laser light acts as a transverse electromagnetic wave pump (t). The excited mode in this case may be another transverse mode (t'), a longitudinal Langmuir electron wave (ℓ) or an ion sound wave (s). The possible mode couplings are:

$t \rightarrow \ell + s$	"Parametric" Instability
$t \rightarrow t' + s$	Stimulated Brillouin Scattering
$t \rightarrow t' + \ell$	Stimulated Raman Scattering
$t \rightarrow \ell + \ell$	Two Plasmon Instability .

In parametric and two-plasmon instabilities laser light energy can be absorbed efficiently under certain conditions. On the other hand, in stimulated Raman and Brillouin scattering part of the laser energy will be lost in the form of sideband electromagnetic waves (t'). The loss is more pronounced in Brillouin scattering simply because the ion wave mode involved in this process takes up only a small fraction of the laser energy.

Since we are merely concerned with parametric processes which can excite low frequency ion modes, it will be assumed that the high frequency electromagnetic pump (\bar{k}_0, ω_0) decays into a low frequency ion mode (\bar{k}, ω) and two high frequency waves ($\bar{k}_\pm = \bar{k} \pm \bar{k}_0, \omega_\pm = \omega \pm \omega_0$). The sideband modes (\bar{k}_\pm, ω_\pm), which are generated through a nonlinear coupling between the low frequency perturbation (\bar{k}, ω) and the pump wave, will interact in turn with the pump wave and therefore create a ponderomotive force which reacts back with the (\bar{k}, ω) mode. According to Drake, *et al.* [15], the ponderomotive force is given by

$$\bar{F}_p = \frac{e^2}{m\omega_0^2} (\bar{E}_{0+} \cdot \bar{E}_- + \bar{E}_{0-} \cdot \bar{E}_+) , \quad (2.2)$$

where $\bar{E}_{0\pm} = \bar{E}_0(\pm\omega_0, \pm\bar{k}_0)$ refer to the components of the pump electric field and \bar{E}_\pm refer to the sideband waves' electric field.

The derivation of the generalized dispersion relation is based on the insertion of Eqn.(2.2) into the Maxwell-Vlasov equations of the

plasma [16,17]. This dispersion relation for $(\omega_0/\omega_{pe}) \gg 1$ and $k\lambda_0 \ll 1$ is given by

$$\frac{1}{\chi_e(\bar{k})} - \frac{1}{\chi_i(\bar{k},\omega)+1} = k^2 \left[\frac{|\bar{k}_- \times \bar{v}_0|^2}{k_-^2 D_-} - \frac{(\bar{k}_- \cdot \bar{v}_0)^2}{k_-^2 \omega^2 \epsilon_-} + \frac{|\bar{k}_+ \times \bar{v}_0|^2}{k_+^2 D_+} - \frac{(\bar{k}_+ \cdot \bar{v}_0)^2}{k_+^2 \omega^2 \epsilon_+} \right], \quad (2.3)$$

where χ_e and χ_i are the usual electron and ion susceptibilities defined for a Maxwellian plasma by

$$\chi_{e,i}(\bar{k},\omega) = \frac{T_e}{T_{e,i}} \frac{1}{k^2 \lambda_D^2} \left| 1 + \frac{\omega}{k v_{e,i}} Z\left(\frac{\omega}{k v_{e,i}}\right) \right|, \quad (2.4)$$

$$Z(y) = \frac{1}{\sqrt{\pi}} \int_{-\infty}^{\infty} \frac{dx \exp(-x^2)}{x-y}$$

where $v_{e,i}$ are electron and ion thermal speeds, λ_D is the Debye length,

$$v_0 = eE_0/m\omega_0, \quad \epsilon_{\pm} = 1 - \frac{\omega^2}{\omega_{\pm}^2} \text{ and } D_{\pm} = c^2 k^2 \pm 2\bar{k} \cdot \bar{k}_0 c^2 + 2\omega_0 \omega - \omega^2.$$

The $\bar{k} \times \bar{v}_0$ terms in Eqn. (2.3) arise from the electromagnetic components of the sideband modes and the $\bar{k}_0 \cdot \bar{v}_0$ terms from the electrostatic components. Inspection of Eqn. (2.3) reveals the following:

i) The $\bar{k}_0 \cdot \bar{v}_0$ terms will be predominant only near the critical plasma density region ($\epsilon_{\pm} \approx 0$ or $\omega_{pe} \approx \omega_0$) which is of no interest in this study.

ii) For $D_{\pm} \approx 0$, the permittivity $\epsilon_{\pm} \neq 0$ and the $\bar{k}_0 \cdot \bar{v}_0$ terms (the electrostatic terms) in the equation are nonresonant and therefore negligible. In this case, the high frequency sidebands are predominantly electromagnetic and the problem is essentially that of

stimulated scattering where the pump wave excites an electrostatic wave and an electromagnetic wave at a shifted frequency. If we limit ourselves to the case where the electrostatic mode is an ion mode ($\omega \ll \omega_0$) then $\omega_{\pm} \approx \omega_0$ and $\bar{k}_{\pm} \cdot \bar{E}_0 \sim 0$ and Eqn. (2.3) can be reduced to the form

$$\epsilon(\bar{k}, \omega) D_+(\bar{k}, \omega) D_-(\bar{k}, \omega) = \frac{k^2 v_0^2}{4} \chi_e(\bar{k}, \omega) |1 + \chi_i(\bar{k}, \omega)| \times |D_+(\bar{k}, \omega) + D_-(\bar{k}, \omega)|, \quad (2.5)$$

where ϵ is the plasma dielectric constant ($= 1 + \chi_e + \chi_i$).

Equation (2.5) describes stimulated Brillouin scattering (SBS) and filamentation (a purely growing mode, i.e., Real $\omega = 0$, occurs when D_+ and D_- are comparable).

2.2.1 Numerical solutions of the Drake dispersion relation.

The numerical technique used for the solution of the Drake dispersion relation as given by Eqn. (2.5) is discussed in Appendix A. Results showing the dependence of both the growth rate γ and the real frequency component ω_R (where $\omega_0 = \omega_R + i\gamma$) of the ion wave instability in an oxygen plasma ($z = 6$, $T_i = T_e = 100$ eV) on k/k_D , with n/n_e and v_0/v_e taken as parameters are given in Figs. 2.2-2.5 (where $k_D = 1/\lambda_D$). As shown in the figures results are plotted for three different angles θ , where θ is the angle between \bar{k} and \bar{E}_0 (where $\theta = 0, 45$ and 90° correspond respectively to Brillouin backscattering, Brillouin side-scattering and filamentation).

According to Fig. 2.2 ($n/n_e = 0.5$, $v_0/v_e = 2$) for $\theta < 90^\circ$ there are narrow regions of k/k_D where γ and ω_R are approximately of the same order and within which the maximum growth rates are located. For

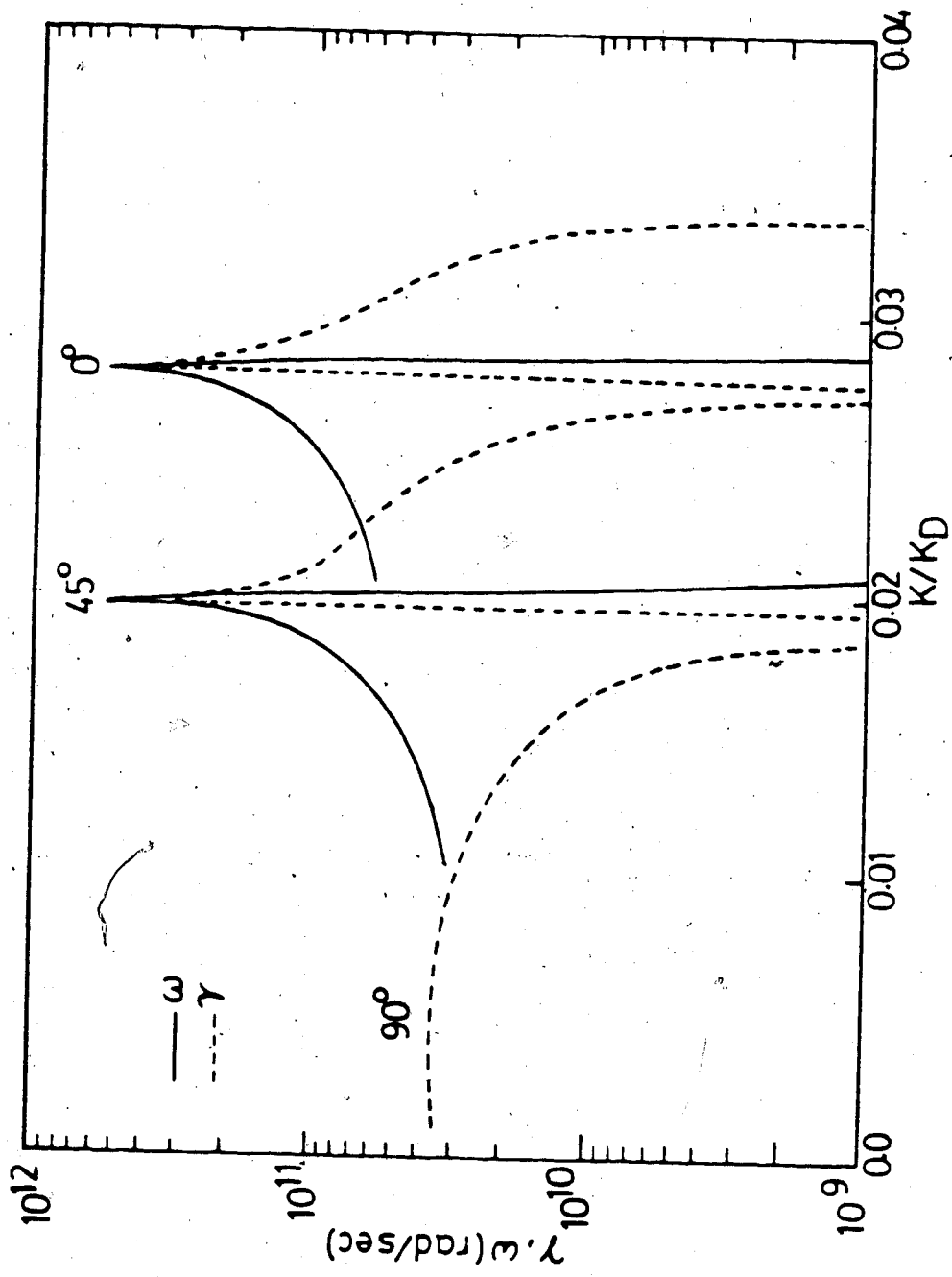


Fig. 2.2 Growth rate and real frequency of ion wave for Brillouin scattering and filamentation versus k/k_D for oxygen plasma ($z=6$) with $T_e = T_i = 100$ eV, $n/n_c = 0.5$ and $v_0/v_e = 2$.

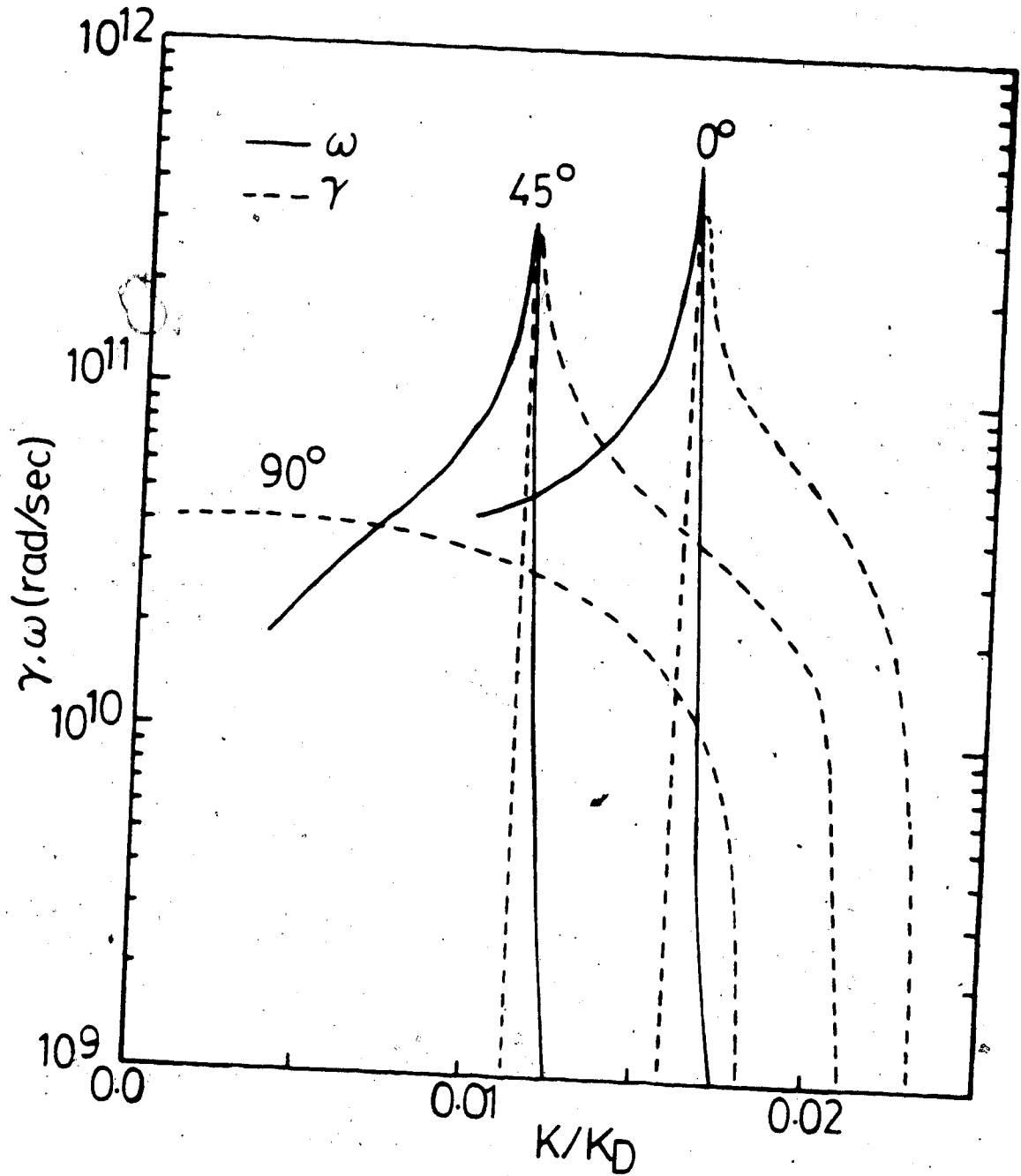


Fig. 2.3 Growth rate and real frequency of ion wave for Brillouin scattering and filamentation versus k/k_D for oxygen plasma ($z=6$) with $T_i = T_e = 100$ eV, $n/n_e = 0.75$ and $v_0/v_e = 2$.

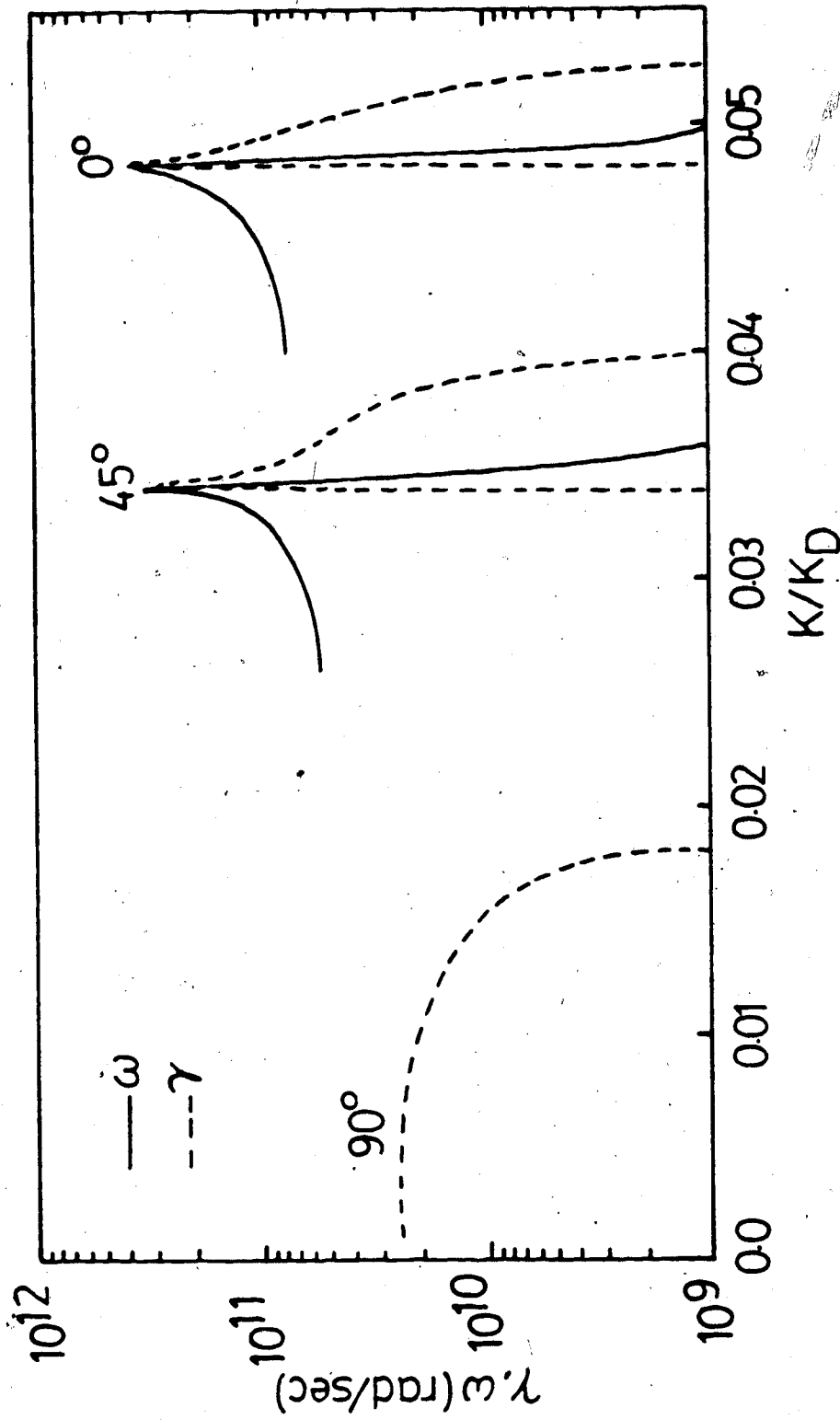


Fig. 2.4 Growth rate and real frequency of ion wave for Brillouin scattering and filamentation versus k/k_D for oxygen plasma ($z=6$) with $T_i = T_e = 100$ eV, $n/n_c = 0.25$ and $v_0/v_e = 1.2$.

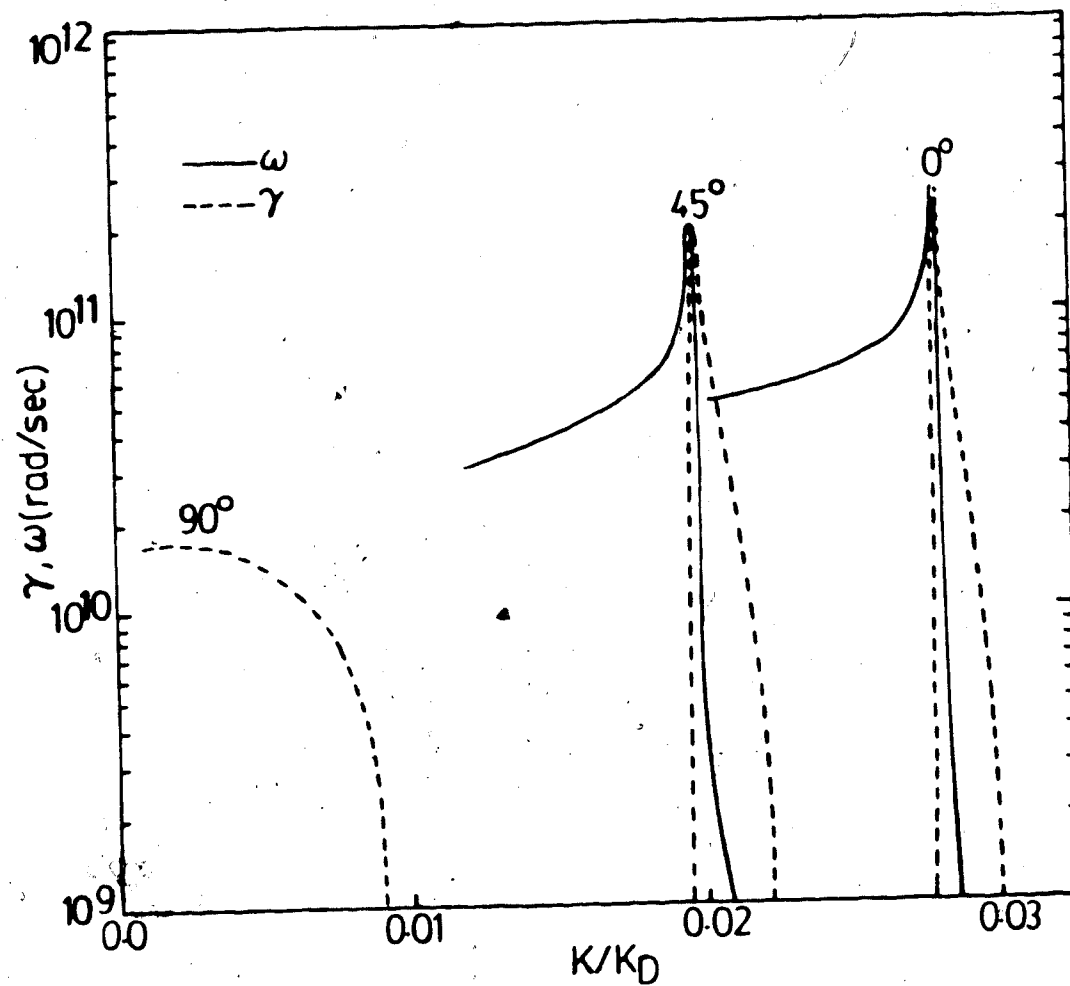


Fig. 2.5 Growth rate and real frequency of ion wave for Brillouin scattering and filamentation versus k/k_D for oxygen plasma ($z=6$) with $T_i = T_e = 100$ eV, $n/n_c = 0.5$ and $\nu_0/\nu_e = 1$.

smaller values of k , i.e., to the left of these maxima γ drops to zero rapidly while w_R decreases slowly before it approaches a constant value. To the right of the maxima w_R drops faster than γ and the mode grows almost purely ($w_R = 0$) for certain ranges of k/k_D . The growth rate decreases slowly over an order of magnitude range as θ increases from 0° to 90° . The instability covers a small range of k/k_D (~ 0 to 0.03) whereas the filamentation alone covers the range $0 \leq k/k_D \leq 0.02$. This range of k/k_D has no significant role in enhancing absorption.

Figures 2.3 and 2.4 are similar to Fig. 2.2 but with $n/n_c = 0.75$ and 0.25 respectively. It is clear from these figures that the k/k_D range of the ion fluctuations induced by SBS increases with density, while the k/k_D range of the ion fluctuations induced by filamentation stays constant. In addition, the growth rate of the most highly excited mode of Brillouin scattering remains fairly constant in the range $0.2 \leq n/n_c \leq 0.75$, while for filamentation the growth rate increases slowly with density.

Figure 2.5 is similar to Fig. 2.2 except v_0/v_e was reduced to 1. This figure shows, in addition to the reduction in the growth rate, shorter ranges of k/k_D for each individual angle with the most highly excited modes remaining at the same k/k_D .

It was also found from the solution of Eqn. (2.5) that both the Brillouin scattering and filamentation were insensitive to the temperature ratio, at least in the range $0.25 \leq T_i/T_e \leq 4$. Furthermore, these instabilities have v_0/v_e thresholds of order 0.1 for the lowest threshold excited modes for $n/n_c = 0.5$. This threshold value is quite low and is equivalent to a CO_2 laser intensity of $\sim 2.4 \times 10^{10}$ W/cm².

As was pointed out earlier and noticed in Figs. 2.2 to 2.5, the above instabilities (SBS and filamentation) occur in the $k\lambda_D \ll 1$ region which is of no interest for absorption. However, these instabilities may be important in explaining the general ion fluctuation k -spectrum.

2.2.2 Off-resonance oscillating two stream and parametric decay instabilities.

Other parametric instabilities, which involve ion modes and may occur in the subcritical density region, are the oscillating two stream (OTS) and the parametric (electron-ion) decay (PE) off-resonance instabilities. In these instabilities, similar to the usual OTS and PE instabilities, the pump decays into ion wave (with $\omega_D = 0$ in the case of OTS) and electron wave. Unfortunately, for the parameters $\omega_{pe} \lesssim \omega_0$, $k\lambda_D \approx 0(1)$ and a pump strength $\omega_0/v_e \approx 0(1)$, the general dispersion relation of Drake *et al.* (2.3) is not valid. However, another approach to the problem of the instability induced by a strong pump field has been carried out by Silin [17] and Sanmartin [18]. Although the solution is limited to electrostatic sidebands, it is quite general for an arbitrary pump strength and a wide range of k/k_D . The general dispersion relation for a uniform high frequency field and a homogeneous and an unmagnetized plasma, derived from Vlasov-Poisson equations using Fourier and Laplace transformations, is given by [18]

$$1 + \chi_i^0 \left| \frac{\nu_0^2}{D_e^0} + \sum_{j=1}^{\infty} \nu_j^2 \left(\frac{1}{D_e^j} + \frac{1}{D_e^{-j}} \right) \right| = 0, \quad (2.6)$$

where $D_\alpha^j = 1 + \frac{\nu_\alpha^j}{\omega_\alpha}$ (α stands for i or e),

$$X_{\alpha}^j = \frac{\omega^2}{k^2} \int \frac{\bar{k} \cdot (\partial F_{\alpha 0} / \partial \bar{u}_{\alpha}) d\bar{u}_{\alpha}}{\omega - j\omega_0 - \bar{k}_0 \cdot \bar{u}_{\alpha}} \quad (2.7)$$

where $F_{\alpha 0}$ is the zero order velocity distribution function of the particle α , $\omega_{p\alpha}^2 = (4\pi q_{\alpha}^2 n_{\alpha} / m_{\alpha})$ and q_{α} and n_{α} are the α particle charge and density respectively. In Eqn. (2.7) the particle velocity \bar{v} in the distribution function has been transformed to $\bar{u} = \bar{v} + \omega_0 \bar{\epsilon}_{\alpha} \cos \omega_0 t$ with $\bar{\epsilon}_{\alpha} = q_{\alpha} \bar{E}_0 / m_{\alpha} \omega_0^2$. The argument of the Bessel function J in (2.6) is $\bar{k} \cdot \bar{\epsilon} = k \frac{\partial E_0}{m_e \omega_0^2} \cos \theta$ where θ is the angle defined before.

Equation (2.6) was previously analyzed by many authors [17-19] and its solution for unstable ion modes in the underdense plasma was obtained for certain conditions. For a more general case and for conditions of relevance to our experiment, Eqn. (2.6) has been solved numerically (Appendix A) for different values of the parameters (k/k_D) , (n/n_e) , (v_0/v_e) , (T_i/T_e) and $\cos \theta$.

The solution of Eqn. (2.6) as illustrated in Figs. 2.6-2.10 indicates that an unstable ion wave can be excited for $n/n_e \geq 0.5$ and $k/k_D \lesssim 0.5$. These figures also reveal the existence of two types of ion waves, the off-resonance PD and OTS. As can be seen in these figures, at low densities ($0.5 \lesssim n/n_e < 0.7$) only PD off-resonance can be excited. However, at $n/n_e \sim 0.7$ the OTS off-resonance starts to take place and then dominates at higher densities (Fig. 2.10). The thresholds of the most easily excited modes of these instabilities vary between $v_0/v_e \sim 0.02$ to ~ 1.1 , depending on the density region (Fig. 2.11). These thresholds are considered low and could be easily achieved in our experiment. Figure 2.11 also shows that the off-resonance PD instability has a lower threshold than the OTS instability.

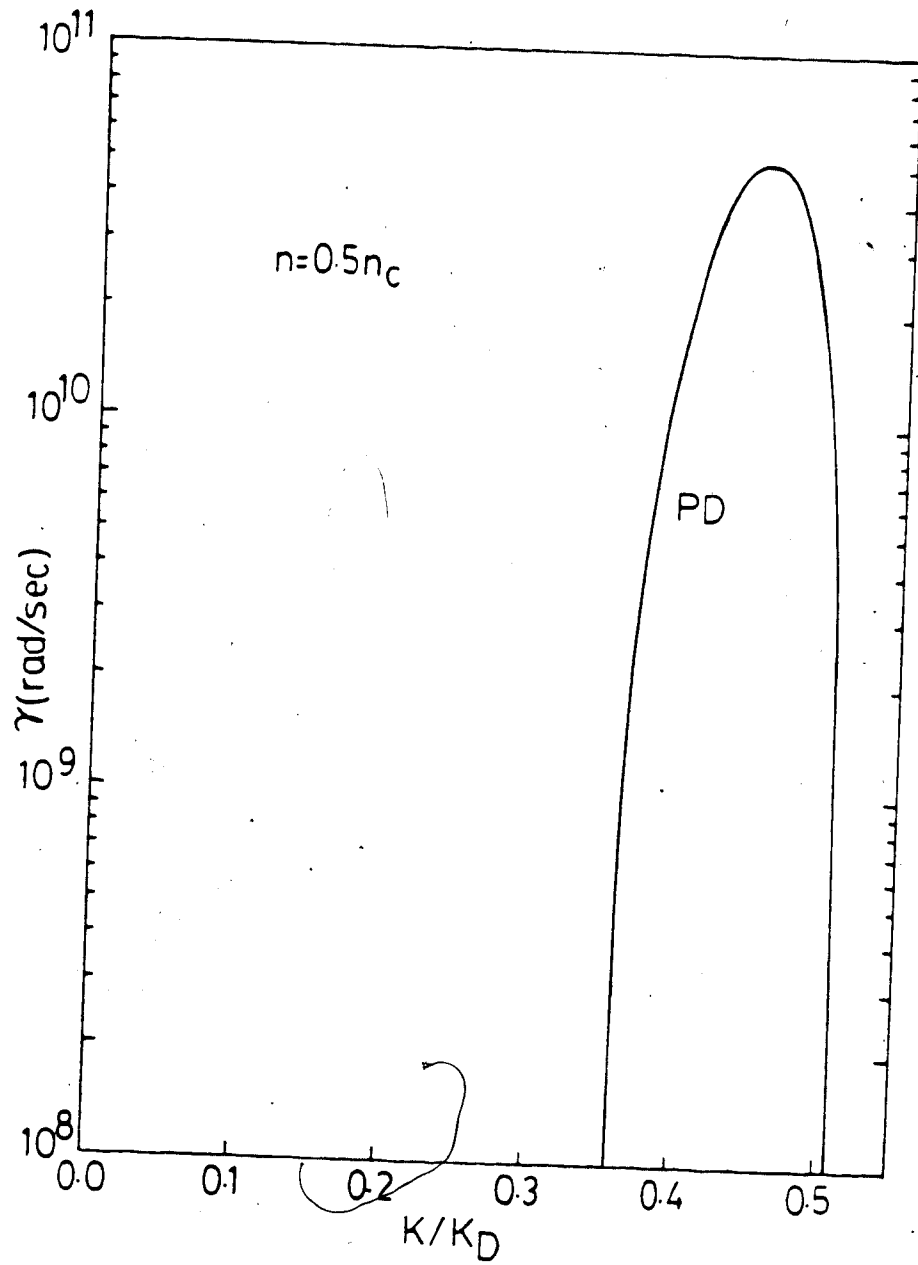


Fig. 2.6 Growth rate of ion wave instability (PD off-resonance) versus k/k_D for oxygen plasma ($z=6$) with $T_i = T_e = 100$ eV, $n/n_c = 0.5$ and $v_0/v_e = 1$.

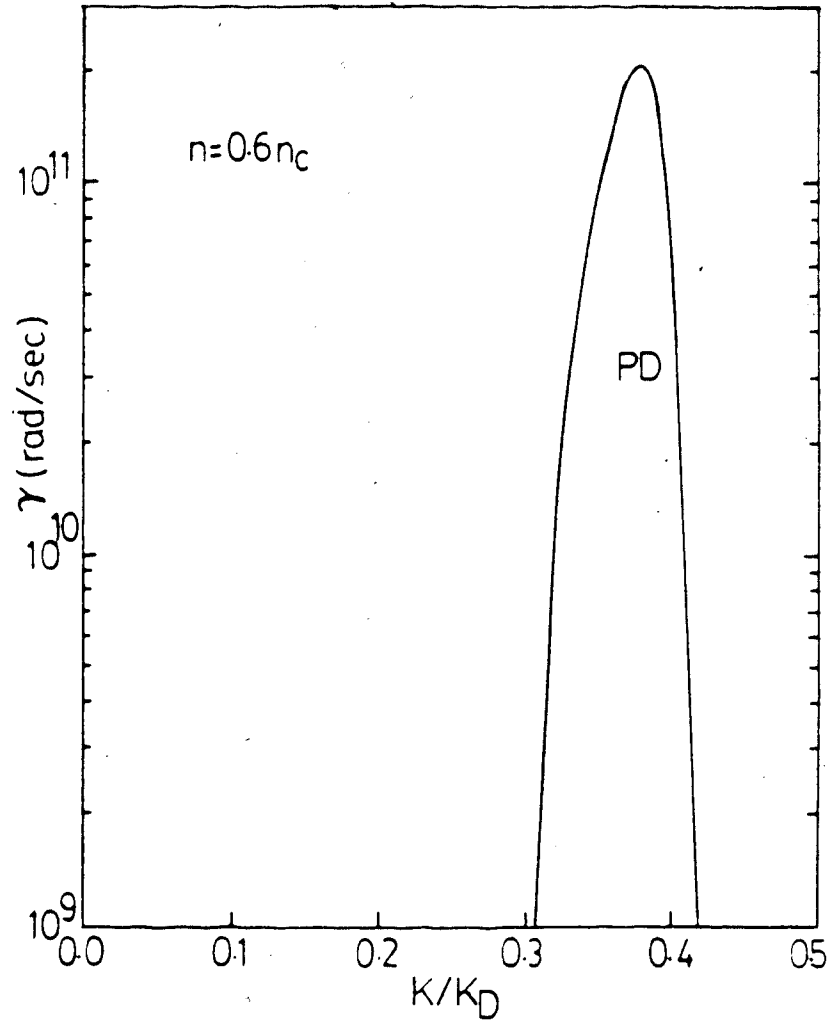


Fig. 2.7 Growth rate of ion wave instability (PD off-resonance) versus k/k_D for oxygen plasma ($z=6$) with $T_i = T_e = 100$ eV, $n/n_c = 0.6$ and $v_0/v_e = 1$.

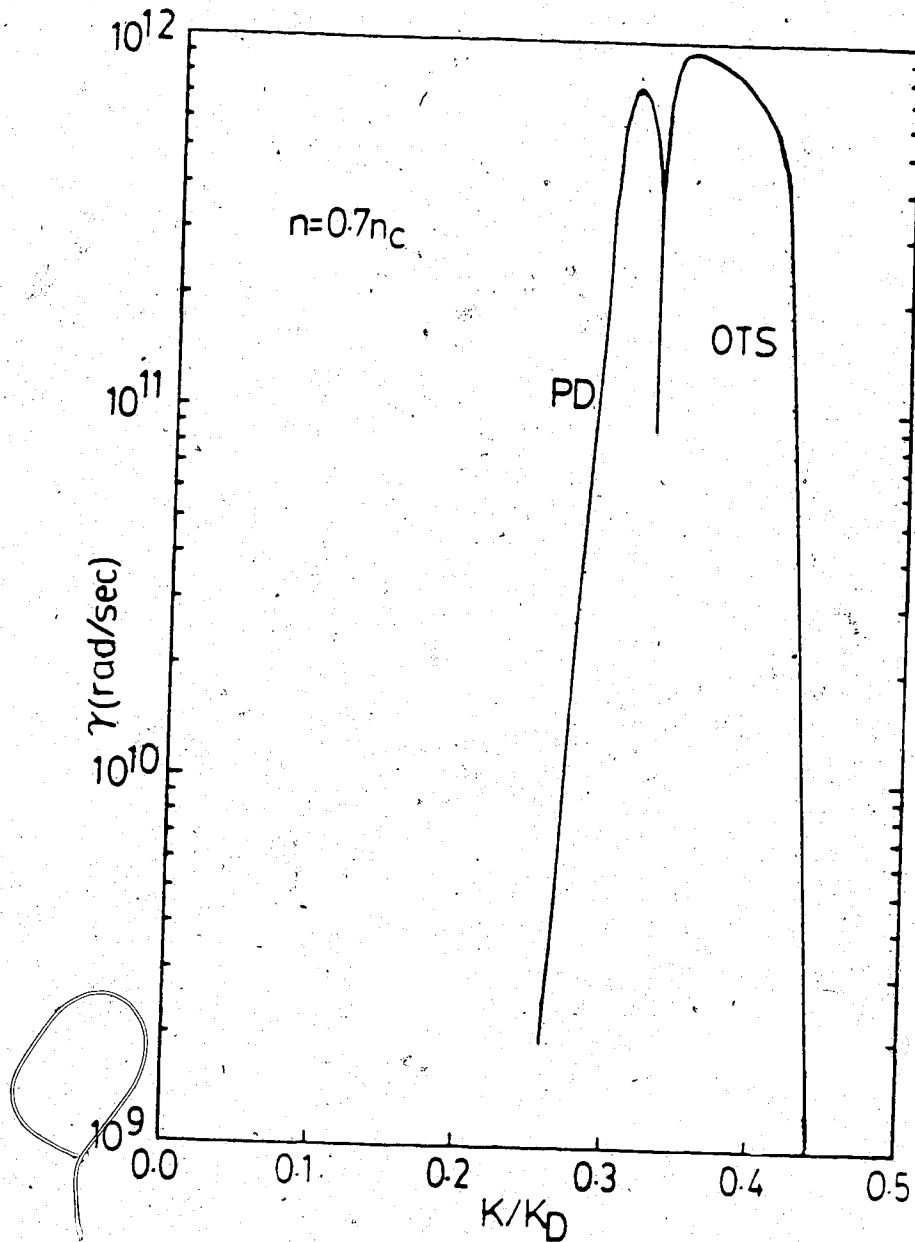


Fig. 2.8 Growth rate of ion wave instability (PD and OTS off-resonance) versus k/k_D for oxygen plasma ($z=6$) with $T_i = T_e = 100$ eV, $n/n_c = 0.7$ and $v_0/v_e = 1$.

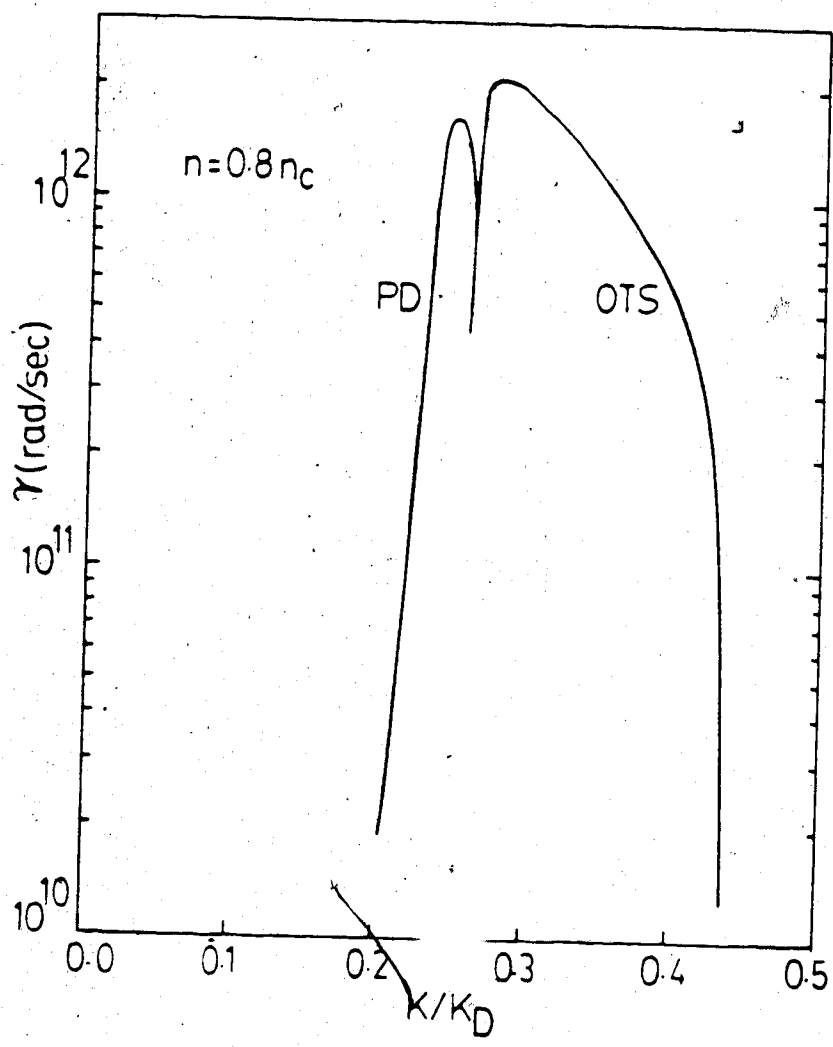


Fig. 2.9 Growth rate of ion wave instability (PD and OTS off-resonance) versus k/k_D for oxygen plasma ($z=6$) with $T_i = T_e = 100$ eV, $n/n_e = 0.8$ and $v_0/v_e = 1$.

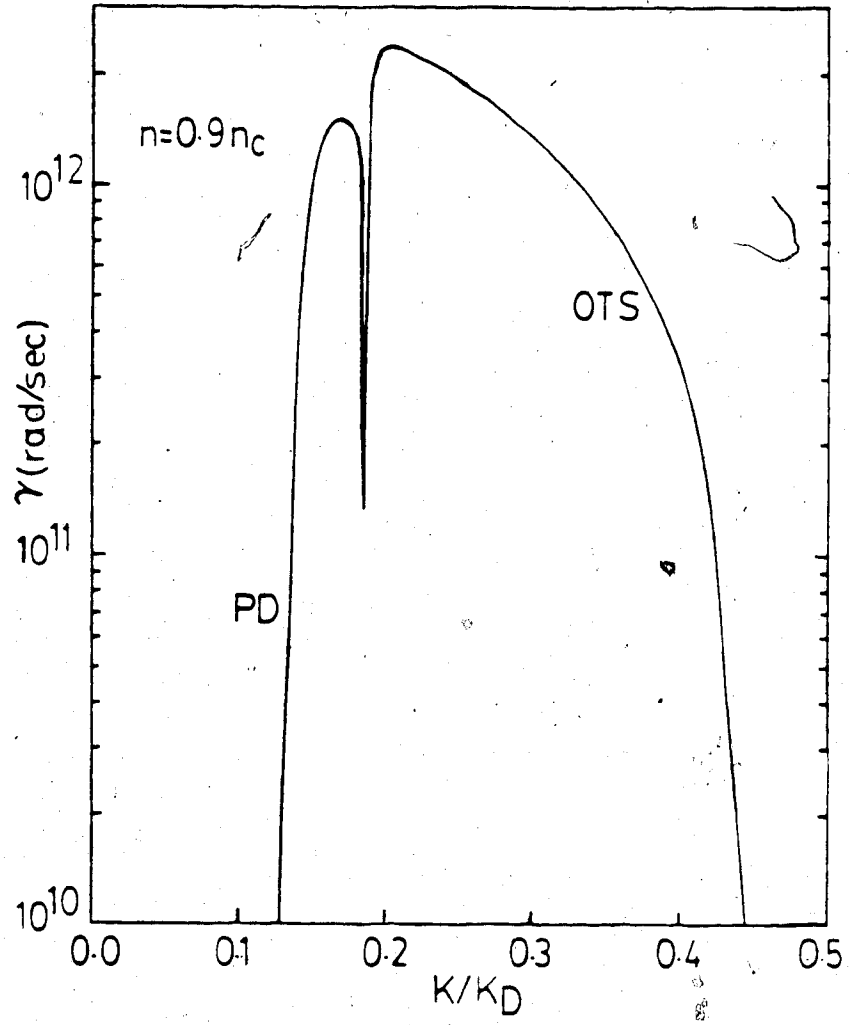


Fig. 2.10 Growth rate of ion wave instability (PD and OTS off-resonance) versus k/k_D for oxygen plasma ($z=6$) with $T_i = T_e = 100$ eV, $n/n_c = 0.9$ and $\nu_0/\nu_e = 1$.

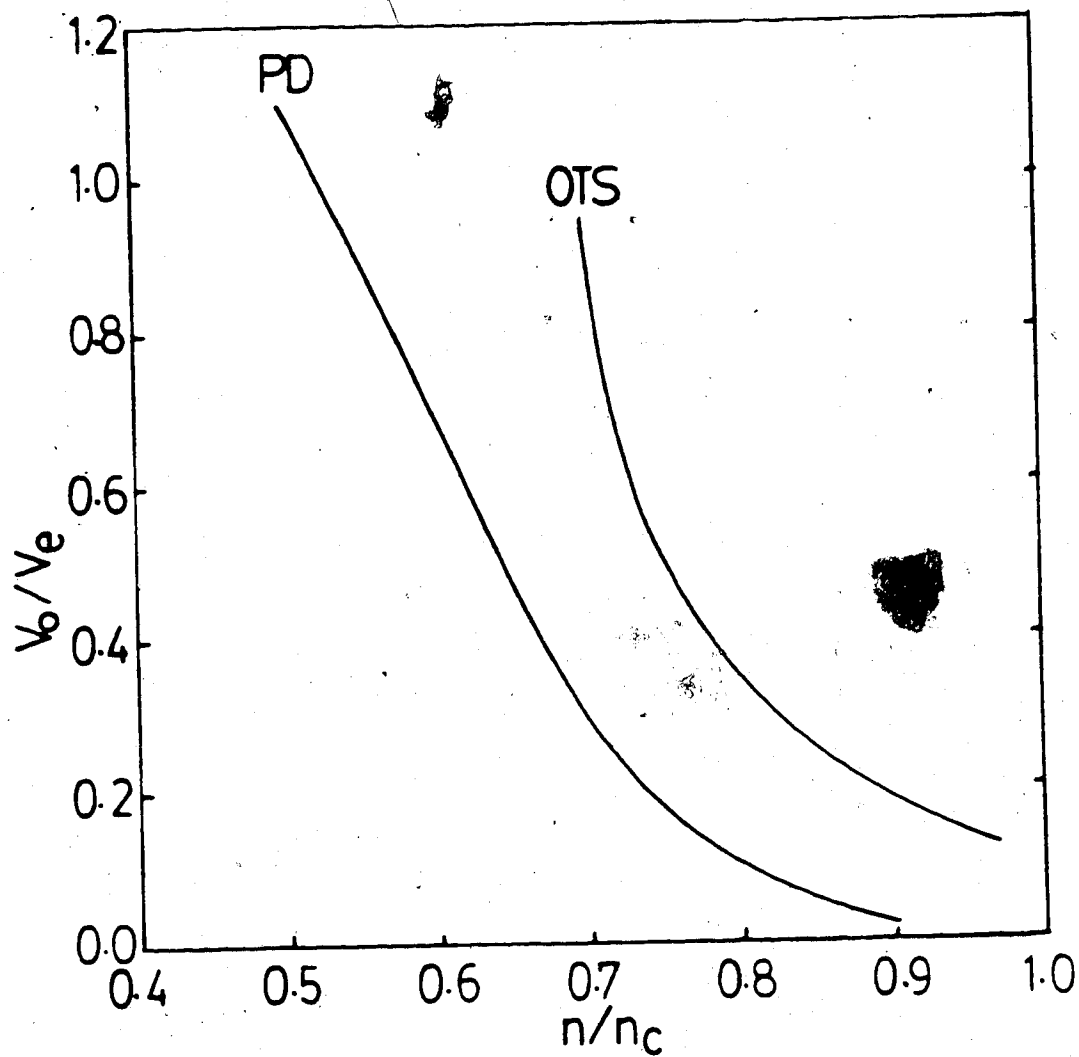


Fig. 2.11 Thresholds of the most easily excited modes of PD and OTS off-resonance instabilities as a function of density for oxygen plasma ($z=6$) with $T_i = T_e = 100$ eV.

The range of k for which these instabilities can be excited varies with the pump strength ν_0/ν_e as indicated in Fig. 2.12 for $n/n_e = 0.8$. The growth rates of the PD and OTS off-resonance as well as the frequency of the ion acoustic mode (in the PD) as a function of the angle θ are shown in Fig. 2.13. Similar dependence is expected with respect to ν_0/ν_e since both $\cos \theta$ and ν_0/ν_e enter Eqn. (2.6) through the argument of the Bessel function only. The angular frequency of the ion waves generated from the PD instability varies quite significantly with k/k_D (Fig. 2.14). The peak of the frequency occurs very close to the most unstable k mode.

It is clear from Figs. 2.6 to 2.10 that the growth rates of the PD and OTS off-resonance instabilities are quite high for $\nu_0/\nu_e \sim 0(1)$, particularly for densities close to n_e , which makes one wonder about the possible mechanisms for saturating these instabilities. However, before discussing possible saturation mechanisms, it should be emphasized here that the above discussion is only valid for homogeneous plasmas. Inhomogeneity of the plasma could severely increase the thresholds and reduce the growth rates.

2.2.3 Saturation of ion waves.

Numerical simulations and theory indicate that the ion wave instability may be stabilized by ion trapping [20-22]. According to this nonlinear mechanism an ion will be trapped in the electrostatic potential ϕ of the ion wave as the wave grows and stabilization only occurs when a large number of ions are trapped. The ion trapping velocity ν_{tr} is given (assuming a waterbag distribution function for the ions) by [23]

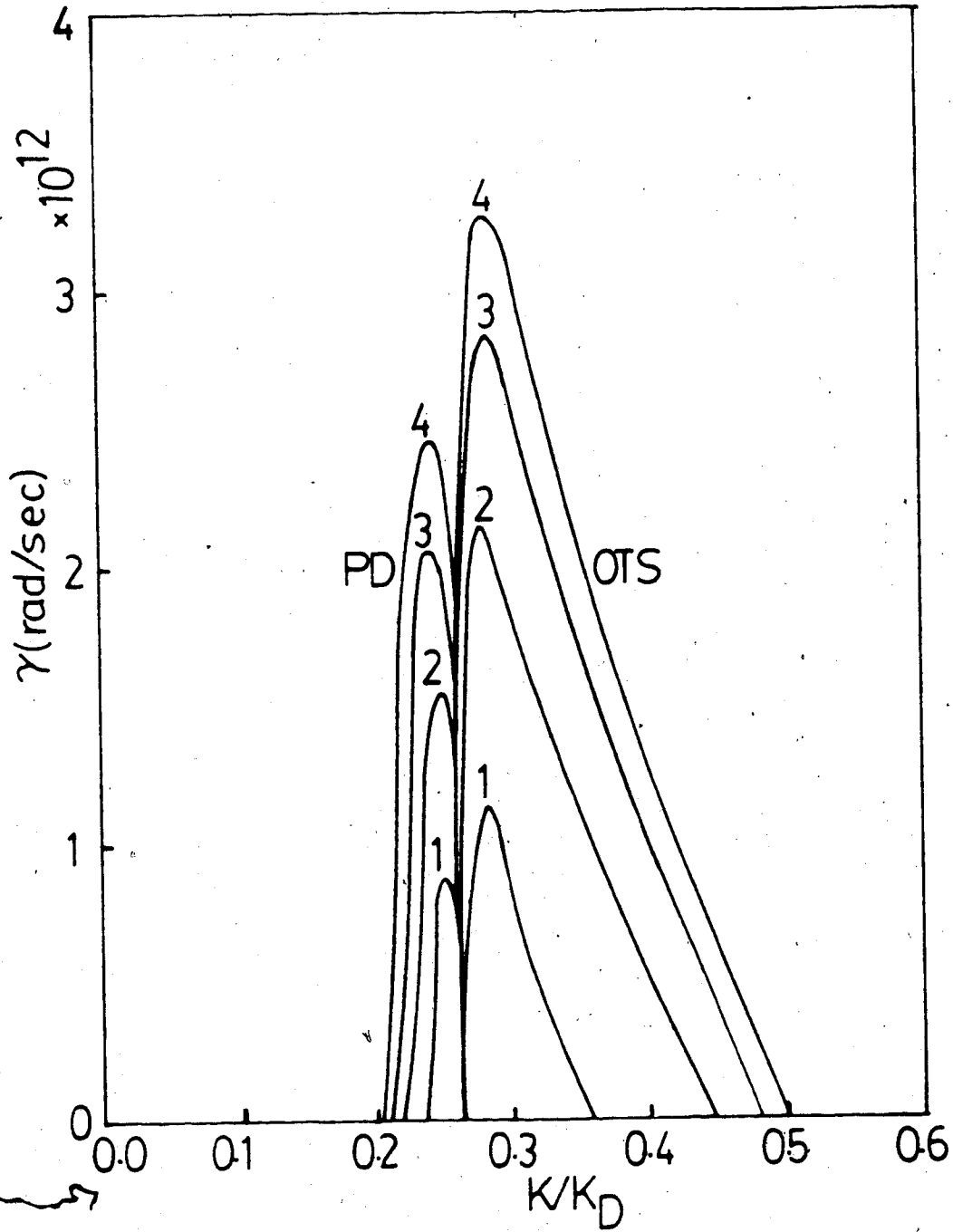


Fig. 2.12 Growth rate of PD and OTS off-resonance instabilities versus k/k_D for oxygen plasma ($z=6$) with $T_i = T_e = 100$ eV, $n/n_e = 0.8$ and $v_0/v_e = 1, 2, 3$ and 4.

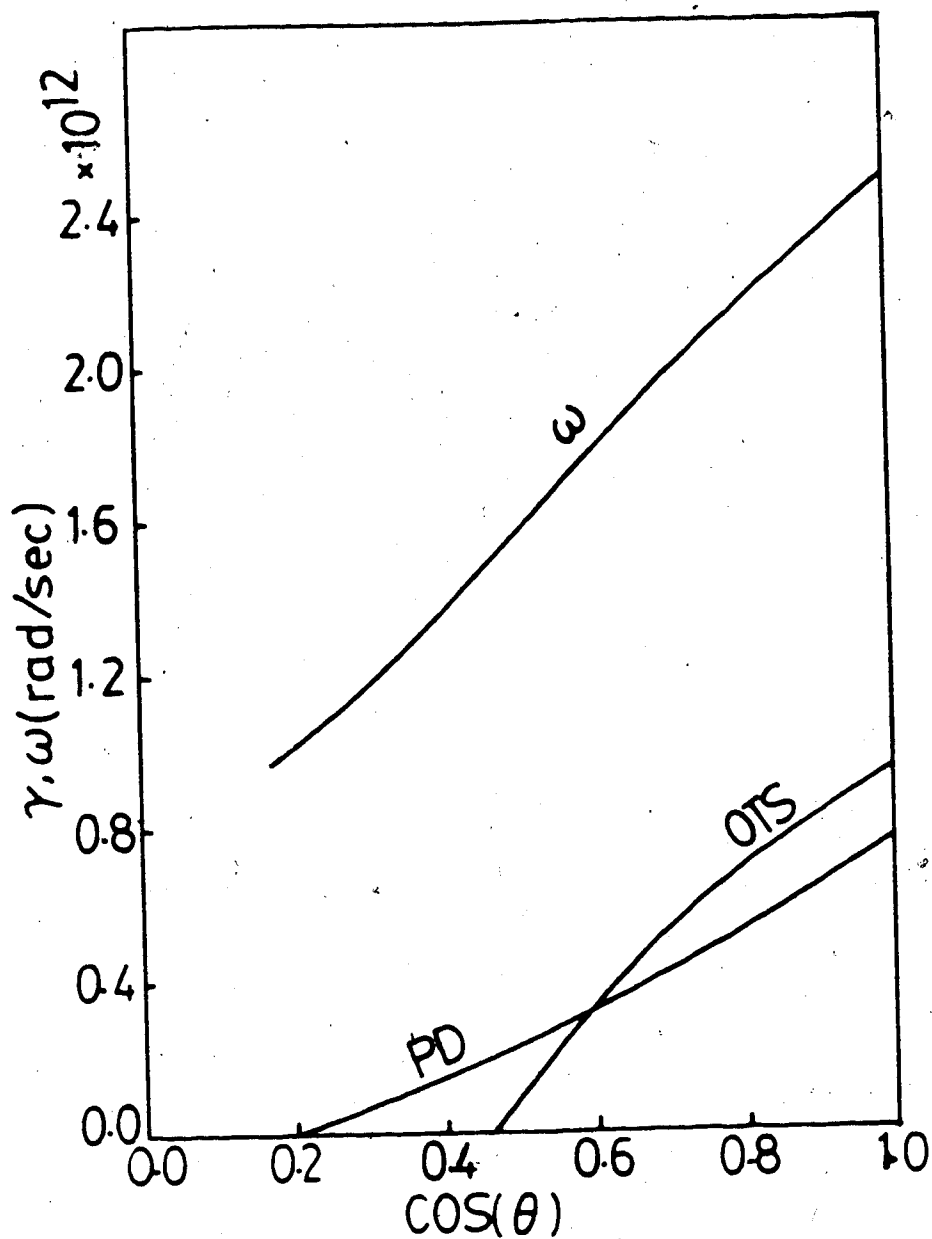


Fig. 2.13 The growth rate (of PD and OTS off-resonance) and the real frequency (of PD off-resonance) of the most easily excited ion modes in an oxygen plasma ($z=6$) with $T_i = T_e = 100$ eV, $n/n_e = 0.7$ and $v_0/v_e = 1$.

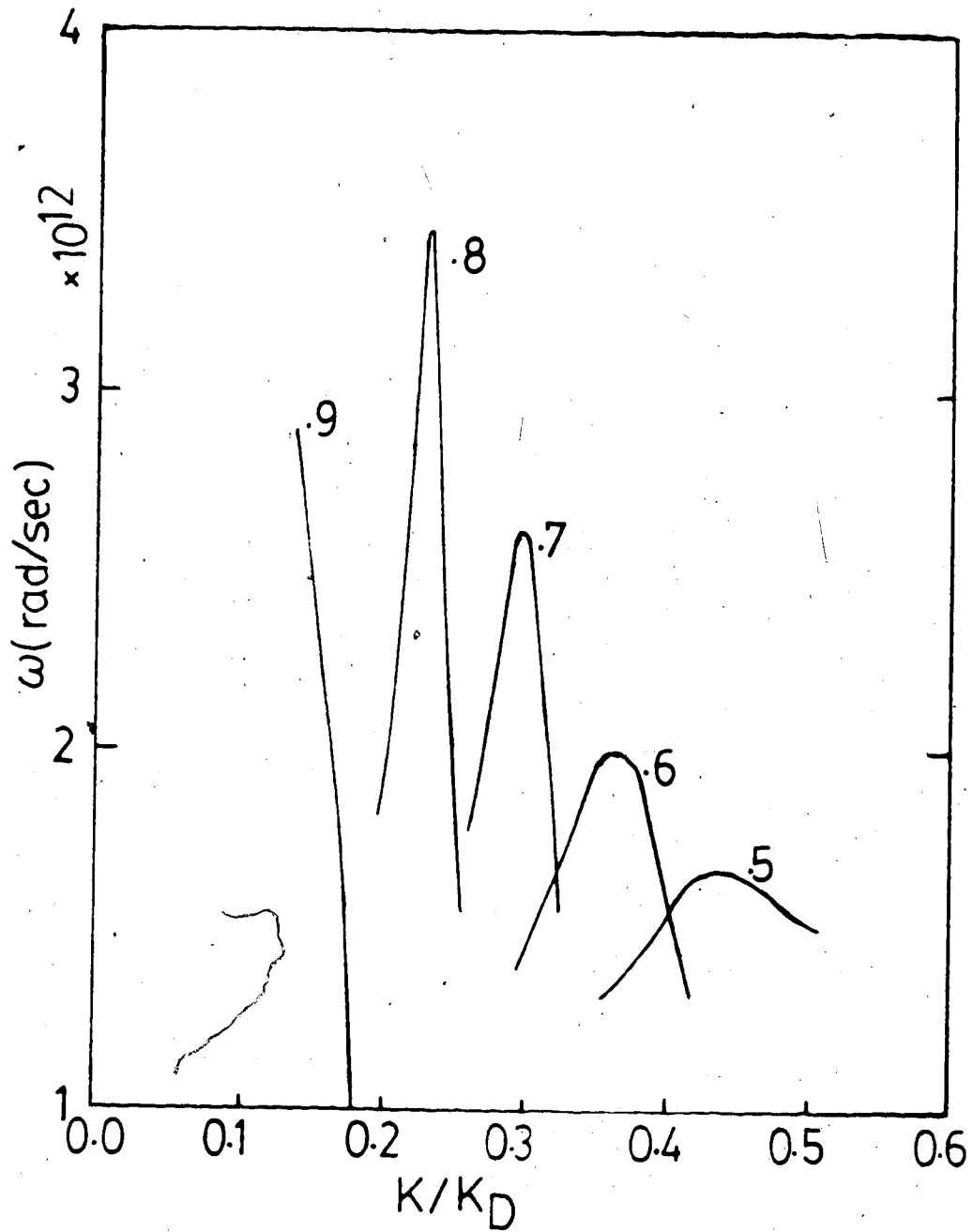


Fig. 2.14 Real frequency of the ion waves of PD off-resonance instability as a function of k/k_D for an oxygen plasma ($z=6$) with $T_i = T_e = 100$ eV, $v_0/v_e = 1$ and $n/n_0 = 0.5$, 0.6, 0.7, 0.8 and 0.9.

$$v_{tr} \approx v_p - \sqrt{3}v_i, \quad (2.8)$$

where v_p is the phase velocity of the wave and v_i is the ion thermal velocity. The energy that the wave loses to the trapped particle is then given by

$$e\phi = \frac{1}{2} Mv_{tr}^2 = \frac{1}{2} M|v_p - \sqrt{3}v_i|^2. \quad (2.9)$$

Substituting for v_p (using the ion dispersion relation), v_i and $e\phi (= T_e \delta n/n)$ Eqn. (2.9) takes the following form

$$\frac{\delta n}{n} \leq \frac{1}{2} \left| \left(\frac{z + T_i/T_e}{1 + k^2 \lambda_D^2} \right)^{\frac{1}{2}} - (3T_i/T_e)^{\frac{1}{2}} \right|^2. \quad (2.10)$$

Equation (2.10) sets the upper limit for the density fluctuation when saturation occurs. For $T_i = T_e$, $k\lambda_D = 0.5$ and $z = 6$ (for oxygen plasma), $\delta n/n \sim 0.2$.

Numerical simulations [21,22] show that saturation takes place in a very short time ($\omega_{pe} t \approx 2200$) while ion heating is still occurring. For a plasma with $n = 0.5n_c$ (where $n_c = 10^{19} \text{ cm}^{-3}$) the saturation time is only ~ 20 ps.

Electron heating could be another mechanism for saturating the PD and OTS off-resonance instabilities. Although the temperature ratio T_i/T_e has a very limited effect on the growth rates of these instabilities as illustrated in Fig. 2.15, heating of electrons to high temperatures, on the other hand, will decrease v_0/v_e , possibly below the instability threshold.

The PD off-resonance may also be stabilized by harmonic generation where coupling of the form

$$\omega_1 \pm \omega_2 = \omega_3, \quad \bar{\kappa}_1 \pm \bar{\kappa}_2 = \bar{\kappa}_3,$$

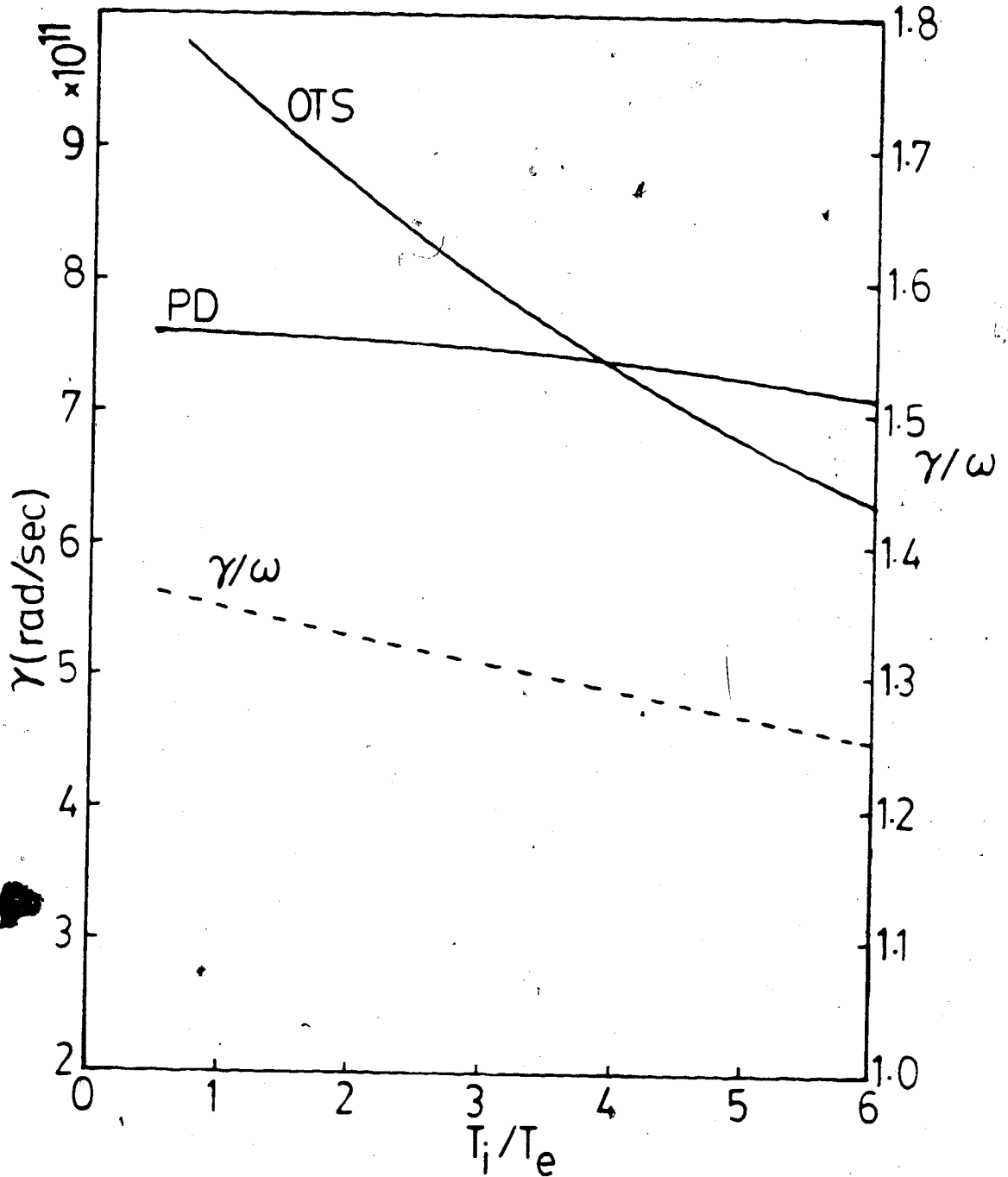


Fig. 2.15 The growth rate (of PD and OTS off-resonance) and the ratio γ/ω (of PD off-resonance) of the most easily excited ion modes versus T_i/T_e for an oxygen plasma ($z=6$) with $v_0/v_e = 1$ and $n/n_c = 0.7$.

may take place (where 1 and 2 refer to two unstable ion modes and 3 to a new ion wave). Saturation occurs in this process when the new wave is heavily damped. It should be pointed out that the general dispersion relation of Sanmartin predicts the existence of stable ion modes with a wide range of k and ω (Fig. 2.14). These stable modes could be a product of wave couplings described above. Recently, the importance of harmonic generation as a saturation mechanism for PD off-resonance instability has been confirmed theoretically and by simulation [24].

2.3 Absorption Due to Ion Turbulence

Central to the problem of heating plasmas to **temperatures of interest** for fusion by means of lasers is the absorption of radiation. The simplest absorption process is that of inverse bremsstrahlung or resistive absorption due to electron-ion collisions (discussed in Chapter 1). For the case of a linear density profile of scale length L and $\omega \leq \omega_p$, the absorbed fraction of light is given by [25]

$$A = 1 - \exp\left[-\frac{32}{15}(\nu_{ei}^c L/c)\right], \quad (2.11)$$

where ν_{ei}^c is the electron-ion collision frequency at critical density. Equation (2.11) indicates that in order to have strong absorption, a short wavelength laser is preferred (simply because $\nu_{ei}^c \propto n_e \propto 1/\lambda^2$) and the plasma conditions must be such that both L and ν_{ei}^c are sufficiently large. The latter assumption, however, means low temperature plasmas are needed (since ν_{ei}^c varies with $T_e^{-3/2}$). It is therefore expected that the inverse bremsstrahlung absorption is very inefficient in hot plasmas where a high temperature is required for fusion. The situation is even worse when a long wavelength laser is used, such as the CO_2 laser.

In addition to inverse bremsstrahlung absorption, light can also be absorbed via wave-particle interactions. Such a process may occur when high intensity radiation excites plasma waves, which, in turn, transfer their energy to the particles by damping. In this case Eqn. (2.11) will be approximately valid but with v_{ei}^c replaced by the effective damping rate of the effective collision frequency ν_{eff} . An example of this process, which is of relevance to us and will be discussed here, is the absorption due to ion turbulence.

The effect of ion turbulence on absorption was first discussed by Dawson and Oberman [26] and subsequently by Faehl and Kruer [3]. It is necessary, in order to understand absorption via ion turbulence, to briefly review the theory that was given by Faehl and Kruer [3].

A simple interpretation can be obtained by considering the pump field of the form $E_0 \cos \omega_0 t$ propagating in a plasma whose density is

$$n = n_0 + \sum \delta n_{ki} \cos \vec{k}_i \cdot \vec{x} . \quad (2.12)$$

As the electrons oscillate with velocity $\vec{v} = \frac{e\vec{E}_0 \sin \omega_0 t}{m\omega_0}$ between low and high-density regions a source term of high frequency density fluctuations $n_s(\vec{x}, t)$ is produced

$$n_s(\vec{x}, t) = n(\vec{x} + \vec{r}_0) - n(\vec{x}) = \vec{r}_0 \cdot \nabla n , \quad (2.13)$$

where $\vec{r}_0 = \frac{e\vec{E}_0 \cos \omega_0 t}{m\omega_0^2}$. The self-consistent field resulting from this

source term is given by Poissons' equation whose Fourier transform is

$$i\epsilon(\vec{k}, \omega) \vec{k} \cdot \vec{E}(\vec{k}, \omega) = -4\pi e n_s(\vec{k}, \omega) , \quad (2.14)$$

where $\epsilon(\vec{k}, \omega)$ is the plasma dielectric function.

The energy transferred to the plasma via this field is

$$W = - \int \bar{\mathbf{J}} \cdot \bar{\mathbf{E}} dt d\bar{\mathbf{x}}, \quad (2.15)$$

in which the current density $\bar{\mathbf{J}}$ is given by the continuity equation $\nabla \cdot \bar{\mathbf{J}} - e \partial n'_s(\bar{\mathbf{x}}, t) / \partial t = 0$. By taking the Fourier transform of Eqn. (2.15) and substituting Eqn. (2.14) for $\bar{\mathbf{E}}(\bar{\mathbf{k}}, \omega)$ we obtain

$$W = \frac{4e^2}{(2\pi)^3} \int d\bar{\mathbf{k}} d\omega \frac{\omega}{k^2} \text{Im} \left[\frac{1}{\epsilon(\bar{\mathbf{k}}, \omega)} \right] |n'_s(\bar{\mathbf{k}}, \omega)|^2, \quad (2.16)$$

where the imaginary term is used for a real energy. The time averaged power per unit volume transferred to the plasma can be obtained by substituting $n'_s(\bar{\mathbf{k}}, \omega)$ (by Fourier transforming Eqn. (2.13)) into Eqn. (2.16), then integrating with respect to $\bar{\mathbf{k}}$ and taking the derivative with respect to ω . This gives

$$P = \frac{\omega_0}{2} \frac{E_0^2}{8\pi} \sum_{ki} \left(\frac{\delta n_{ki}}{n_c} \right)^2 \text{Im} \left[\frac{1}{\epsilon(\bar{\mathbf{k}}, \omega)} \right] \cos^2 \theta_i, \quad (2.17)$$

where n_c is the critical density and θ_i is the angle between $\bar{\mathbf{k}}_i$ and $\bar{\mathbf{E}}_0$.

The effective collision frequency is then calculated from the energy balance equation $P = \nu_{eff} (E_0^2 / 8\pi)$, yielding

$$\nu_{eff} = \frac{\omega_0}{2} \sum_{ki} \left(\frac{\delta n_{ki}}{n_c} \right)^2 \text{Im} \left[\frac{1}{\epsilon(\bar{\mathbf{k}}, \omega)} \right] \cos^2 \theta_i. \quad (2.18)$$

If we assume an ion turbulence which is broad in angle and extends over a region of wavenumbers, then the average effective collision frequency can be written in the form

$$\nu_{eff} = \frac{\omega_0}{4} \left| \frac{\bar{y}}{n_c} \right|^2 < \text{Im} \frac{1}{\epsilon(\bar{\mathbf{k}}, \omega_0)}, \quad (2.19)$$

where \bar{y} is the mean density fluctuation and the brackets denote an

average over the range of wavenumbers. In order to estimate ν_{eff} , the function $\langle \text{Im} \frac{1}{\epsilon(\bar{k}_i, \omega_0)} \rangle$ averaged over a uniform spectrum of ion fluctuations between $k/k_D = 0.1$ and $k/k_D = 1$ was plotted and is shown in Fig. 2.16 as a function of density for a Maxwellian oxygen plasma. As shown in Fig. 2.16, the function $\langle \text{Im} \frac{1}{\epsilon(\bar{k}_i, \omega_0)} \rangle$ is significant near $n/n_e = 0.8$. At this density and for plasma conditions $T_e = 100$ eV, $z = 6$ and $\Delta n/n_e = 0.1$, the effective collision frequency $\nu_{eff} \approx \nu_{cl}$ (where ν_{cl} is the classical inverse bremsstrahlung collision frequency). For higher temperatures and low z , $\nu_{cl} \gg 1$.

Evidently, the above analysis assumed a given ion fluctuation level regardless of the source or cause of the fluctuations. As we have discussed earlier ion fluctuations can be generated by parametric processes in the presence of a strong electromagnetic pump. The presence of such a pump could by itself modify the plasma dispersion function and thus the collision frequency. The effect of a high frequency electromagnetic field on electron collisionality was considered recently by Rozmus *et al.* [27] for a stable plasma. An approximate expression for ν_{eff} is given by

$$\nu_{eff}/\omega_0 \approx - \left(\frac{n_e}{n_c} \right) \frac{8\pi z e}{E_0^2} \int \frac{d\bar{k}}{(2\pi)^3} (\bar{k} \cdot \bar{E}_0) \int d\bar{u}_i \frac{F_i(\bar{u}_i)}{|\Gamma(\bar{k}, \bar{k} \cdot \bar{u}_i)|^2} .$$

$$\sum_j \prod_j \prod_{j^{-1}} \text{Im} \frac{\chi_e^{-j}(\bar{k}, \bar{k} \cdot \bar{u}_i)}{D_e^{-j}(\bar{k}, \bar{k} \cdot \bar{u}_i) D_e^{-j+1}(\bar{k}, \bar{k} \cdot \bar{u}_i)} \quad (2.20)$$

where $\Gamma(\bar{k}, \bar{k} \cdot \bar{u}_i)$ is the general dispersion function given by Eqn. (2.6) and the remaining terms are as defined previously.

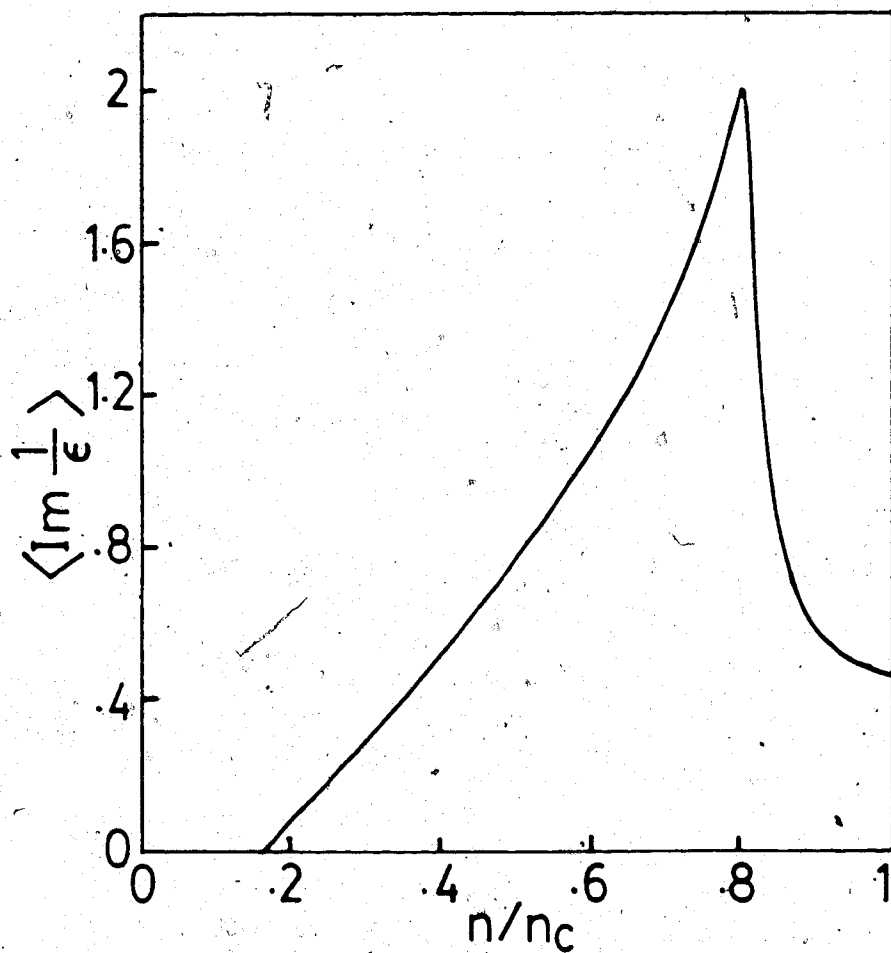


Fig. 2.16 $\text{Im} \frac{1}{\epsilon(\bar{k}_i, \omega_0)}$ averaged over a uniform spectrum of ion fluctuations as a function of density for a Maxwellian oxygen plasma with $T_i = T_e = 100$ eV.

The factor under summation can be reduced to $\text{Im} \frac{1}{D_e(\vec{k}, \vec{k} \cdot \vec{u}_e)}$ for the small pump limit which is similar to the factor appearing in the Faehl-Kruer expression (2.19). In addition, enhancement can occur when $D_e(\vec{k}, \vec{k} \cdot \vec{u}_e)$ is at or near resonance. It is important to note that even for sub-threshold pump, significant enhancement of collision frequency is possible. For example, for an oxygen target ($s=c$, $A=10$), $T_e = T_i = 100$ eV and $n = 0.8n_c$, expression (2.20) gives $\nu_{eff} = 3\nu_{cl}$ for the \vec{k} vector along the direction of \vec{E}_0 , corresponding to the least damped mode and ν_0/ν_e below the threshold for the PI off-resonance instability. For lower densities where the threshold is relatively high, even larger enhancements over classical are obtained. However, Eqn.(2.20) is only valid for stable plasma and represents the lowest enhancement limit. With a pump strength above the threshold for PI and OIS off-resonance the enhancement can be substantially larger. Consequently, plasma absorption may be expected to be enhanced over classical when large ion fluctuations are induced. This may occur even for pump waves below threshold for parametric instabilities, though clearly significant effects may be anticipated for above-threshold pump waves.

CHAPTER 3

EXPERIMENTAL METHODS AND TECHNIQUES

3.1 Introduction

A number of experimental methods and techniques were applied in the course of studying ion turbulence in laser plasma interactions. It is in this chapter of the thesis that these methods and techniques will be outlined in some detail. The experimental setup consists mainly of two sections. First, the part in which the plasma was formed and heated using an intense pulsed CO₂ laser and a μm target is presented. The second section is concerned with the diagnostic techniques which in turn is divided into two; one for measuring the different plasma parameters such as density and temperature and the other for studying ion turbulence by means of a ruby laser Thomson scattering technique. In the forthcoming sections the laser plasma experiment, the diagnostic techniques and related topics will be discussed in this order.

3.2 Laser Plasma Experiment

3.2.1 Carbon dioxide laser.

A transverse excited atmospheric TEA type pulsed CO₂ laser was used for studying laser plasma interactions. This laser was designed and built at the University of Alberta. A brief description will be given here.

The laser consisted of six pairs of opposing aluminum and graphite electrodes. Each electrode was 50 cm long with a Rogowski

cross sectional profile modified by a 3.7 cm wide flat area in the centre. The electrodes were connected to a two stage Marx circuit using 1.1 μF capacitors. UV preionization was provided by means of a series of 500 pf capacitors driving pin discharges. These pins were situated 2 cm apart along both sides of the main discharge volume and were electrically conductively parallel with the main discharge electrodes. The total gain length of the laser was 8 m and the unstable optical cavity resonator was formed using two mirrors of 34.57.

An unstable resonator has many advantages over a stable one. Two of these advantages are the high energy extraction for having a larger mode volume, the discrimination against high order transverse modes and the most important of all is the low output beam divergence. The cavity mirrors for this laser were designed to give a positive type unstable resonator with 10% output coupling. The system was operated at a charge voltage of 85 kV and a gas mixture of approximately 10% He, 10% Ne_2 and 80% Ne . It should be noted that the system was designed for a more favorable gas ratio: 10% He, 40% Ne_2 and 50% Ne . However, because of deterioration in the electrodes and electrical components the He concentration in the gas mix was increased to maintain arcs between the electrodes.

Table 1.1 lists the main features of this laser. The output laser pulse consisted of a multi-transverse mode structure, approximately 1.5 cm in cross section, with an initial gain switched spike peak of approximately 700 MW in a pulse duration of 0.50 ns. Figure 1.1 shows a typical pulse shape of the existing Ne_2 laser as recorded by a Geiger detector. Similar results were obtained theoretically by

Wavelength	10.6 μm
Storage Capacitance	1.2 μf
Preionization	50 discharge pins, each side
Active Volume	3.7 cm \times 3.7 cm \times 300 cm
Electrical Efficiency	\approx 2.7%
Cavity Length	4 meter unstable resonator
Rear Mirror	12.8 m radius of curvature, 5 cm diameter
Front Mirror	-5 m radius of curvature, 1 cm diameter
Gas Mixture ($\text{CO}_2:\text{N}_2:\text{He}$)	22:8:70
Output Energy	\approx 40 Joules
Peak Power	\approx 500 MW gain switched
Pulse Duration	\approx 50 ns
Beam Divergence	\approx 0.5 mrad

Table 3.1 Features of the CO_2 TEA Laser

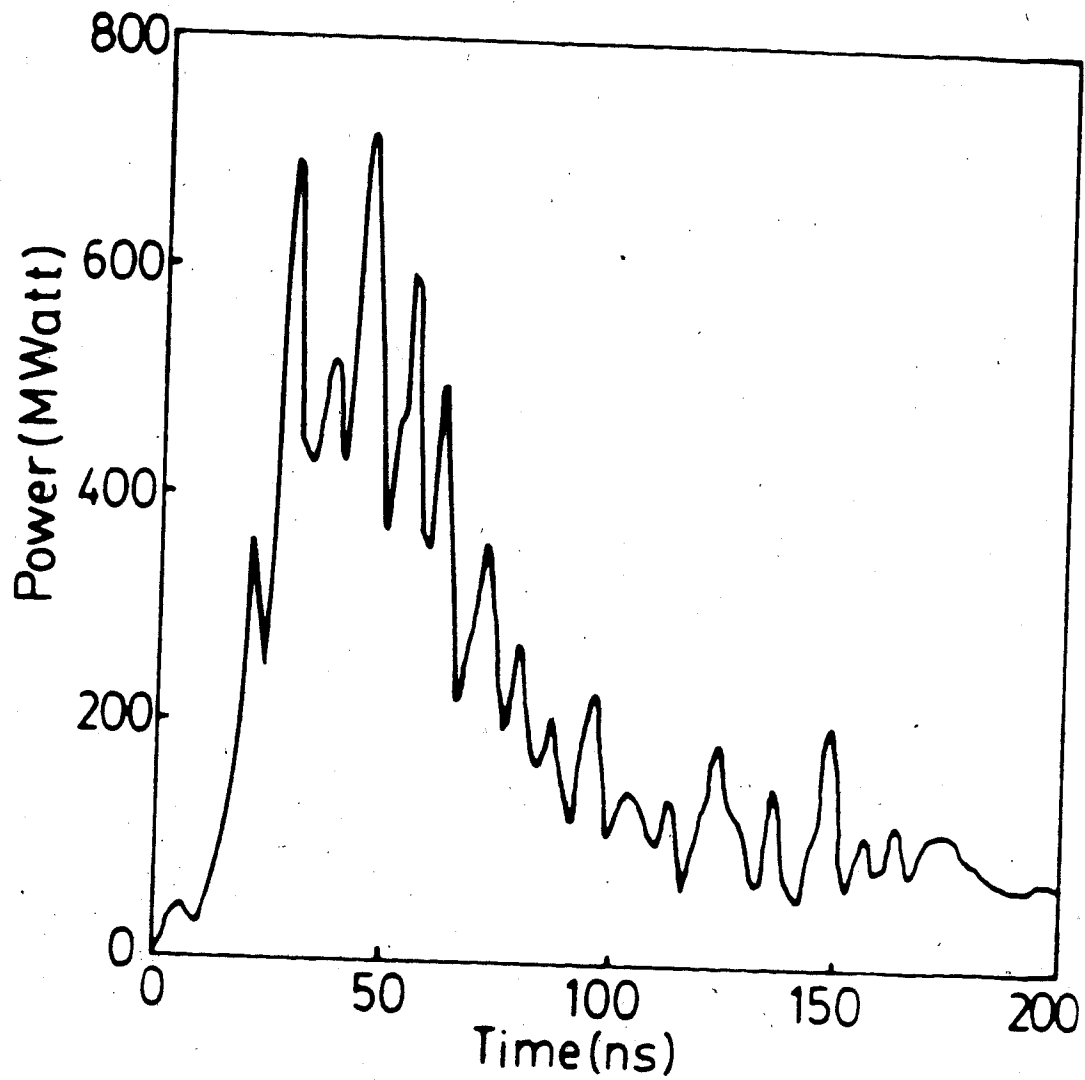


Fig. 3.1 Gain switched CO₂ laser output.

numerically solving the kinetic rate equations (Appendix F). These results are shown in Fig. 3.2. It should be noted, however, that the calculations do not predict the temporal fine structure shown in Fig. 3.1. This structure is usually observed due to the presence of several modes in the cavity.

3.2.2 Gas jet and target chamber.

A cylindrical aluminum target chamber 75 cm in diameter and 30 cm in height with a 1 cm thick plexiglas top was used throughout the course of these investigations. A base pressure of $\sim 10^{-3}$ torr inside the chamber was readily achieved using a 15 cfm rotary pump. A NaCl window was fitted onto the chamber in such a way that normal incidence of the CO_2 laser beam could be avoided. In addition, the chamber was provided with seven large ports which could be used for windows in order to attain optical access for different diagnostics (Fig. 3.3). The chamber pressure was maintained at ~ 10 torr of He pressure during the experiment and was evacuated and back-filled with fresh He every time the laser was focused on the gas target.

A gas jet assembly, developed by Burnett *et al.* [1], was used to provide the gas target. Near laminar flow was obtained by pulsing high pressure (~ 2000 torr) oxygen gas through a convergent-divergent (supersonic) nozzle into the 20 torr He background. Equilibrium between O_2 and He was reached in ~ 2 ms, forming a ~ 1.5 mm thick and a 1 cm wide target. The optimal oxygen to helium pressure ratio (2000:10) was necessary in order to produce a laminar O_2 flow with a minimum He background. The oxygen flow was measured utilizing ruby laser shadowgraphy. The neutral O_2 density of the jet was not uniform but rather varied as a function of both height and thickness.

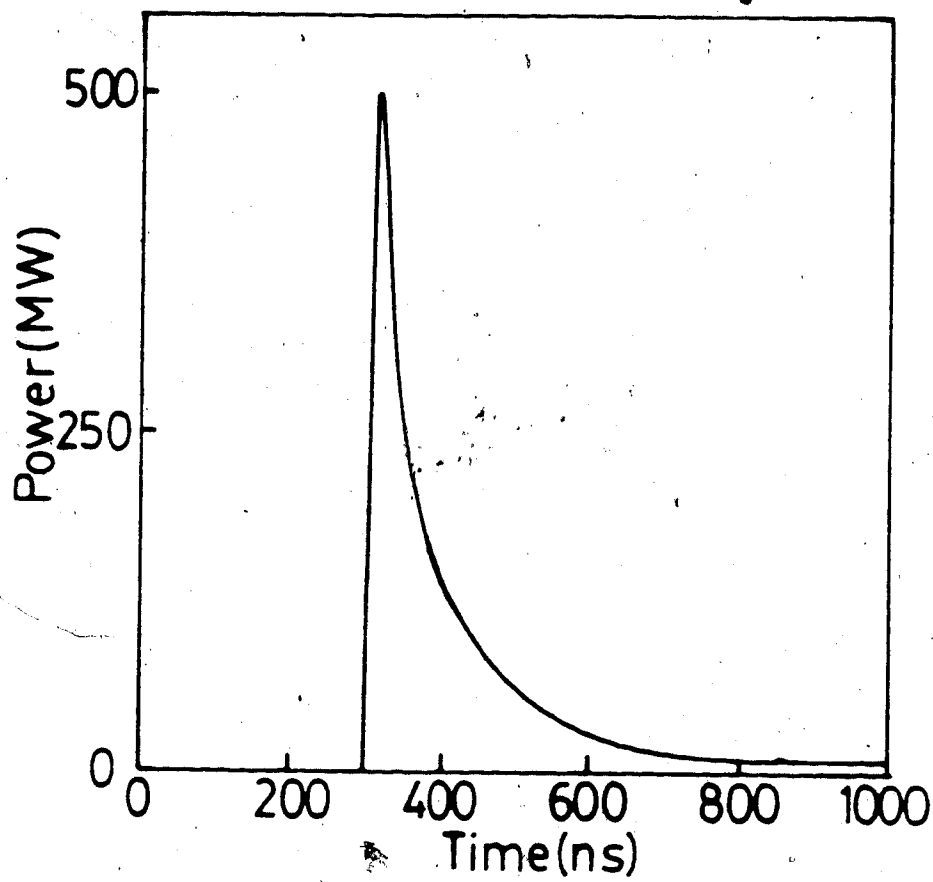


Fig. 3.2 Output of the CO₂ laser model.

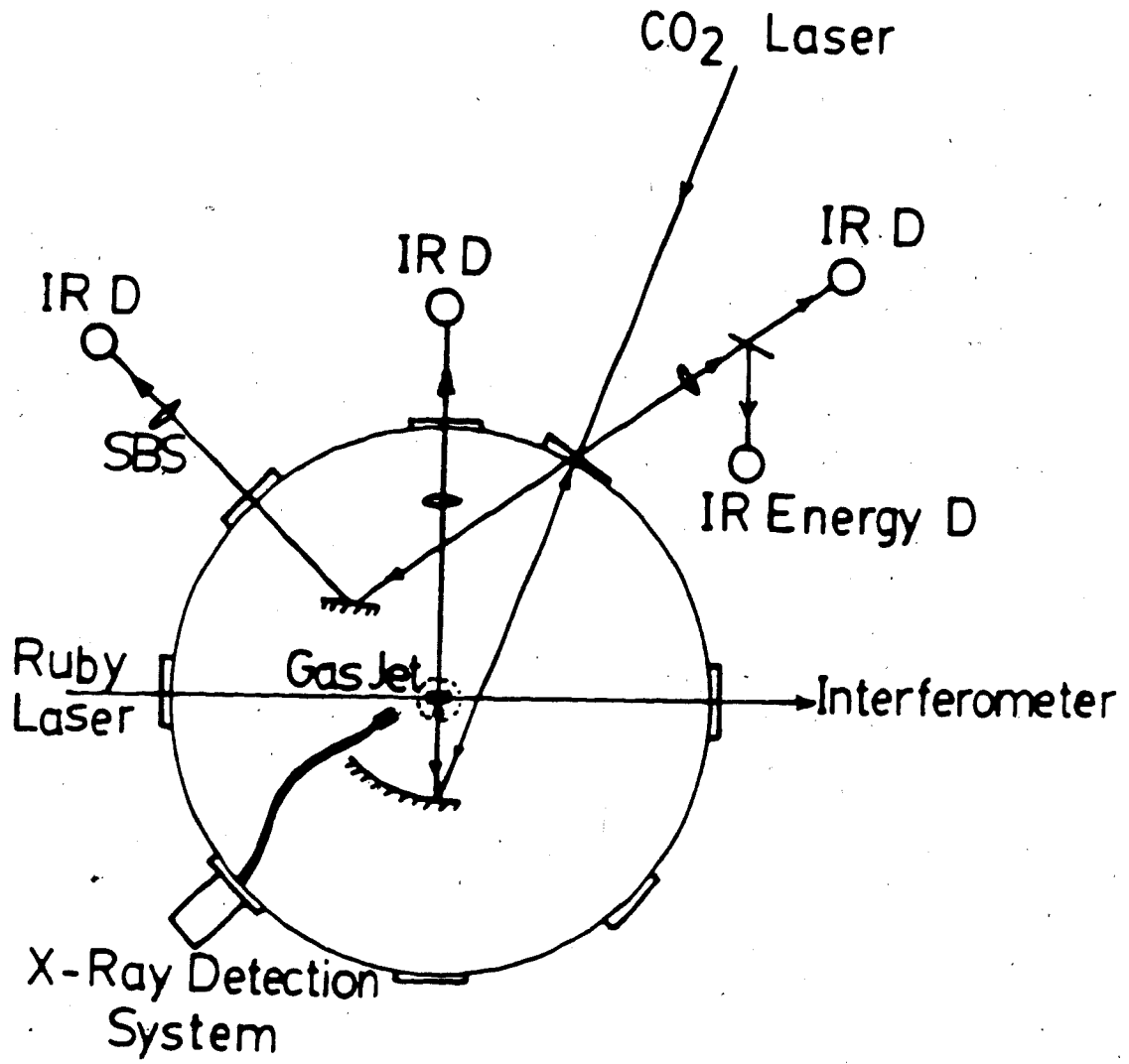


Fig. 3.3 Schematic diagram of the CO₂ laser plasma interaction experiment and various diagnostics.

The hydrodynamic calculations of Giles [2] showed that at 4 mm above the nozzle (where the laser beam was focused) the jet consisted of a high density region ($\sim 1.5 \times 10^{18}$ molecule/cm³) located between an inner low density region ($\sim 2.2 \times 10^{17}$ molecule/cm³) and an outer low density region ($\sim 6.5 \times 10^{17}$ molecule/cm³). If it is assumed that all the oxygen atoms in these regions were ionized to the degree $z = 6$ then this will lead to the formation of a nonuniform electron density distribution (between $1.75 n_c$ and $0.29 n_c$). This nonuniformity, however, is very transient because of heating and subsequent hydrodynamic expansion.

Many unique features of the gas target make it preferable to a solid target. These features include: (a) variable density n from above critical to below critical; (b) nondestructive target which can be pulsed just prior to the firing of the laser; (c) good diagnostic access for measurements.

In brief, the laser firing operation can be explained with the help of Figs. 3.4 and 3.5 as follows:

- (a) The high pressure oxygen gas is admitted into reservoir B and to the chamber upon triggering the solenoid valve between reservoirs A and B.
- (b) A delayed signal from a piezoelectric pressure transducer B is used to trigger an EG&G high voltage pulse generator which in turn triggers the main laser spark gaps. The delay time is adjusted so that the laser is fired at the peak of the pressure transducer. Typical time jitter in the laser system was 10-20 ns.

The only remaining component in the target chamber to be described here is the parabolic mirror used for focusing the 3μ laser beam. This mirror was made of an aluminum substrate (13 cm in diameter and 10 cm

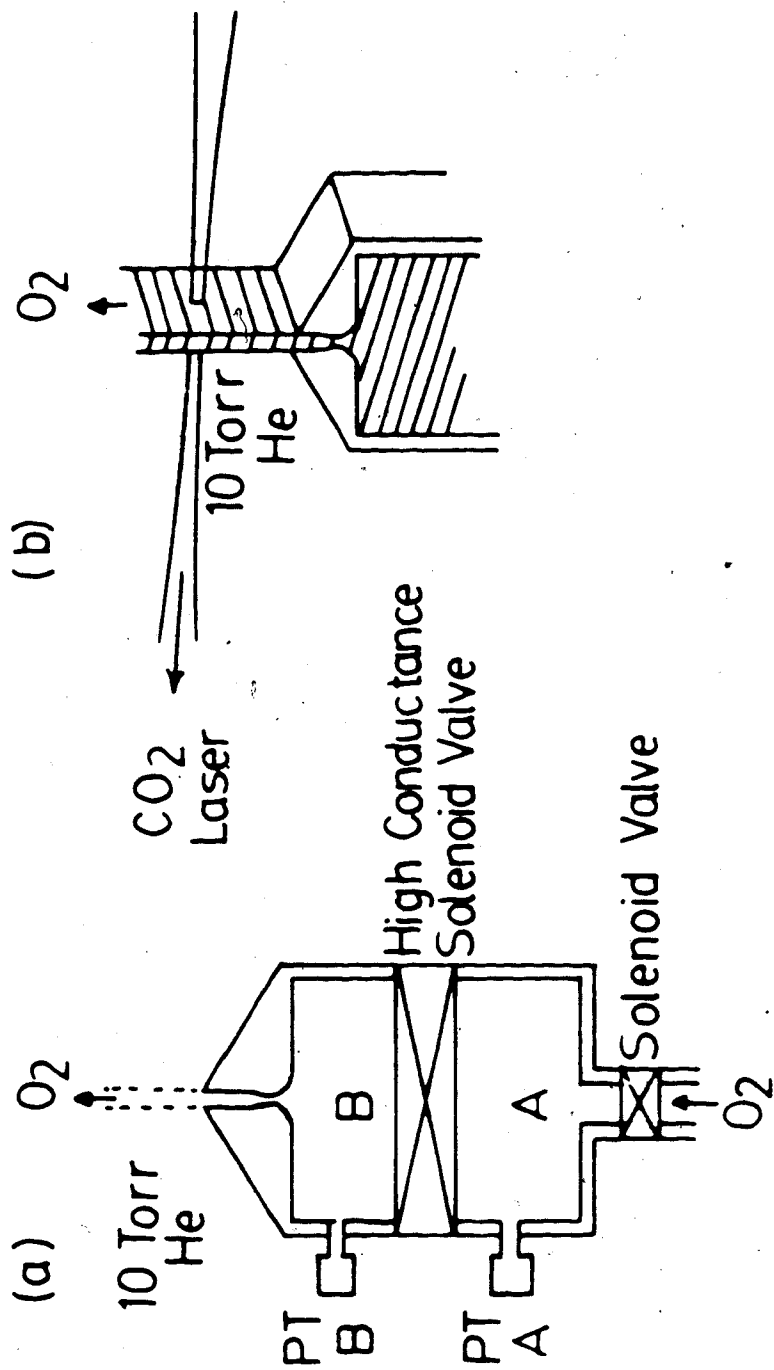


Fig. 3.4 Gas target design.

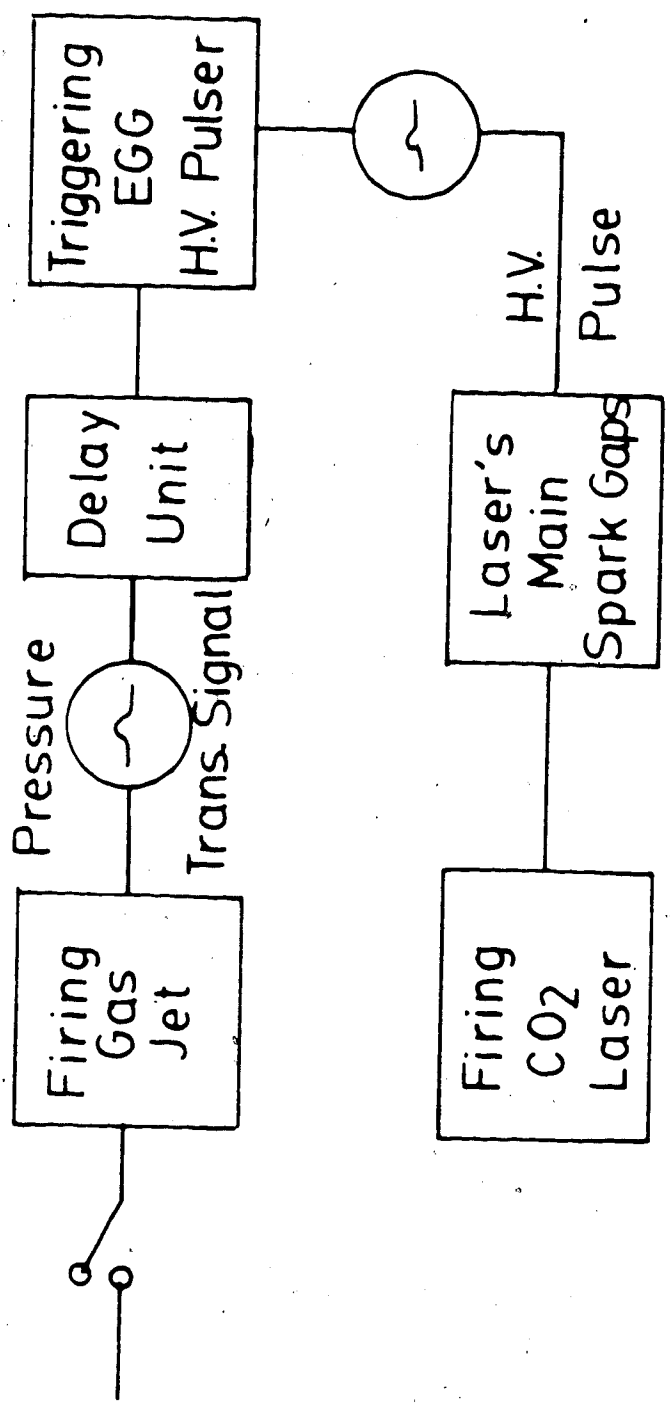


Fig. 3.5 Firing sequence of the CO₂ Laser.

focal length), coated with copper and then overcoated with gold for high reflectivity and high damage threshold [3]. A mirror holder that could be moved or tilted in order to focus the laser beam anywhere within the gas target was used. The approximate focal point size was determined by measuring accurately the laser beam divergence and mirror focal length. The beam divergence was determined from a focal beam waist calculation for the half power point using a long focal length mirror. For a 10 cm mirror focal length and a 0.5 mrad beam divergence the minimum focal spot size is 50 μm . However, this is only true for an ideal, aberration free mirror. More accurate calculations were done by Giles [2] for the mirror used in these studies using a ray tracing technique and a direct measurement of the mirror radius of curvature. These calculations showed that 60 to 70 percent of the incident laser power was within a 100 μm diameter spot (this corresponds to a focal point laser beam intensity of $\leq 5 \times 10^{12} \text{w/cm}^2$). The approximate focal length of the system was 800 μm . At the axial position of peak intensity, the beam FWHM was 60 μm . It should be noted, however, that according to Giles' calculations, a significant fraction of the incident power could be off-axis by 150 microns.

3.3 Measurements of Plasma Parameters

3.3.1 Infrared (IR) radiation detection.

Different types of IR power detectors were used during the course of these investigations. For relatively strong signals such as the incident and transmitted laser radiation, simple photo-voltaic or photo-conductive type detectors were adequate. Examples

of these types of detectors are the Rofin photon drag model 7400 (~ 1 ns risetime and responsivity of $0.5 \mu\text{V/W}$ into 50Ω resistor) or its duplicate homemade detector. For weaker signals such as the $2\omega_0$ and $3/2\omega_0$ harmonics and stimulated Brillouin scattering (SBS), a liquid nitrogen cooled, gold doped germanium (Ge:Au) photo-detector (~ 1 ns risetime and responsivity of $\sim 0.2 \text{ V/W}$ into 50Ω at $\lambda \sim 10 \mu$) was used. In both cases the output was amplified by Avantek preamplifiers (0.9 ns risetime and 30-40 dB gain). For even faster response detection a pyroelectric detector (~ 100 ps risetime and responsivity of $\sim 5 \mu\text{V/W}$ into 50Ω) directly coupled to a Tektonix 7104 oscilloscope (1 GHz) was used.

The laser output energy was measured with a pyroelectric thermal detector (Gen-Tec ED-200) which has a time response of 5 ms and a known responsivity of 5 volts per joule into a 1 megohm resistance. The power was then determined from $\int P(t)dt = E$, where $P(t)$ is the instantaneous power and E is the total energy. This allowed many power detectors to be calibrated by simultaneously monitoring the power and energy using NaCl etalons of known reflectivity and Mylar of different thickness for attenuation.

As was mentioned before, an NaCl window was mounted to the chamber in such a way that normal incidence of the laser beam was avoided. This arrangement not only prevented damaging the laser optics but it also made the simultaneous measurements of both the SBS back-scattering and the incident radiation possible (Fig. 3.3).

The SBS, transmitted and incident radiation signals were accurately synchronized by carefully measuring the optical paths and adjusting the cable lengths so that any of these signals could be used as a reference for timing other signals.

3.3.2 Electron temperature (T_e).

The electron temperature was measured using the x-ray foil absorption method. To apply this technique the x-ray spectrum was assumed to be dominated by free-free bremsstrahlung radiation (the recombination and line radiation must be very small). This is true for a single species plasma with no impurities present and for temperature in the range $T_e \geq 100$ eV [4]. The above assumption was checked and was found valid for our O_2 target using a computer code developed by Salzmann [5].

For bremsstrahlung radiation, the intensity (I_λ) at a particular wavelength λ (and energy $E = hc/\lambda$) transmitted through an absorbing foil of thickness x with an absorption coefficient μ , is given by [6]

$$I_\lambda \propto \lambda^{-2} (kT_e)^{-3/2} \exp(-E/kT_e - \mu x) \quad (3.1)$$

In practice, Eqn. (3.1) must be integrated over a wide range of wavelengths to give the total intensity I . The simplest way of determining the electron temperature is to plot the ratio of two x-ray intensities I_1/I_2 for two different foil thicknesses according to Eqn. (3.1) after being integrated versus temperature. The intensity ratio I_1/I_2 is then measured experimentally for the two foil thickness. The corresponding electron temperature is finally obtained from I_1/I_2 versus temperature curve. Since the error in using only one pair of foils can be large, a more reliable method - a multifoil technique - was used as explained in Fig. 3.3. Equation (3.1) was first integrated numerically over a photon range (up to 1000 eV) with a fine mesh near the absorption edges for different foil thicknesses and different electron temperatures. Secondly, the resultant intensities were normalized to the intensity at a foil thickness $x = 3 \mu\text{m}$.

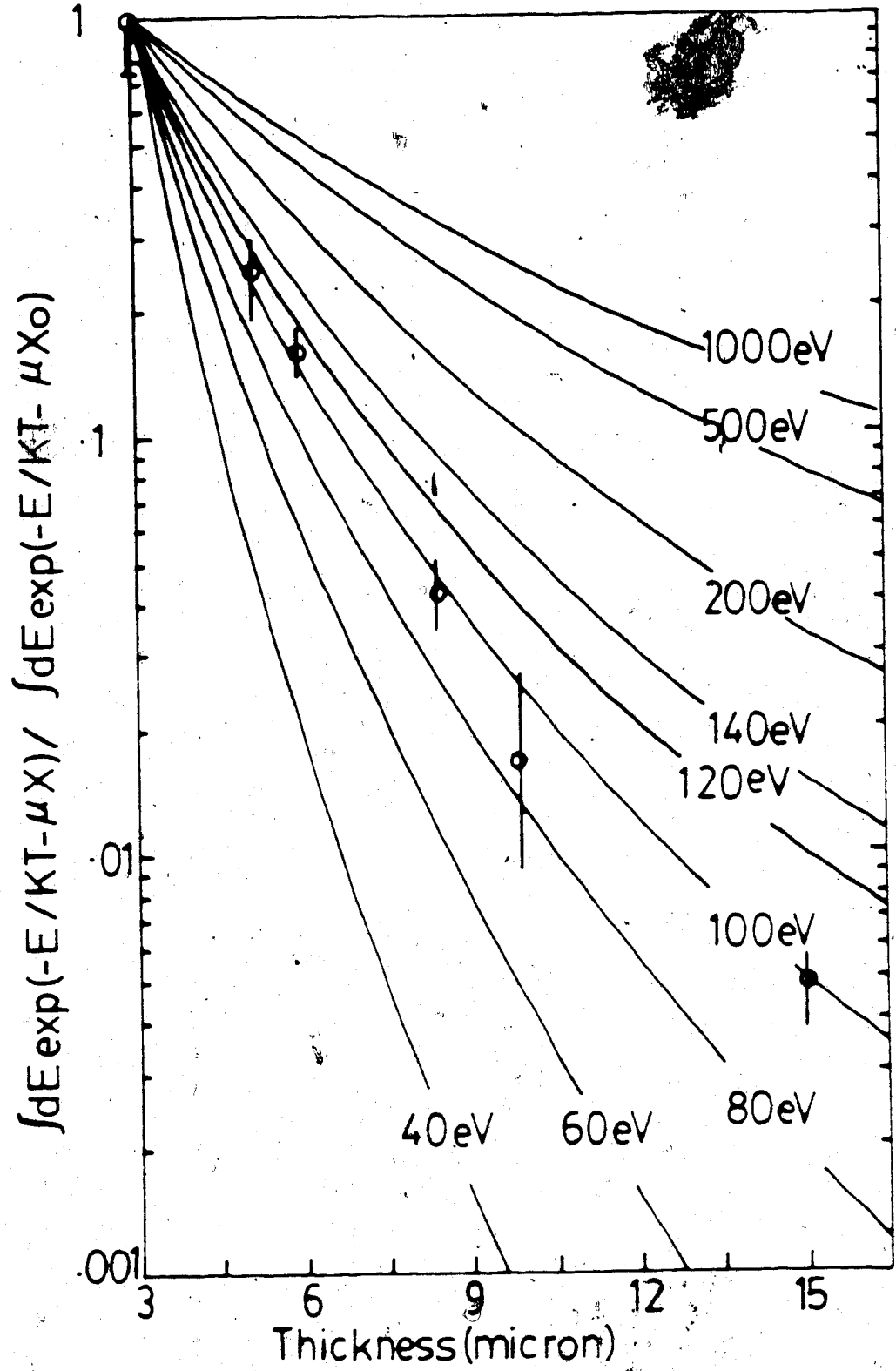


Fig. 2. Normalized x-ray intensity as a function of aluminum thickness. The circles are experimental data from the 10 laser plasma experiment.

thinnest foil used experimentally). To determine the electron temperature the x-ray intensity was measured for different foil thicknesses, normalized to the measured intensity at $x = 3\mu\text{m}$ and plotted in the same figure. Comparison with the experimental data shows that an electron temperature of $\sim 100\text{ eV}$ provides a best fit. The total uncertainty in temperature measurement using this technique is $\pm 20\%$ (limited by the shot to shot variation in the x-ray signal) including the uncertainty due to variation in the plasma condition.

To utilize the above method a simple x-ray detector was designed (Fig. 3.7a). It consisted of an aluminum cylinder (2.5 cm long and 1.5 cm diameter) arranged to accommodate one end of a low loss fiber optic bundle and to hold a plastic scintillator type NE102A (2.1 cm in diameter and 2 mm thick). Aluminum foils ($\geq 3\mu\text{m}$ thick) were used to filter the x-rays reaching the scintillator. These filters were held adjacent to the detector body by a plastic ring and a screw mounted, thin aluminum cap. The other end of the fiber optic bundle was connected to an RCA-8645 photomultiplier tube. The scintillator, fiber optic bundle and photomultiplier tube were all sealed against any light leakage from outside. The photomultiplier signal was amplified by an Avantek preamplifier and recorded on a Tektronix 7834 storage oscilloscope with 7A12 vertical amplifiers (400 MHz bandwidth). A typical x-ray signal ($T_e \ll 1\text{ keV}$ emission) is shown in Fig. 3.7b. For higher energy x-rays, the pulse width is considerably shorter (characteristic of a nonthermal electron distribution).

3.3.3 Electron density (n_e) measurement.

A reliable method for measuring electron density in high temperature and high density plasmas is laser interferometry. This technique

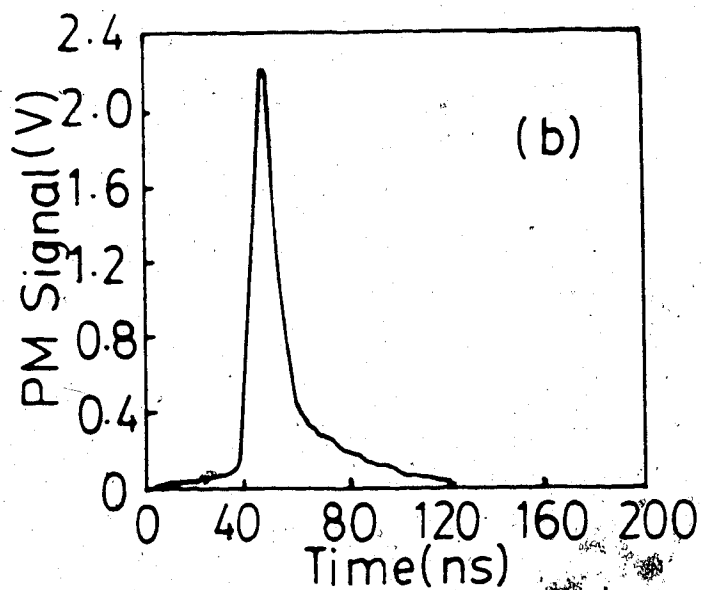
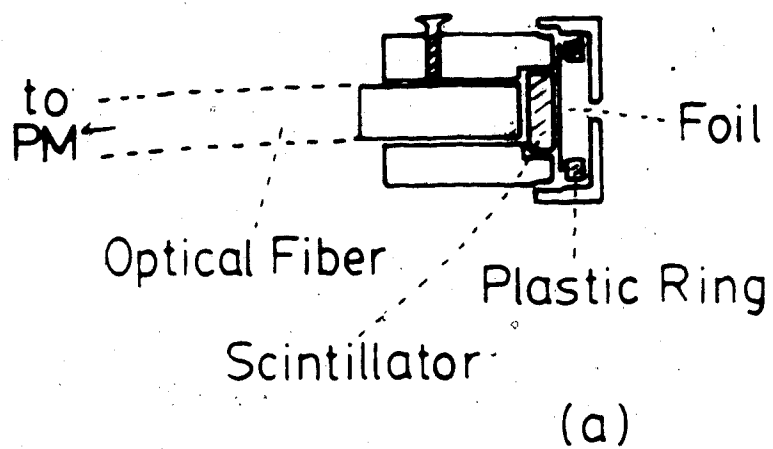


Fig. 3.7 a) X-ray detector b) Typical x-ray signal for $T_e \ll 1$ keV.

is based on measuring the density from the phase shift of an electromagnetic wave as it propagates along two paths of equal length, one through the plasma and the second through a vacuum. In a fully ionized plasma with zero external magnetic field and collisionless electrons the plasma refractive index n , defined as the ratio of phase velocity in vacuum to phase velocity in the plasma, for light frequencies $\omega_0 \gg \omega_{pe}$, is given by [7]

$$n = \left(1 - \frac{\omega_{pe}^2}{\omega_0^2} \right)^{1/2} \approx 1 - \frac{e^2 N_0}{4\pi m^2 \omega_0^2}$$

where λ_0 is the probing laser light wavelength. The phase difference $\Delta\phi$ between the two paths of equal length L as the electromagnetic wave propagates through is given by

$$\Delta\phi = \int_0^L \left(1 - \frac{e^2 N(y)}{4\pi m^2 \omega_0^2} \right) \frac{2\pi}{\lambda_0} dy = \frac{2\pi}{\lambda_0} \int_0^L n(y) dy$$

where y is the abscissa along the path L . The shift in units of fringes ΔN is then given by

$$\Delta N = \frac{\Delta\phi}{2\pi} = \frac{e^2 N_0}{4\pi m^2 \omega_0^2} \int_0^L n(y) dy$$

If the plasma is assumed to be cylindrically symmetric, it is possible to deduce, from the value of ΔN as given by Eq. (1), the electron density along the path L using an Abel inverse algorithm [Appendix C].

For our experiment, different interferometers were tried to measure the electron density. Mach-Zehnder and Michelson interferometers were found to be inadequate because of the low probe laser intensity resulting from the sub-nanosecond pulse, extended over

and the resolution of the light intensity arising from the wave-front
 Hologram. Best results were obtained by using the Fresnel's mirror
 interferometer. This interferometer has two advantages over the other
 interferometers: (a) minimum optical path length (single pass) and
 (b) smaller variation in light intensity (only due to the mirrors'
 reflectivities).

The optical system for Fresnel's mirror interferometer is shown
 in fig. 1. The expanded and collimated ruby laser light passes
 through the plane of S_1 . Only half the beam is used as the test beam
 while the other half is the reference beam. The two lenses L_1 and L_2
 separated by $f_1 + f_2$ (f_1 and f_2 are focal lengths of L_1 and L_2 respec-
 tively), were used for magnifying and localizing the fringes at the
 camera. The Fresnel's mirrors were placed close to L_2 with the contact
 point at the centre of the beam and were incident at an angle α to each
 other. The two halves of the beam were overlapped at S_2 and viewed by
 a camera of magnification M . From the thin lens equation

$$\frac{1}{S_1} + \frac{1}{S_2} = \frac{1}{f_1}$$

$$\frac{1}{S_2} + \frac{1}{S_3} = \frac{1}{f_2}$$

the magnifying lens L_1 is a real object S_1 with $S_1 = f_1 + f_2$
 and $S_2 = f_1$ and the magnifying lens L_2 is a real object S_2 with $S_2 = f_2$
 and $S_3 = f_2$.

$$\frac{1}{S_1} + \frac{1}{S_3} = \frac{1}{f_1} + \frac{1}{f_2}$$

the magnifying lens L_1 is a real object S_1 with $S_1 = f_1 + f_2$
 and $S_3 = f_2$ and the magnifying lens L_2 is a real object S_2 with $S_2 = f_2$
 and $S_3 = f_2$.

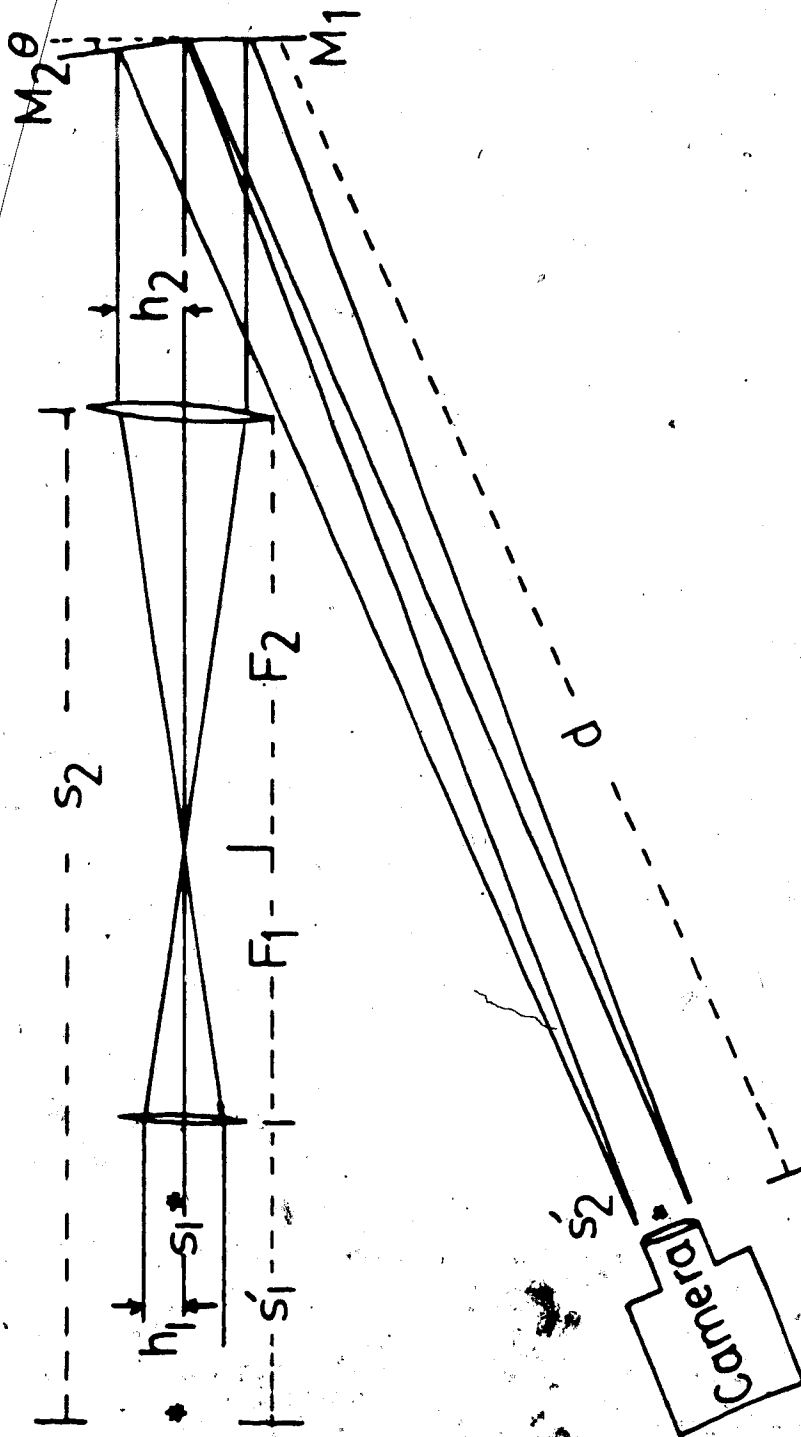


Fig. 3.8 Optical system for Fresnel's bimirror interferometer.

From the interference theory of Fresnel's bimirron, the fringe spacing at s' is [8]

$$\delta = \lambda_0 d / h_2, \quad (3.11)$$

where λ_0 is the ruby laser wavelength and $h_2 (= h_1 \frac{f_2}{f_1})$ is the width of the overlapping area. The corresponding fringe spacing in the plane (resolution) is

$$\delta_p = \delta / M. \quad (3.12)$$

With $f_1 = 30$ cm, $f_2 = 67$ cm, $h_1 = 3$ mm, $s_1 = 26.5$ cm and $l = s_1'$, Eqns. (3.9) to (3.12) give $\delta_p = 27$ μ m. This value agreed very well with the resolution measured by counting the number of fringes across the field of a 200 micron diameter microsphere positioned at the 30_2 laser focus. The total magnification of the telescope and the camera was $M_T = M \times M_c \approx 50$.

The main light source used in this interferometer was a Q-switched Korad K1 ruby laser whose features are given in Table 1.1. An optical shutter was used to slice a 300 ps pulse from the 20 ns FWHM ruby laser pulse. The shutter consisted of three major components: the Lasermetric KD*P Pockels cell, the crossed Glan-Thomson polarizers and the laser triggered spark gap (LTS).

In order to achieve the fastest switching, a solid dielectric laser triggered spark gap was employed. This permitted the earliest avalanche breakdown. The solid dielectric material used in the LTS was a piece of mylar. The switching procedure was done as follows (Fig. 1.2). The Pockels cell was aligned so that the x crystallographic axis was parallel to the polarization axis of the laser light (shown vertical). Further, the Glan-Thomson polarizer was

Wavelength	6943 Å
Polarization	Plane polarized by interscavity Brewster stack polarizer.
Pumping	With helical Xe flash lamp.
Output Energy	1 Joule with full beam size, 0.2 J with 2.5 mm diameter inside cavity iris.
Pulse Duration	10 ns FWHM with λ -switching alone.
Beam Divergence	1.5 mrad.
Pulsed Switching	With Pockels cell, Han-Thomson polarizers and laser triggered spark gap.
Output Pulse Width	10 ns FWHM.

Table 3.1 Features of the λ -switched Ruby Laser

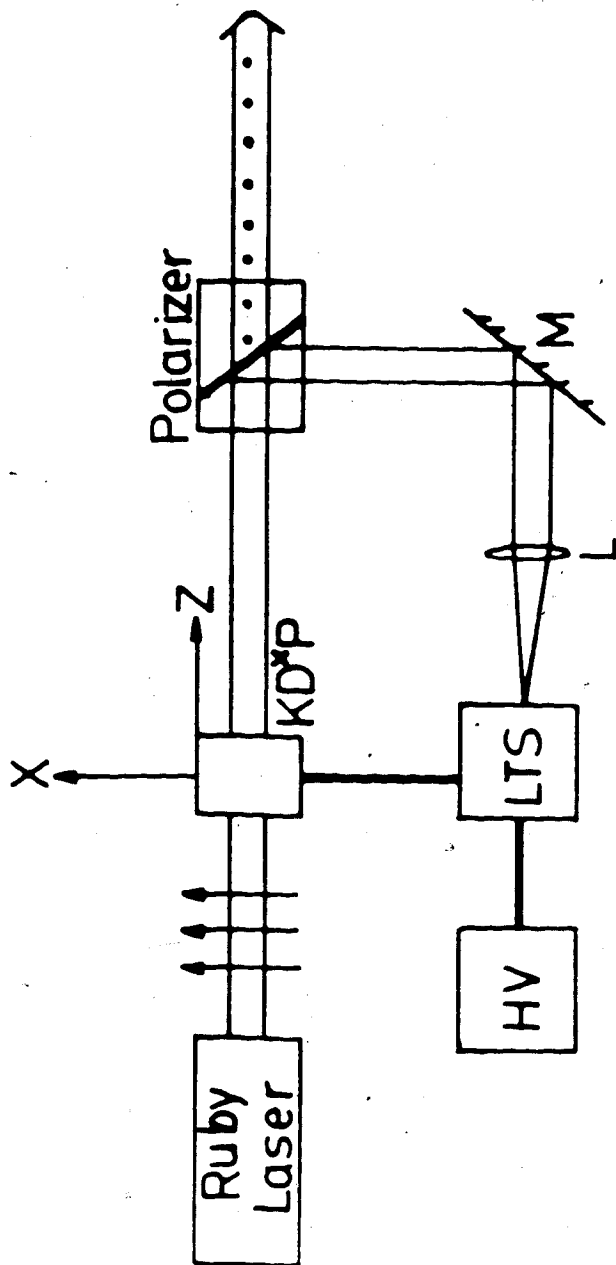


Fig. 3.9 Subnanosecond switching of the ruby laser.

← CO₂

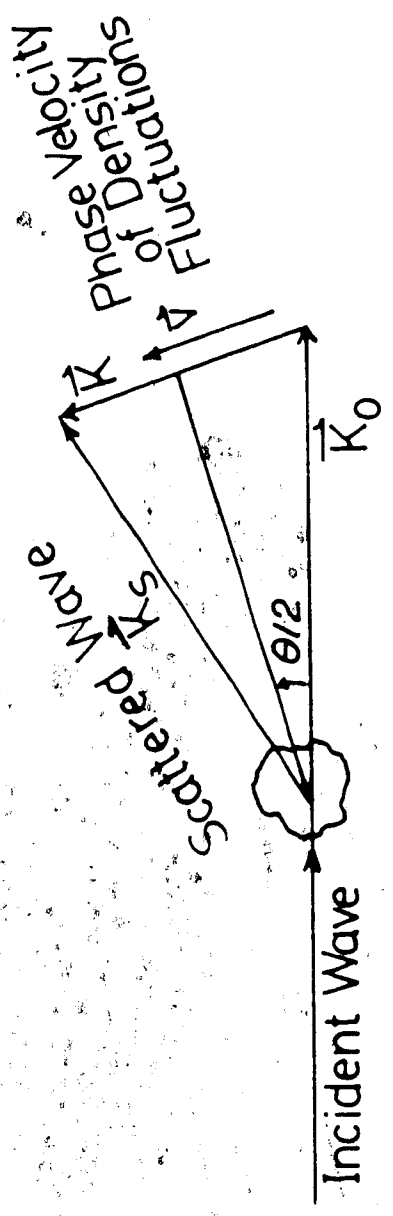


Fig. 1. Interference pattern with CO_2 in the center.

fluctuations in the density distribution. The scattered wave vector is
 $\vec{k}_s = \vec{k}_i - \vec{k}_p$ where \vec{k}_i and \vec{k}_p are the incident wave vector and
 plasma wave vector respectively. The scattered wave vector is
 $\vec{k}_s = \vec{k}_i - \vec{k}_p$ where \vec{k}_i and \vec{k}_p are the incident wave vector and
 plasma wave vector respectively.

It is well known that only dipole distributed charge particles
 scatter electromagnetic waves only in the forward direction. It is the
 inhomogeneity in the density distribution (fluctuations) which do
 the scattering in other directions. The scattering cross
 section is inversely proportional to the square of the particle
 size and therefore light scattering is mainly due to the plasma
 electrons. Random density fluctuations give rise to incoherent scattering
 from individual electrons where the scattered power is simply the
 sum of the individual contributions. If the inhomogeneity is due to
 large scale correlated effects such as plasma waves or nonthermal
 fluctuations, the scattering will be coherent or cooperative and
 substantially enhanced over that caused by thermal fluctuations. For
 a plasma in thermal equilibrium these two regimes can be distinguished
 experimentally by matching the fluctuation wave vector \vec{k}_p to the typical
 shielding distance (coherence length) or the Debye length λ_D . When
 $k_p \lambda_D \ll 1$ the probe's scale length will be much smaller than λ_D , hence
 we deal with individual electron motions. If, on the other hand,
 $k_p \lambda_D \gg 1$ collective effects caused by coherent electron motion
 will be observed. In the following discussion, the parameter $k_p \lambda_D$
 will be introduced, for convenience, to classify the two regimes.

Figure 3.11 shows a schematic diagram of the scattering geometry.
 A plane polarized electromagnetic wave (\vec{k}_i, \vec{e}_i) is incident on a plasma
 and a scattered wave (\vec{k}_s, \vec{e}_s) is observed at an angle θ . If the density
 fluctuation velocity is \vec{v} the observed signal will be Doppler shifted



$$\vec{k} = \vec{k}_s - \vec{k}_0$$

$$k^2 = k_s^2 + k_0^2 - 2k_s k_0 \cos \theta$$

$$\approx 2k_0^2 (1 - \cos \theta)$$

$$k = 2k_0 \sin \frac{\theta}{2}$$

Fig. 3.11 Schematic diagram of the scattering geometry.

in frequency by ω where

$$\omega = \omega_0 - \omega_s = \vec{k} \cdot \vec{v},$$

and

$$\vec{k}_s = \vec{k}_0 - \vec{k}_s.$$

(3.13)

For small frequency shifts, $k_0 \approx k_s$ and the wave number k can be approximated by

$$k = 2k_0 \sin \frac{\theta}{2}. \quad (3.14)$$

Hence, Eqn. 3.13 can be rewritten as

$$\omega = 2k_0 v \sin \frac{\theta}{2}, \quad (3.15)$$

and in this case the shift in wavelength $\Delta\lambda$ will be given by

$$\Delta\lambda = \frac{2\lambda_0}{c} v \sin \frac{\theta}{2}. \quad (3.16)$$

where λ_0 is the wavelength of the incident wave.

In the individual scattering regime, $\alpha \ll 1$, v represents the phase velocity of the electron fluctuations while in the collective scattering regime, $\alpha \gg 1$, v can be related to the phase velocity of both the ion and electron fluctuations. The role of the ions in this case (although they do not scatter) arises from the collective interaction between electrons and ions -- the ions impose fluctuations on the electrons and these fluctuations move with the characteristic ion velocity. Thus the Doppler shift observed corresponds to the ion rather than the electron velocity. Since the velocity depends on the particle temperature, the $\alpha \ll 1$ regime gives information about electron temperature while the ion temperature can be determined in the $\alpha \gg 1$ regime.

Experimentally, the angle of scattering θ and the beam wavelength λ determine the scattering vector \vec{k} . The Raman shift ω , Eq. 3.19, is given by

$$\omega = \omega_0 - \vec{k} \cdot \vec{v}$$

The scattered power as a function of frequency, ω , and angle θ is explicitly related to the electron density through the electron density through the relation

$$P_s(\omega, \theta) = V \sigma(\omega, \vec{k}) S(\omega, \vec{k})$$

where V is the scattering volume, $\sigma(\omega, \vec{k})$ is the scattering cross section and $S(\omega, \vec{k})$ is the density spectrum defined by

$$S(\omega, \vec{k}) = \int_{-\infty}^{\infty} \langle \delta n(\vec{r}, t) \delta n(\vec{r}, t') \rangle e^{-i\omega(t-t')} e^{-i\vec{k} \cdot (\vec{r} - \vec{r}')} dt dt'$$

where σ is the scattering cross section.

The density spectrum $S(\omega, \vec{k})$ is a very important function in scattering, from which the full spectrum of the scattered light can be determined and consequently plasma parameters such as T_e , T_i and fluctuation level can be inferred. With Maxwellian velocity distributions for both ions and electrons in a collisionless plasma, the density spectrum is given by [9]

$$S(\omega, \vec{k}) = \frac{N}{k} \left\{ \left| \frac{1-G_i}{1-G_e-G_i} \right|^2 F_e \left(\frac{-\omega}{k} \right) + z \left| \frac{G_e}{1-G_e-G_i} \right|^2 F_i \left(\frac{-\omega}{k} \right) \right\}, \quad (3.20)$$

where F_e and F_i are the Maxwell-Boltzmann velocity distribution functions for the electrons and ions respectively, z is the ionization degree, N is the total number of electrons in the scattering volume and

The first two terms in Eq. (3.1) are the electron and ion components, respectively. Examining Eq. (3.1) shows that:

the scattered radiation has a dispersion curve dependent on the electron temperature which can be deduced from its width.

The dispersion curve of a central narrow peak in the spectrum whose shape is a function of the electron temperature which can be determined from its width and the dispersion of the electron component symmetrically displaced on either side of the central peak by a frequency $\omega = \omega_0 \pm \omega_p$.

A computer program was written to determine the full spectrum $S(\omega, \vec{k})$ as given by Eqn. (3.2) (Appendix D) in order to be compared with that obtained experimentally.

Another important, measurable quantity in a scattering experiment is the total scattering spectrum $S(\vec{k})$ defined as the integral of $S(\omega, \vec{k})$ over all frequencies. $S(\vec{k})$ can be given approximately (for

The total scattering spectrum is given by

$$S(\omega) = \frac{1}{2\pi} \int_{-\infty}^{\infty} \langle \delta n(t) \delta n(t+\tau) \rangle e^{-i\omega\tau} d\tau$$

$$= \frac{1}{2\pi} \int_{-\infty}^{\infty} \int_{-\infty}^{\infty} \langle \delta n(t) \delta n(t+\tau) \rangle e^{-i\omega\tau} d\tau dt$$

where the brackets denote an average over all frequencies and wave numbers. The total scattering spectrum is given by



The total scattering spectrum $S(\omega)$ is related to the plasma density fluctuations. By definition, the density $\delta n(t)$ and the total scattering spectrum are given by

$$S(\omega) = \lim_{T \rightarrow \infty} \frac{1}{2\pi T} \int_{-T}^T \delta n(t) e^{-i\omega t} dt$$

where t is the time and δn is the Fourier transformed density fluctuation. According to this definition the total spectral density fluctuation averaged over all frequencies and wave numbers can be written in the form [13]

The main effect is that the unstable spectrum will be very small, and for these cases the spectral density function is measured well above the thermal level. Consequently, this will increase the scattering level as indicated in Eqn. (3.3). The direction will depend on the type of instability generated in the plasma.

As an example, the current driven ion instability gives a total spectrum of the ion feature $S_i(\omega)$ of the form $k^{-3} \ln(2/\pi) \omega^3$.

3.1.2 Scattering arrangements:

The Q-switched ruby laser, described previously, with full pulse of ~ 20 ns FWHM was used for Thomson scattering measurements of ion fluctuations. Further slicing of the ruby pulse was avoided to minimize jittering and to save ~ 50% of the output power which may be

...at 50 degrees. In addition, the ruby laser was filtered into three
...lines either at a 90 degree angle...
...between the incident intensity and...
...particularly at small scattering angles.

The scattered light was collected by an F 5-f... 22-15...
...with...
...in a horizontal plane, defined...
...of a vertical plane...
...and the direction of the scattered...
...directed by a set of mirrors
...towards the optical detectors. Precise alignment was
...by placing a microscope...
...illuminating...
...ruby laser alignment...
...laser beam. The HeNe light scattered from the microspheres and col-
...lected by the collecting lens was easily seen by the eye and could
...through the optical system up to the detectors.

The stray light was severely reduced by using a variety of
...techniques including spatial filtering, proper dumping of the ruby
...laser probe beam, blackening and covering of surfaces and using
...Brewster windows instead of normal incidence windows. For some inte-
...grated spectral measurements and, in particular, when the full
...scattered spectrum was under investigation, a Kerr shutter with 5 ns
...gating was used to reduce detection of the relatively long-lived
...bremsstrahlung radiation. In addition, approximately 50% of the
...unpolarized bremsstrahlung radiation was always eliminated by insert-
...ing an optical polarizer in the path of the scattered light.

The timing between scattering and plasma formation was determined by simultaneously monitoring the incident ruby laser beam and either the SRS or the transmitted CO_2 laser signal with a 1 ns resolution.

Spectral measurements of $S(\omega, \vec{q})$ were made by focusing the scattered light into a 100 μm wide entrance slit of an 0.50 m Perkin-Elmer E-1 monochromator (Fig. 3.12). The spectrally dispersed light of the monochromator was imaged onto the target plate of a PARC type 1255 optical multichannel analyzer (OMA) using a relay lens. The OMA target plate consisted of a silicon intensified vidicon with an S-20 photocathode. The 14.5 mm target plate was a linear array of 500 channels, each with 10-20 photon/count sensitivity. The OMA signal was digitized by a PARC type 1206A console, stored in a solid state memory and then recorded as an X-Y display on an oscilloscope. The overall monochromator-OMA dispersion and resolution were 0.24 \AA per channel and 0.1 \AA respectively. This resolution was quite adequate for studying the 0.5-2 \AA range ion feature in the Thomson scattered spectrum. In addition, a deconvolution routine was used in cases where the spectral width of the ion feature and the instrumental resolution were comparable (Appendix E).

Measurement of the electron component of the scattered spectrum was made in two different ways: either by scanning the Perkin-Elmer monochromator at ~ 10 \AA intervals, or by using a 0.25 m Jarrel-Ash monochromator with a 1180 lines/mm grating to give a full spectral scan of ~ 1500 \AA on the OMA (~ 3 \AA per channel). It is for this latter case that a Kerr shutter was used.

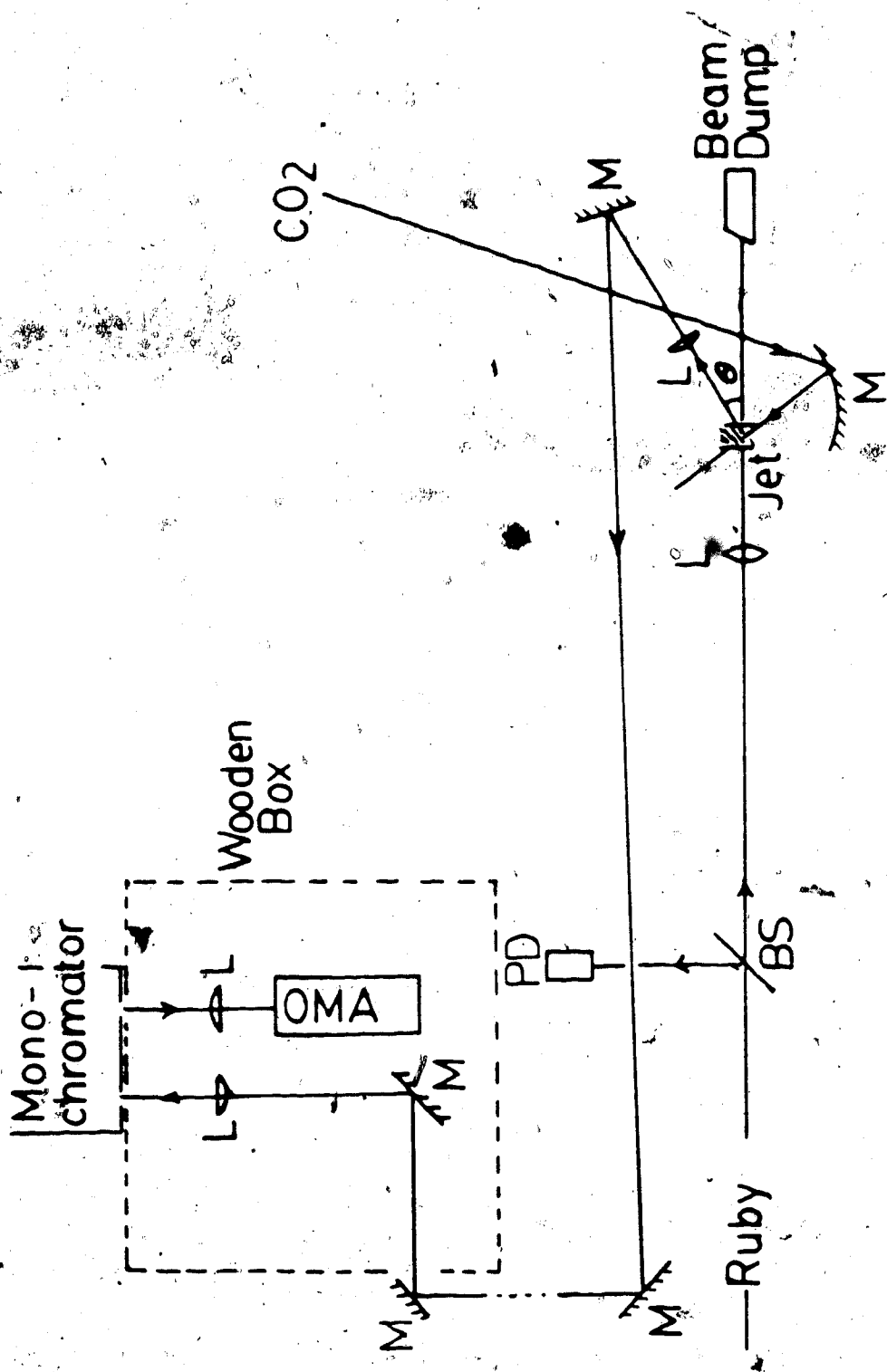


Fig. 3.12 Thomson scattering experiment for $S(\omega, \theta)$ measurements. M: mirror, L: lens, BS: beam splitter, PD: photodiode.

The scattered signal was received in a vertical plane parallel to the target, i.e., in the $\theta = \pi/2$ plane with respect to the parallel polarization plane of the incident ruby laser radiation, which was of particular interest for this system. In addition, measurements were also made in a horizontal plane in order to compare results for different directions but with the same wave vector.

The collection optics were used, the light at a $\theta = \pi/2$ angle of scattering was collected by a lens and the other part at an angle $\theta = \pi/4$ by a lens. The beams collected from the collection were imaged onto the photomultiplier using two more lenses. The lens used for the main beam, separated at angle θ matched the collecting lens in focal length and magnification. An interference filter of the bandwidth was used in front of the photomultiplier to discriminate the component from the rest of the spectrum.

For absolute measurements of $I(\theta)$, the scattered power (as defined by Eq. (1.3)) was calibrated by comparing the Thomson scattered signal with that from emission from a blackbody source.

Because of the high stray light levels detection of Rayleigh scatter was not possible and therefore the parameters I_0 , α , β and Ω were carefully determined: the scattering volume was determined from the intersection dimensions of the focused ruby laser, the CO_2 laser, and the collecting optics - the latter being limited by the 100 μm iris size in front of the photomultiplier; the ruby light intensity was determined by measuring the ruby laser power, energy and spatial distribution with a pinhole and long focal length lens; the solid angle was determined from the f number of the collecting lens; the electron plasma density was measured with Fresnel's bimirror

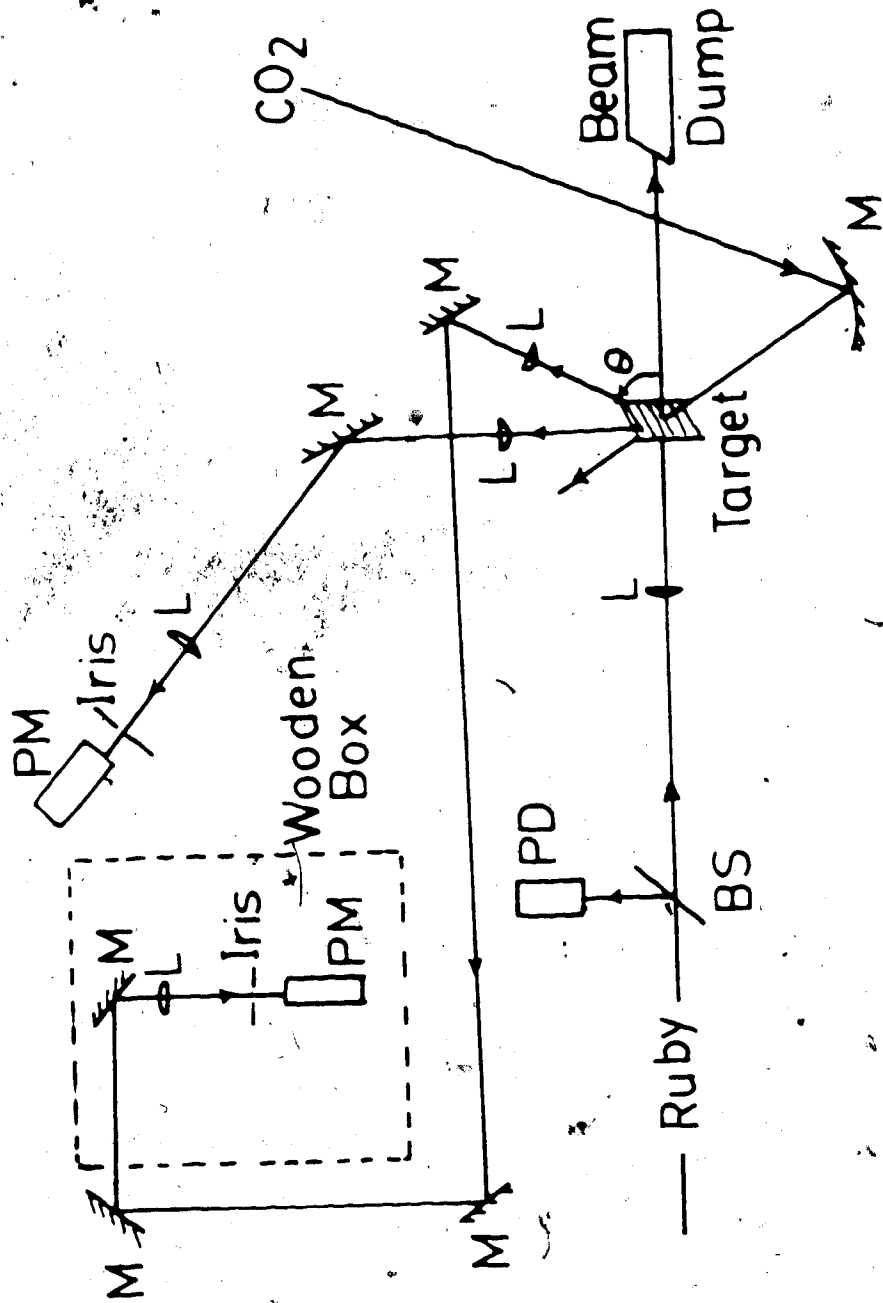


Fig. 3.13 Thomson scattering experiment for atomic measurements. M: mirror, L: lens, BS: beam splitter, PD: photodiode, BB: beam dump, IR: optical filter.

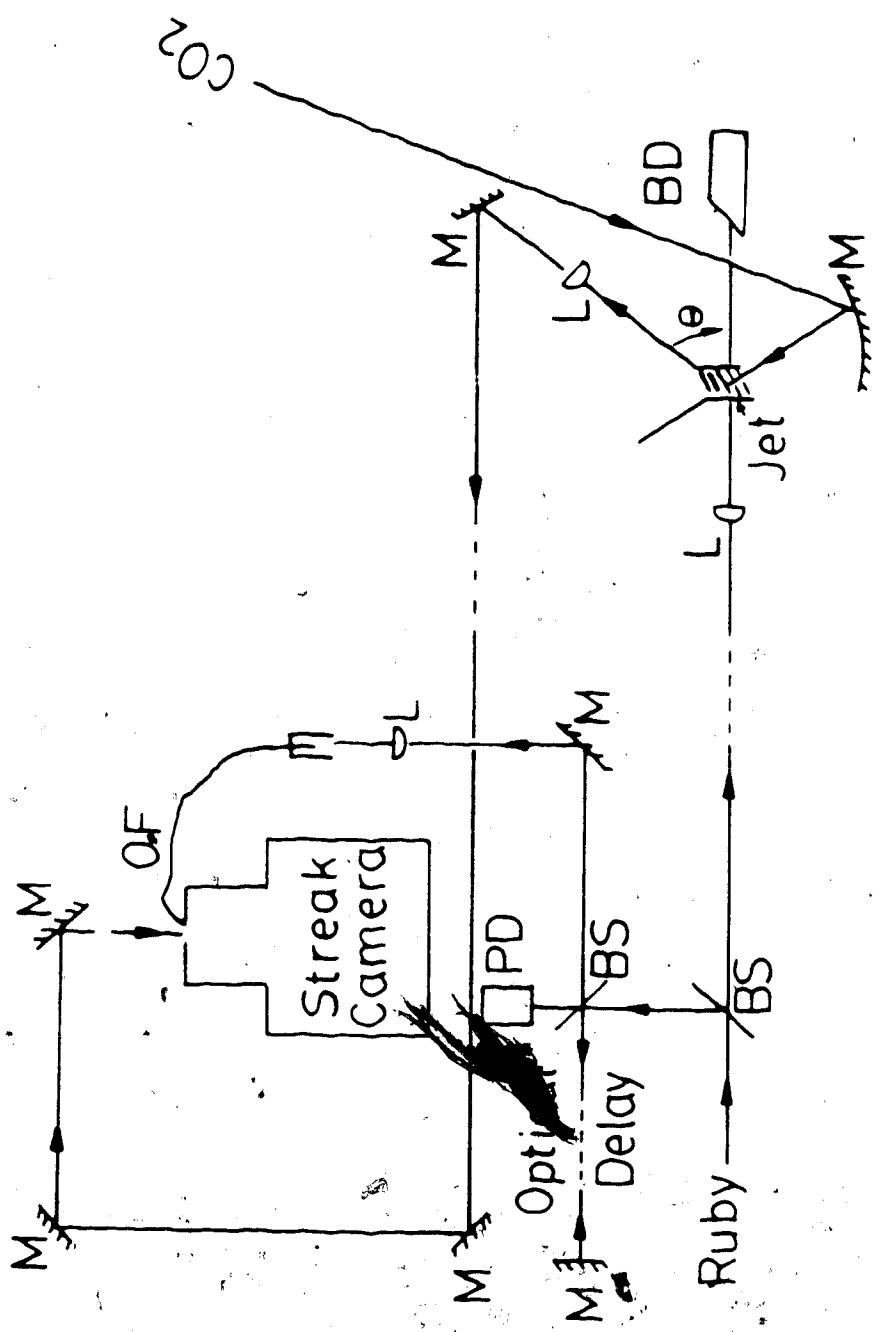


FIG. 3.14 Thomson scattering experiment for CO_2 laser. BS: beam splitter, PD: photodetector, M: mirror, L: lens.

...microscopic ... of the ... layer ... from ...
... was ...

EXPERIMENTAL RESULTS AND DISCUSSION

4.1 Introduction

In this chapter the results of various laser plasma interaction experiments are presented. Plasma parameters such as electron temperature and density as well as the general behaviour of the laser plasma interaction are first discussed. In later sections the results of the Thomson scattering experiment are given.

Temporal characteristics of plasma formation are shown in Fig. 4.1. Figure 4.1a shows the CO₂ laser power transmitted (as collected with an f/2 lens and detected by a photon drag). It can be seen from the figure that gas breakdown occurs approximately 10-15 ns before the peak of incident CO₂ laser beam is reached. In the breakdown region the transmitted signal level approaches a noise level as a result of absorption and scattering. As can be seen in Fig. 4.1b the transient phase of stimulated Brillouin backscattering takes place only once in the laser plasma interaction. This happens 0.4 ns after gas breakdown regardless of the fluctuation in the incident laser power. The measured SBS level was approximately 10% at an operating laser intensity of $5 \times 10^{12} \text{ W/cm}^2$. In a time interval of 0.2 ns following SBS, nearly complete absorption occurs. This is followed by target burn through and refraction domination (not seen in the figure because it takes place outside the focal cone of the

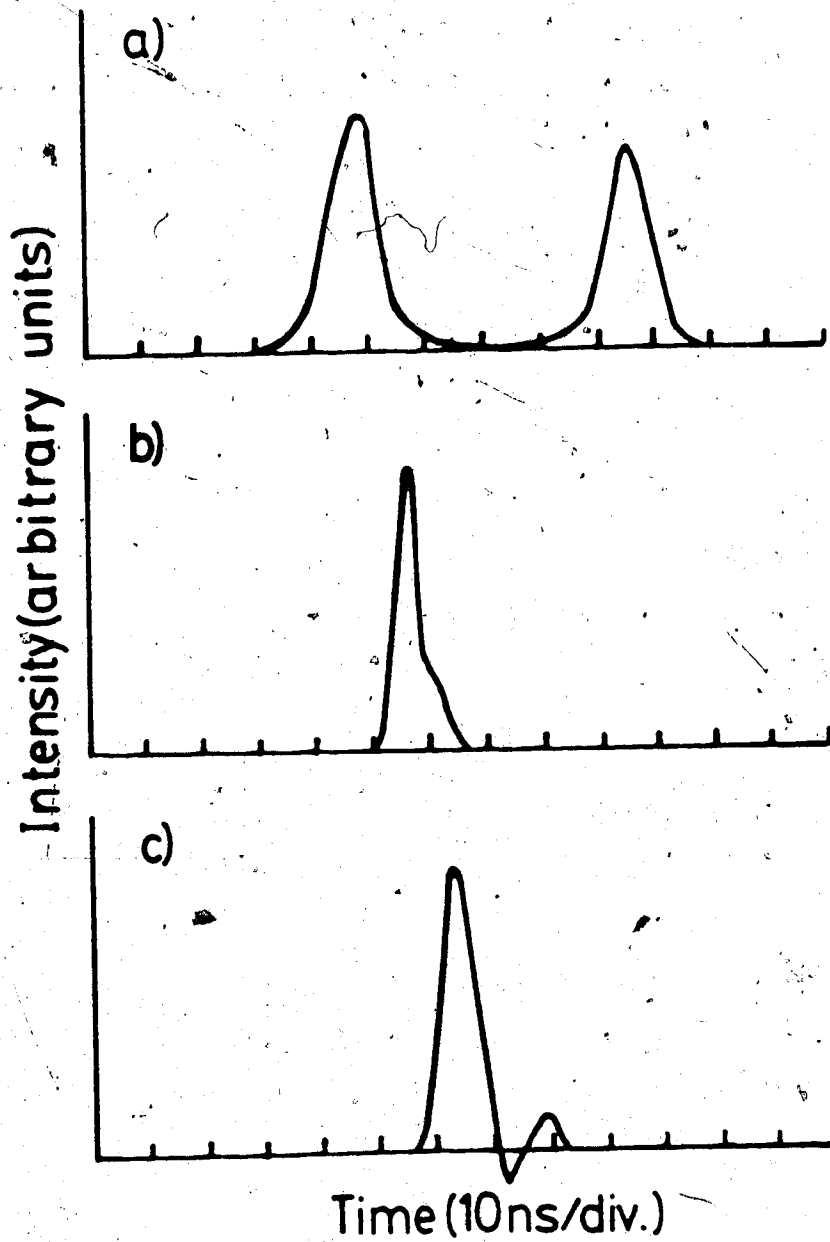


Fig. 4.1 General behaviour of the CO_2 laser plasma interaction;
a) Transmitted CO_2 laser signal, b) SBS signal,
c) Thomson scattering signal.

forward transmission). Throughout these experiments the SBS signal was used as a reference for timing purposes simply because of being strong, fast and well characterized (details of the SBS instability generated in the present experiment can be found in [1] and [2]). Finally, Fig. 4.1c shows a typical Thomson scattering signal from ion fluctuations (recorded by a photomultiplier), the characteristics of which will be discussed later in this chapter.

Results from x-ray measurements indicate a single electron temperature of 100 ± 20 eV (Fig. 3.2) at focused CO_2 laser intensity. This is quite different from those reported in [1] where two Maxwellian electron temperatures were identified at ~ 160 eV and ~ 1 keV. The absence of the hot electron component in the present experiment may be due to the many changes in laser plasma interaction conditions such as lower electron plasma density and reduced laser intensity level compared to the earlier work, though the precise reasons are not understood. The absence of the critical density layer (as will be seen later) eliminated the role of many mechanisms responsible for producing energetic electrons such as two stream instability and $e-i$ decay instability at critical and resonance absorption.

Since the x-ray detection risetime was ~ 3 ns, temporal evolution of the x-ray emission could not be examined in detail. However, a pulse duration of < 20 ns was observed in all the x-ray measurements. This was much less than the CO_2 laser pulse duration and could be related to the low density plasma that results at a later time due to plasma expansion.

Direct measurement of ion temperature in this experiment was not possible. This is because Faraday cups and electrostatic ion analyzers require a low pressure environment ($< 10^{-4}$ torr) and therefore could not be used in our experiment since we were operating with a 10 torr helium background (necessary to stabilize the oxygen jet). Optical spectroscopy also was not possible since, for a high degree of ionization, most of the oxygen lines are in the x-ray region. Thomson scattering, on the other hand, would potentially be an excellent technique for measuring the ion temperature if scattering from thermal plasma fluctuations could be observed. But because the measured Thomson scattering spectra were dominated by enhanced scattering from nonthermal ion fluctuations, isolation of the thermal component from the nonthermal one was again impossible. However, the ion temperature was roughly estimated from enhanced Thomson scattering of the nonthermal ion fluctuations as will be discussed later.

The other very important parameter to be determined accurately is the plasma electron density. Unfortunately, due to a number of changes in the operating conditions of the CO_2 laser plasma interaction, particularly focused laser conditions (a new focusing mirror with larger aberrations was used), different regimes were observed and therefore must be dealt with separately. Thus, according to the density regime prevailing during different periods in the thesis research, two cases will be considered for analysis (Case I and Case II).

In discussing the plasma density throughout these analyses, we will refer to Fig. 4.2 where a schematic diagram of the experiment is shown. In this figure it is assumed that the CO_2 laser light is

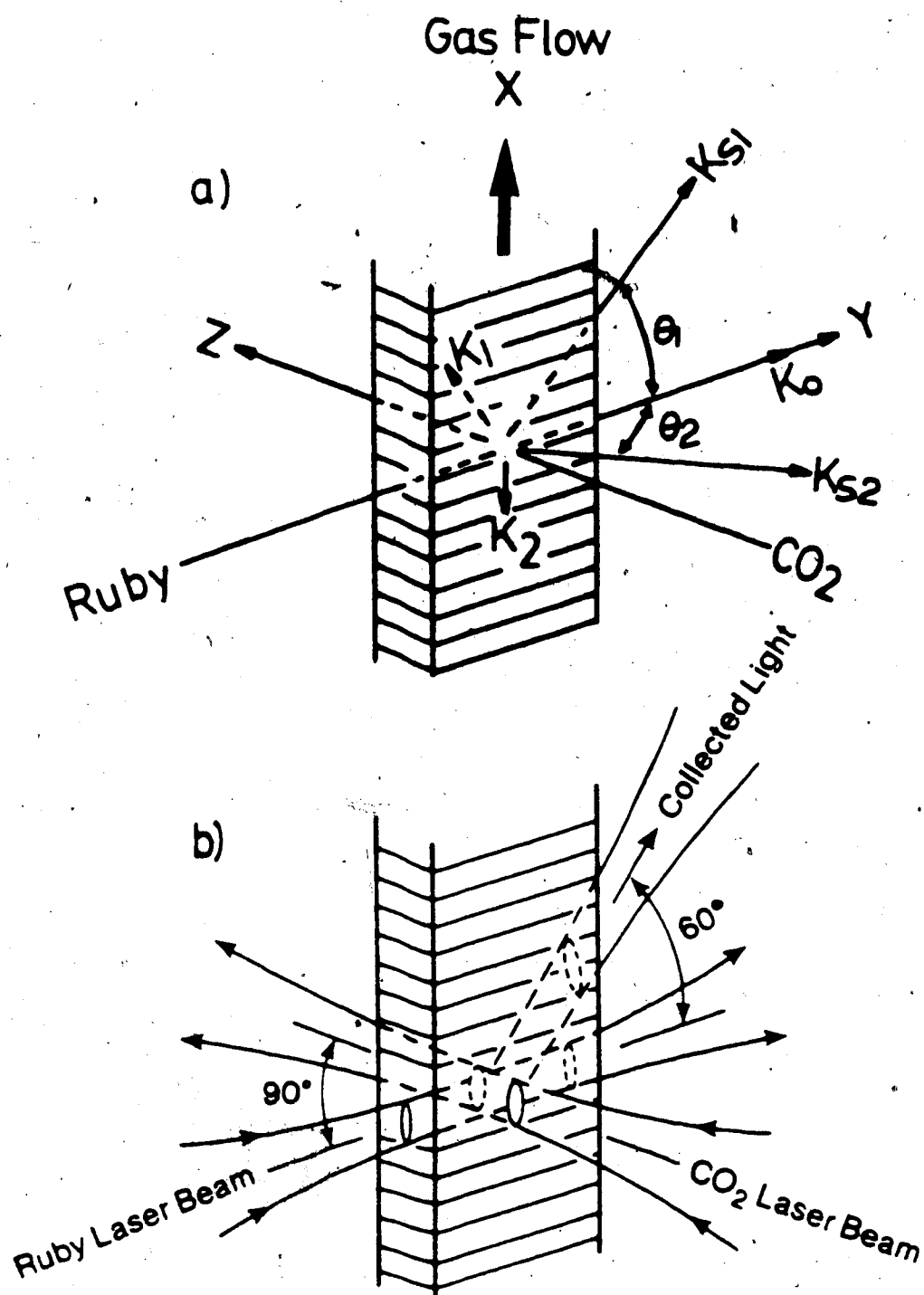


Fig. 4.2 Schematic diagram of the Thomson scattering experiment;
 a) Wave vectors at the incident, scattered and ion waves,
 b) Intersection of the CO₂ laser beam, ruby laser beam and
 the collected light.

propagating in the X-direction and the O_2 gas is flowing in the Y-direction. Because a symmetric CO_2 laser beam was used, the plasma density was assumed cylindrically symmetric (i.e., in the X-Y plane). The Abel transform could therefore be used to invert plasma interferograms in order to obtain the radial electron density profiles at different axial positions in the plasma. Following SBS, the highest density for both cases (I and II) is reached and maintained for a period of ~ 8 ns. The density then decays due to hydrodynamic expansion.

Figures 4.3-4.5 illustrate the behaviour of the electron density in Case I. Figure 4.3 shows a typical unfolded interferogram of the plasma density ~ 3 ns after the start of SBS. The radius of the plasma spot size at the leading edge is ~ 100 μm . The radius then increases almost linearly with Z according to the relation $R \approx 100 + 0.5Z$ where both R and Z are in μm . As can be seen in the figure, the plasma density is maximum at $r = 0$ and then decays sharply towards zero at the plasma edge ($r = R$). Figure 4.4 shows the variation of the peak density ($r = 0$) with the axial distance Z. According to the figure, this variation is almost linear and the plasma density at the focal point is approximately equal to $0.8 n_c$. In this early stage of the plasma formation, the plasma was found to extend ~ 0.3 mm in the axial direction with a scale length $L = n \frac{dz}{dn}$ of 200 ± 30 μm at the focal point. At a later time ($t > 8$ ns) the plasma density decreased and occupied a larger volume (Fig. 4.5) with $L \geq 800$ μm and $n \approx 0.25 \cdot n_c$ at the focal point.

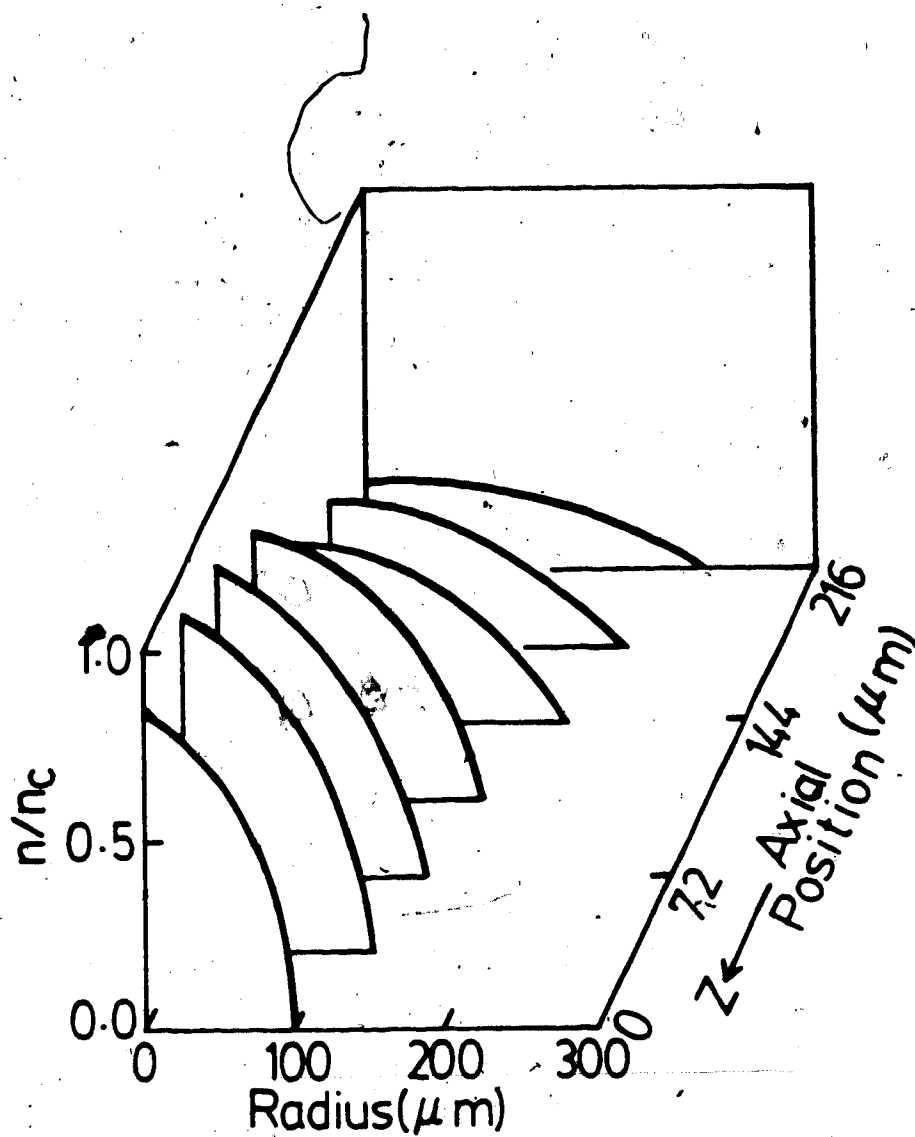


Fig. 4.3 Unfolded interferogram of the plasma density of Case I at ~ 3 ns after the start of SBS.

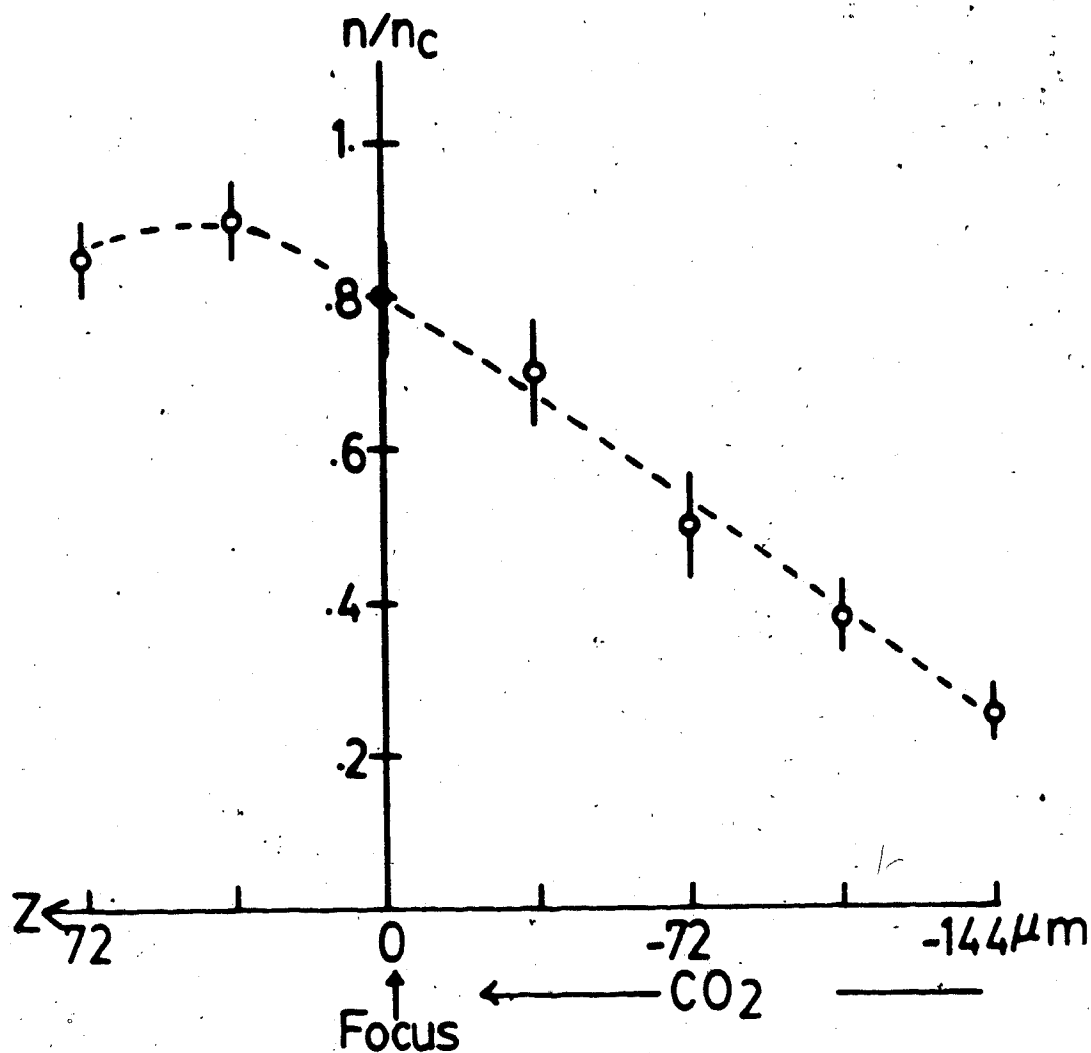


Fig. 4.4 Variation of the peak density ($r=0$) with the axial distance Z at ~ 3 ns after the start of SBS for Case I.

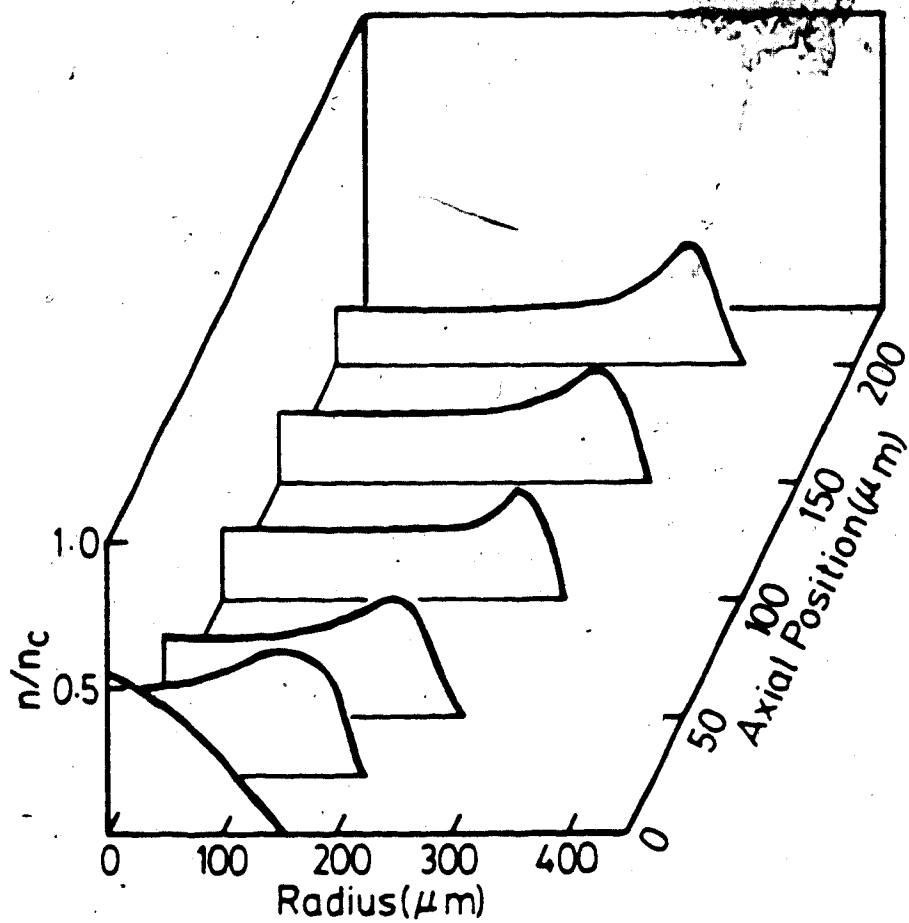


Fig. 4.5 Unfolded interferogram of the plasma density of Case I at ~ 10 ns after the start of SBS.

Figures 4.6 and 4.7 show the measured plasma density profiles for Case II at a time 2 - 8 ns after the start of SBS. In comparison with the previous figures, it can be seen that the density is generally lower than in Case I ($\sim 0.7 n_c$ at the focus) with larger axial extension (0.8 - 1 mm with $L = 450 \pm 50 \mu\text{m}$). Similar behaviour to that shown in Fig. 4.5 (for Case I) was observed at a later time, $t > 8$ ns for Case II.

It should be mentioned that although the change in the focused laser conditions affected the plasma density, no significant change was seen in the electron plasma temperature (taking into account the 20% error associated with the x-ray results). The temporal evolution of the electron temperature might not have been identical for Case I and Case II, however, the time integrated x-ray measurements did not show any difference.

Thomson scattering results and ion turbulence measurements obtained in Case I will be presented in Section 4.2, while those results obtained in Case II will be discussed later in Section 4.3.

4.2 Ion Turbulence and Absorption

4.2.1 Laser light absorption in CO_2 laser plasma interaction.

It has been shown in an experiment very similar to the present one [1,3] that inverse bremsstrahlung absorption cannot fully account for the nominal 100% absorption observed. With a scale length as small as $100 \mu\text{m}$ and with the existence of a critical density layer the fraction of light which should be absorbed is only 40% (as calculated from Eqn. (2.10)), assuming a linear density profile. Similarly, we

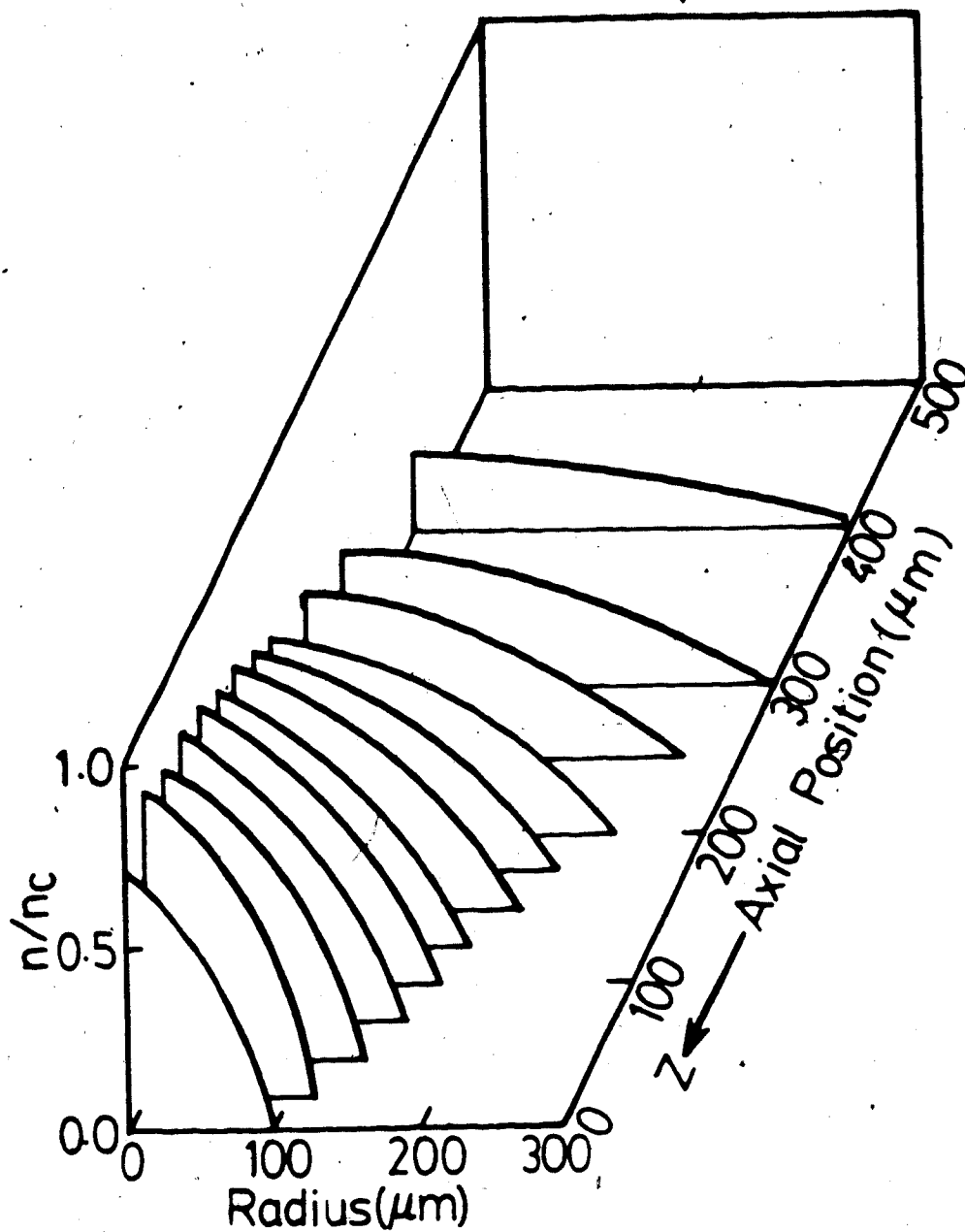


Fig. 4.6 Unfolded interferogram of the plasma density of Case II at ~ 3 ns after the start of SBS.

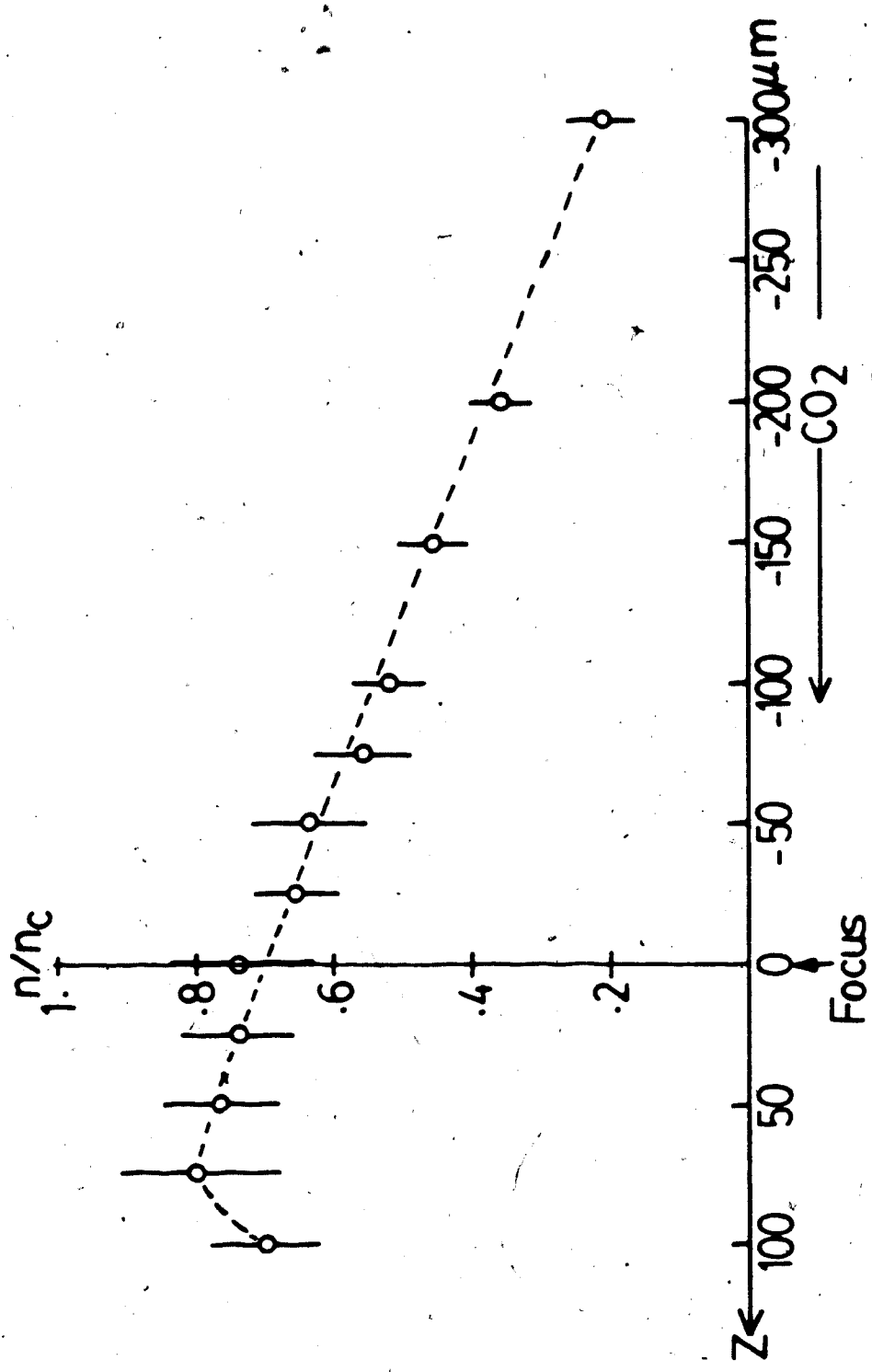


Fig. 4.7 Variation of the peak density ($n=0$) with the axial distance Z at $t = 3$ ns after the start of SRS for Case II.

can estimate the fraction of the incident light being absorbed by the inverse bremsstrahlung mechanism in the present experiment. Since Eqn. (2.16) is only valid for a linear density profile reaching critical density, it cannot be used to estimate the fraction of the absorbed light in the present experiment ($n < n_c$). However, the original formula of Dawson, *et al.* [4] can still be applied:

$$A = 1 - e^{-K_T}$$

$$K_T = \int_{-\infty}^{\infty} \frac{\nu_{ei}(z) \omega_{pe}^2(z) dz}{\omega_0 |\omega_0^2 - \omega_{pe}^2(z)|^2} \quad (4.1)$$

By using the measured density profile (Fig. 4.4) and assuming a uniform plasma temperature over the entire region (with $T_e = 100$ eV), Eqn. (4.1) can then be integrated numerically to give an absorbed fraction $A \leq 0.7$. It should be noted, however, that in calculating the collision frequency ν_{ei} , the charge state was taken to be $Z = 6$ which is the maximum value estimated by comparing the neutral oxygen density and the maximum plasma density ($n = 0.9 n_c$). This is consistent with the ionization state calculated for $T_e = 100$ eV [5]. Thus, the above calculated absorption fraction represents an upper limit. At a later time ($t > 8$ ns after the start of SBS) where the density is very low or for $Z < 6$ the absorption is even lower. Therefore, at least 30% of the incident light has to be absorbed by a mechanism or mechanisms other than inverse bremsstrahlung.

A variety of processes may account for the discrepancy between the observed and classical calculated absorption. Parametric processes are one possibility. Stimulated Brillouin scattering has the lowest

threshold among the possible parametric instabilities. However, the SBS scattered power was not more than 10% of the incident power. Moreover, the anomalous absorption and SBS occurred at different times during the interaction (i.e., absorption follows SBS). Thus, while SBS can influence subsequent events by enhancing fluctuations, it cannot account directly for absorption.

The absence of the critical density and high energy electrons eliminate resonance absorption and absorption due to OTS and $e-i$ decay instabilities (at critical density) as candidates for the observed absorption. Equally, the Raman instability cannot contribute to absorption in the present experiment because of its high threshold ($> 10^{13} \text{ W/cm}^2$). Moreover, it was sought for but was not detected experimentally.

Other parametric instabilities such as two-plasmon decay and off-resonance OTS and PD may contribute to absorption directly but in a very limited way because of relatively high thresholds (compared to SBS, e.g.). However, the OTS and PD off-resonance may contribute to absorption indirectly by generating ion turbulence which, in turn, can enhance the absorption as explained in Chapter 2. This mechanism will be discussed here in light of the experimental data.

4.2.2 Scattering parameters.

To study ion turbulence and its effect in laser light absorption, both spectral and temporal behaviour along with the turbulence level of the ion fluctuations were measured using ruby laser Thomson scattering techniques (described previously in Chapter 3). The scattered light was detected at an angle θ with respect to the incident ruby laser direction (Fig. 4.2a) either in the plane of the target (\vec{k}_{s1} in the

X-Y plane) or in a plane defined by the ruby laser and CO₂ laser propagation directions (\bar{k}_{s2} in the Y-Z plane). This probed the ion fluctuation wave vector \bar{k}_1 perpendicular to the CO₂ laser light wave vector (parallel to the unpolarized electric field of the CO₂ laser radiation, i.e., in the X-Y plane) and \bar{k}_2 in the Y-Z plane at an angle $90^\circ \pm \theta_2/2$ with the electric vector of the CO₂ laser light. Both \bar{k}_1 and \bar{k}_2 are related to the incident ruby laser wave vector \bar{k}_0 through the equation $\bar{k}_{1,2} = \bar{k}_{s1,2} - \bar{k}_0$ and the wave numbers are given by

$$k_{1,2} \approx 2k_0 \sin \theta_{1,2}/2 = 1.81 \times 10^5 \sin \theta_{1,2}/2 \text{ cm}^{-1}. \quad (4.2)$$

We will refer to these two cases (scattering in the X-Y and Y-Z planes) as Case a and Case b respectively.

If we realize that the plasma density varies from 0 - 0.9 n_c the range of $k\lambda_D$ will be very wide ($k\lambda_D \geq 0.224$ for $\theta = 60^\circ$, e.g.). However, the scattering volume was defined by the intersection of the focused ruby laser, the CO₂ laser, and the collecting optics (Fig. 4.2b) and was approximately $150 \times 150 \times 100 \mu\text{m}^3$ around the laser focal point. Therefore, by referring to Figs. 4.3 to 4.5, the density of the probed plasma roughly ranged between 0.4 and 0.9 n_c . The corresponding $k\lambda_D$ and α (for $\theta = 60^\circ$) were $0.224 \leq k\lambda_D \leq 0.336$ and $3.0 \leq \alpha \leq 4.5$.

The scattered light was detected simultaneously with a photomultiplier for temporal behaviour and an optical multichannel analyzer-monochromator system for (time integrated) spectral measurements.

4.2.3 Temporal measurements of ion fluctuations.

The power of the scattering signal (recorded by the photomultiplier) was many orders of magnitude greater than that which would arise from scattering from thermal fluctuations. A simple calculation

shows that refraction of a ruby laser beam caused by electron density gradients cannot account for the enhanced scattering at the observed angles. This is because the deflection angle is small compared to the scattering angles used (22° - 90°). The deflection angle is given by $|\delta|$

$$\Delta\alpha = \int_0^L \left(\frac{1}{\mu} \right) \bar{\nabla} \mu \times dz, \quad (4.3)$$

where the refractive index $\mu \approx 1 - 2\pi n e^2 / m\omega_0^2$ and ω_0 is the ruby laser light frequency. By approximating the density n by $0.9 n_c \left[1 - \frac{Z}{0.3} \right]$, where Z is in mm, Eqn. (4.3) gives $\Delta\alpha \sim 10^{-3}$ rad ($= 0.057^\circ$). This indicates that simple refraction is not expected to be important and that the observed scattering is due to enhanced fluctuations. In addition, the fact that no electron features were observed (at large ω shifts such as the scattering theory predicts for $\alpha > 1$) demonstrates that the enhancement is most likely due to ion fluctuations. This will be discussed in more detail when spectral results are analyzed.

Temporal characteristics of the enhanced Thomson scattering in relation to the transmitted CO_2 laser radiation and strong stimulated Brillouin backscattering are shown in Fig. 4.1c. This time behaviour (although limited by the 3 ns photomultiplier risetime) is important in identifying (a) that the ion turbulence follows rather than occurs simultaneously with strong parametric interaction such as SBS, (b) that the duration of enhanced ion fluctuations (≤ 10 ns) is consistent with the observed period of anomalous absorption. The first observation clearly shows that while subsequent mode coupling or other plasma transport mechanisms may be responsible for ion turbulence, the fluctuations associated with the prompt SBS could not be directly responsible. Likewise, two-plasmon decay is coincident

with SBS. Ruby laser scattering from ion fluctuations induced by SBS, however, did not show any time delay between the SBS signal and the Thomson scattering signal. These observations, together with the density measurements discussed before, indicate that enhanced ion fluctuations were generated in a relatively high density plasma. This is supported by the fact that, on average, the strongest scattering signal was observed only when the peak of the incident ruby laser took place within 10 ns of the start of SBS.

Scattering in the X-Y plane (Case a) and in the Y-Z plane (Case b) showed similar behaviour for a 60° scattering angle. The same characteristics were also observed for other angles, e.g. 22° in Case a and 90° in Case b. Details of the temporal structure of the ion fluctuations could not be obtained because of the limited resolution of the photomultiplier. Better information of the fine structure of the ion turbulence was obtained by employing a streak camera; the results will be discussed in Section 4.3.

4.2.4 Spectral measurements.

The main goal of the spectral measurements was to identify the source of the enhanced scattering. In the case of thermal plasma the spectra (theoretical) of the scattered light $S(k, \omega)$ for $\theta = 60^\circ$ are shown in Figs. 4.8 and 4.9 for $n = 0.8 n_c$ and $n = 0.4 n_c$ respectively (for $T_i = T_e = 100$ eV). As can be seen, the central region of each spectrum consists of an ion feature (solid) and an electron feature (dashed), the intensity of the ion component being approximately an order of magnitude higher than that of the electron. In addition, the electron component has two side components located at wavelengths $\Delta\lambda$, where $\Delta\lambda$ is given by

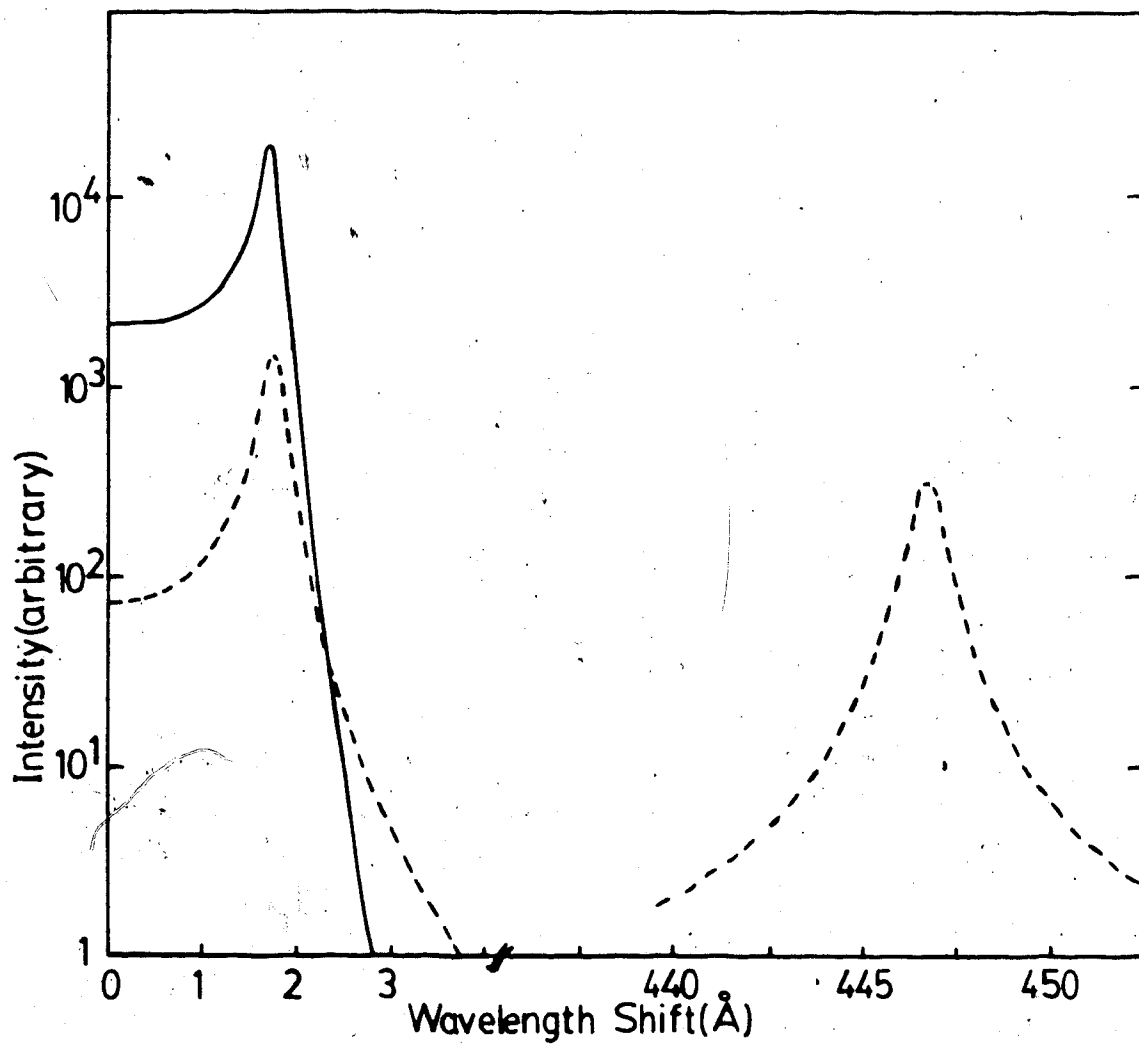


Fig. 4.8 Calculated ruby Thomson scattering spectrum for a thermal oxygen plasma with $T_i = T_e = 100$ eV, $n = 0.8 n_c$, $z = 6$ and $\theta = 60^\circ$ showing the ion component (solid) and the electron component (dashed).

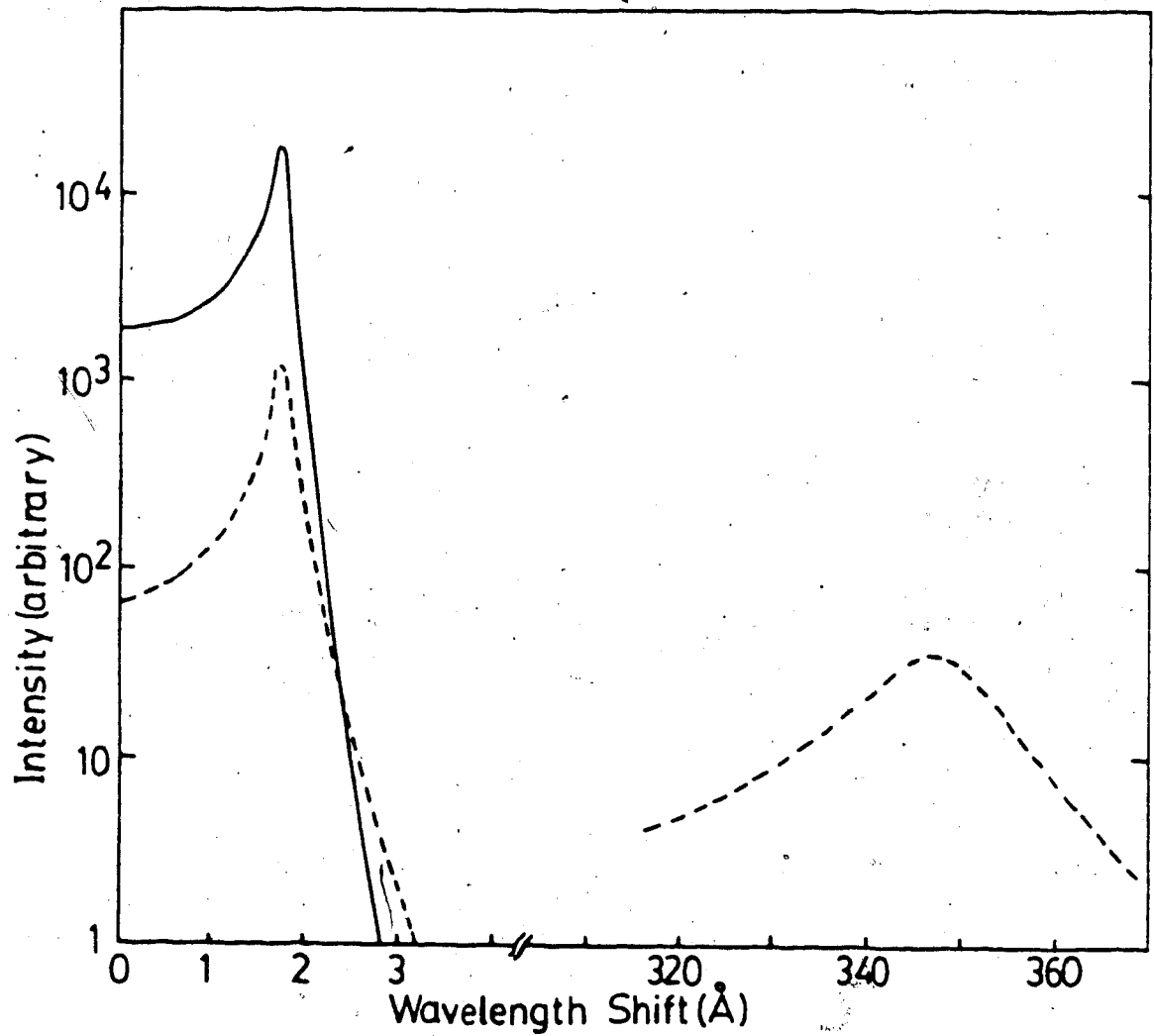


Fig. 4.9 Calculated ruby Thomson scattering spectrum for a thermal oxygen plasma with $T_i = T_e = 100$ eV, $n = 0.4 n_c$, $z = 6$ and $\theta = 60^\circ$ showing the ion component (solid) and the electron component (dashed).

$$\Delta\lambda = \lambda_0 \pm \frac{\lambda_0}{\omega_0} \Delta\omega, \quad (4.4)$$

in which λ_0 and ω_0 are the ruby laser wavelength and frequency respectively and $\Delta\omega$ is the frequency of the electron plasma waves determined approximately from the Bohm-Gross dispersion relation

$$\Delta\omega^2 \approx \omega_{pe}^2 (1 + 3k^2 \lambda_D^2). \quad (4.5)$$

Experimentally, only the central parts of the scattered spectra were observed. Attempts were made to observe any electron plasma wave components by scanning over $\pm 600 \text{ \AA}$ around λ_0 (corresponding to $n \leq 1.5 n_c$) but were not successful. This indicated that there was no significant enhancement in the electron feature and that the observed scattering was mainly due to strong ion fluctuations. For nearly thermal plasma the electron wave feature, of course, is very weak for the values of $k\lambda_D$ experimentally probed and will therefore be buried in the bremsstrahlung radiation.

Comparison of Figs. 4.8 and 4.9 shows that due to the relatively small change in the plasma density, only the side components of the electron feature change materially (location and shape). The shape of the ion component, on the other hand, varies significantly with ion temperature and degree of ionization z as shown in Fig. 4.10 (for $n = 0.8 n_c$ and $T_e = 100 \text{ eV}$ oxygen plasma).

Since the measured spectra correspond to time-integrated scattering, precise structure is lacking because of frequency smearing effects associated with time varying plasma conditions. Nevertheless, considerable information can be obtained from these spectra. Spectral measurements of the scattered ruby light are summarized in Fig. 4.11

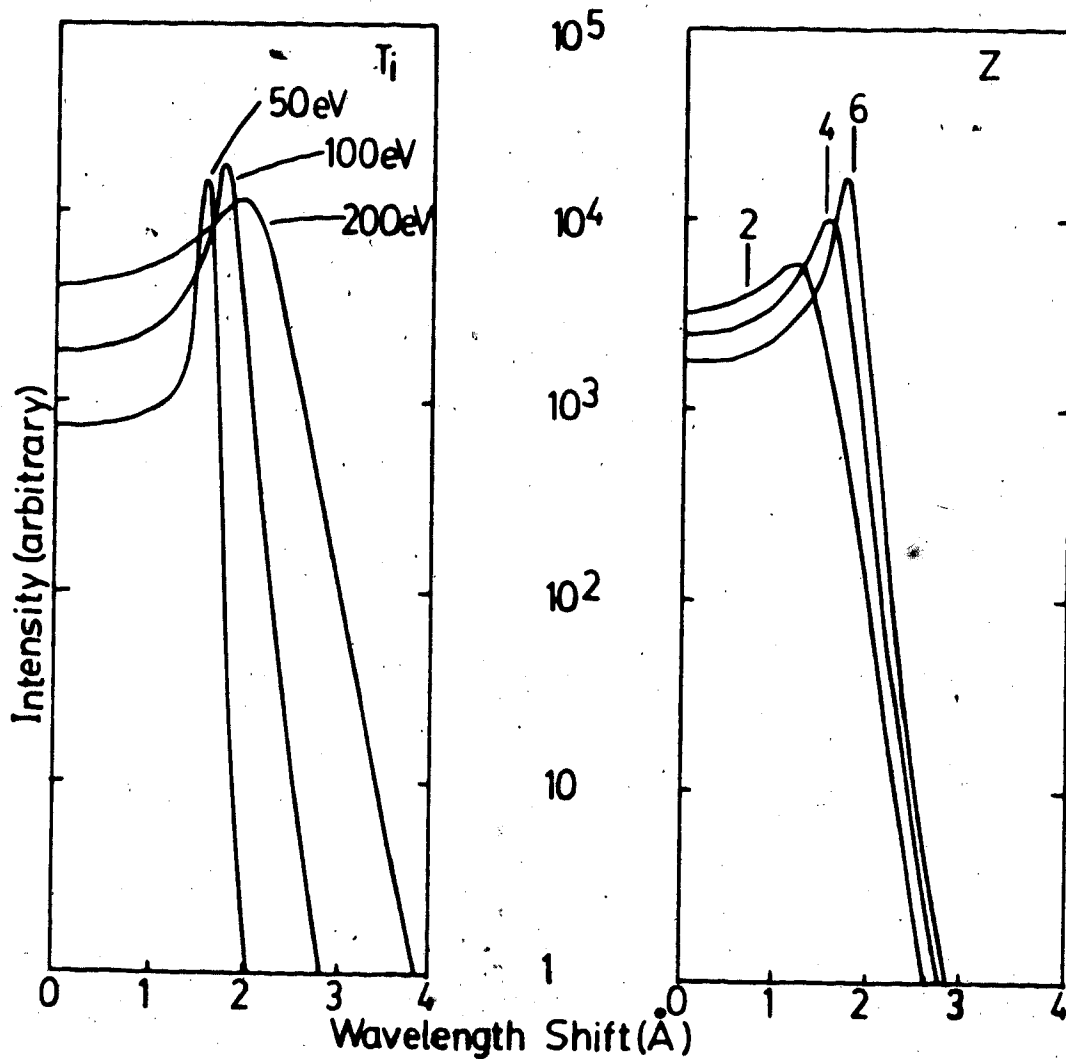


Fig. 4.10 Variation of the ion component (of the ruby Thomson scattering spectrum) with ion temperature and degree of ionization for a thermal oxygen plasma with $T_e = 100$ eV, $n = 0.8 n_c$ and $\theta = 60^\circ$.

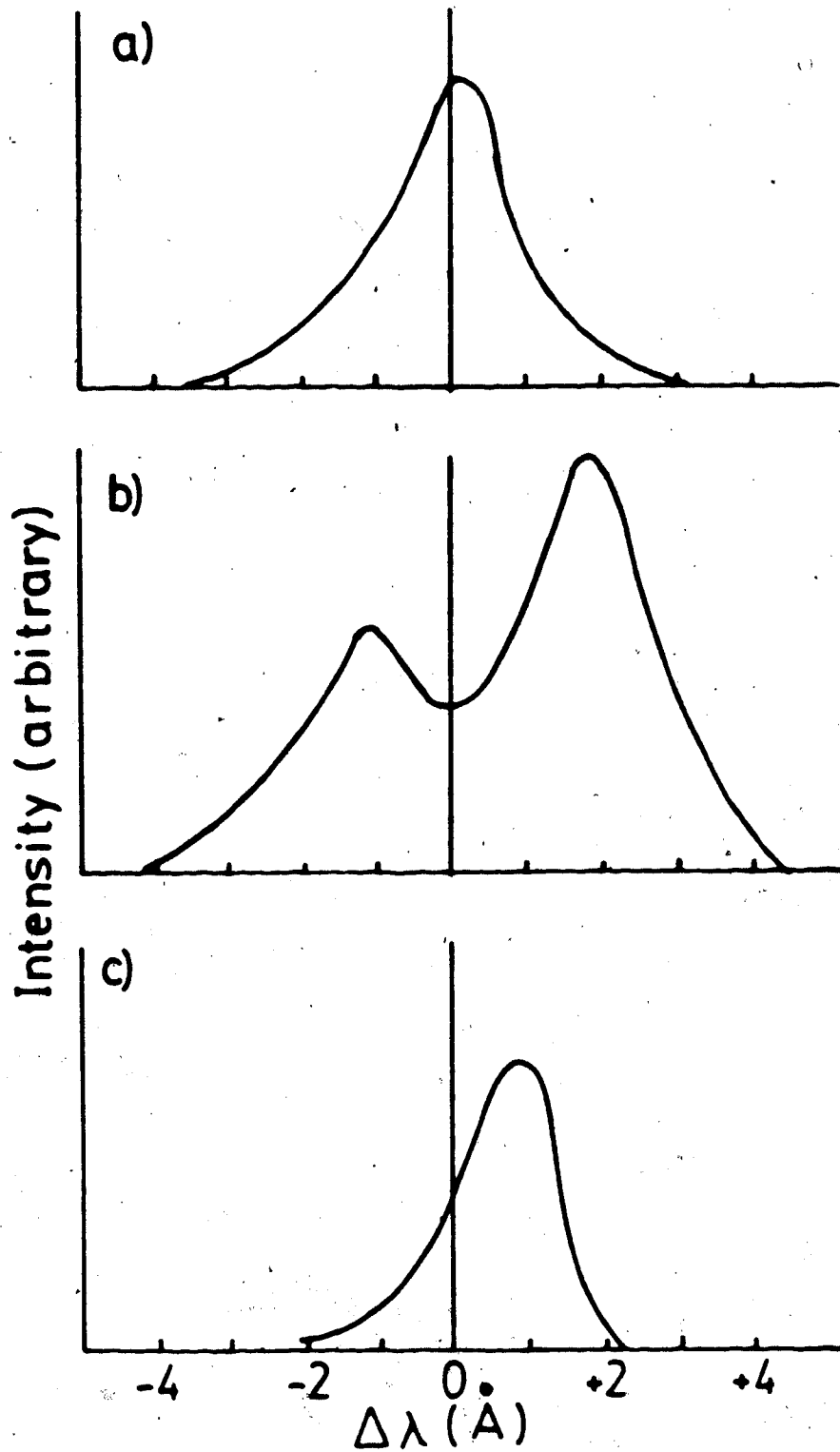


Fig. 4.11 Spectra of scattered ruby laser light for $\theta = 60^\circ$ showing a) and b) unshifted typical spectra (for scattering in the X-Y plane), c) shifted spectrum for scattering in the Z-Y plane.

for $\theta = 60^\circ$. The upper two spectra (Case a) show a symmetric behaviour with respect to λ_0 , i.e., no discernable Doppler shift due to a superimposed plasma drift. This is most likely expected for ion waves (\bar{k}_1, ω) propagating perpendicular to the CO_2 laser beam. Moreover, the lower spectrum (Case b) shows a distinct red shift for the case of \bar{k}_2 lying in the plane containing the incident CO_2 and the ruby laser beams (Fig. 4.2a). The magnitude and direction of this shift $\Delta\lambda/\lambda_0 = -\Delta\omega/\omega_0 = -(\bar{k}_2, \bar{u})/\omega_0$ implies a motion \bar{u} in the direction of propagation of the incident CO_2 laser beam of magnitude $u \approx 5 \times 10^6$ cm/sec. It should be pointed out that Fig. 4.11c shows an extreme case for the maximum shift observed. Results such as those seen in Fig. 4.11a and 4.11b were observed frequently for Case b as well as Case a. The detailed behaviour depended on the statistical variation of the plasma conditions and on the timing between the CO_2 and ruby laser beams.

If we assume that the peak of the enhanced ion spectrum is unchanged from that of a thermal plasma, then we may calculate T_i from the following equation [7],

$$T_i = \left\{ 77.63 \left(\frac{\Delta\lambda}{\sin \theta/2} \right)^2 - zT_e \right\} / 3 . \quad (4.6)$$

In this case, the peak of the scattered spectrum for Fig. 4.11b at $\Delta\lambda \approx 1.7 \text{ \AA}$ corresponds to an ion temperature $T_i \approx 100 \text{ eV}$ (for $T_e = 100 \text{ eV}$, $\theta = 60^\circ$ and $z = 6$). Similar results were obtained for other angles, e.g., $\theta = 22^\circ$ and are shown in Fig. 4.12 (in which $\Delta\lambda \approx 0.7 \text{ \AA}$). The average temperature obtained from different spectra was $T_i = 100 \pm 50 \text{ eV}$. This temperature, as indicated before, has been inferred for ions that are a small percentage of the total ions; i.e., those which

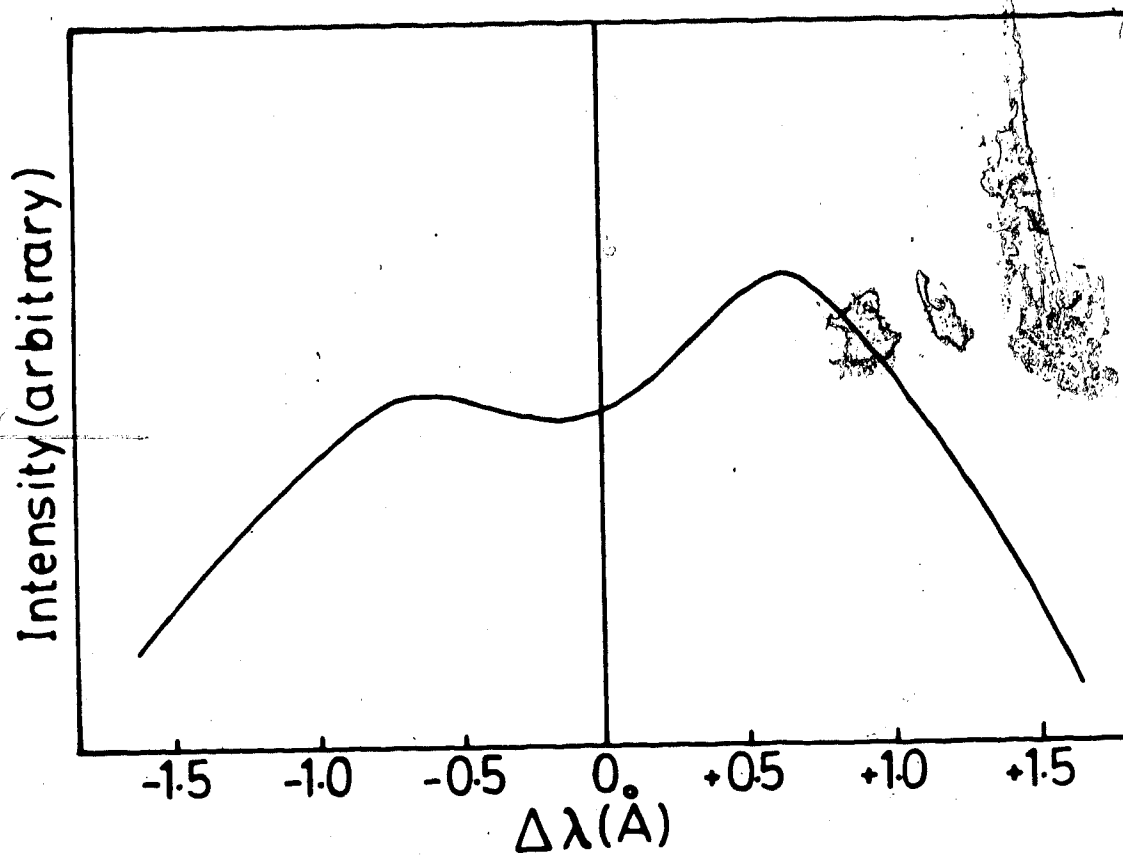


Fig. 4.12 Spectrum of the scattered ruby laser light for $\theta = 22^\circ$ in the X-Y plane.

cause the enhanced scattering. It should be noted, however, that it is reasonable to assume that the background (bulk) ions have ~ 100 eV temperature as well for two reasons: 1) Even without taking into consideration the heating effect due to parametric instabilities, hydrodynamic calculations of the laser heated gas target experiment give an average ion temperature of not less than 50% of electron temperature [3] ($T_i \geq 50$ eV in this experiment) and 2) The time required for the ions (nonthermal) to relax to a Maxwellian distribution of thermal temperature $T_i = 100$ eV using the formula given in [2] is ~ 0.4 ns which is very short compared to the total enhanced fluctuation time observed, at least in this part of the experiment (Case I).

The other information which can be deduced from the shift in wavelength of the ion feature is the ion wave frequency. Simple calculations from Figs. 4.11b and 4.12 give $\omega = 5.6 \times 10^{11}$ and $2.4 \times 10^{11} \text{ sec}^{-1}$ for $\theta = 60^\circ$ and 22° respectively. These same calculations, when made using the ion wave dispersion relation

$$\omega = \frac{\omega_{pi} k \lambda_D}{(1 + k^2 \lambda_D^2)^{1/2}}, \quad (4.7)$$

give $\omega = 5.3 \times 10^{11}$ and $2.1 \times 10^{11} \text{ sec}^{-1}$ (where $\omega_{pi} \sim 2.3 \times 10^{12} \text{ sec}^{-1}$ for $n = 0.8 n_0$). These values agree well with the experimental calculations.

In addition, from the spectral width ($\Delta\lambda = 1.5 \text{ \AA}$) of Figs. 4.11a and 4.11c one can estimate an ion wave damping rate of $\sim 3 \times 10^{11} \text{ sec}^{-1}$ which implies a mode lifetime of ~ 3 ps. Numerical simulations of current driven ion turbulence [10] show that enhanced ion fluctuations rapidly build up and decay, accompanied by electron heating, ion tail formation, and Landau damping of the instability.

Thus, if quasilinear effects (rather than nonlinear effects) are important in stabilizing the ion instability (regardless of how the instability is produced) in this experiment, many periods of growth and decay of ion fluctuations are occurring, as is indeed inferred from the short mode lifetime.

4.2.5 Ion fluctuation level.

The study of ion fluctuations produced by a laser beam is important because such fluctuations can affect energy absorption (discussed in Chapter 3) and transport processes [11,12] in the plasma. The presence of large amplitude ion fluctuations near critical density has often been invoked as a source of anomalous resistivity in order to explain the strongly reduced thermal conductivity seen in laser heated plasma experiments [13]. Therefore, many experiments have been made to measure the ion fluctuation levels. The first Thomson scattering measurements of ion turbulence in a high intensity laser heated hydrogen plasma target were reported by Offenberger *et al.* [14] where enhancement of $\sim 10^4$ over the thermal level was observed for $k\lambda_D \approx 0.5$. Similar measurements have been carried out on oxygen [3] and carbon [15] plasmas in the presence of a critical density layer. In both experiments substantial enhancement over the thermal level was observed.

In the present experiment the main goal was to study the effect of ion turbulence on absorption of laser radiation in underdense plasma ($n < n_c$). Because it has already been established that the observed enhanced scattering is due to enhanced ion fluctuations, the turbulence level has to be determined in order to estimate the amount of absorption that enhanced ion fluctuations can contribute.

The density fluctuation level was determined from the absolute measurements of $S(k)$ which is related to the scattered power, P_s , through Eqn. (3-25). The absolute value of P_s was obtained by calibrating the photomultiplier detection system with a tungsten filament blackbody source positioned at the point where the CO_2 laser beam is focused. Light from the filament was collected through the same optics used for collecting the scattered light from the plasma. The blackbody light power P reaching the photomultiplier is given by [16]

$$P = \frac{2hc^2}{\lambda^5} \frac{\epsilon A}{\exp(hc/kT\lambda) - 1} d\lambda d\Omega, \quad (4.8)$$

where A is the emitting area, ϵ is the emissivity of the filament (≈ 0.43 for tungsten at 6943 \AA and $\sim 1700^\circ \text{ K}$), $d\Omega$ is the collection solid angle and $d\lambda$ is the detection bandwidth (defined by the bandwidth of the interference filter in front of the photomultiplier). The absolute value of both P_s and $S(k)$ were determined by comparing the scattered signal with the signal generated by the tungsten filament. The error associated with $S(k)$, due to the uncertainty in determining parameters relating P_s to $S(k)$, as discussed in Chapter 3, was $\sim 26\%$ (uncertainty in I_0 , n , ν and $\Delta\Omega$ were $\sim 5\%$, 15% , 5% and 1% respectively).

The scattering form factor $S(k)$ calculated for a measured CO_2 laser intensity $I \sim 5 \times 10^{12} \text{ W/cm}^2$ and assumed $T_e = 100 \text{ eV}$ and $n = 0.8 n_e$ (taking the density at the focal point as the average density), is shown in Fig. 4.13 as a function of k for $\theta = 22^\circ$, 60° and 90° . The scattering at $\theta = 60^\circ$ shows almost equal $S(k)$ values for both the X-Y and the Z-Y planes (Case a and Case b). The $S(k)$ value for $\theta = 60^\circ$ is $\sim 6.8 \times 10^2$ which is approximately 10^3 times the thermal scattering form factor $S_{Th}(k)$. The $S(k)$ measurements for $\theta = 22^\circ$ and $\theta = 90^\circ$ were made in the Z-Y and X-Y planes respectively.

for convenience. It should be pointed out that the observed $S(k)$ spectrum is not a Kadomtsev like spectrum (shown dashed in Fig. 4.13 as normalized to $S(k)$ at $\theta = 60^\circ$) which is theoretically predicted for current driven ion turbulence [17].

The corresponding average spatial ion fluctuation level was calculated from a numerical integration of Eqn. (3.25) over k space using a second degree polynomial of the form

$$\ln S_i(k\lambda_D) = 10.89 - 22.14(k\lambda_D) + 15.9(k\lambda_D)^2$$

to represent the $S_i(k)$ spectrum given in Fig. 4.13. The result of this integration gave a value of $(\frac{\delta n}{n}) \approx 0.036$. Although this fluctuation level is lower than the one reported previously in a similar experiment [3] it is still very high. This may be due to the changes in focused laser conditions and plasma parameters such as T_e , n (lower in this experiment), L and/or the absence of the critical density layer. If the reduction in $(\delta n/n)$ is due to a change in plasma density then this would indicate that the instability (responsible for generating the ion turbulence) favours high plasma densities. This is expected for the off-resonance OTS and PD instabilities discussed in Chapter 3.

To calculate the effective collision frequency ν_{eff} required to determine the enhanced absorption, we assume a broad isotropic turbulence spectrum (as indicated experimentally). The calculation then

gives $\sum_{ki} \left(\frac{\delta n_{ki}}{n_c} \right)^2 \approx 0.05$ where from Eqn. (2.19) we get

$$\nu_{eff} \approx \frac{\omega_0}{4} \left\langle \text{Im} \frac{1}{\epsilon(k, \omega)} \right\rangle \sum_{ki} \left(\frac{\delta n_{ki}}{n_c} \right)^2, \quad (4.9)$$

gives $\nu_{eff} \approx 4.7 \times 10^{12} \text{ sec}^{-1}$, using Fig. 2.16 for the value of

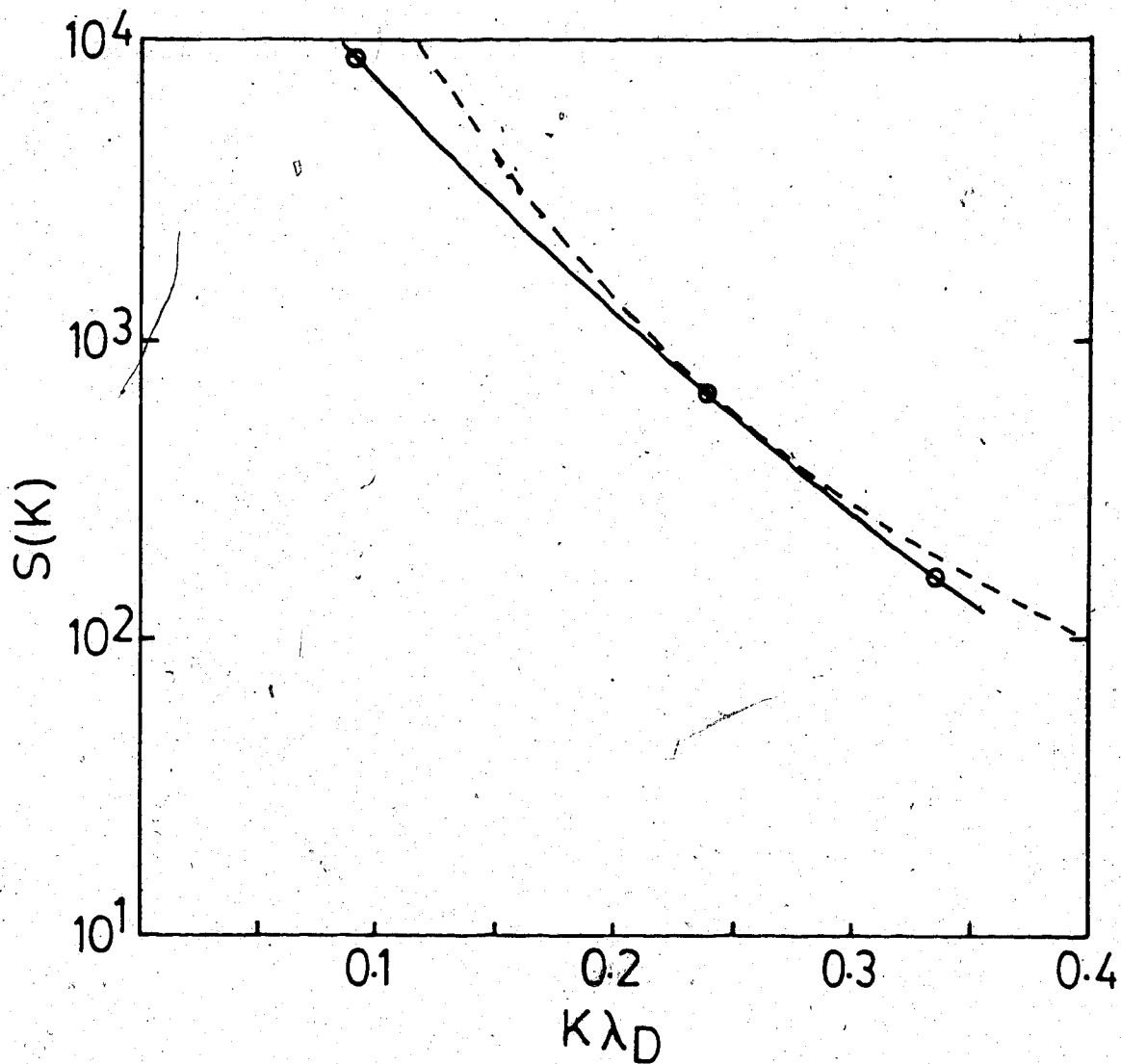


Fig. 4.13 Ion turbulence spectrum $S(k)$ as a function of $k\lambda_D$ assuming an average density $n = 0.8 n_e$ and $T_e = 100$ eV (Case I). The Kadomtsev type spectrum is also shown (dashed).

$\langle \text{Im} \frac{1}{\epsilon(k, \omega)} \rangle$. In comparison, this frequency is approximately 3.6 times the classical collision frequency and it is more than enough to account for the remaining 30% absorption factor.

In summary, the ruby laser Thomson scattering measurements which probed directly the ion fluctuations for $k\lambda_D = 0.091, 0.238$ and 0.336 , corresponding to scattering angles $\theta = 22^\circ, 60^\circ$ and 90° respectively (assuming $n = 0.8 n_c$), confirmed three important turbulence related points; (a) spectrally resolved measurements showed the enhanced ion feature, (b) temporally resolved measurements showed enhanced fluctuations for $t \lesssim 10$ ns, consistent with the period of strong absorption, and (c) measurements of the scattering form factor $S(k)$ showed that the enhancement in ion density fluctuations could account for the anomalous absorption observed. In addition, it was shown that the turbulence levels for $k\lambda_D \approx 0.238$ ($\theta=60^\circ$) in and out of the plane of the target are almost equal, indicating an isotropic turbulence.

Discussion of the source of the ion turbulence is delayed to the next section where further experimental information will be provided.

4.3 Results of Case II

4.3.1 Introductory remarks.

The results presented here belong to the second regime of the experiment in which the plasma density is lower and the density scale length ($\sim 450 \pm 50 \mu\text{m}$ at the focal point) is larger than that observed in Case I. Spectral analysis of the ruby laser Thomson scattering measurements showed unequivocal enhanced ion features. Some of the characteristics of the plasma in Case I were also seen here,

particularly those shown in Fig. 4.1. The fraction of light which should be absorbed, as given by Eqn. (4.1), is $\sim 85\%$ assuming the density profile of Fig. 4.7. Thus, ion turbulence need only to contribute 15% of the total to account for full laser absorption in this case.

More importantly, ion turbulence generated in laser plasma experiments has not been systematically investigated. It was, therefore, the aim of this experiment to study comprehensively the characteristics of the ion fluctuations, particularly those propagating in the direction of the electric field of the CO_2 laser radiation (in the X-Y plane). These fluctuations are potentially important for laser light absorption by plasmas ($\cos \theta = 1$ in Eqn. (2.17)). Section 4.3.2 will present the results of the $S(k)$ measurements. Section 4.3.3 will discuss the results of the streak camera measurements. Finally, the possible mechanism or mechanisms for generating ion turbulence in this experiment will be discussed in Section 4.3.4.

It should be noticed that the average plasma density throughout this part of the experiment will be taken as being equal to the density at the point where the CO_2 beam is focused, which prevails for $t \approx 2 - 8$ ns after the start of the SBS (i.e., $n = 0.7 n_c$). The electron temperature and the ionization state will be taken as 100 eV and 6 respectively.

4.3.2 $S(k)$ measurements.

The principal experimental result in this section is the determination of the $S(k)$ spectrum. The time integrated $S(k)$ was measured by a photomultiplier for six scattering angles ($\theta = 10^\circ, 16^\circ, 30^\circ, 45^\circ, 60^\circ$ and 90°) in the X-Y plane (Fig. 4.2). Thus, the wave number of the scanned ion fluctuations was in the range $1.6 \times 10^4 \text{ cm}^{-1} \leq k \leq 1.3 \times$

10^5 cm^{-1} or $0.045 \leq k\lambda_D \leq 0.36$ ($2.8 \leq \alpha \leq 22.2$) assuming $n = 0.7 n_c$.

Figure 4.14 shows the experimentally derived $S(k)$ as a function of $k\lambda_D$ for ion fluctuations in the X-Y plane. As will be seen in the following section, there is considerable modulation in the str of the scattered light; consequently, peak fluctuation levels can be substantially higher than the average values shown. In the region $k\lambda_D < 0.2$, $S(k)$ is strongly dependent on k (almost exponentially); however, for larger values of $k\lambda_D$ it varies more slowly. Over the experimental range of $k\lambda_D$, the time-integrated turbulence level varies from as little as 2 to $\approx 10^4$ times the thermal fluctuation level. It should be pointed out that, in general, this level is almost an order of magnitude below the previously reported value (Case I), probably due to a number of changes in the operating conditions of the CO_2 laser plasma interaction, particularly, focused laser conditions and the resultant change in the plasma density. This again illustrates the plasma density dependence of the ion wave instability generating the ion turbulence. It is apparent that no cutoff in the small $k\lambda_D$ region - such as that predicted for current driven ion instability and seen experimentally [18] - is found (at least for $0.045 \leq k\lambda_D < 0.36$).

We were unable to measure $S(k)$ in the X-Y plane for $\theta > 90^\circ$ because of geometrical limitations that resulted in overlapping of the focusing and collecting optics. However, no enhanced scattering over that expected from thermal fluctuations was seen for $\theta = 120^\circ$ ($k\lambda_D \approx 0.44$) in the Y-Z plane. Furthermore, no change in the turbulence level was observed as we changed the direction of observation from the X-Y plane to the Y-Z plane for $\theta = 45^\circ$ and 60° . This suggests an approximately isotropic spectrum. Moreover, we probed the $S(k)$ of the

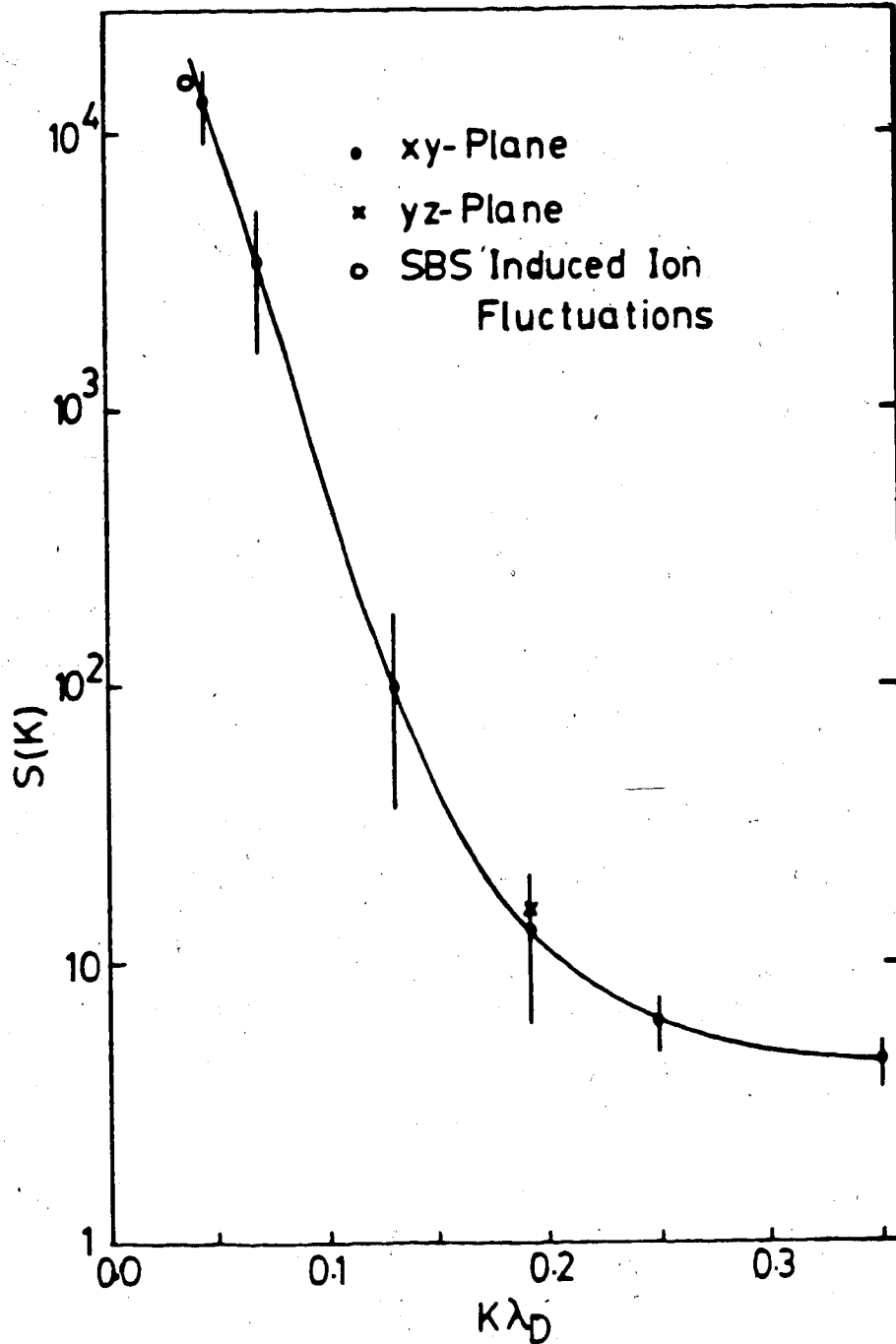


Fig. 4.14 Ion turbulence spectrum $S(k)$ as a function of $k\lambda_D$ assuming an average density $n = 0.7 n_c$ at $T_e = 100$ eV (Case II). The error bars indicate the standard deviations for thirty shots at average focused CO_2 laser intensity of $5 \times 10^{12} \text{ W/cm}^2$.

ion fluctuations induced by stimulated Brillouin backscattering (Fig. 4.15) and the result, shown as a circle in Fig. 4.14, fits well on the curve. It is significant that, although the SBS induced ion fluctuations (along the Z axis) and the ion turbulence in the X-Y plane are undoubtedly generated by different instability mechanisms, they are of comparable levels.

In order to estimate the absorption fraction due to ion turbulence in this experiment the $S(k)$ spectrum of Fig. 4.14 was approximated by a polynomial of the form

$$\ln S(k\lambda_D) \approx 12.25 - 69.03(k\lambda_D) + 1.0789(k\lambda_D)^2 \quad (4.10)$$

Integrating Eqn. (3.25) numerically (using Eqn. (4.10)) leads to a

fluctuation level $(\delta n/n) \sim 0.008$ which, in turn, gives $\sum_{ki} \left(\frac{\delta n_{ki}}{n_{cl}} \right)^2 \approx 0.013$.

and $v_{eff} \approx 0.9 v_{cl}$. This value is 1/4 of the v_{eff} calculated in Case I. The lower value may be related to the longer density scale-length and greater classical absorption prior to focus. However, it is still adequate to account for the remaining 15% absorption. It should be pointed out that the density fluctuation levels, observed in Cases I and II, are much lower than the maximum level predicted by the ion trapping saturation mechanism discussed in Chapter 2.

4.3.3 Streak measurements.

In order to follow the rapid temporal variations of the enhanced ion fluctuations, a streak camera was employed to record the Thomson scattered ruby laser light. While the timing between SBS and the Thomson scattering signal was the same as in Case I (Fig. 4.1), the duration of the Thomson scattering signal for this interaction regime (Case II), when recorded by a photomultiplier, was found to be shorter than that

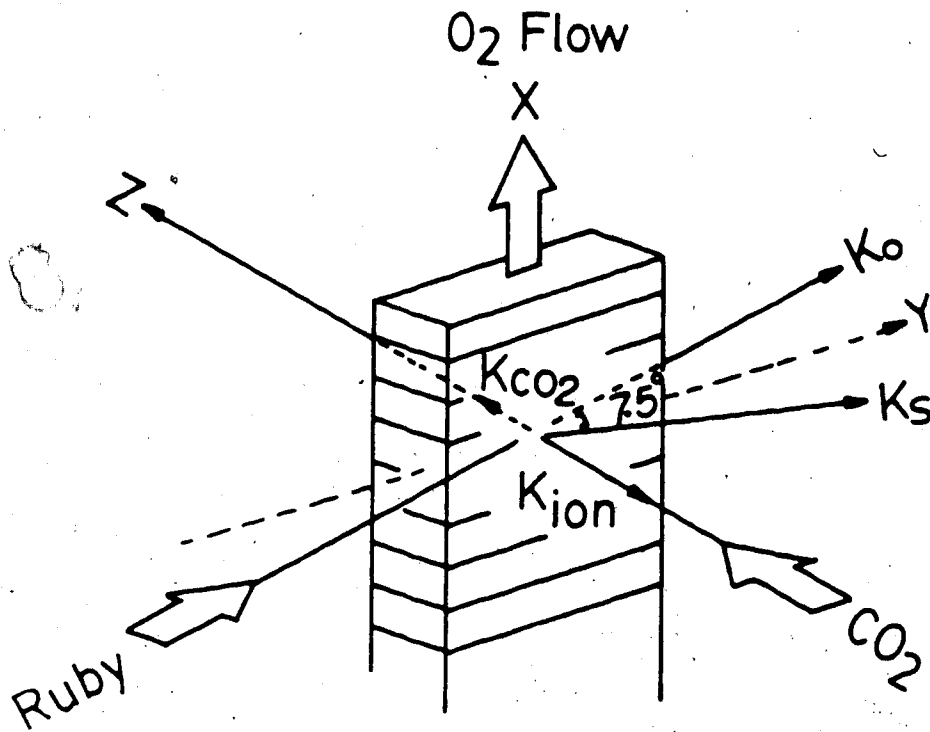


Fig. 4.15 Schematic diagram of the Thomson scattering experiment to probe ion fluctuations induced by stimulated Brillouin backscattering.

observed in Case I (~ 6 ns FWHM). But, since this was approaching the limit of the detector resolution, it is possible that the pulse duration could be much shorter than the observed one.

This was confirmed when a high speed streak camera was used. The results are summarized in Fig. 4.16 for varying observations with three different streak speeds. In general, the duration of the ion turbulence varied from approximately 3 ns (Fig. 4.16a) to as short as 250 ps (Fig. 4.16b). The risetime also varied from ~ 1 ns to 50 ps. Highly modulated pulse structures (~ 50 ps duration) were observed in 50% of the shots using the very high speed mode (Fig. 4.16c). This fine structure in the scattered light could not be fully accounted for by the fine structure of the incident ruby laser (monitored simultaneously with the scattered light and shown at the top of Figs. 4.16), nor to the modulations of the incident CO_2 laser beam which has a structure of ≥ 3 ns duration. Similar behaviour was observed in the scattered light of the ion fluctuations driven by SBS. In addition, changing the scattering direction from the X-Y plane to the Y-Z plane showed no systematic changes in the temporal behaviour of the scattered light. Indeed, the observed temporal changes were merely statistical and not related to the probing of CO_2 lasers. If the parametric instabilities discussed in Chapter 2 were responsible for generating the ion turbulence, then the observed variation in risetime could be explained by the wide range of the growth rate ($\gamma > 1$ ps) predicted for these instabilities when the density values accessible and the variation in v_0/v_e are taken into account. The ion trapping mechanism also predicts that saturation takes place in as short as 15 ps for $n = 0.7 n_c$ [19].

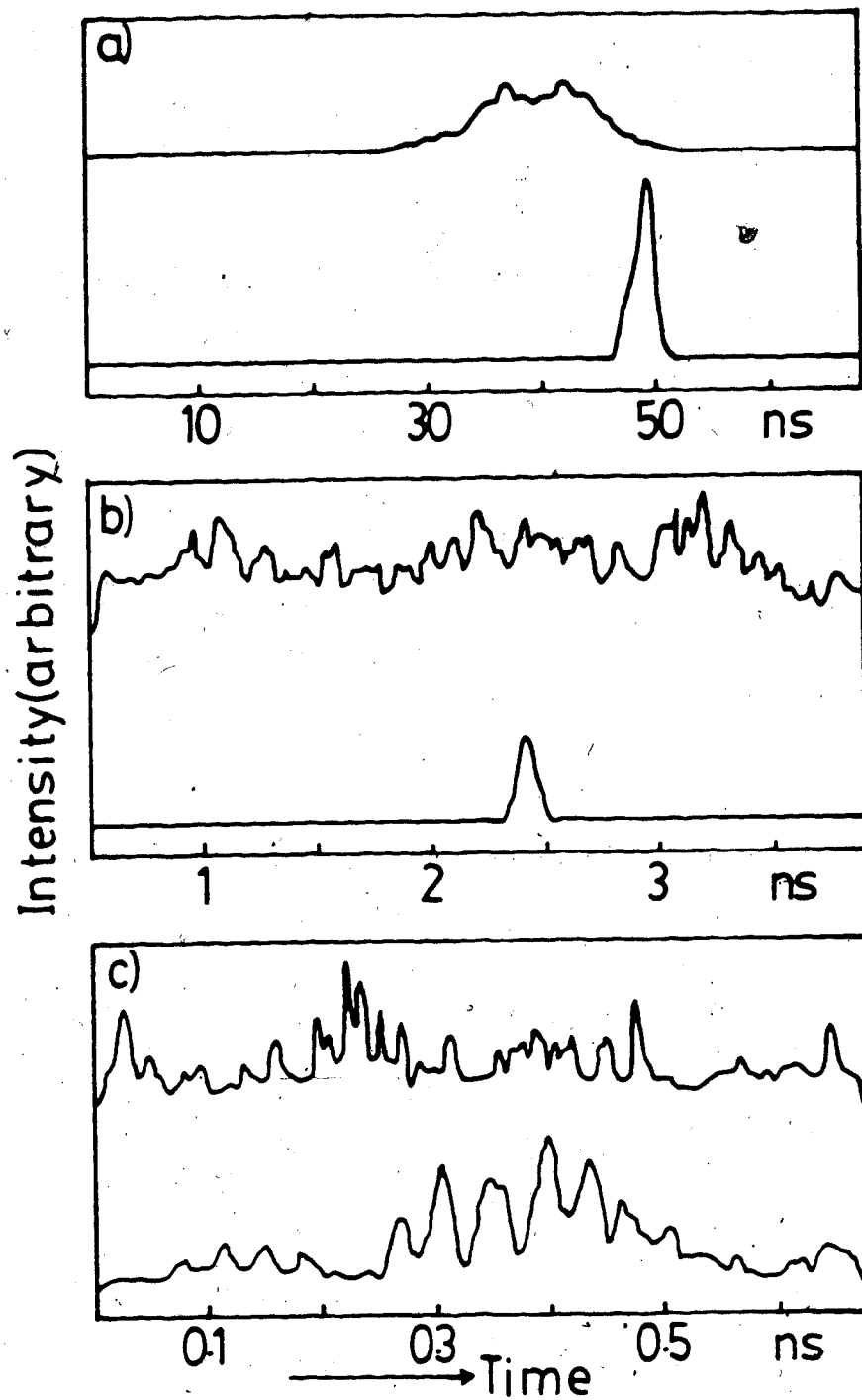


Fig. 4.16 Temporal behaviour of the Thomson scattered light (bottom) and the incident ruby light (top) in each plate showing different features; a) long lived fluctuations (~ 3 ns), b) short lived fluctuations (250 ps) with no structure, c) ~ 0.7 ns long fluctuations with fine structure.

The nanosecond duration of the enhanced fluctuations is significantly shorter than the value reported in Case I (FWHM $\lesssim 10$ ns). The very real difference is not attributable to the difference between the temporal resolution of the camera and the photomultiplier. For the photomultiplier (risetime ≈ 3 ns), the true duration Δ_t , the measured one, Δ_m , and the instrumental width Δ_s ($= 3$ ns) are governed by the relation $\Delta_t^2 = \Delta_m^2 - \Delta_s^2$. This gives, for $\Delta_m = 8 - 10$ ns, a true duration of at least 7.4 - 9.5 ns.

Thus, the difference between turbulence duration of Case I and Case II is obvious. This again is probably due to changes in the laser plasma interaction conditions (including n and L) which might affect the growth rate and damping of the ion wave instability responsible for the observed ion fluctuations.

The streak camera also proved to be a useful tool for measuring the size of the turbulence region and for following the temporal evolution of the ion fluctuation spatial distribution. This was realized by relaying the scattering region onto the 3 mm wide slit of the camera. The measurements were done with an effective spatial resolution of as high as 33 μm (depending on the magnification of the relay lens). Figure 4.17 shows a streak photograph of a 1.2 ns duration enhanced scattering event from ion turbulence. The observed turbulence region is only 170 μm in size. The variation in the spatial extent of the measured ion turbulence was found to lie between 100 to 250 μm . The 250 μm length plasma near the CO_2 beam focus corresponds to a density which varies, for early time near breakdown, between $0.43 n_c$ and $0.8 n_c$ (Fig. 4.7). This indicates that the instability responsible for generating ion turbulence takes

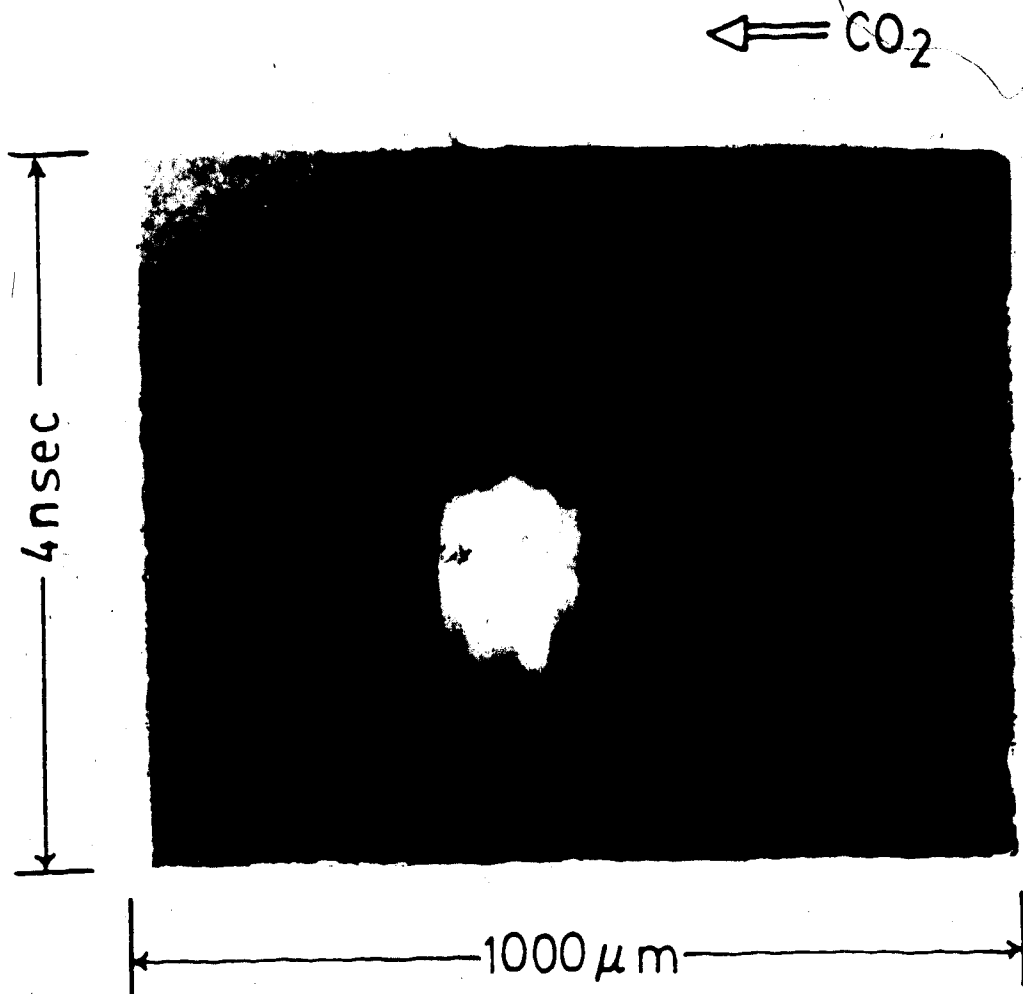


Fig. 4.17 Streak photograph of ~ 1.2 ns duration enhanced scattering event from ion turbulence occupying ~ 170 μm region.

place in relatively high density regions. This is consistent with the density region required to induce off-resonance OTS and PD ($n \geq 0.5 n_c$).

The time evolution of the spatial distribution of turbulence is shown (for $\theta = 16^\circ$ in the X-Y plane) for 3 ns duration (Fig. 4.18) and 0.8 ns duration (Fig. 4.19) cases. Both figures show a build-up of turbulence in two regions. This could be due to spatial inhomogeneity of the original incident CO₂ or ruby laser beams or due to filaments occurring in the plasma that make the CO₂ laser beam intensity distribution irregular. Such filaments have been observed previously in a CO₂ laser heated hydrogen plasma [20]. Figure 4.20 shows, for the same scattering angle, a rare case of turbulence in which the peak fluctuation region moves with a speed of $\sim 2 \times 10^7$ cm/sec in the direction of propagation of the CO₂ laser beam. This behaviour is similar to that observed spectrally in Case I although the speed here is 4 times higher. The measured speed is very high compared to the ion thermal speed ($\sim 2.4 \times 10^6$ cm/sec assuming $T_i = 100$ eV) and ion acoustic speed ($\sim 7.3 \times 10^5$ cm/sec for $T_e = T_i = 100$ eV and $z = 6$) and could be due to cases when higher intensity CO₂ radiation ionizes the gas ahead and then propagates towards the focus. This is characteristic of ionization (propagation) speeds observed in laser induced breakdown experiments [20].

4.3.4 Sources of ion turbulence.

Three mechanisms for generating ion turbulence in laser produced plasma experiments were suggested in Chapter 2. These mechanisms are heat flow driven ion acoustic instability, ion-ion streaming instability and ion waves induced by parametric instabilities. The first instability is expected to take place when $zT_e > T_i$ and $u > c_s$ where u is the

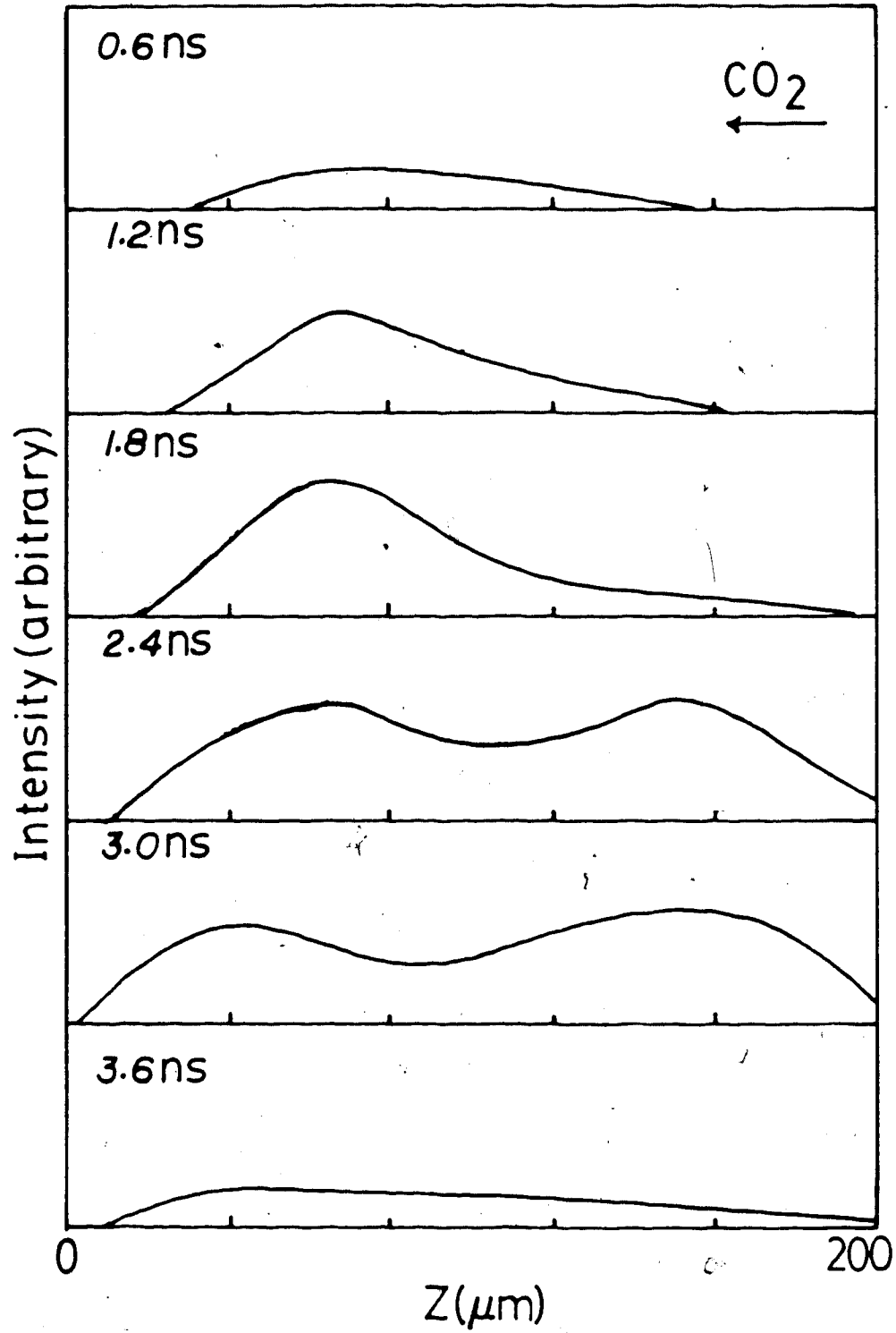


Fig. 4.18 Time evolution of the spatial distribution of turbulence for a 3 ns duration ($\theta = 16^\circ$ in the X-Y plane).

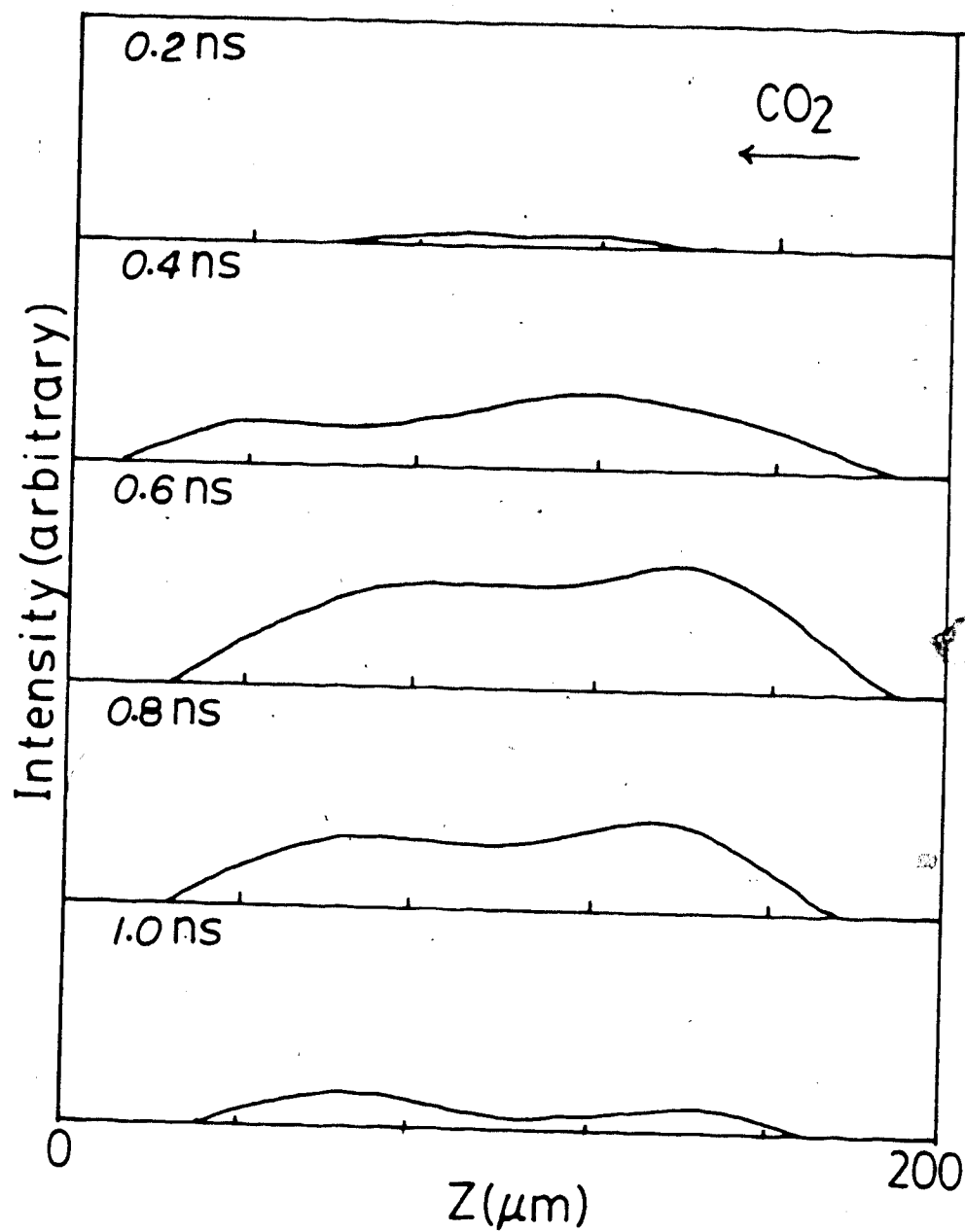


Fig. 4.19 Time evolution of the spatial distribution of turbulence for a 0.8 ns duration ($\theta = 16^\circ$ in the X-Y plane).

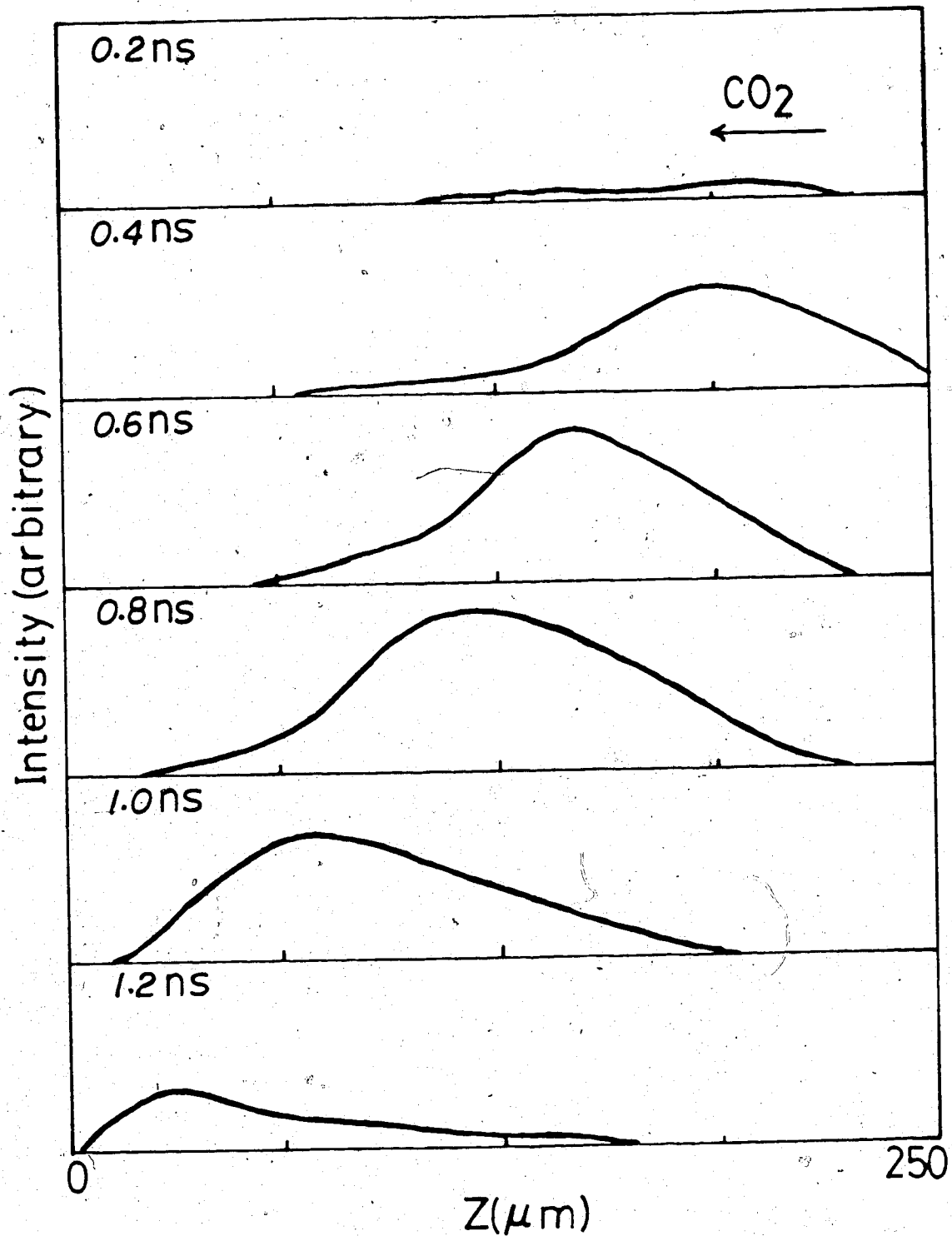


Fig. 4.20 Time evolution of the spatial distribution of turbulence showing a motion of the fluctuation region in the direction of the propagation of the CO_2 laser beam ($\theta = 16^\circ$ in the X-Y plane).

electron drift speed and c_s is the ion acoustic speed. In the current¹²² experiment the first condition was satisfied, while the second one was not always fulfilled. The spectral measurements of Case I, for example, occasionally showed a drift speed of $\leq 5 \times 10^6$ cm/sec which is close to the ion acoustic speed ($\sim 7.3 \times 10^6$ cm/sec for $T_i = T_e = 100$ eV and $z = 6$). On other occasions no drift was observed although enhanced scattering was substantial.

Now, even if the above conditions were satisfied, the current driven ion instability is not likely to be the source of the observed ion fluctuations for three reasons: (a) the observed $S(k)$ spectrum, particularly that shown in Fig. 4.14, is not a Kadomtsev type spectrum [17] nor is it similar to any of the modified spectra that the current driven instability suggests [21]; (b) no hot electrons were observed in the present experiment which could provide a return current source for a current driven instability; (c) the ion fluctuations driven by this instability are mainly distributed within a cone (of angle $\theta \sim 50 - 60^\circ$) around the direction of the return current (CO_2 laser beam) [22]; therefore the observed fluctuations in the X - Y plane cannot be explained by this instability.

The second possible candidate, the ion-ion streaming instability, suggested by Faehl and Kruer [23] only occurs when different ion species are available (different masses or z). For species of different masses only helium ions are potentially available in the background. However, in the oxygen target where the interactions take place, their number is insignificant. It may be possible to have oxygen ions with different z but again, near the focus of the CO_2 laser beam (≤ 250 μm diameter sphere) most of the oxygen atoms will be ionized to $z = 6$. Moreover,

the condition $v_{ii} < v_i/L$, required for this instability (Chapter 2), must be satisfied. If we assume in the best conditions that $T_e = T_i \approx 100$ eV, $L \approx 100$ μm and v_i is equal to the thermal ion speed, a value of $v_i/L \approx 2.4 \times 10^9 \text{ sec}^{-1}$ will be obtained. For the same plasma parameters, one can also calculate that

$$v_{ii} = v_{ei} \left(\frac{m_e}{m_i} \right)^{1/2} \approx 9 \times 10^9 \text{ sec}^{-1}.$$

Thus, the above instability condition is not satisfied and it will be even worse if $T_i < T_e$. In addition, it should be pointed out that high level ion turbulence was also observed in a pure hydrogen plasma target where species of different masses and z are negligible [14]. In conclusion, it is unlikely that the ion-ion streaming instability is a significant mechanism for generating the observed enhanced ion fluctuations in this experiment.

Before discussing the last suggested mechanism (ion waves produced by parametric instabilities) we should first mention that under the conditions of the present experiment, particularly with $v_0/v_e \leq 1.4$ (corresponding to CO_2 laser intensity $I \leq 5 \times 10^{12} \text{ W/cm}^2$ and $T_e = 100$ eV), many nonlinear processes are expected to take place. Indeed, among the processes that have been detected previously are stimulated Brillouin and Compton scattering, filamentation, two-plasmon decay, and harmonic generation [21].

Although much emphasis has been given by many investigators to ion waves produced by parametric instabilities of importance to scattering (stimulated Brillouin) or direct absorption (PD and OTS near critical density) processes, not much attention has been given to other instabilities such as off-resonance PD and OTS which may play

an important role in underdense plasmas. Our analysis in Chapter 2 showed clearly that PD and OTS off-resonance and filamentation can easily be excited for our experimental conditions (Figs. 2.2 - 2.11).

The correlations between experimental results and the theory of ion waves generated by parametric instabilities (discussed in Chapter 2) can be summarized as follows: (a) The off-resonance PD and OTS and filamentation have higher growth rates for higher densities that may explain the existence of ion turbulence in the high density region and the reduction in the turbulence level when moving from a high density regime to a low density one; (b) The wave number of the observed turbulence could be explained by PD and OTS off-resonance in the high $k\lambda_D$ range and by filamentation and mode coupling in the low $k\lambda_D$ range; (c) Both PD and OTS off-resonance and filamentation can produce ion waves in the direction of the electric field of the incident CO_2 laser beam (indeed, the growth rate γ is maximum in this direction) which are of interest for absorption; (d) Calculated values of γ for these instabilities agree qualitatively with the observed risetime range of the ion turbulence when the density values accessible and the variations in v_0/v_e , due to temporal modulation in CO_2 laser power, are taken into account; (e) The narrow spectral features of ion turbulence, observed in some shots, could also be explained by the purely growing modes of these instabilities (i.e., $\omega_R = 0$).

In addition, recent theoretical studies of PD and OTS off-resonance [25] have shown that $S(k)$ can be enhanced in the presence of a strong pump, even for stationary stable plasmas. The results are summarized in Fig. 4.21 for oxygen plasma with $z = 6$, $T_i = T_e = 100$ eV and $n = 0.6 n_c$ and $0.8 n_c$ for (a) and (b) respectively. The values of

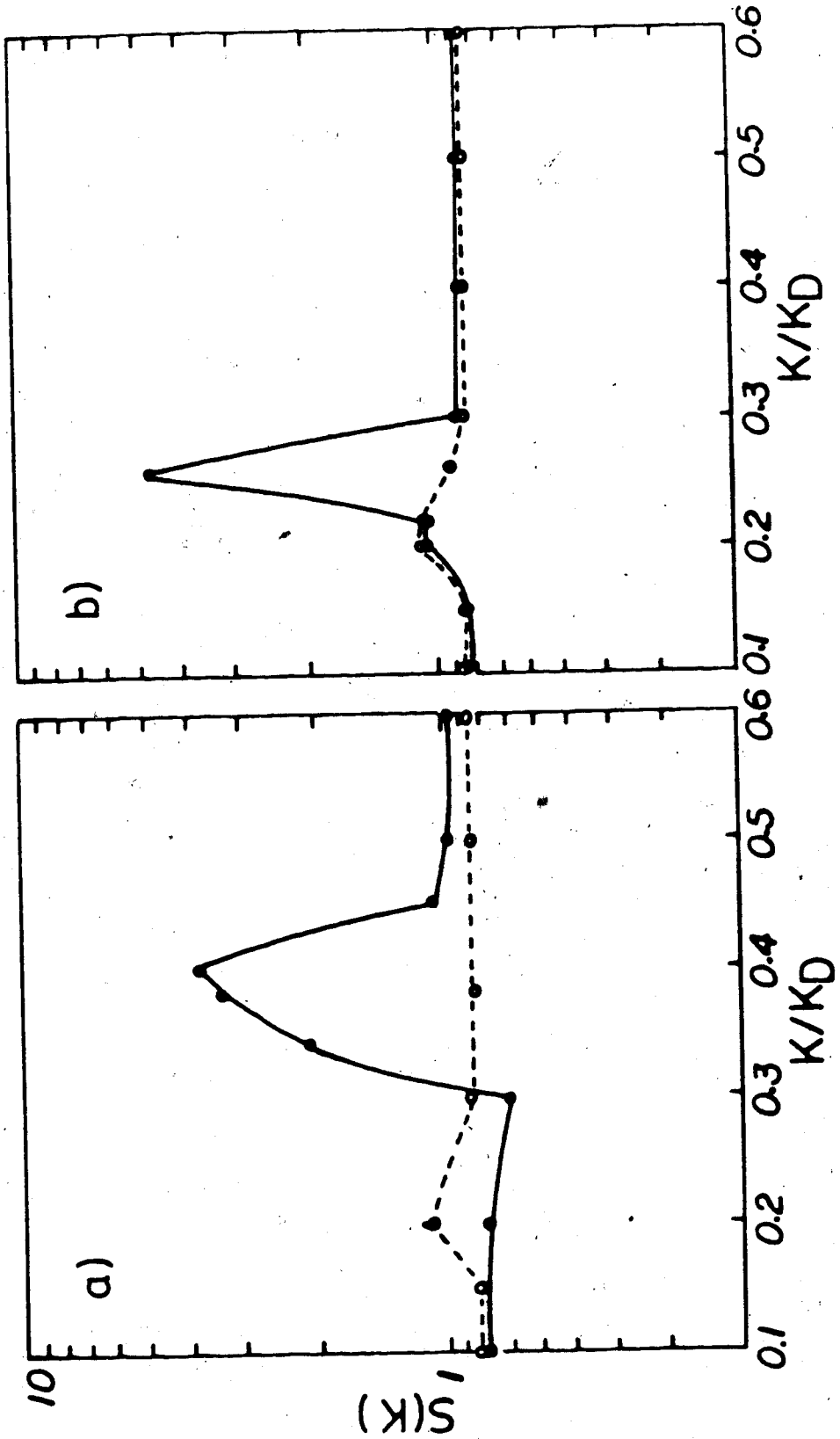


Fig. 4.21 Numerical values of $S(k)$ versus k/k_D for oxygen plasma with $z = 6$ and $T_i = T_e = 100$ eV. Dashed curves correspond to $v_0/v_e = 0$. a) $v_0/v_e = 0.6$, $n/n_c = 0.6$, b) $v_0/v_e = 0.05$, $n/n_c = 0.8$.

v_0/v_e were taken just below the threshold values of PD and OTS off-resonance. Considerable enhancement over thermal fluctuation levels (shown dashed) can be seen. Moreover, these calculations set a lower bound for enhancement. With higher values of v_0/v_e (above threshold) the enhancement is naturally expected to be substantially higher (ultimately limited by some nonlinear saturation processes).

Before coming to any final conclusions, three observational facts must be recalled: first, the shape of $S(k)$ observed in Case I and Case II; second, the isotropy of the ion turbulence; third, the absence of electron features in Thomson scattering when electron waves can be generated by PD and OTS off-resonance. Regarding the first point, although no theoretical knowledge is available concerning $S(k)$ for ion turbulence generated by off-resonance PD and OTS and filamentation above the instability threshold, if the observed $S(k)$ spectrum is related to the above instabilities, then it should be a function of density and pump strength. This may be the reason for observing different $S(k)$ spectra in Case I and Case II. In addition, because of the considerable variation in plasma density available in the scattering volume, the observed spectrum is an integration of the form factor over that density range.

With respect to the second point, the low $k\lambda_D$ part of the spectrum can be explained by filamentation and mode coupling (Brillouin) instabilities, which cover a wide angle \bar{E} between the electric field of the CO_2 laser beam and the \bar{k} vector of the ion waves (Fig. 2.2 - 2.5). For the other instabilities (off-resonance PD and OTS), although they are functions of θ , the reduction in the growth rate at $\theta = 60^\circ$, compared to $\theta = 0$, is $\sim 60-70\%$ (Fig. 2.13). This reduction is even lower if we take

into account the angle subtended by the CO_2 laser beam ($\sim 29^\circ$) when focused by an $f/2$ mirror. Finally, the electron waves generated by off-resonance instability are heavily damped. A simple calculation for $T_e = 100$ eV, $k\lambda_D = 0.3$ and $n = 0.7 n_0$, shows that Landau damping is $\sim 8 \times 10^{12} \text{ sec}^{-1}$. This is somewhat greater than the growth rate of the instabilities mentioned above. Therefore, the instability is a heavily damped driven mode.

In conclusion, it appears more likely that our observations may be related to strong pump induced ion instabilities generated by off-resonance PD and OTS, filamentation and mode coupling processes. However, in order to convincingly demonstrate this, considerable additional experimental and theoretical study is required. This will be discussed in the next chapter.

Importantly, the experimental work presented here is a beginning towards understanding complicated non-equilibrium phenomena which can be induced in plasma under the influence of a strong external electromagnetic wave.

CHAPTER 5

SUMMARY AND CONCLUSION

A successful attempt has been made to study ion turbulence in high intensity CO_2 laser produced plasmas both theoretically and experimentally. In the theoretical aspect of the study, emphasis has been given to ion waves generated by parametric instabilities in an underdense plasma ($n < n_c$). The Drake dispersion equation for $k\lambda_D \ll 1$ and the Silin dispersion equation for $k\lambda_D = 0(1)$ have been solved numerically for a homogeneous oxygen plasma in the presence of a high intensity CO_2 laser beam. The results have shown that ion waves can be easily induced by stimulated Brillouin scattering, filamentation and off-resonance electron-ion decay and oscillating two stream instabilities. The latter instabilities are believed to be possible candidates for generating the enhanced ion fluctuations associated with the anomalous absorption reported previously.

On the experimental side the Thomson scattering technique was applied to study ion turbulence in the CO_2 laser heated oxygen gas target plasma. Measurements were made for two density regimes, short scale length and long scale length. Both regimes showed strong enhancement of the ion fluctuations over thermal levels. The turbulence level along with temporal and spectral analysis of the ion fluctuations indicates that the measured ion turbulence could account for the anomalous absorption observed in this experiment.

In the long scale length regime the $S(k)$ spectrum was measured for ion turbulence in and out of the plane of the target. The approximately isotropic spectrum that was observed suggests that the enhancement in the plane of the target could be as strong as that induced by low threshold parametric instabilities such as SBS (for the same wave number). High speed streak camera measurements of the scattered light show short lived and fast rising ion fluctuations. These measurements also indicate that ion turbulence is generated in a high density region in the vicinity of the focal point of the laser beam. This is consistent with the density regime at which OTS and PD off-resonance can be excited.

The possible mechanisms for generating ion fluctuations were discussed in the light of the experimental results. It is believed that the observed ion fluctuations are likely related to strong pump induced parametric instabilities (particularly off-resonance PD and OTS instabilities and filamentation) rather than the current driven ion instability or ion-ion streaming instability.

Since the analyses performed in this thesis were carried out only for thermal homogeneous plasma, the effect of the inhomogeneity in a plasma on off-resonance PD and OTS instabilities, together with the different $S(k)$ spectra produced by these instabilities in a non-thermal plasma, are worth further analysis.

Experimental verification of the existence of both PD and OTS off-resonance instabilities is needed. This could be achieved by two different methods; first by varying the intensity of the incident laser beam in order to determine the thresholds of these instabilities, and second by the proper selection of a plasma density region, it might

be possible to simultaneously observe features of both electrons and ions in a Thomson scattering experiment.

The effect of ion fluctuations on energy transport in laser/plasma interactions is another interesting topic for investigation. A possible way of doing it is through the measurement of the particle energy distribution and determining the corresponding energy flux limit.

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APPENDIX A

SOLUTION OF THE DISPERSION EQUATION

The reduced form of the Drake dispersion function, Eqn. (2.5), and the general dispersion function of Sanmartin, Eqn. (2.6), were solved graphically by finding the values of $z = \alpha + j\beta$ which satisfy the conditions

$$\operatorname{Re} F(z) = 0 \quad (A.1)$$

and

$$\operatorname{Im} F(z) = 0 \quad (A.2)$$

separately, where $F(z)$ is the dispersion function. The intersection (or intersections) of the curves of z that satisfy A.1 and A.2, in the $\alpha\beta$ diagram, is the solution to the function $F(z) = 0$. Separation of the real and imaginary parts of $F(z)$ was not practical analytically without a proper approximation, and therefore the procedure was done numerically. Neither Newton's method nor its modifications were successful in finding the roots of Eqns. A.1 and A.2 because the roots are so close to each other and convergence could not be achieved. Instead, a simple technique called the bisection or half-interval method [1] was used. Suppose, as an example, that the roots of $S(z) = \operatorname{Re} F(z) = 0$ are required. We first keep one variable fixed, say β , and evaluate $S(\alpha)$ at equal intervals in a certain range of α . Fig. A.1. The bisection method gives a root if $S(\alpha_1, \beta_1)$ and $S(\alpha_2, \beta_1)$ are opposite in sign. Next we evaluate $S(\alpha)$ at α_3, β_1 where $\alpha_3 = (\alpha_1 + \alpha_2)/2$. $S(\alpha_3, \beta_1)$ will be either zero or have the sign of $S(\alpha_1, \beta_1)$ or the sign

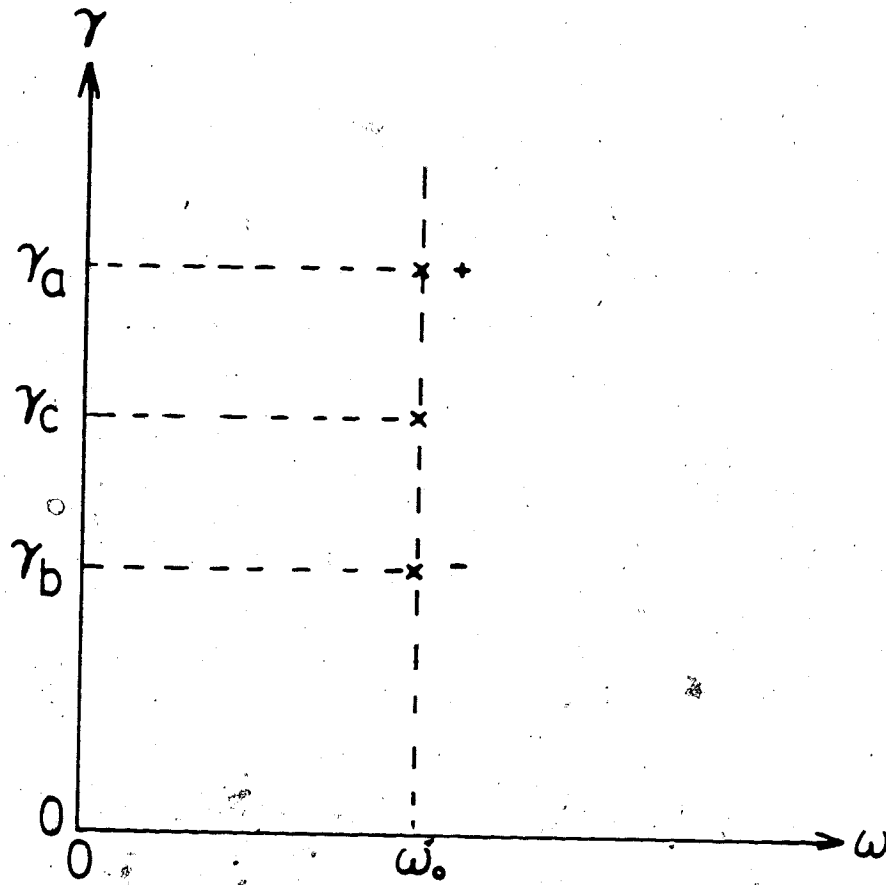


Fig. 1. Points found using the bisection method.

of $S(\omega_0, \gamma_3)$. If $S(\omega_0, \gamma_c)$ is not zero $S(z)$ will be evaluated at point (ω_0, γ_d) , where $\gamma_d = (\gamma_c + \gamma_a)/2$ or $= (\gamma_c + \gamma_b)/2$, whichever point, γ_a or γ_b has an opposite sign to γ_c . Continuing in this manner, there is always a point in the interval for which $F(z) = 0$. Ten to fifteen operations were found to be adequate enough to find fairly accurate roots in any pair of points that have opposite signs.

Next, we ~~will~~ look at other roots. The same procedure was applied to the ~~imaginary~~ part of the function $F(z)$.

Programs FILAM and DISPL were written in FORTRAN to solve Eqs. (2.5) and (2.6) respectively. It is more convenient for understanding the programs to write the variables of these equations in terms of parameters which are of interest to us. We define first the following normalized complex frequencies

$$s = \frac{\omega}{\omega_0} + j\frac{\gamma}{\gamma_0} \tag{A.3}$$

and

$$s_1 = \frac{\omega}{\omega_0} + j\frac{\gamma}{\gamma_0} = s \left(\frac{\omega_0}{\omega} + j\frac{\gamma_0}{\gamma} \right) \tag{A.4}$$

where $\omega_0 = \gamma_0^{-1}$ is the complex frequency which appeared in the general susceptibility of Eqn. 2.7. With Maxwellian distribution it is easy to show that χ_1 can be written in the form

$$\chi_1 = \frac{\chi_0}{1 - s_1^2} = \chi_0 \frac{1 - s_1^2}{1 - s_1^2} \tag{A.5}$$

$$\chi_1 = \frac{\chi_0}{1 - s_1^2} = \chi_0 \frac{1 - s_1^2}{1 - s_1^2} \tag{A.6}$$

where s_1 is the ionization degree,

$$\Omega = \frac{\omega_0}{\sqrt{2} k (T_e/m_e)^{1/2}} = \frac{1}{\sqrt{2}} \frac{\omega_0}{\omega_{pe}} \frac{k}{k_D}$$

and the function Z is called the plasma dispersion function and is defined by the integral

$$Z(z) = \pi^{-1/2} \int_{-\infty}^{\infty} dx \exp(-x^2) / (x-z) \quad (A.7)$$

The function $Z(z)$ was solved by series expansion in the subroutine ZPI following the method of [2]. The argument of the Bessel function in Eqn. (2.6), $k \frac{eE_0}{m_e \omega_0^2} \cos(\theta)$, which will be called α , can be written as follows:

$$\alpha = \frac{k}{\omega_0} \frac{eE_0}{m_e \omega_0^2} \cos(\theta) = \frac{v_0}{\omega_0} k \cos(\theta) = \left(\frac{v_0}{v_e} \right) \left(\frac{k}{k_D} \right) \left(\frac{k_D v_e}{\omega_0} \right) \cos(\theta) =$$

$$\left(\frac{v_0}{v_e} \right) \left(\frac{k}{k_D} \right) \left(\frac{k_D v_e}{\omega_0} \right) \cos(\theta) = \left(\frac{v_0}{v_e} \right) \left(\frac{k}{k_D} \right) \left(\frac{v_e}{v_0} \right) \cos(\theta)$$

Thus, the important parameters in Eqn. (2.6) are $\frac{v_0}{v_e}$, $\frac{k}{k_D}$, $\frac{v_e}{v_0}$, $\frac{v_e}{\omega_0}$, ω_0 , and θ .

Similarly the functions J_{\pm} of Eqn. (2.5) can be written in the form

$$J_{\pm} = \omega_0^2 \left(\frac{k}{k_D} \right)^2 \left(\frac{v_e}{v_0} \right)^2 = \omega_0^2 \left(\frac{k}{k_D} \right) \left(\frac{v_0}{v_e} \right) \left(\frac{v_e}{v_0} \right)^2 \cos^2 \theta =$$

$$\omega_0^2 \left(\frac{k}{k_D} \right) \left(\frac{v_0}{v_e} \right)^2 \cos^2 \theta = \omega_0^2 \left(\frac{k}{k_D} \right) \cos^2 \theta$$

with $\beta = 1 + \frac{v_0}{v_e} + \frac{v_e}{v_0}$ while β_1 and β_2 are given by Eqs. (A.5) and (A.6) with $\kappa = 1$.

The list of the computer programs and their subroutines are given below. The searched roots are given in the normalized form

$$z = x + iy \quad \text{where } x = \frac{\omega_R}{\sqrt{2} k(T_i/m_i)^{1/2}} \quad \text{and } y = \frac{Y}{\sqrt{2} k(T_i/m_i)^{1/2}} .$$

```

1 C .....
2 C * PROGRAM DISPI FOR SOLVING THE GENERAL DISPERSION EQUA--
3 C * TION OF SILIN & SANMARTIN. THE FIRST INPUT TO THIS
4 C * PROGRAM CONTAINS TE & TI (ELECT. & ION TEMP., MI (ION
5 C * MASS NUMBER) & ZI (IONIZATION DEGREE). THE SECOND
6 C * INPUT CONTAINS K (IN UNITS OF KD), N (IN UNITS OF NC)
7 C * VE (IN UNITS OF VO) & CTH (COS OF THE ANGLE BETWEEN E
8 C * & K. THE PROGRAM REQUIRES ONE SUBROUTINE (BESJ) & TWO
9 C * EXTERNAL FUNCTIONS (F & ZP1).
10 C .....
11 C COMPLEX Z,ZO,Z1,Z2,F,S(200)
12 C REAL K,N,MI
13 C DIMENSION Y(200),S1(200),Y1(25),Y2(25),Y3(25),Y4(25),S2(200)
14 C COMMON K,A,ZI,T,OM,BJ(20)
15 C READ(5,1)TE,TI,MI,ZI
16 C 1 FORMAT(4F6.2)
17 C T=TI/TE
18 C WRITE(3,20)TE,T,MI,ZI
19 C READ(5,2)K,N,V,CTH
20 C 2 FORMAT(4F6.3)
21 C WRITE(3,21)K,N,V,CTH
22 C OM=0.7071/(K*SQRT(N))
23 C ALPHA=K*V*CTH*SQRT(N)
24 C CALCULATION OF 20 TERMS OF BESSLE FUNCTION
25 C DO 3 I=1,20
26 C J=I-1
27 C CALL BESJ(ALPHA,J,H,IRE)
28 C BJ(I)=H*H
29 C 3 CONTINUE
30 C A=T/(1836.2*MI)
31 C A=SQRT(A)
32 C THIS INPUT DETERMINES THE RANGE OF THE NORMALIZED REAL (X)
33 C & THE IMAGINARY (Y) PARTS OF THE FREQUENCY. N1 & N2 ARE
34 C THE NUMBER OF STEPS IN X & Y. DX & DY ARE STEPS AND DXO &
35 C DYO ARE THE STARTING POINTS.
36 C READ(5,22)N1,N2,DX,DY,DXO,DYO
37 C K=K*K
38 C SEARCHING FOR POSSIBLE ROOTS OF THE REAL & IMAG. PARTS OF
39 C THE DISPERSION EQUATION.
40 C DO 16 I=1,N1
41 C X=DX*I+DXO
42 C WRITE(3,23)X
43 C DO 4 J=1,N2
44 C Y(J)=DY*J+DYO
45 C Z=CMPLX(X,Y(J))
46 C S(J)=F(Z)
47 C S1(J)=REAL(S(J))
48 C 4 S2(J)=AIMAG(S(J))
49 C IR=0
50 C II=0
51 C DO 6 J=2,N2
52 C F1=S1(J)/S1(J-1)
53 C IF(F1.GT.0.0)GO TO 5
54 C IR=IR+1
55 C Y1(IR)=Y(J-1)
56 C Y2(IR)=Y(J)
57 C 5 F2=S2(J)/S2(J-1)
58 C IF(F2.GT.0.0)GO TO 6
59 C II=II+1
60 C Y3(II)=Y(J-1)
61 C Y4(II)=Y(J)
62 C 6 CONTINUE
63 C USING BISECTION METHOD TO DETERMINE THE REAL ROOTS OF THE
64 C DISPERSION EQUATION
65 C IF(IR.EQ.0)GO TO 11
66 C DO 10 K1=1,IR
67 C Y11=Y1(K1)
68 C Y22=Y2(K1)
69 C Z1=CMPLX(X,Y11)
70 C Z2=CMPLX(X,Y22)
71 C F1=REAL(F(Z1))

```

```

72      F2=REAL(F(Z2))
73      DO 8 KO=1,20
74      YO=(Y22+Y11)/2.DO
75      ZO=CMLPX(X,YO)
76      FO=REAL(F(ZO))
77      FF=FO/F1
78      IF(FF.GT.O.O)GO TO 7
79      F2=FO
80      Y22=YO
81      GO TO 8
82      7   F1=FO
83      Y11=YO
84      8   CONTINUE
85      WRITE(3,9)YO,FO
86      9   FORMAT(10X,'Y='.E12.5,5X,'RE(D)='.E12.5)
87      10  CONTINUE
88      C   USING BISECTION METHOD TO DETERMINE THE IMAG ROOTS
89      11  IF(II.EQ.O.O)GO TO 16
90      DO 15 K1=1,II
91      Y33=Y3(K1)
92      Y44=Y4(K1)
93      Z1=CMLPX(X,Y33)
94      Z2=CMLPX(X,Y44)
95      F1=AIMAG(F(Z1))
96      F2=AIMAG(F(Z2))
97      DO 13 KO=1,20
98      YO=(Y44+Y33)/2.DO
99      ZO=CMLPX(X,YO)
100     FO=AIMAG(F(ZO))
101     FF=FO/F1
102     IF(FF.GT.O.O)GO TO 12
103     F2=FO
104     Y44=YO
105     GO TO 13
106     12  F1=FO
107     Y33=YO
108     13  CONTINUE
109     WRITE(3,14)YO,FO
110     14  FORMAT(10X,'Y='.E12.5,5X,'IM(D)='.E12.5)
111     15  CONTINUE
112     16  CONTINUE
113     20  FORMAT(5X,'TE='.F6.1,5X,'TI/TE='.F6.3,5X,'MI='.F5.2,5X,
114     'I/Z='.F5.2)
115     21  FORMAT(5X,'K/KD='.F5.3,5X,'N/NC='.F5.3,5X,'VO/VT='.
116     'F5.2,5X,'COS( )='.F6.2)
117     22  FORMAT(2I3,4F6.3)
118     23  FORMAT(15X,'***X='.F7.4,'****')
119     STOP
120     END
121     C   .....
122     C   * FUNCTION F IS THE GENERAL DISPERSION EQUATION THE *
123     C   * COMPLEX ARGUMENT Z IS DETERMINED FROM X & Y IN THE *
124     C   * MAIN PROGRAM. *
125     C   .....
126     COMPLEX FUNCTION F(Z)
127     COMPLEX Z,Z1,ZP1,XO,OO,G,O1,O2,OP,ON,G1
128     REAL K
129     COMMON K,A,ZI,T,OM,BJ(20)
130     Z1=A*Z
131     XO=ZI*(1.O+Z*ZP1(Z))/(K*T)
132     DO=1.O+(1.O+Z1*ZP1(Z1))/K
133     G=CMLPX(O.O,O.O)
134     DO 10 I=2,20
135     J=I-1
136     O1=J*OM+Z1
137     O2=-J*OM+Z1
138     OP=1.O+(1.O+O1*ZP1(O1))/K
139     ON=1.O+(1.O+O2*ZP1(O2))/K
140     G1=G+BJ(I)*(1.O/OP+1.O/ON)
141     IF(REAL(G1).EQ.O.O.AND.AIMAG(G1).EQ.O.O)GO TO 20
142     X1=(REAL(G1)-REAL(G))/REAL(G1)
143     1   X2=(AIMAG(G1)-AIMAG(G))/AIMAG(G1)

```



```

144      2  IF(ABS(X1) GT 0.001 OR ABS(X2) GT 0.001)GO TO 5
145      GO TO 20
146      5  G=G1
147      10 CONTINUE
148      20 F=1.0+X0*(BJ(1)/DO+G1)
149      RETURN
150      END
1      C
2      C
3      C
4      C
5      SUBROUTINE BESJ(X,N,BJ,IER)
6      IER=0
7      BJ=0.
8      IF(N)10,20,25
9      10  IER=1
10     RETURN
11     20  IF(X)30,21,31
12     21  BJ=1
13     RETURN
14     25  IF(X)30,27,31
15     27  BJ=0
16     RETURN
17     30  IER=2
18     RETURN
19     31  IF(X-15.)32,32,34
20     32  NTEST=20.+10.*X-X**2/3
21     GO TO 36
22     34  NTEST=90.+X/2
23     36  IF(N-NTEST)40,38,38
24     38  IER=4
25     RETURN
26     40  IER=0
27     NI=N+1
28     BPREV=0.
29     C
30     C
31     C
32     IF(X-5.)50,60,60
33     50  MA=X+6
34     GO TO 70
35     60  MA=1.4*X+60./X
36     70  MB=N+IFIX(X)/4+2
37     MZERO=MAXO(MA,MB)
38     C
39     C
40     C
41     MMAX=NTEST
42     100 DO 190 M=MZERO,MMAX,3
43     C
44     C
45     C
46     FM1=1.0E-28
47     FM=0.
48     ALPHA=0.0
49     IF(M-(M/2)*2)120,110,120
50     110  JT=-1
51     GO TO 130
52     120  JT=1
53     130  M2=M-2
54     DO 160 K=1,M2
55     MK=M-K
56     BMK=2.*FLOAT(MK)*FM1/X-FM
57     FM=FM1
58     FM1=BMK
59     IF(MK-N-1)150,140,150
60     140  BJ=BMK
61     150  JT=-JT
62     S=1+JT
63     160  ALPHA=ALPHA+BMK*S
64     BMK=2.*FM1/X-FM
65     IF(N)180,170,180

```

```

66      170  BJ=BMK
67      180  ALPHA=ALPHA+BMK
68      BJ=BJ/ALPHA
69      IF(ABS(BJ-BPREV)-ABS(0.0001*BJ))200,200,190
70      190  BPREV=BJ
71      IER=3
72      200  RETURN
73      END
1      C
2      C *****
3      C * FILAM IS A PROGRAM TO SOLVE DRAKE DISPERSION EQUATION *
4      C * FOR BRILQUIN & FILAMENTATION. IT USES TWO EXTERNAL *
5      C * FUNCTIONS ZP1 & F. THE INPUTS TO THIS PROGRAM ARE THE *
6      C * SAME AS IN DISP1. THE SAME BISECTION METHOD IS ALSO *
7      C * USED HERE. NO BESSLE FUNCTION IS USED & THE FUNCTION F *
8      C * HERE IS NOT THE SAME F BEEN USED IN DISP1 BUT ZP1 IS *
9      C * THE SAME. *****
10     C
11     C COMPLEX Z,Z0,Z1,Z2,F,S(200)
12     * REAL K,N,MI,KD,KO,KN
13     DIMENSION Y(200),S1(200),Y1(25),Y2(25),Y3(25),Y4(25),S2(200)
14     COMMON K,R,A,Q1,Q2,Q3,Q4,Q5,QP,QM
15     1 READ(5,1)TE,II,MI,ZI
16     FORMAT(4F6.2)
17     T=TI/TE
18     R=ZI/T
19     WRITE(3,19)TE,T,MI,ZI
20     2 READ(5,2)K,N,V,CTH
21     FORMAT(4F6.4)
22     A=T/(1836.2*MI)
23     A=SQRT(A)
24     VT=4.191E7*SQRT(TE)
25     VC=3.0E10/VT
26     KO=4.256E6*SQRT(N/TE)
27     KO=1.0-N
28     KO=5.9426E3*SQRT(KO)/KO
29     KN=K/(2.0*KO)
30     WRITE(3,20)N,CTH,K
31     WRITE(3,21)V,VC,KN
32     VC=VC*VC
33     V=V*V
34     Q2=2.0*K*KO*VC*CTH
35     Q3=2.82843*K/SQRT(N)
36     K=K*K
37     R=R/K
38     Q1=K*VC
39     Q4=0.5*K*V
40     Q5=2.0*K
41     QP=Q1+Q2
42     QM=Q1-Q2
43     READ(5,22)N1,N2,DX,DY,DXO,DYO
44     DO 16 I=1,N1
45     X=DX*I+DXO
46     WRITE(3,23)X
47     DO 4 J=1,N2
48     Y(J)=DY*J+DYO
49     Z=CMPLX(X,Y(J))
50     S(J)=F(Z)
51     S1(J)=REAL(S(J))
52     S2(J)=AIMAG(S(J))
53     IR=0
54     II=0
55     DO 6 J=2,N2
56     F1=S1(J)/S1(J-1)
57     IF(F1.GT.0.0)GO TO 5
58     IR=IR+1
59     Y1(IR)=Y(J-1)
60     Y2(IR)=Y(J)
61     5 F2=S2(J)/S2(J-1)
62     IF(F2.GT.0.0)GO TO 6
63     II=II+1
64     Y3(II)=Y(J-1)
65     Y4(II)=Y(J)

```

```

65      6      CONTINUE
66      IF(IR.EQ.O)GO TO 11
67      DO 10 K1=1,IR
68      Y11=Y1(K1)
69      Y22=Y2(K1)
70      Z1=CMPLX(X,Y11)
71      Z2=CMPLX(X,Y22)
72      F1=REAL(F(Z1))
73      F2=REAL(F(Z2))
74      DO 8 KO=1,20
75      YO=(Y22+Y11)/2.DO
76      ZO=CMPLX(X,YO)
77      FO=REAL(F(ZO))
78      FF=FO/F1
79      IF(FF.GT.O.O)GO TO 7
80      F2=FO
81      Y22=YO
82      GO TO 8
83      7      F1=FO
84      Y11=YO
85      8      CONTINUE
86      WRITE(3,9)YO,FO
87      9      FORMAT(10X,'Y=',E12.5X,'RE(D)=',E12.5)
88      10     CONTINUE
89      11     IF(II.EQ.O)GO TO 16
90      DO 15 K1=1,II
91      Y33=Y3(K1)
92      Y44=Y4(K1)
93      Z1=CMPLX(X,Y33)
94      Z2=CMPLX(X,Y44)
95      F1=AIMAG(F(Z1))
96      F2=AIMAG(F(Z2))
97      DO 13 KO=1,20
98      YO=(Y44+Y33)/2.DO
99      ZO=CMPLX(X,YO)
100     FO=AIMAG(F(ZO))
101     FF=FO/F1
102     IF(FF.GT.O.O)GO TO 12
103     F2=FO
104     Y44=YO
105     GO TO 13
106     12     F1=FO
107     Y33=YO
108     13     CONTINUE
109     WRITE(3,14)YO,FO
110     14     FORMAT(10X,'Y=',E12.5X,'IM(D)=',E12.5)
111     15     CONTINUE
112     16     CONTINUE
113     19     FORMAT(5X,'TE=',F6.15X,'TI/TE=',F6.35X,'MI=',F5.25X,
114     1'Z=',F5.2)
115     20     FORMAT(5X,'N/NC=',F5.35X,'COS( )=',F6.25X,'K/KO=',F5.31)
116     21     FORMAT(5X,'VO/VT=',F5.25X,'C/VT=',E12.55X,'K/2KO=',E12.5)
117     22     FORMAT(2I3,4F6.3)
118     23     FORMAT(15X,'***X=',F7.4,'***')
119     STOP
120     END
121     COMPLEX FUNCTION F(Z)
122     C .....
123     C * THIS FUNCTION IS REQUIRED BY FILAM PROGRAM TO DETER-
124     C * MINE THE DRAKE DISPERSION FUNCTION FOR A GIVEN COMPLEX
125     C * VARIABLE Z.
126     C .....
127     COMPLEX Z,Z1,ZP1,XI,XE,ZS,DP,DM
128     REAL K
129     COMMON K,R,A,Q1,Q2,Q3,Q4,Q5,QP,QM
130     Z1=A*Z
131     XE=(1.O+Z1*ZP1(Z1))/K
132     XI=R*(1.O+Z*ZP1(Z))
133     ZS=Z1*Z1*Q5
134     DP=QP-Q3*Z1-ZS
135     DM=QM+Q3*Z1-ZS
136     F=(1.O+XE+XI)*DP*DM-Q4*XE*(1.O+XI)*(Q1-ZS)

```

```

137 RETURN
138 END
1 C
2 C * THIS FUNCTION IS REQUIRED BY DISP1 & FILAM TO EVALUATE
3 C * THE PLASMA DISPERSION INTEGRAL USING POLYNOMIAL EXPAN-
4 C * SION METHOD.
5 C
6 COMPLEX FUNCTION ZP1(Y)
7 COMPLEX*16 U,Z,U2,AZP,AZPOLD,USQM,ZP
8 COMPLEX Y
9 U=Y
10 NA=10
11 IF(CDABS(U) GE 4 DO)GO TO 3
12 USQM=-U**2
13 IF(DREAL(USQM) GT 150 DO)USQM=DCMPLX(150 DO,0 DO)
14 ZP=DCMPLX(0 DO,1 DO)*1.772453850905516*COEXP(USQM)
15 U2=-2 DO*U**2
16 AZP=-2 DO*U
17 DO 2 N=1,100
18 ZP=ZP+AZP
19 2 AZP=AZP*U2/(2 DO*N+1 DO)
20 ZP=ZP+AZP
21 GO TO 30
22 3 Z=1 DO/U
23 IF(DIMAG(U) LE 0 DO)GO TO 10
24 ZP=0 DO
25 GO TO 20
26 10 CONTINUE
27 USQM=-U**2
28 IF(DREAL(USQM) GT 150 DO)USQM=DCMPLX(150 DO,0 DO)
29 ZP=DCMPLX(0 DO,1 DO)*1.772453850905516*COEXP(USQM)
30 IF(DIMAG(U) LT 0 DO)ZP=2 DO*ZP
31 20 AZP=Z
32 U2=0.5DO*Z**2
33 DO 25 N=1,NA
34 ZP=ZP-AZP
35 AZPOLD=AZP
36 AZP=(2 DO*N-1 DO)*AZP*U2
37 IF(CDABS(AZP) GE CDABS(AZPOLD))GO TO 30
38 25 CONTINUE
39 ZP=ZP-AZP
40 30 CONTINUE
41 ZP1=ZP
42 RETURN
43 END

```

APPENDIX B

MODELING OF THE CO₂ LASER

B.1 Gas Kinetics of the CO₂-N₂-He Mixture

The physics of the CO₂ laser is well known and can be found in most laser textbooks. Only a simple description is given here. The CO₂ molecule undergoes three vibrational modes. The energy of excitation of a molecule in any one mode can have only discrete values which are integral multiples of some fundamental value. At any time a carbon dioxide molecule can vibrate in a linear combination of the three modes. The energy state of the molecule can be represented by three energy quantum numbers (v_1, v_2, v_3) . Some of the low-lying vibrational levels of the CO₂ molecule that are responsible for the laser action are shown in Fig. B.1. In the CO₂ laser, molecules are excited from the ground state to higher energy states where they lose their energy and reach a long-lived state (001) by radiative and nonradiative processes. With sufficient pumping, a population inversion is produced between the (001) state and the (100) state. If the losses in the laser cavity surrounding the medium are sufficiently low, laser oscillation begins. To improve the laser output, nitrogen and helium are added to the CO₂ gas. The excited nitrogen molecules transfer energy to the CO₂ molecules by near-resonance collisional transfer, while the helium speeds up the transition from the (100) state to the ground level, thereby maintaining a large population inversion. In

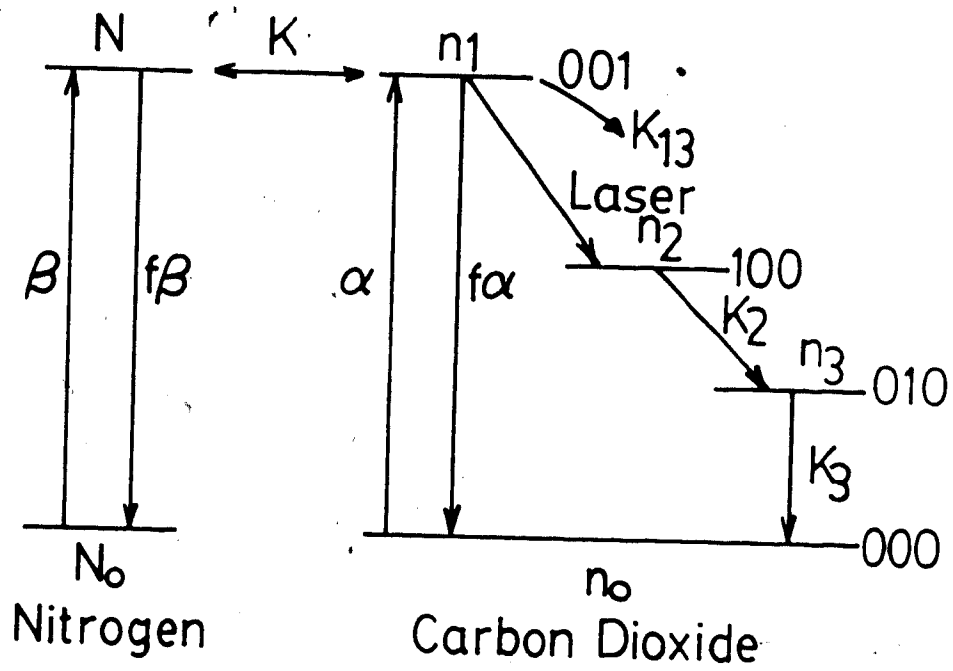


Fig. B.1 Some of the low lying levels of the CO_2 and N_2 molecules.

addition, helium helps to eliminate arcs between the electrodes so that efficient pumping can be provided. The output power and energy as well as the time delay between the electrical discharge and the peak of the laser emission depend on gas mixture, mirror reflectivities and pumping level. Several models have been developed by various workers [1-4] to study the effect of the operating conditions on the output pulse characteristics. For the purpose of comparison between theory and experiment, the model described by Andrews *et al.* [3] was found to be simple and adequate. The following set of rate equations was used to describe the CO₂-N₂ energy level scheme adopted by them:

$$\begin{aligned} \frac{dn_1}{dt} &= \alpha(n_0 - \beta n_1) + \beta(n_0 - n_1)N_0 - k_{13}n_1 - S_0(n_1 - n_2), \\ \frac{dn_0}{dt} &= \beta(N_0 - N) - k_1(N_0 - n_1)N_0, \\ \frac{dn_2}{dt} &= S_0(n_1 - n_2) - k_2n_2, \\ \frac{dn_3}{dt} &= k_2n_2 - k_3n_3 + k_{13}n_1, \\ \frac{dq}{dt} &= S_0(n_1 - n_2) - \omega q - \beta n_1, \end{aligned} \quad (B.1)$$

where n_1 , N , n_2 and n_3 are the electron population densities of the levels shown in Fig. A.1, k_1 and k_2 are the effective decay rates of the (100) and (010) levels respectively, and α and β are the effective electron excitation rates to the (001) level of the CO₂ molecule and the first excited level of the N₂ molecule respectively. All other transitions of level (001) were assumed to go directly to the (010) level with the k_{13} rate since the intermediate states relax rapidly to the (010) level. The last equation in B.1 describes the growth of

photon density q in the laser cavity where ω is the cavity decay rate, D defines the rate of spontaneous emission into the cavity mode of interest and Sq is the rate of stimulated emission. For a laser with cavity mirror reflectivities R_1 and R_2 , cavity length L , gain length l and gas partial pressures x , y and z for N_2 , He and CO_2 respectively, the following parameters are obtained for our laser conditions 3 :

$$\omega = -c \ln R_1 R_2 / 2L = 7 \times 10^7 \text{ sec}^{-1},$$

$$S = cl_0 / L = 7.5 \times 10^{-20} \text{ cm}^{-1} \text{ sec}^{-1},$$

$$k_{13} = (85x + 110y + 265z) = 1.129 \times 10^5 \text{ sec}^{-1},$$

$$k_2 = 1.4 \times 10^5 (x + 0.40y + 0.056z) \times 760 = 3.1 \times 10^7 \text{ sec}^{-1},$$

$$k_3 = (4 \times 10^3 x + 40y + 200z) = 2.16 \times 10^6 \text{ sec}^{-1},$$

$$\alpha = 6 \times 10^{-9} n_e \text{ sec}^{-1},$$

$$\beta = 2 \times 10^{-8} n_e \text{ sec}^{-1},$$

$$D = 10^{-10} \text{ sec}^{-1},$$

where $c (= 10^{-19} \text{ cm}^{-2})$ is the cross section for stimulated emission and n_e is the time dependent discharge electron density. A good estimate of the electron density, based on the discharge current waveform measurements of Burnett and Offenberger 5, is given by

$$n_e(t) = 3 \times 10^{13} \sin\left(\frac{\pi}{500} t\right) \text{ cm}^{-3} \quad (0 \leq t \leq 500 \text{ ns}),$$

$$n_e(t) = 0 \text{ otherwise},$$

where t is in ns and the 500 ns shown in Eq. 4 is the current duration. The power is determined from the partial densities using the following relation 6 :

$$P_0 \approx -h\nu_0 \frac{e \ln R_1}{2L}$$

where P_0 is the power per unit mode volume and $h\nu_0$ is the photon energy of the laser transition.

3.2 Numerical Solution of the Rate Equations

In modeling the TEA CO_2 laser the system of ordinary differential equations (3.1) had to be solved. The classical (fourth order) Runge-Kutta method was applied with a time interval of $\Delta t = 2$ ns. To simplify the method, we assume the N ordinary differential equations of N variables are:

$$\frac{dx_i}{dt} = f_i(t, x_1, x_2, \dots, x_N), \quad i = 1 \text{ to } N.$$

The variable $x_{i,j+1}$ at time $t_j + \Delta t$ is related to $x_{i,j}$ at time t_j by the following equation (4):

$$x_{i,j+1} = x_{i,j} + \Delta t \left(k_{i,1} + 2k_{i,2} + 2k_{i,3} + k_{i,4} \right),$$

where

$$k_{i,1} = \Delta t \cdot f_i(t_j, x_{1,j}, x_{2,j}, \dots, x_{N,j}),$$

$$k_{i,2} = \Delta t \cdot f_i(t_j + \frac{\Delta t}{2}, x_{1,j} + \frac{\Delta t}{2} k_{1,1}, x_{2,j} + \frac{\Delta t}{2} k_{2,1}, \dots, x_{N,j} + \frac{\Delta t}{2} k_{N,1}),$$

$$k_{i,3} = \Delta t \cdot f_i(t_j + \frac{\Delta t}{2}, x_{1,j} - \frac{\Delta t}{2} k_{1,1}, x_{2,j} - \frac{\Delta t}{2} k_{2,1}, \dots, x_{N,j} - \frac{\Delta t}{2} k_{N,1}),$$

$$k_{i,4} = \Delta t \cdot f_i(t_j + \Delta t, x_{1,j} + \Delta t k_{1,1}, x_{2,j} + \Delta t k_{2,1}, \dots, x_{N,j} + \Delta t k_{N,1}).$$

The results of the above analysis is shown in Fig. 3.2 where the laser power is plotted as a function of time. As seen in the

figure, lasing starts about 30 ns after the start of the discharge current and the power reaches maximum in less than 20 ns. The laser pulse is seen to consist of a high amplitude gain-switched spike of duration ≈ 50 ns (FWHM), followed by a relatively low amplitude tail extending for about 1 μ s.

A program listing the above numerical technique for modeling the laser written for the ADAM computer is given below.

```

10  REM      SET THE INITIAL CONDITIONS
20  D1=6E-09
30  D2=2E-08
40  K=5 4E-13
50  K1=112900
60  K2=31380000
70  K3=21640000
80  S=7 5E-20
90  F=1 157
100 C=3E+10
110 W=690000000
120 D=1E-10
130 M=0
140 M1=0
150 M2=0
160 M3=0
170 Q=0
180 T=0
190 NC=5 91E+18
200 NN=2 15E+18
210 N=M
220 N1=M1
230 N2=M2
240 N3=M3
250 Q=00
260 NQ=NC-M1-M2-M3
270 NG=NN-M
280 IF T<500 GOTO 310
290 NE=0
300 GOTO 320
310 NE=3E+13*SIN(6 2832E-03*T)
315 REM      EVALUATE U1,V1,W1,X1,Y1
320 F1=D1*NE*(NO-F*N1)+K*(N*NO-N1*NG)-K1*N1-S1*(N1-N2)
330 F2=D2*NE*(NG-F*N1)+K*(N*NO-N1*NG)
340 F3=S*C*Q*(N1-N2)+K2*N2
350 F4=K2*N2-K3*N3+K1*N1
360 F5=S*C*Q*(N1-N2)-W*Q+D*N1
370 U1=2E-09*F1
380 V1=2E-09*F2
390 W1=2E-09*F3
400 X1=2E-09*F4
410 Y1=2E-09*F5
420 T=T+1
430 N1=M1+U1/2
440 N=M+V1/2
450 N2=M2+W1/2
460 N3=M3+X1/2
470 Q=Q0+Y1/2
480 IF T<500 GOTO 510
490 NE=0
500 GOTO 520
510 NE=3E+13*SIN(6 2832E-03*T)
520 NQ=NC-N1-N2-N3
530 NG=NN-N
535 REM      EVALUATE U2,V2,W2,X2,Y2
540 F1=D1*NE*(NO-E*N1)+K*(N*NO-N1*NG)-K1*N1-S1*(N1-N2)
550 F2=D2*NE*(NG-F*N1)+K*(N*NO-N1*NG)
560 F3=S*C*Q*(N1-N2)+K2*N2
570 F4=K2*N2-K3*N3+K1*N1
580 F5=S*C*Q*(N1-N2)-W*Q+D*N1
590 U2=2E-09*F1
600 V2=2E-09*F2
610 W2=2E-09*F3
620 X2=2E-09*F4
630 Y2=2E-09*F5
640 N1=M1+U2/2
650 N=M+V2/2
660 N2=M2+W2/2
670 N3=M3+X2/2
680 Q=Q0+Y2/2

```

```

690 NO=NC-N1-N2-N3
700 NG=NN-N
710 REM T=T+1 NANOSECOND WILL BE USED LATER
720 T=T+1
730 REM EVALUATE U3,V3,W3,X3,Y3
740 F1=D1*NE*(NO-F*N1)+K*(N*NO-N1*NG)-K1*N1-S*C*Q*(N1-N2)
750 F2=D2*NE*(NG-F*N)-K*(N*NO-N1*NG)
760 F3=S*C*Q*(N1-N2)-K2*N2
770 F4=K2*N2-K3*N3+K1*N1
780 F5=S*C*Q*(N1-N2)-W*Q*D*N1
790 U3=2E-09*F1
800 V3=2E-09*F2
810 W3=2E-09*F3
820 X3=2E-09*F4
830 Y3=2E-09*F5
840 N1=M1+U3
850 N=M+V3
860 N2=M2+W3
870 N3=M3+X3
880 Q=Q0+Y3
890 IF T<500 GOTO 920
900 NE=0
910 GOTO 930
920 NE=3E+13*SIN(6.2832E-03*T)
930 NO=NC-N1-N1-N3
940 NG=NN-N
950 REM EVALUATE U4,V4,W4,X4,Y4
960 F1=D1*NE*(NO-F*N1)+K*(N*NO-N1*NG)-K1*N1-S*C*Q*(N1-N2)
970 F2=D2*NE*(NG-F*N)-K*(N*NO-N1*NG)
980 F3=S*C*Q*(N1-N2)-K2*N2
990 F4=K2*N2-K3*N3+K1*N1
1000 F5=S*C*Q*(N1-N2)-W*Q*D*N1
1010 U4=2E-09*F1
1020 V4=2E-09*F2
1030 W4=2E-09*F3
1040 X4=2E-09*F4
1050 Y4=2E-09*F5
1060 REM CALCULATE THE VARIABLE AT T=T+2
1070 M1=M1+(U1+2*U2+2*U3+U4)/6
1080 M=M+(V1+2*V2+2*V3+V4)/6
1090 M2=M2+(W1+2*W2+2*W3+W4)/6
1100 M3=M3+(X1+2*X2+2*X3+X4)/6
1110 Q0=Q0+(Y1+2*Y2+2*Y3+Y4)/6
1120 REM CALCULATE THE POWER
1130 P=1.3186E-12*Q0
1140 PRINT T,P
1150 IF T<1000 GOTO 210,
1160 END

```

APPENDIX C

ABEL INVERSION

In cylindrically symmetric plasma one can determine the plasma density from the phase shift measured by an interferometer. Figure C.1 represents a cross sectional view of the plasma. The plasma density is assumed to be symmetric in any azimuthally symmetric circle of radius r . The change in fringes that an observer sees due to the variation in density along the path AA' is given by Eqn. (3.7)

$$N = 2G \int_0^{y_0} n(r) dy, \quad (C.1)$$

where $G = \frac{e^2 \lambda_0}{mc^2}$. Equation (C.1) can be written in a more convenient form after making the substitution, $y = (r^2 - x^2)^{\frac{1}{2}}$

$$N(x) = 2G \int_x^R \frac{n(r) r dr}{(r^2 - x^2)^{\frac{1}{2}}}. \quad (C.2)$$

Abel inversion of Eqn. (C.2) yields

$$n(r) = -\frac{1}{\pi G} \int_r^R \frac{N'(x) dx}{x (x^2 - r^2)^{\frac{1}{2}}}, \quad (C.3)$$

where $N'(x)$ is the first derivative of $N(x)$ with respect to x .

Several numerical methods have been used to solve the integral in Eqn. (C.3). A summary of these methods is given in reference [1]. The method of Nestor and Olsen [2] was found to be simple and adequate. This method is based on linear expansion of the integral in (C.3) assuming equal intervals along x and R . Accordingly, the plasma

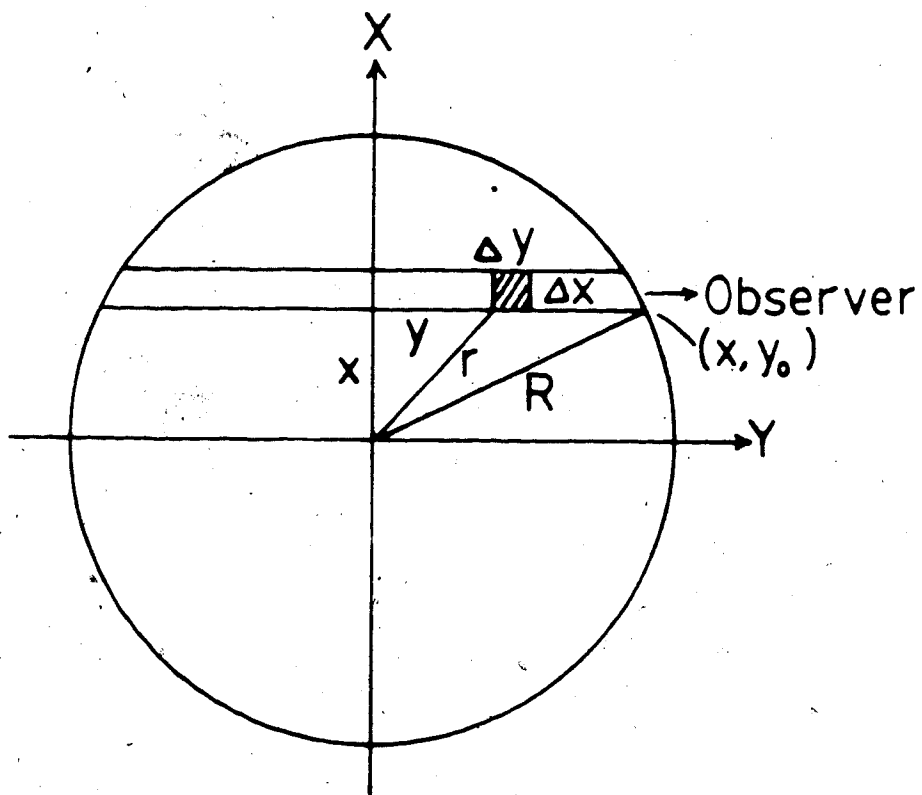


Fig. 3.1 Cross sectional view of a cylindrically symmetric plasma.

at a radius of $(k-1) x$ is

$$n_k(r) = - \frac{2}{\pi G \Delta x} \sum_{n=k}^{N_0} n(x) B_{k,n}, \quad (3.4)$$

where

$$B_{k,n} = -A_{k,k} \text{ for } n = k$$

$$B_{k,n} = A_{k,n-1} - A_{k,n} \text{ for } n \geq k+1$$

$$A_{k,n} = \frac{|n^2 - (k-1)^2|^{\frac{1}{2}} - |(n-1)^2 - (k-1)^2|^{\frac{1}{2}}}{2n - 1}.$$

Here k and n represent the respective radial and lateral points under consideration, Δx is the interval width, and N_0 is the number of points used.

A computer program was written for the HP-85 computer to do Abel inversion. The program digitizes the fringe pattern from an enlarged photograph, calculates the density $n(r)$ and plots it as a function of r . A list of the program is given below.

```

10 PLOTTER IS 705
20 OPTION BASE 1
30 DIM A(6),B(6),X(100),Y(100)
40 DIM F(100)
50 DISP "CALCULATE THE BASE LINE"
60 DISP "DIGITIZE 3 POINTS EACH SIDE OF THE LINE"
70 BEEP
80 FOR I=1 TO 6
90 DIGITIZE B(I),A(I)
100 NEXT I
110 DISP "GIVE THE RESOLUTION IN MICRON/FRINGE"
120 BEEP
130 INPUT R
140 C1=C2=C3=C4=0
150 FOR I=1 TO 6
160 C1=C1+B(I)*A(I)
170 C2=C2+A(I)
180 C3=C3+B(I)
190 C4=C4+A(I)*A(I)
200 NEXT I
210 A2=(6*C1-C2*C3)/(6*C4-C2*C2)
220 A1=(C3-A2*C2)/6
230 DISP "DETERMINE THE CENTRE OF THE PROFILE"
240 BEEP
250 DISP "DIGITIZE TWO POINTS AT THE EDGES OF THE
PROFILE STARTING FROM THE BOTTOM"
260 DIGITIZE X1,Y1
270 DIGITIZE X2,Y2
280 YO=(Y1+Y2)/2
290 D=(Y2-Y1)/2
300 XO=(X2+X1)/2
310 MOVE XO,YO
320 DISP "SPECIFY THE # OF DATA INPUT"
330 BEEP
340 INPUT N
350 DISP "WRITE 1 FOR MOVING UP OR 2 FOR MOVING DOWN"
360 BEEP
370 INPUT L
380 IF L=1 THEN 410
390 D=-D/(N-1)
400 GOTO 420
410 D=D/(N-1)
420 X(1)=XO
430 BEEP
440 DISP "START DIGITIZING"
450 FOR I=1 TO N
460 Y(I)=YO+(I-1)*D
470 MOVE X(I),Y(I)
480 DIGITIZE X(I),Y(I)
490 X(I+1)=X(I)
500 NEXT I
510 DISP "CALCULATE THE DATA IN UNITS OF FRINGE SHIFT"
520 DISP "DIGITIZE TWO POINTS ON TWO STRAIGHT LINES"
530 BEEP
540 DIGITIZE X1,Y1
550 DIGITIZE X2,Y2
560 DISP "WRITE # OF FRINGES BETWEEN THESE LINES"
570 INPUT M
580 Z=ABS(X2-X1)/M
590 FOR I=1 TO N
600 X(I)=(X(I)-A1-A2*Y(I))/Z
610 Y(I)=ABS(Y(I)-YO)*R/Z
620 Y(I)=ABS(Y(I))
630 DISP X(I),Y(I)
640 NEXT I
650 D=0.001*ABS(D)*R/Z
660 FOR I=K TO N-1
670 F(K)=0
680 FOR M=K TO N-1
690 X1=(K-1)*(M-1)
700 X2=(M-1)*(M-1)
710 X3=(M-2)*(M-2)

```



```
720 IF M=K THEN 770
730 A1=SQRT(M*M - X1) - SQRT(X2 - X1) / (2*M - 1)
740 A2=(SQRT(X2 - X1) - SQRT(X3 - X1)) / (2*M - 1)
750 B=A2 - A1
760 GOTO 780
770 B=-1/SQRT(2*K - 1)
780 F(K)=F(K)+X(M)*B
790 NEXT M
800 F(K)=F(K)*2.0479E17/D
810 PRINT Y(K), F(K)
820 NEXT K
830 FOR I=1 TO N-1
840 F(I)=F(I)/1.E18
850 Y(I)=Y(I)/100
860 NEXT I
870 Y7=CEIL(Y(N-1))
880 Y6=Y7/8
890 PLOTTER IS 1
900 PEN 1 • GCLEAR
910 SCALE -Y6, Y7, 0, 10
920 XAXIS 1, 1, 0, 7
930 YAXIS 0, 1, 1, 10
940 LDIR 0
950 FOR X7=0 TO 10
960 MOVE X7, 0
970 LABEL VAL$(X7)
980 NEXT X7
990 FOR Y5=1 TO 10
1000 MOVE -Y6, Y5
1010 LABEL VAL$(Y5)
1020 NEXT Y5
1030 PENUP
1040 FOR I=1 TO N-1
1050 PLOT Y(I), F(I)
1060 NEXT I
1070 END
```

APPENDIX D

NUMERICAL SOLUTION OF THE $S(\omega, \bar{k})$ FUNCTION

The numerical solution of the $S(\omega, \bar{k})$ function as defined in Eqs. (3.20) and (3.21), including the complex functions $G_e(\omega)$ and $A(\omega)$, is straight forward except for the integral part in the definition of the function f . To solve this part, the function f may be rewritten in the following form

$$f(x) = 2x \int_0^x \exp(t^2 - x^2) dt, \quad (D.1)$$

where the variable x was moved inside the integral. The $\exp(t^2 - x^2)$ function is only significant when t is close to x . Therefore, the integral in (D.1) is evaluated for the interval

$$t = 0 \text{ to } x, \quad \text{when } |x| < 5$$

and

$$t = (x^2 - 1)^{1/2} \text{ to } x, \quad \text{when } |x| \geq 5.$$

The above range of t is then divided into 100 intervals where Simpson's rule is applied to evaluate the integral. The above method is proved to be fast and fairly accurate.

Below is listed a Fortran IV program for the AMDAHL 470 computer.

The inputs for this program are T_0 and T_1 in eV, M_0 (required to calculate λ_D), the scattering angle, the ion mass number, the spectrum range in k and the step in k at which $S(\omega, \bar{k})$ is to be calculated.

```

1 C .....
2 C * PROGRAM SCATT CALCULATES THOMSON SCATTERING SPECTRUM. *
3 C * S(K,W) FOR A GIVEN TEMPERATURES (TE & TI IN eV). *
4 C * PLASMA DENSITY (N IN CM-3), ANGLE (THETA IN DEGREES). *
5 C * ION MASS (A IN ATOMIC UNITS) AND IONIZATION DEGREE (Z) *
6 C * IN A WAVELENGTH RANGE (RANGE IN ANGSTROMS) WITH A STEP *
7 C * D IN ANGSTROMS). *
8 C .....
9 REAL N,K
10 COMPLEX GE,GI,GEE,GII,E
11 READ(5,1)TE, TI, N, THETA, A, RANGE, D, Z
12 1 FORMAT(2F5.1, E7.2, 5F7.2)
13 TR=TE/TI
14 K=1.8099E5*SIN(0.00873*THETA)
15 ALPHA=SQRT(N/TE)/(743.4*K)
16 WRITE(6,5)
17 5) FORMAT('1')
18 C WRITING THE PLASMA & SCATTERING PARAMETERS
19 WRITE(6,10)TE, TI, A, Z, N
20 10) FORMAT(1X, 'TE=' ,F6.1,5X, 'TI=' ,F6.1,5X, 'A=' ,F5.1,5X, 'Z=' ,F4.1,5X,
21 'N=' ,E7.2)
22 WRITE(6,16)TR, THETA, ALPHA
23 16) FORMAT(1X, 'RATIO=' ,F5.2,5X, 'ANGLE=' ,F5.1,5X, 'ALPHA=' ,E7.2)
24 WRITE(6,17)
25 WRITE(6,20)
26 17) FORMAT(////)
27 20) FORMAT(' WLENGTH' ,5X, 'ION COMP' ,9X, 'ELECTRON COMP' ,4X, 'S(W,K)')
28 WRITE(6,21)
29 21) FORMAT(1X, '-----')
30 1)-----)
31 C CALCULATE THE COMPLEX FUNCTIONS GE & GI
32 R=A/TI
33 A1=9.5126E-9/SQRT(TE)
34 A2=-2.8428E-16/TE
35 B1=4.076E-7*SQRT(R)
36 B2=-5.2197E-13*R
37 WE=5.931E7*K*SQRT(TE)
38 WI=1.384E6*K/SQRT(R)
39 ALP=ALPHA*ALPHA
40 D1=-RANGE
41 25) D1=D1+D
42 IF(D1.GT.0.0)GO TO 150
43 VE=3.9103E11*ABS(D1)
44 VE=(W/K)**2
45 X=W/WE
46 Y=W/WI
47 GI1=1-F(Y)
48 GE1=1-F(X)
49 X2=X*X
50 Y2=Y*Y
51 IF(Y2.GT.30.0)GO TO 30
52 GI2=1.7725*Y*EXP(-Y2)
53 GO TO 40
54 30) GI2=0.0
55 40) IF(X2.GT.30.0)GO TO 50
56 GE2=1.7725*X*EXP(-X2)
57 GO TO 60
58 50) GE2=0.0
59 60) GE=CMPLX(GE1,GE2)
60 GI=CMPLX(GI1,GI2)
61 GE=-ALP*GE
62 GI=-Z*TR*ALP*GI
63 E=1-GI-GE
64 GEE=(1-GI)/E
65 GII=GE/E
66 C1=VE*A2
67 IF(C1.LT.-50.0)GO TO 70
68 FE=A1*EXP(C1)*CABS(GEE)**2
69 GO TO 80
70 70) FE=0.0
71 80) C2=VE*B2

```

```

72      IF(C2.LT.-50.0)GO TO 90
73      FI=Z*B1*EXP(C2)*CABS(GII)**2
74      GO TO 95
75      90  FI=0.0
76      95  FEI=FE+FI
77      WRITE(6,100)D1,FI,FE,FEI
78      100 FORMAT(1X,F7.2,5X,3(E12.5,5X))
79      GO TO 25
80      150 STOP
81      END
82      C *****
83      C * FUNCTION F IS USED TO EVALUATE THE INTEGRAL PART IN *
84      C * S(K,W) USING THE METHOD DESCRIBED IN THIS APPENDIX *
85      C *****
86      FUNCTION F(X)
87      REAL*8 S(101),Y,Y1,T,T1,D,F1,F2,FN
88      IF(X.EQ.0.0)GO TO 100
89      Y=DBLE(X)
90      Y1=Y*Y
91      T=0.0
92      F1=0.0
93      F2=0.0
94      IF(Y.LT.5.0)GO TO 10
95      T=DSQRT(Y1-16.0)
96      10  D=(Y-T)/100.0
97      DO 20 I=1,101
98      T1=T+(I-1)*D
99      20  S(I)=DEXP(T1*T1-Y1)
100     DO 50 I=2,100,2
101     F1=F1+S(I)
102     DO 60 I=3,100,2
103     F2=F2+S(I)
104     FN=2.0*(4.0*F1+2.0*F2+S(1)+S(101))*D*Y/3.0
105     F=SINGL(FN)
106     GO TO 200
107     100 F=0.0
108     200 RETURN
109     END

```

End of file

APPENDIX E

DECONVOLUTION OF SPECTRA

Accurate measurement of the scattered spectrum is possible only if the spectral bandwidth of the instrument is negligible in comparison with the halfwidth of the spectral shape. If the spectral width and the instrumental width are of comparable magnitude, the recorded distribution $A(\lambda)$ will be a complicated function of the true spectrum $B(\lambda)$ and the instrumental function $S(\lambda)$. Mathematically the relation can be expressed by the convolution integral

$$A(\lambda) = \int_{-\infty}^{\infty} S(\lambda - \lambda') B(\lambda') d\lambda' \quad (E.1)$$

In measuring the ion feature of the scattered spectra the width of the ion feature is so often small that the effect of the spectrometer slit cannot be ignored. To recover the true spectral shape $B(\lambda)$ from the recorded one, a deconvolution routine similar to the one suggested by Jones *et al.* [1] was used. This method may be explained by first writing Eqn. (E.1) in a more convenient form

$$A_i = \sum_{j=1}^N S_{ij} B_j, \quad i = 1 \text{ to } N, \quad (E.2)$$

or in a matrix form

$$A = SB, \quad (E.3)$$

where N is the number of data input. The technique used by Jones *et al.* [1] is to recover B from A and S using the following iterative method

where the iterations can cease when ϵ approaches to within a certain limit. The elements of the slit matrix were of the form

$$c_{ij} = \begin{matrix} c_{11} & c_{12} & \dots & c_{1N} \\ c_{21} & c_{22} & & c_{2N} \\ \vdots & \vdots & & \vdots \\ c_{N1} & c_{N2} & & c_{NN} \end{matrix}$$

with rows normalized to unity,

For our application, each row was taken as a truncated normalized Gaussian function; its center was taken at $(N+1)/2$ (where N is an odd number) and its FWHM was made equivalent to the FWHM of a monochromatic line seen by the spectrometer (usually the He-Ne laser). A five point smoothing routine

$$A'_i = [17A_i + 12(A_{i-1} + A_{i+1}) - 3(A_{i-2} + A_{i+2})] / 35 \quad (E.7)$$

was used to smooth the experimental data A and the calculated data A^n after each iteration.

A FORTRAN program was written for the AMDAHL 470 capable of handling up to 50 equally spaced data points. The program plots the

deconvoluted data of the pattern and provides the result of the
 the recovery of the original data. Usually three to four iterations
 are required to deconvolute the matrix A of 100-200 \times data. Figure
 7.1 shows a convoluted spectrum a constructed from a true spectrum,
 b , and a truncated Gaussian slit function of FWHM = 13 using Eqs.
 7.1 and 7.5. The deconvoluted spectrum of Fig. 7.1 is shown in
 Fig. 7.2 after three iterations. A listing of the program codes
 is given below.

11

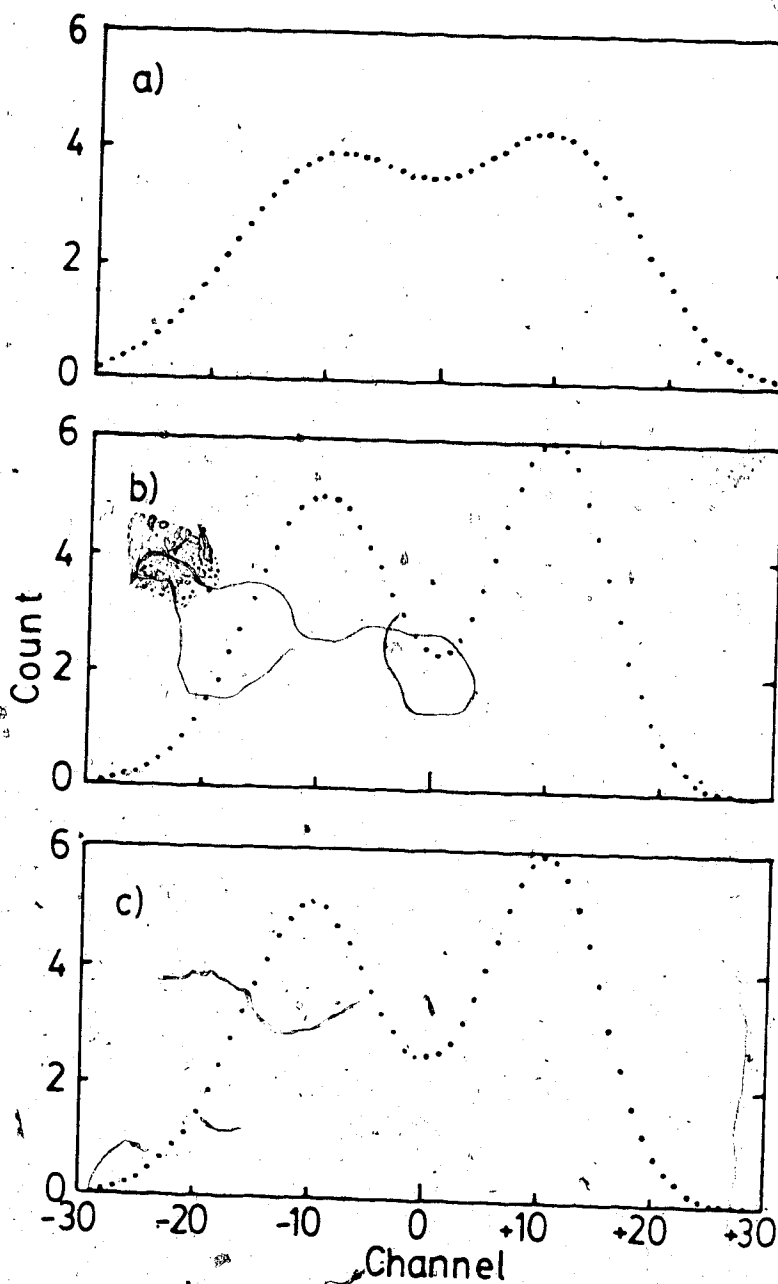


Fig. E.1 Effect of a finite slit on a spectrum.
a) The spectrum as observed through a Gaussian slit of FWHM = 13 channels. b) The true spectrum. c) The recovery of the true spectrum from the observed one using the deconvolution technique.


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.....
PROGRAM DEC FOR DECONVOLVING EXPERIMENTAL DATA USING
TRUNCATED GAUSSIAN SLIT FUNCTION AND PLOTTING THE
RESULTS ON THE PRINTER THE INPUTS TO THIS PROGRAM ARE
FW THE FWHM OF THE SLIT FUNCTION IN CHANNELS; N, THE
NO OF DATA TO BE DECONVOLVED; A(N), THE MEASURED
DATA
.....
DIMENSION A(500),B(500),C(500),S(500),A1(500),DELTA(16)
DIMENSION AA(80),LL(500)
DATA X1,X2/
READ(5,1)FW,N
FORMAT(F4,1,13)
DETERMINING THE SLIT FUNCTION
S(1)=0.93944/FW
DO 5 I=2,N
X=I-1
Y=2.7726*(X/FW)**2
IF(Y.GT.100.0)GO TO 2
S(I)=S(1)*EXP(-Y)
GO TO 5
S(I)=0.0
CONTINUE
READ(2,10)(A(I),I=1,N)
FORMAT(10F5,2)
WRITE(6,20)(A(I),I=1,N)
FORMAT(5(1X,E10,3))
DO 25 I=1,16
DELTA(I)=0
ITERATIONS START TO CALCULATE THE DECONVOLUTION MATRIX
IT=1
CALL SMOOTH(N,A)
WRITE(6,20)(A(I),I=1,N)
CALL MATM(N,A,B,S)
CALL RATIO(N,A,B,C,A)
CALL SMOOTH(N,C)
DO 30 I=1,N
DELTA(IT)=DELTA(IT)+C(I)-A(I)
A1(I)=C(I)
DELTA(IT)=DELTA(IT)/N
CALL MATM(N,A1,B,S)
CALL RATIO(N,A1,B,A)
DO 40 I=1,N
DELTA(IT+1)=DELTA(IT)+C(I)-A1(I)
DELTA(IT+1)=DELTA(IT+1)/N
WRITE(6,42)DELTA(IT)
FORMAT(5X,E12,5)
DEL=(DELTA(IT+1)-DELTA(IT))/DELTA(IT)
IF(ABS(DEL).LT.0.05)GO TO 60
IF(IT.EQ.15)GO TO 50
IT=IT+1
CALL SMOOTH(N,C)
DO 45 I=1,N
A1(I)=C(I)
GO TO 35
WRITE(6,55)
FORMAT(10X, '***CONVERGENCE COULD NOT BE REACHED***')
WRITE(6,65)IT
FORMAT(15X, 'IT=',12)
WRITE(6,20)(C(I),I=1,N)
CALL MATM(N,C,B,S)
WRITE(6,20)(B(I),I=1,N)
XX=C(1)
DO 70 I=2,N
IF(C(I).GT.XX)XX=C(I)
CONTINUE
DO 75 I=1,N
LL(I)=80.0*C(I)/XX
CONTINUE
DO 100 I=1,N
DO 85 J=1,80
IF(J.EQ.LL(I))GO TO 80

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73      AA(J)=X1
74      GO TO 85
75      80      AA(J)=X2
76      85      CONTINUE
77      WRITE(6,90)I,(AA(K1),K1=1,80)
78      90      FORMAT(1X,I3,' ',BOA1)
79      100     CONTINUE
80      STOP
81      END
82      C
83      C * SUBROUTINE SMOOTH IS USED TO SMOOTH THE INPUT DATA AND
84      C * THE DECONVOLVED DATA AFTER EACH ITERATION AS WELL.
85      C * USING FIVE-POINT SMOOTHING TECHNIQUE
86      C
87      SUBROUTINE SMOOTH(N,A)
88      DIMENSION A(500),F(500)
89      M=N-2
90      DO 5 I=3,M
91      F(I)=(17 *A(I)+12 *(A(I-1))+3 *(A(I-2))+A(I+2)))/35 0
92      5      CONTINUE
93      DO 10 I=3,M
94      10     A(I)=F(I)
95      A(1)=(17 *A(2)+12 *A(3))/35 0
96      A(2)=(17 *A(1)+12 *(A(1)+A(3))-3 *A(4))/35 0
97      A(N)=(17 *A(N-1)+12 *(A(N-2)+A(N))-3 *A(N-3))/35 0
98      A(N)=(17 *A(N-1)+3 *A(N-2))/35 0
99      RETURN
100     END
101     C
102     C * SUBROUTINE MATM PERFORMS THE MULTIPLICATION OF THE SLIT
103     C * MATRIX AND THE DATA MATRIX
104     C
105     SUBROUTINE MATM(N,A,B,S)
106     DIMENSION A(500),B(500),S(500)
107     DO 10 I=1,N
108     B(I)=0.0
109     DO 5 J=1,I
110     B(I)=B(I)+S(I-J+1)*A(J)
111     5      CONTINUE
112     IF(I EQ.N)GO TO 10
113     J2=N-I+1
114     DO 10 J=2,J2
115     B(I)=B(I)+S(J)*A(J+I-1)
116     10     CONTINUE
117     RETURN
118     END
119     C
120     C * SUBROUTINE RATIO CALCULATES THE NEW DECONVOLVED
121     C * MATRIX C FROM A AND B.
122     C
123     SUBROUTINE RATIO(N,A1,B,C,A)
124     DIMENSION A1(500),B(500),C(500),A(500)
125     DO 10 I=1,N
126     IF(B(I).EQ.0.0)GO TO 5
127     C(I)=A1(I)*A(I)/B(I)
128     GO TO 10
129     5      C(I)=0.0
130     10     CONTINUE
131     RETURN
132     END

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