

**Practical considerations for body composition assessment of adults with class II/III obesity  
using bioelectrical impedance analysis or dual-energy X-ray absorptiometry.**

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## **Abstract**

**Purpose of Review:** To explore the practical considerations for body composition assessment of adults with class II/III obesity. Studies assessing adults (18-64 years) with a body mass index (BMI)  $\geq 35$  kg/m<sup>2</sup> with bioelectrical impedance analysis (BIA) and/or dual-energy X-ray absorptiometry (DXA) were included. **Recent Findings:** Twelve studies met inclusion criteria. Five considerations were identified: variances in equipment and technology; equipment weight capacity; subject positioning; tissue penetration; and total body hydration. In subjects with BMI  $\geq 35$  kg/m<sup>2</sup>, BIA overestimated fat free mass with scaling errors as BMI increased. DXA provided accurate and reliable body composition measures, but equipment-related barriers prevented assessment of some taller, wider and heavier subjects. **Summary:** BIA is an unreliable method to assess body composition in class II/III obesity. Advancements in DXA technology (e.g. iDXA), methodology (e.g. subject positioning, longer scan times) and more inclusive testing criteria (e.g. use equipment limits not just BMI) may improve access and understanding of body composition in this cohort.

**Key Words:** bioelectrical impedance analysis; body composition; fat mass; dual energy X-ray absorptiometry; lean soft tissue; obesity.

## 1 **Introduction**

2 Obesity defined as a body mass index [BMI]  $\geq 30$  kg/m<sup>2</sup> affects one in three adults in the United  
3 States (US) [1] and Canada [2]. There are three classes of obesity: class I (BMI 30-34.9 kg/m<sup>2</sup>),  
4 class II (BMI 35-39.9 kg/m<sup>2</sup>) and class III (BMI  $\geq 40$ kg/m<sup>2</sup>) [3]. Class III obesity affects 2.5% of  
5 Canadian and 6.4% of American adults, impacting more women (3% Canada, 8.3% US) than  
6 men (2% Canada, 4.4% US), and is associated with the highest level of health risk [1, 2].

7 BMI is commonly used to identify those at increased health risk and as referral criteria for  
8 obesity treatment, including bariatric surgery (e.g. BMI  $\geq 35$  kg/m<sup>2</sup>) [4]. Although quick and easy  
9 to determine, BMI is a proxy measure for adiposity; it cannot estimate or quantify fat mass nor  
10 determine the presence of conditions such as sarcopenia (lower muscle mass and function).  
11 Sarcopenia is most associated with older adults [5], but it can occur across all age and BMI  
12 categories [6] and in healthy middle-aged adults [7]. Body composition analysis is needed to  
13 quantify fat mass (FM) and fat-free mass (FFM), including the components of FFM, specifically  
14 bone, lean soft tissue (LST) and total body water (TBW). Although there is great emphasis on  
15 FM in obesity, the amount of FFM is essential to health. A desirable outcome of obesity  
16 treatment is to not just reduce total body weight, but to achieve a reduction in FM while  
17 preserving FFM. Lower FFM combined with higher FM, known as sarcopenic obesity, is linked  
18 with increased morbidity and mortality [8].

19 Validated methods and tools for the assessment of body composition have been developed to  
20 objectively quantify FM and FFM. The two most commonly used tools for body composition  
21 analysis in clinical and research settings are bioelectrical impedance analysis (BIA) and dual-  
22 energy X-ray absorptiometry (DXA), respectively. Clinicians and researchers are increasingly  
23 interested in the assessment of body composition as part of the obesity treatment plan to help  
24 inform treatment decisions and optimize patient outcomes. To provide some background on  
25 these methods in the context of obesity, a brief overview of BIA and DXA is included. Interested  
26 readers may want to review the following tutorials: a two-part series on BIA published by Kyle  
27 et al. [9, 10], LST imaging including BIA and DXA by Prado & Heymsfield [11], and body  
28 composition tools for assessment of adult malnutrition by Earthman [12].

29

## 30 **Bioelectrical Impedance Analysis**

31 BIA is commonly used in a clinical setting because the equipment is small, portable, affordable,  
32 and relatively easy to use requiring minimal training. BIA utilizes a mild electrical current  
33 (single or multi-frequency waves) to measure differences in resistance and reactance between  
34 tissue types based upon water and electrolytes content. Population-specific regression equations  
35 are used to estimate FM and FFM, usually based in the relation between TBW and FFM. If  
36 normal-weight regression equations are used for subjects with obesity, measurement errors from  
37 abnormal tissue density and hydration can result.

38 Foundational to the technology, two important assumptions are made: 1) the body is a consistent  
39 cylinder [9, 10] and 2) tissue hydration status is constant (73.2%) [13] and the ratio of  
40 extracellular water (ECW) to intracellular water (ICW) is a consistent proportion (1/3). Obesity  
41 challenges both of these assumptions. With obesity, there can be variance in FM distribution  
42 (e.g. central vs. peripheral, android vs. gynoid [14]), and fluid distribution (e.g. edema,  
43 lymphedema) or altered body shape (e.g. shortened limbs or amputees [10]), resulting in body  
44 segments not shaped as a consistent cylinder.

45 For the second assumption, tissue hydration status is not a constant across BMI categories.  
46 Obesity is associated with a state of general “overhydration”, with excess TBW and an increased  
47 ratio of ECW relative to ICW. The hydration status of FFM is elevated, one study measured  
48 75.6% using isotope dilution [15]. Elevated TBW and ECW will result in errors of  
49 overestimation of FFM and thereby underestimation of FM, with lower accuracy at higher levels  
50 of obesity [14, 16].

51 Another challenge with BIA and obesity is the fact that single frequency (50 kHz) waves cannot  
52 fully penetrate the cell membrane. Only some of the ICW is included in the TBW values,  
53 resulting in an overestimation of TBW and FFM and underestimation of FM [17]. Although  
54 multiple-frequency waves can improve tissue penetration, the altered ratio of ECW:ICW and  
55 increased resistance of ICW still result in overestimation of FFM in subjects with obesity [17,  
56 16]. A summary of the measurement errors using BIA in subjects with obesity is presented in  
57 **Table 1**. In the 2004 European Society for Parenteral and Enteral Nutrition (ESPEN) Guidelines,

58 BIA assessment was determined to have questionable validity for FFM and FM when BMI > 34  
59 kg/m<sup>2</sup> [10].

### 60 **Dual-energy X-ray Absorptiometry**

61 DXA utilizes x-rays (photons with two different energy levels) to measure the attenuation (i.e.  
62 energy absorbed) by each tissue type. FM and FFM (which includes separate measures for bone  
63 and LST) are measured for the whole body or segments of interest such as appendicular skeletal  
64 muscle mass (ASM= sum of the LST from the limbs, a surrogate measurement of total muscle  
65 mass). DXA provides an accurate and safe assessment of body composition, with minimal  
66 radiation exposure and provides measurement of more components than BIA. DXA is commonly  
67 used in research or clinical diagnostic settings (e.g. bone density), as it requires trained  
68 technicians, a large dedicated room space and capital expenditure. The precision and reliability  
69 of DXA leads it to be the reference method for body composition analysis [11].

70 Although BIA and DXA have been extensively used in “healthy” populations (i.e. normal BMI  
71 18.5-24.9 kg/m<sup>2</sup>) and older adults (e.g. for bone density studies), these tools are less commonly  
72 used in adults with class II/III obesity. One of the benefits of DXA over BIA is the ability to  
73 assess bone density, which is now recommended for patients after bariatric surgery [4].  
74 Measurements of body composition in this cohort can enhance assessment and risk stratification  
75 of the complex and diverse chronic disease of obesity, including identification of sarcopenic  
76 obesity (i.e. low muscle mass and high adiposity) and osteosarcopenic obesity [6, 18, 7, 19].  
77 Understanding the barriers to body composition assessment can support patient care management  
78 with evidence-based practice tools and identify opportunities for future research.

### 79 **Literature Search Methodology**

80 The purpose of this review was to identify recent studies assessing body composition in adults  
81 (18-64 years) with class II /III obesity (BMI ≥ 35 kg/m<sup>2</sup>) and explore practical considerations for  
82 use of the two most commonly used body composition methods, BIA and DXA. A literature  
83 search was conducted using Medline, Scopus and Web of Science databases for studies  
84 published from 01 October 2005 to 31 October 2015 that measured body composition with BIA  
85 and/or DXA of adults (18-64 years) with a BMI ≥35 kg/m<sup>2</sup>. Studies including children (17 years

86 or less), older adults (65 years or more), subjects with a BMI  $<35$  kg/m<sup>2</sup> or cancer, were  
87 excluded.

88 Twelve studies published met inclusion criteria; nine studies used a single method, either BIA  
89 (five studies) [20-24] or DXA (four studies) [25-28], while three of the 12 studies compared BIA  
90 to DXA [29-31]. Of the eight BIA studies, five utilized a single frequency wave (50 kHz) [29,  
91 20-23] and three utilized multi-frequency waves [30, 24, 31]. Of the seven DXA studies, five  
92 used the standard DXA technology [29-31, 26, 28] and two studies used newer iDXA technology  
93 [25, 27]. In total, there were 920 subjects (77.7% female) and six of the 12 studies included post-  
94 bariatric surgery subjects (n=500, 69.2% roux-en-Y gastric bypass).

### 95 **Defining Obesity: Comparing BMI to Percentage of Fat Mass (%FM)**

96 Obesity can be defined by %FM based upon body composition analysis. There are several  
97 published cut-points for %FM that are sex-specific [32]. Frankenfield et al. [21] used BIA  
98 (n=141, BMI 15.9-93.4 kg/m<sup>2</sup>) to explore the accuracy and specificity of BMI to identify  
99 subjects that exceeded the %FM cut-points. All subjects with obesity (approximately 40% of the  
100 sample) exceeded the %FM cut-points ( $>25\%$  for males and  $>30\%$  for females), showing BMI  $\geq$   
101 30 kg/m<sup>2</sup> had a high sensitivity and accuracy to identify excess adiposity. For subjects with a  
102 BMI  $<30$  kg/m<sup>2</sup>, 46% of females and 30% of males exceeded %FM cut points. The authors noted  
103 alterations in FM and FFM were identified across all BMI categories, supporting the notion that  
104 BMI alone can misclassify subjects at increased health risk due to unfavourable body  
105 composition [21].

### 106 **Barriers to Assessment of Adults with Class II/III Obesity**

107 Methodological and equipment-related limitations for the assessment of adults with class II/III  
108 obesity were identified. These barriers to assessment of body composition in this clinical cohort  
109 were clustered into five key areas: differences in equipment and technology; equipment weight  
110 capacity; subject positioning; total body water; and tissue penetration.

#### 111 *Differences in equipment and technology*

112 In the selected studies, five countries (Brazil, Canada, France, Italy, United States of America)  
113 were represented. Eight different BIA models and four different DXA models from two

114 manufacturers (Hologic, GE Healthcare) were used. The software versions were not often  
115 reported, which is important as software upgrades occur more often than hardware. The  
116 difference in equipment is inevitable, considering the number of countries, different  
117 manufacturers, product advancements, different times of procurement and publication. It is  
118 important to keep in mind there are differences in technique, measurement and study samples,  
119 impacting the outcome data and comparisons of studies [12].

### 120 *Equipment weight capacity*

121 Both BIA and DXA require measured total body weight to determine body composition. A  
122 weigh scale is often integrated into the equipment, with weight capacity limits in place by the  
123 manufacturer. A summary of weight capacity limits for different full body DXA models is found  
124 in **Table 2**. A separate or “stand-alone” scale may also be used to measure body weight. All  
125 reviewed studies reported measured weights. Only four of the eight BIA studies indicated the use  
126 of a stand-alone weigh scale and no BIA studies reported the scale weight capacity. Compared to  
127 the DXA studies, subjects with the highest weights were included in the BIA studies (maximum  
128 214.0 kg [20]). Weight and BMI were used as exclusion criteria from DXA studies due to  
129 equipment weight capacity limits. Five of the seven DXA studies reported the weight capacities  
130 from 120 to 160 kg [29, 30, 26] with the recently commercialized iDXA limits of 182 kg [27] up  
131 to 204 kg [25]. Two of the seven DXA studies did not report equipment weight capacity, instead,  
132 used  $BMI > 40 \text{ kg/m}^2$  as a surrogate marker for exclusion [31, 28].

133 Equipment weight capacity limits the available data on subjects with class II/III obesity and  
134 validation of body composition tools in this cohort. Due to individual variance in height and  
135 weight, use of BMI alone may unnecessarily exclude some subjects from DXA. Assessment and  
136 reporting subject anthropometrics for each limiting factor may improve inclusion criteria and  
137 access to those excluded from DXA measurements based upon BMI alone. In addition, reporting  
138 exclusion criteria based upon anthropometrics could help clinicians determine if body  
139 composition analysis is feasible for their patient.

### 140 *Subject positioning*

141 For segmental BIA models, the electrodes are contact points integrated into the standing scale  
142 and handgrips. Subjects are required to stand with legs separated (45 degrees) and hold the

143 handgrips with arms extended (30 degrees) to ensure limb separation while maintaining adequate  
144 skin contact with the electrodes [10]. Utilization of the two-point method to measure impedance  
145 of the lower (i.e. foot-to-foot) or upper body (i.e. hand-to-hand) segments can produce estimation  
146 errors for whole body composition. Four- to eight-point electrode placements are required for  
147 whole body BIA assessment. With this method, individual electrodes are placed directly upon  
148 the skin, permitting measurement in either a standing or supine position.

149 Any skin contact, either between the legs or the arms and torso, results in measurement errors  
150 (up to 18% [33]). For some subjects with obesity, it may not be possible to achieve leg  
151 separation while maintaining foot contact with the electrodes on a narrow standing platform. The  
152 reviewed BIA studies provided limited methodology or descriptions for subject positioning, with  
153 one exception. Frankenfield et al. provided details to achieve limb separation, including  
154 placement of a dry towel to avoid skin-to-skin contact [21]. No study reported on the subjects'  
155 ability to stand or sustain the required body position for the BIA test.

156 For DXA, subjects are required to lie still in a supine position while the scan arm moves across  
157 the subject for the length of the instrument bed. The subject's supine length (height), width and  
158 depth must fit within the DXA scan area limits. Dimension limits of different full body DXA  
159 models are summarized in **Table 2**. In the reviewed DXA studies, scan arm height and supine  
160 body depth were not reported. Just one study measured body depth, with supine  
161 anterior/posterior thickness  $>25$  cm used as exclusion criteria [31]. Although waist  
162 circumference was reported in one study [29], this measure is taken from a standing position, it  
163 could not be substituted for *supine* width or depth. Although the supine length dimension of  
164 DXA models (198 cm) is sufficient to accommodate most North American adults (95<sup>th</sup>  
165 percentile, age 20 years and older for women=173.7 cm, men=188.2 cm [34]), some taller  
166 subjects may still be excluded. Validated techniques for scanning taller subjects (e.g. bent knees)  
167 within normal BMI ranges could be explored for class II/III subjects [35].

168 To assess wider subjects, "reflection positioning" has been used [36]. The subject is positioned  
169 off-center (typically towards the right-side of the scan bed) to include the torso and right arm,  
170 with the lower portion of the left arm positioned outside of the scan area. Based upon the  
171 bilateral symmetry of the human body, the values of the right arm are used to "reflect" the left  
172 arm values. This alternative method was validated by Tataranni et al. (n=183, BMI 17.7-52.8

173 kg/m<sup>2</sup>) with low predictive errors for %FM ( $r^2=.90$  [SEE=4.1%]), FFM ( $r^2=.89$  [SEE=3.72 kg])  
174 and FM ( $r^2=.95$  [SEE=3.57 kg]) [37]. Similar values were recorded for all three measurements  
175 between right and left sides. In the reviewed studies, only one discussed this method. Carver et  
176 al. examined 65 subjects with class III obesity (BMI  $49 \pm 6$  kg/m<sup>2</sup>); 51% required reflection  
177 positioning for whole body composition analysis despite wider scan bed limits with iDXA [25].

178 Rothney et al. used an alternative method for assessment of wider subjects. This study explored  
179 measuring either the left or right half of the body (i.e. half-body scans also called hemi-scans) as  
180 a proxy for a full body measurement by iDXA. The half-body scans of 52 subjects (BMI 30.4 -  
181 41.0 kg/m<sup>2</sup>) were validated against their whole-body scans for within-subject (>97%) and within-  
182 group (> 99.9%) variances for total FM, %FM and LST (all  $r^2 < 0.033$ ). A small variance with  
183 increased bone mass consistent with handedness (+30 g, 1%) was measured [27]. In this study,  
184 half-body scans provided a valid method using DXA to assess subjects that exceed supine width  
185 limits. The maximum BMI in this study was 41 kg/m<sup>2</sup>, only representing the lowest range of  
186 class III obesity. Both studies utilized iDXA, with larger scan bed area and weight capacity,  
187 permitting imaging of subjects with wider, thicker, and heavier body dimensions [25, 27].  
188 Further validation is required of the half-body scan method with class III subjects.

### 189 ***Total body water***

190 Two of the eight BIA studies reviewed reported %TBW. De Freitas et al. compared single-  
191 frequency (50 kHz) BIA (Quantum II, RLJ Systems) for 36 patients before and 6 months after  
192 bariatric surgery. Before surgery, TBW was  $36.1 \pm 4.8$  % (29-48%) with an increase to  $45.0 \pm$   
193  $5.8$  % (36-58%) at 6 months after surgery [20]. Nicoletti et al. used single-frequency (50 kHz)  
194 BIA (101-Q, RLJ Systems) for 43 women before and annually for four years after bariatric  
195 surgery. The %TBW was  $33.1 \pm 3.8$  % before surgery, with an increase to  $48.5 \pm 6.7$  % at one  
196 year and  $46.6 \pm 6.7$  % at year four. Both studies showed a reduced hydration status both before  
197 and after bariatric surgery, with trends for %TBW to increase after bariatric surgery. Studies on  
198 body composition of adults with class II/III obesity without bariatric surgery are needed.

### 199 ***Tissue penetration***

200 For DXA, the x-ray beams must be able to penetrate (attenuate) the body in order to differentiate  
201 the tissues measured. Tissue depth is important; attenuation errors occur when tissue depths

202 exceed 25 cm, resulting in an underestimation of FM. To account for this, some DXA models  
203 can increase scan time (i.e. use “slow” or “thick” mode) to improve attenuation and scan  
204 accuracy. No study reviewed specifically discussed wave frequency or attenuation in context of  
205 their results. For one iDXA study, longer scan modes (13 vs. 7 minutes) were reported to  
206 enhance tissue penetration and reduce measurement errors although the type of errors were not  
207 specified [25]. Due to increased DXA scan time, subjects have a small but increased radiation  
208 exposure and may become too uncomfortable to sustain a still, supine position. This may present  
209 a barrier for assessment in some subjects.

210

### 211 **Comparing BIA to DXA**

212 Three of the reviewed studies completed cross-sectional validations of BIA to DXA  
213 measurements [29-31]. Bedogni et al. compared measures of FM using single-frequency (50 Hz)  
214 BIA to DXA (GE Lunar Prodigy) and utilized an obese-specific regression equation (validated  
215 by Jimenez et al. using iDXA n=159, 79% female) to determine FM from impedance values in  
216 women (n= 57, BMI 37.3- 55.2 kg/m<sup>2</sup>) [29]. The BIA measurements were not reliable (%FM  
217 levels of agreement -4.9% to 8.2%), leading investigators to conclude that BIA, even with an  
218 obese-specific equation, was not interchangeable with DXA. The use of a different BIA device  
219 from the Jimenez’s validated equation can justify the lack of accuracy found in Bedogni’s study.

220

221 The second study by Faria et al. compared FM measurements of 73 subjects (89% female, BMI  
222  $40.17 \pm 4.08$  kg/m<sup>2</sup>) using a multi-frequency BIA (InBody 720) and DXA. Both methods to  
223 measure FM produced an “almost perfect correlation”, however BIA significantly  
224 underestimated FM (-2.05 kg [p<0.0001]) or -1.16% [p<0.0001]) and overestimated FFM (1.28  
225 kg [p=0.0007], or 1.61% [p<0.0001]) compared to DXA. These results, in contrast to the  
226 authors’ conclusions, suggest that BIA was not accurate enough for research or application to  
227 clinical practice in an obese cohort or for individual assessment [30].

228

229 In the third study, Shafer et al. utilized an eight-point, segmental, multi-frequency BIA (Inbody  
230 320) to compare %FM measures to those obtained from DXA (Hologic QDR Delphi-W) in 132  
231 subjects (n= 42 with BMI 30-39 kg/m<sup>2</sup>, class III obesity excluded). In subjects with class I/II  
232 obesity, BIA overestimated %FM ( $3.40 \pm 0.39$ ) with increased error as %FM increased (r=0.424,

233  $P < 0.0001$ ) with limits of agreement ranging from -5.7% to 7.2% FM. This study concluded BIA  
234 was not a reliable tool to measure body composition in an adult cohort with class I/II obesity  
235 [31].

236  
237 In all three studies, BIA results were not consistent with DXA with the rate of error increasing  
238 with higher adiposity, with the maximum BMI studied was  $55.2 \text{ kg/m}^2$ . Although the Bland-  
239 Altman analysis reported for each study demonstrated agreement between BIA and DXA for  
240 some individuals, there was overall high variability and estimation bias, making individual  
241 measurements unreliable [31, 30, 29]. Each study excluded subjects due to equipment weight  
242 limitations (120 kg[30], 130 kg[29], BMI  $< 40 \text{ kg/m}^2$  [31]), restricting the data available from  
243 subjects with class III obesity.

#### 244 **Additional Considerations**

245 A few considerations are highlighted to inform future research and clinical practice.

246 ***Males are underrepresented.*** Male subjects often represent less than 20% in both clinical obesity  
247 practice and research obesity literature. Compared to females, males have more FFM and  
248 potentially are at increased risk of greater FFM loss during weight loss [38]. Further research is  
249 required not only for body composition of men with obesity, but the possible barriers leading to  
250 underrepresentation in treatment and research.

251 ***Data on subjects with higher BMI in Class III obesity is limited or lacking.*** Many studies either  
252 collate results for all class III subjects or exclude subjects with BMI  $> 40 \text{ kg/m}^2$  or who exceed  
253 equipment limits. Our understanding of body composition at higher levels of obesity as a result is  
254 very limited. Unlike other BMI categories with a narrow five-point range, class III obesity has  
255 the widest range, with no upper limit above  $40 \text{ kg/m}^2$ . Stratifying results within class III could  
256 enhance the understanding of body composition within class III and at extremes of BMI.

257 ***The % FFM can increase, despite loss of FFM (kg).*** Reporting of body composition results can  
258 be misleading; for the same subject or group, FFM could be reported as both a loss and a gain.  
259 For example, Ciangura et al. examined the body composition of patients before and after  
260 bariatric surgery. In the first three months post surgery, subjects lost LST mass [a component of  
261 FFM] at a rate of  $-2.3 \pm 1.2 \text{ kg/month}$ , however when reported as a percentage relative to FM,

262 %FFM increased by 2.8% [26]. This study demonstrated post-surgical subjects lost FFM at a  
263 specific rate and the time frame. However, it could be misinterpreted that subjects *increased* lean  
264 tissue after surgery because %FFM increased. Subjects who are actually losing FFM mass may  
265 not be identified as at risk, impacting assessment and treatment decisions. The reported  
266 preservation or increase in %FFM after weight loss is confounded by a possible elevation in  
267 TBW, contributing to a *relative* change compared to %FM [21]. The rate of error was  
268 proportional to body weight, increasing at higher body weights. When examining the outcome  
269 data for body composition during weight loss, the absolute changes in FFM independent of FM  
270 and BMI (e.g. appendicular skeletal muscle by height [m<sup>2</sup>], from DXA) may better marker of  
271 FFM changes.[8]

## 272 **Conclusions**

273 Although extensive research and reviews on body composition are reported in the literature,  
274 relatively few studies using BIA and/or DXA including subjects with class II/III obesity were  
275 identified. In general, both BIA and DXA can provide relatively safe, quick and non-invasive  
276 measures of body composition.

277 It is easy to understand the interest of clinicians in BIA; it is inexpensive, portable, low risk, able  
278 to accommodate people with larger body dimensions and requires minimal training or expertise.  
279 Anthropometric measures and BMI are important but have limited value for body composition.  
280 BIA can estimate adiposity better than BMI when BMI < 35 kg/m<sup>2</sup>, but there are methodological  
281 problems for subjects with class II/III obesity limiting the reliability for body composition.

282 DXA can provide accurate and reliable measures of body composition, yet equipment-related  
283 barriers have limited assessment of heavier, taller and wider subjects. As demonstrated with  
284 iDXA and half-body scans, advancements with equipment, technology and methodology permit  
285 assessment of more people with class II/III obesity. Accurate and reliable assessments of body  
286 composition in this cohort are important to help determine health risk in more adults with class  
287 III obesity and contribute to understanding of the longer-term effects of this disease and  
288 treatment. Further studies are needed to measure body composition with DXA at initiation and at  
289 several points during interventions to support individualized obesity treatment, risk reduction and  
290 outcome optimization. Longitudinal studies of body composition across interventions and phases  
291 of treatment (loss, maintenance, gain/regain) should also be considered, to optimize patient care.

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294 **Compliance with Ethics Guidelines**

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**Table 1.** Summary of errors associated with assessing body composition using bioelectrical impedance analysis (BIA) in subjects with class II/III obesity.

	<b>Fat Mass</b>	<b>Fat-Free Mass</b>
Increased TBW (>73.2%)	Underestimated	Overestimated
Increased ECW	Underestimated	Overestimated
Use of normal-weight regression prediction formulas	Underestimated	Overestimated
Use of two vs. eight electrodes	Underestimated	Overestimated
Use of single (50 kHz) vs. multi-frequency waves	Underestimated	Overestimated

TBW: total body water; ECW: extracellular water; kHz: kilohertz

**Table 2.** Scan area dimensions and subject weight capacity of full body dual-energy X-ray absorptiometers (DXA) from two manufacturers.

Scan area	Hologic, Inc[39-41]		GE Healthcare[42]	
			DXP, Prodigy	Lunar iDXA
Length, cm	All models	195	DXP 195 Prodigy 197.7	197.7
Width, cm	All models	65	60	66
Weight capacity, kg	Delphi, QDR, Explorer:	136	DXP: 136	204
	Discovery A,W:		Prodigy: 160	
	Prior to 03/05	159		
	03/05 to 04-07	182		
	After 04/07	204		
	Horizon:	204		