Assessment of body composition in indian adults: Comparison between dual-energy X-ray absorptiometry and isotope dilution technique

Bharati Kulkarni, Queensland University of Technology
Hannah Kuper
Amy Taylor, University of Bristol
Jonathan Wells
Radhakrishna, Radhakrishna, et al.

Available at: https://works.bepress.com/nuala_byrne/27/
Assessment of body composition in Indian adults: comparison between dual-energy X-ray absorptiometry and isotope dilution technique

Bharati Kulkarni1,2*, Hannah Kuper3, Amy Taylor4, Jonathan C. Wells5, K. V. Radhakrishna2, Sanjay Kinra2, Yoav Ben-Shlomo4, George Davey Smith4, Shah Ebrahim3,6, A. V. Kurpad7, Nuala M. Byrne1 and Andrew P. Hills8

1School of Exercise and Nutrition Sciences and Institute of Health and Biomedical Innovation, Queensland University of Technology, Brisbane, QLD, Australia
2Clinical Division, National Institute of Nutrition, Jamai Osmania PO, Hyderabad 500 007, India
3Department of Non-communicable Disease Epidemiology, London School of Hygiene and Tropical Medicine, London, UK
4School of Social and Community Medicine, University of Bristol, Bristol, UK
5Childhood Nutrition Research Centre, UCL Institute of Child Health, London, UK
6South Asia Network for Chronic Disease, Public Health Foundation of India, New Delbi, India
7Saint John’s Research Institute, Bangalore, India
8Mater Mothers’ Hospital, Mater Research Institute – University of Queensland and Griffith Health Institute, Griffith University, Brisbane, QLD, Australia

(Submitted 28 September 2013 – Final revision received 30 April 2014 – Accepted 22 May 2014 – First published online 11 August 2014)

Abstract
Dual-energy X-ray absorptiometry (DXA) and isotope dilution technique have been used as reference methods to validate the estimates of body composition by simple field techniques; however, very few studies have compared these two methods. We compared the estimates of body composition by DXA and isotope dilution (18O) technique in apparently healthy Indian men and women (aged 19–70 years, n 152, 48% men) with a wide range of BMI (14–40 kg/m2). Isotopic enrichment was assessed by isotope ratio mass spectroscopy. The agreement between the estimates of body composition measured by the two techniques was assessed by the Bland–Altman method. The mean age and BMI were 37 (SD 15) years and 23·3 (SD 5·1) kg/m2, respectively, for men and 37 (SD 14) years and 24·1 (SD 5·8) kg/m2, respectively, for women. The estimates of fat-free mass were higher by about 7 (95 % CI 6, 9) %, those of fat mass were lower by about 21 (95 % CI 23, 18) %, and those of body fat percentage (BF%) were lower by about 7·4 (95 % CI 6·6, 8·2) % as obtained by DXA compared with the isotope dilution technique. The Bland–Altman analysis showed wide limits of agreement that indicated poor agreement between the methods. The bias in the estimates of BF% was higher at the lower values of BF%. Thus, the two commonly used reference methods showed substantial differences in the estimates of body composition with wide limits of agreement. As the estimates of body composition are method-dependent, the two methods cannot be used interchangeably.

Key words: Body composition: Dual-energy X-ray absorptiometry: Isotope dilution technique: Indian adults: Bland–Altman analysis

Estimation of body composition is a vital element of nutritional assessment as fat and fat-free compartments of body mass have different health implications. Fat mass (FM) is closely linked with metabolic complications of obesity because the adipose tissue functions as an endocrine organ that releases bioactive substances having pro-inflammatory properties(11). In contrast, fat-free mass (FFM), especially muscle mass, plays a protective role against the risk of chronic diseases including diabetes and osteoporosis(12). Ethnic differences in the relationship between BMI and disease risk have been associated with differences in body composition(2,3,5,13).

A number of techniques are available for the assessment of body composition, and the choice of technique usually depends on precision, accuracy, ease of application as well as the cost. DXA is increasingly used for body composition assessment because of its high precision and low dose of radiation. A number of studies have validated other, less precise, techniques such as anthropometry and bioelectrical

Abbreviations: 4C, four-compartment model; APCAPS, Andhra Pradesh Children and Parents Study; FFM, fat-free mass; FM, fat mass; IMS, Indian Migration Study.

* Corresponding author: B. Kulkarni, fax +91 40 27019074, email dr.bharatikulkarni@gmail.com
Impedance analysis against DXA as a reference method\(^5\)\(^--\)\(^7\). However, DXA is not without limitations. Although studies have shown that estimates of body composition by DXA are highly correlated with those derived using more accurate methods, variations have been reported between the estimates\(^8\)\(^,\)\(^9\).

With increasing recognition of the association between the high prevalence of the metabolic syndrome and ‘thin-fat’ phenotype in Indians, there is enhanced interest in the assessment of body composition\(^10\)\(^,\)\(^11\). A number of studies in India have reported the body composition of different population groups using different techniques including DXA\(^12\)\(^--\)\(^15\). However, different studies that have compared the estimates of body composition using different methods of body composition measurement need to consider the variation in estimates associated with these methods. Moreover, studies comparing different methods of body composition measurement tend to be population-specific due to ethnic variations in body composition\(^16\). Studies comparing the estimates of body composition using DXA with those measured by other reference methods have not so far been reported in India. Therefore, the aim of the present study was to compare the estimates of body composition by DXA with those using the isotope dilution technique.

Participants and methods

The present study was conducted according to the guidelines laid down in the Declaration of Helsinki, and all procedures involving human participants were approved by the ethics committees of the National Institute of Nutrition, Hyderabad, India, the London School of Hygiene & Tropical Medicine, UK and Queensland University of Technology, Australia. Written informed consent was obtained from all participants.

Participants

Healthy volunteers aged 19–70 years were enrolled in the present study from two pre-established cohorts (Andhra Pradesh Children and Parents Study (APCAPS), n 58 and Indian Migration Study (IMS), n 94) living around the city of Hyderabad, India. The APCAPS cohort was established to assess the long-term impact of early nutrition supplementation provided to pregnant women and young children\(^17\), whereas the IMS cohort was established to examine the association between rural to urban migration and cardio-metabolic risk\(^18\). To obtain a representative sample, participants were chosen on the basis of pre-defined age, sex, cohort, intervention group (in the case of the APCAPS) or rural/urban migrants (in the case of the IMS), and BMI categories (see online supplementary Tables S1 and S2). The target enrolment was 160 participants.

Demographic and anthropometric data

Demographic information was collected from all study participants using an interviewer-administered questionnaire. Weight was measured to the nearest 0·1 kg in light clothing without footwear, using a digital Seca scale (www.seca.com). Standing height was measured to the nearest 1 mm using a portable stadiometer (Leicester Height Measure; Chasmors Limited). Anthropometric measurements were taken twice, and the average of the two values for each measurement was used in the analysis. BMI was calculated as weight (kg)/height (m\(^2\)).

Body composition of each participant was assessed by DXA and isotope dilution technique on the same day.

Isotope dilution technique

Participants arrived at the National Institute of Nutrition in the morning after an overnight fast. A baseline urine sample was collected on arrival for the measurement of background isotopic enrichment, followed by the administration of an oral dose of \(^18\)O \((0·2 \text{ g/kg body weight})\) to each participant at about 09.00 hours. The bottle containing the dose was rinsed with 50 ml deionised water before its consumption by the participants. A light breakfast was provided with 50 ml water at about 10.00 hours. Any subsequent oral intake was avoided. Follow-up urine samples were collected 4 and 5 h after the intake of dose to allow complete equilibration of the isotope within the body water compartments\(^19\). Aliquots of all the urine samples were stored in screw-capped glass containers at −20°C until analysis. Isotopic enrichment in the pre- and post-dose urine samples, the dose given and the local tap water was measured using isotope ratio mass spectrometry (Hydra 20-20; SerCon) at St John’s Research Institute, Bangalore, India. The CV calculated using repeated analysis for the natural background samples as well as for the enriched samples was less than 0·01%. Each sample was analysed in duplicate, and the mean was used for the analysis. Total body water was calculated, allowing a correction by 0·7% for \textit{in vivo} exchange\(^20\). FFM was calculated from total body water using a hydration constant of 0·73. FM was calculated by subtracting FFM from body weight.

Dual-energy X-ray absorptiometry scans

Body composition was assessed by a whole-body DXA scan using a fan-beam DXA machine (Hologic Discovery A model, software version 12.5; www.hologic.com). The scanner was calibrated periodically with a phantom, and its performance was monitored according to the manufacturer’s quality assurance protocol. During the scan, the participants were asked to lie supine on the scanning bed with their arms at their sides. Standard software options were used to calculate the total FFM and FM. FFM was the sum of lean soft tissue mass and bone mineral content. Precision estimates (CV%) of body composition by DXA based on repeat measurements in thirty participants were 0·7 and 1·4% for FFM and FM, respectively.

Statistical analyses

All analyses were conducted using Stata (version 11.2; StataCorp). As FFM and FM showed a skewed distribution, these variables were log-transformed before analysis, and, therefore,
the mean differences between the two are expressed as ratios. Other continuous variables were used in the original scale. Differences between the body composition estimates (FFM, FM and BF%) by DXA and isotope dilution technique were assessed using paired $t$ tests. The Bland–Altman method was used to assess the agreement between the estimates of body composition determined by the two techniques\(^\text{(21)}\).

The mean difference in the estimates by the two techniques (bias) and their 95% limits of agreement (2 SD of the mean difference) were calculated. As the bias and limits of agreement for FFM and FM were on a logarithmic scale, these values are presented as ratios. Correlation coefficients were calculated to examine the association between the average values of body composition measurements by the two methods and the difference between these methods, which indicates the proportional bias. All analyses were conducted for the whole sample and additionally stratified by sex.

**Results**

A total of seventy-three men and seventy-nine women participated in the study. Their characteristics are presented in Table 1. The participants were chosen to represent a wide range of BMI varying from 13·8 to 39·7 kg/m\(^2\). The total mass value measured by DXA showed a strong correlation with weight measured by the scale ($0·99$, $P<0·01$). Although there was a strong correlation between the estimates of body composition measured by DXA and isotope dilution technique (FFM: $r=0·95$, FM: $r=0·95$, BF%: $r=0·89$ all $P<0·01$), the estimates of FFM obtained by DXA were higher than those obtained by the isotope dilution technique in the whole sample as well as in the subgroups stratified by sex (Table 2). The estimates of FM and BF% obtained by DXA were lower than those measured by the isotope dilution technique. On average, DXA overestimated the FFM values by 7·4 (95% CI $6·2$, $8·6$%) % (Table 3; Fig. 1(c)). The bias in the estimates of BF% was negatively correlated with the average values of BF%, indicating that the difference between the two methods was higher for the participants with lower values of BF% (Table 3). The estimates of FFM, FM and BF% measured by DXA explained about 89, 85 and 78% of the variation in the respective estimates measured by the isotope dilution technique.

**Discussion**

The present study compared the estimates of body composition measured by two precise techniques – DXA and isotope dilution technique – in apparently healthy, weight-stable Indian men and women with a wide range of BMI. In this sample of participants, the estimates of FFM were higher whereas those of FM and BF% were lower using DXA than using the isotope dilution technique. The agreement between the two methods was not good as indicated by the significant bias between these methods and wide limits of agreement, especially for the estimates of FM and BF%. The bias in the estimates of estimates measured by the two methods for both FFM and FM, indicating that the bias in the estimates of FFM and FM did not change with the amount of FFM and FM, respectively. On average, the estimates of BF% measured by DXA were lower than those measured by the isotope dilution technique by 7·4 (95% CI $8·2$, $6·6$%) % (Table 3; Fig. 1(c)). The bias in the estimates of BF% was negatively correlated with the average values of BF%, indicating that the difference between the two methods was higher for the participants with lower values of BF% (Table 3). The estimates of FFM, FM and BF% measured by DXA explained about 89, 85 and 78% of the variation in the respective estimates measured by the isotope dilution technique.

**Table 1. Characteristics of the study participants**

<table>
<thead>
<tr>
<th></th>
<th>Men (n 73)</th>
<th>Women (n 79)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Mean</strong></td>
<td><strong>sd</strong></td>
<td><strong>Minimum</strong></td>
</tr>
<tr>
<td><strong>Age (years)</strong></td>
<td>37</td>
<td>15</td>
</tr>
<tr>
<td><strong>BMI (kg/m(^2))</strong></td>
<td>23·3</td>
<td>5·1</td>
</tr>
<tr>
<td><strong>Height (cm)</strong></td>
<td>165·5</td>
<td>6·3</td>
</tr>
<tr>
<td><strong>Weight (kg)</strong></td>
<td>64·1</td>
<td>15·1</td>
</tr>
<tr>
<td><strong>TM by DXA (kg)</strong></td>
<td>64·0</td>
<td>15·0</td>
</tr>
</tbody>
</table>

**Table 2. Estimates of body composition by dual-energy X-ray absorptiometry (DXA) and isotope dilution technique**

<table>
<thead>
<tr>
<th></th>
<th>Isotope dilution technique</th>
<th>DXA</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>n</strong></td>
<td>Mean</td>
<td>sd</td>
</tr>
<tr>
<td><strong>Fat-free mass (kg)</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Whole sample</td>
<td>152</td>
<td>37·42</td>
</tr>
<tr>
<td>Men</td>
<td>73</td>
<td>44·18</td>
</tr>
<tr>
<td>Women</td>
<td>79</td>
<td>31·17</td>
</tr>
<tr>
<td><strong>Fat mass (kg)</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Whole sample</td>
<td>152</td>
<td>22·27</td>
</tr>
<tr>
<td>Men</td>
<td>73</td>
<td>19·93</td>
</tr>
<tr>
<td>Women</td>
<td>79</td>
<td>24·43</td>
</tr>
<tr>
<td><strong>Body fat percentage</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Whole sample</td>
<td>152</td>
<td>36·3</td>
</tr>
<tr>
<td>Men</td>
<td>73</td>
<td>29·8</td>
</tr>
<tr>
<td>Women</td>
<td>79</td>
<td>42·3</td>
</tr>
</tbody>
</table>

* $P$ value was obtained from the paired $t$ test of the difference.
BF% measured by the two methods was higher for individuals with lower values of BF%. The present study indicates that these two methods cannot be used interchangeably as systematic differences exist between the estimates of body composition.

Previous studies that have compared the estimates of body composition by DXA and isotope dilution technique have reported inconsistent results. In general, studies that used older equipment (e.g. Hologic QDR 2000, Hologic QDR 1000W, Lunar DPX-L) with scans done in a pencil-beam mode have shown underestimation of FFM and overestimation of FM and BF% by DXA compared with the isotope dilution technique(22,23). In contrast, studies that used newer equipment (e.g. Hologic QDR 4500W, QDR 4500A) have shown overestimation of FFM by DXA compared with the isotope dilution technique24,25. For example, a study in Chinese women in 1999 has shown that DXA (Hologic QDR 2000) underestimated FFM by 0.5 kg and overestimated BF% by 0.8%25. Similarly, a study from the UK (n = 28) in 1992 has also found that DXA (Hologic QDR 1000W) underestimated FFM by 0.2 kg compared with the isotope dilution technique22. However, a later study on Chinese, Malays and Indians living in Singapore has shown overestimation of FM and underestimation of BF% by DXA (Hologic QDR 4500W) compared with the 4H dilution technique. Similarly, a study by Schoeller et al.26 from the USA that compared body composition by DXA with other reference techniques in 1195 men and women (DXA compared with the isotope dilution technique in 395 participants) has also shown that DXA underestimated FFM by 1.8 to 4.7 kg and underestimated BM by about 1.3 to 5.1 kg. The findings of the present study that used a newer model of DXA (Hologic Discovery) are consistent with relatively recent studies that have shown overestimation of FFM by DXA compared with the isotope dilution technique. However, the magnitude of bias in the estimates of FFM (approximately 3 kg) and FM (approximately 4.5 kg) in the present study is larger than the bias reported in other studies.

A number of studies (Table 4) comparing the estimates of body composition by DXA with those by multi-component criterion methods have also reported inconsistent results22,25–29. Although the majority of these studies reported underestimation of BF% by DXA, similar to the present study, a few studies have reported a bias in the opposite direction. For example, a study by Williams et al.26 compared DXA with a four-compartment (4C) model and reported the overestimation of FM and BF% by DXA in non-obese adults. In contrast, a few studies did not detect significant differences in BF% by DXA compared with the 4C model31–33.

Differences in the results of studies comparing the estimates of body composition by DXA with those by other techniques could be related to a number of factors such as the use of DXA machines by different manufacturers and differences in the scan mode or software used for analyses. Machines developed by different manufacturers as well as different models by the same manufacturer, although based on the same physical principles, differ in various aspects such as the generation of high- and low-energy X-ray beams, X-ray detectors, calibration methodology, algorithms used for selective tissue separation, etc.54. Variations in the estimates of body composition with the machines developed by different manufacturers even with different models by the same manufacturer have been reported55–58. In addition, isotope dilution technique has a number of limitations as the estimates of body composition are based on a number of assumptions including the equal distribution of a tracer in body water and constant hydration of FFM.59 Both these techniques are thus error prone, and a lack of agreement between the methods for the estimation of body composition could be related to a number of factors that can lead to inaccuracies in the estimates.

However, limits of agreement between the two methods observed in the present study were wider (FFM: −9, +26%; FM: −46, +17%; BF%: −17.3, 2.6%) than those reported by

### Table 3. Bias and 95% limits of agreement for measures of body composition by dual-energy X-ray absorptiometry (DXA) compared with the isotope dilution technique

<table>
<thead>
<tr>
<th></th>
<th>n</th>
<th>Bias*</th>
<th>95% CI</th>
<th>Limits of agreement†</th>
<th>r‡</th>
<th>P§</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Fat-free mass (kg)</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Whole sample</td>
<td>152</td>
<td>1.07</td>
<td>1.06, 1.09</td>
<td>0.91, 1.26</td>
<td>−0.077</td>
<td>0.35</td>
</tr>
<tr>
<td>Men</td>
<td>73</td>
<td>1.06</td>
<td>1.04, 1.08</td>
<td>0.92, 1.23</td>
<td>−0.127</td>
<td>0.28</td>
</tr>
<tr>
<td>Women</td>
<td>79</td>
<td>1.08</td>
<td>1.06, 1.10</td>
<td>0.91, 1.29</td>
<td>0.083</td>
<td>0.47</td>
</tr>
<tr>
<td><strong>Fat mass (kg)</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Whole sample</td>
<td>152</td>
<td>0.79</td>
<td>0.77, 0.82</td>
<td>0.54, 1.17</td>
<td>0.045</td>
<td>0.58</td>
</tr>
<tr>
<td>Men</td>
<td>73</td>
<td>0.75</td>
<td>0.71, 0.79</td>
<td>0.48, 1.17</td>
<td>0.043</td>
<td>0.71</td>
</tr>
<tr>
<td>Women</td>
<td>79</td>
<td>0.84</td>
<td>0.81, 0.86</td>
<td>0.63, 1.12</td>
<td>−0.181</td>
<td>0.11</td>
</tr>
<tr>
<td><strong>Body fat percentage</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Whole sample</td>
<td>152</td>
<td>−7.4</td>
<td>−8.2, −6.6</td>
<td>−17.3, 2.6</td>
<td>−0.345</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Men</td>
<td>73</td>
<td>−7.5</td>
<td>−8.7, −6.3</td>
<td>−17.7, 2.8</td>
<td>−0.428</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Women</td>
<td>79</td>
<td>−7.3</td>
<td>−8.4, −6.2</td>
<td>−17.0, 2.4</td>
<td>−0.513</td>
<td>0.03</td>
</tr>
</tbody>
</table>

*Mean bias and 95% CI for fat-free mass and fat mass are expressed as the ratio of DXA:isotope dilution technique values. Bias is the difference (DXA – isotope dilution) between the log-transformed values of fat-free mass and fat mass estimated from the two techniques. The values of body fat percentage are given on the original scale.
†95% Limits of agreement (2 std of the mean difference) expressed as the ratio of DXA:isotope dilution technique values of fat-free mass and fat mass. The values of body fat percentage are given on the original scale.
‡r is Pearson’s correlation coefficient between the difference between DXA and isotope dilution technique and the average of DXA and isotope measures of fat-free mass, fat mass and body fat.
§Significance of the correlation coefficient.
other studies, the majority of which have reported the limits of agreement between ±10% of the mean (40). In contrast, a few other studies have reported that DXA could under- or overestimate the FM of an individual by almost 28% (22). One of the reasons for the narrow limits of agreement reported by other studies could be the exclusion of extreme values of the differences. For example, Schoeller et al. (9) excluded observations in which the difference in the estimates of FFM measured by DXA and isotope dilution technique was >6 kg. The present study did not exclude observations with larger differences between the measurements, which may have contributed to a larger bias between the measurements reported herein.

An interesting finding of the present study is that the bias in the estimates of BF% by the two methods was higher at lower values of BF% (r = 0.345, P < 0.001; Table 3). A previous study comparing the estimates of abdominal fat by DXA with those using MRI in this sample has also shown that overestimation of abdominal fat by DXA was greater in individuals with less abdominal fat (41). It is possible that the algorithms used for the estimation of body composition by DXA produce a larger error at very low levels of body fat. A number of studies from other centres have shown that the bias in the estimates of body composition by DXA varied according to a number of factors including age, body size, body fat, sex, health status, type of the instrument, etc. (30).

An important strength of the present study includes enrolment of a large sample representing a broad range of age and BMI. In addition, the present study used 18O as the isotope tracer that may provide a more accurate estimate of total body water than a more commonly used 2H2O as 18O exchanges to a smaller degree with non-aqueous molecules (39). As differences in body composition in relation to ethnicity are well known, population-specific validation studies comparing DXA with other precise methods are required. Therefore, the present study provides much-needed evidence on the comparability of DXA with the isotope dilution technique in a population group that is known to have a high percentage of body fat at a given BMI compared with other ethnic groups (10, 15). A limitation of the present study is the use of the isotope dilution technique for validating DXA measurements of body composition instead of a multi-component criterion technique. However, estimates of body composition using the isotope dilution technique are highly correlated with those using the criterion technique of the 4C model (27). A study comparing the estimates of body composition by densitometry, DXA and isotope dilution technique with those by the 4C model in Asian adults has shown that estimates of BF% by the isotope dilution technique had the lowest bias while those by DXA had the highest bias compared with the 4C model, suggesting that the isotope dilution technique may be the best two-compartment model for measuring body fat (24).

In conclusion, the present study shows that estimates of body composition by two commonly used reference methods such as DXA and isotope dilution technique may be considerably different at the individual level, with particularly larger differences in the estimates of BF%. The two methods are therefore not directly interchangeable. However, the
The present study was funded by the Wellcome Trust, UK (grant no. WT083707AIA). The study sponsor had no role in the study design, in the collection, analysis and interpretation of the data, in the writing of the report, or in the decision to submit the paper for publication.

The authors thank Dr B. Sesikeran, Director, National Institute of Nutrition, Hyderabad for providing the facilities and support system to carry out the study. They thank Mrs K. Usha Rani for conducting and analysing the DXA scans and Pete Shiarly for data management. The authors also thank Ms Sarita, St Johns Research Institute, Bangalore, India for carrying out the isotope analyses. The authors are extremely grateful to the committed and diligent fieldwork team led by Ms Santhi Bhogadi.

The authors' contributions are as follows: B. K., H. K. and A. V. K. conceived and carried out the study; J. C. W. conceived the study and helped in the interpretation of the results; B. K. analysed the data and wrote the manuscript. All authors were involved in the interpretation of the results and writing of the manuscript, and approved the final draft of the manuscript.

The authors declare that they have no competing interests.

Supplementary material

To view supplementary material for this article, please visit http://dx.doi.org/10.1017/S0007114514001718

Acknowledgements

Table 4. Comparison of body fat percentage (BF%) measured by dual-energy X-ray absorptiometry (DXA) and the four-compartment model (4C) (Mean values and standard deviations)

<table>
<thead>
<tr>
<th>Study†</th>
<th>Sex</th>
<th>Mean</th>
<th>SD</th>
<th>DXA system</th>
<th>Type of the X-ray beam</th>
<th>Mean difference in BF% (4C – DXA)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bergsma-Kadijk et al.‡</td>
<td>20 F</td>
<td>22</td>
<td>2</td>
<td>GE Lunar DPX</td>
<td>Pencil</td>
<td>3.1*</td>
</tr>
<tr>
<td>Prior et al.§</td>
<td>18 F</td>
<td>72</td>
<td>4</td>
<td>Hologic QDR</td>
<td>Pencil</td>
<td>5.3*</td>
</tr>
<tr>
<td>Withers et al.¶</td>
<td>81 F</td>
<td>21</td>
<td>2</td>
<td>GE Lunar DPX-L</td>
<td>Pencil</td>
<td>0.8</td>
</tr>
<tr>
<td>Arngrimsson et al.¶¶</td>
<td>22 F</td>
<td>21</td>
<td>2</td>
<td>GE Lunar 1000W</td>
<td>Pencil</td>
<td>0.9</td>
</tr>
<tr>
<td>Deurenberg-Yap et al.¶¶¶</td>
<td>144 M</td>
<td>42</td>
<td>13</td>
<td>Hologic 4500</td>
<td>Fan</td>
<td>2.8%*</td>
</tr>
<tr>
<td>Van Der Ploeg et al.¶¶¶¶</td>
<td>118 M</td>
<td>31</td>
<td>12</td>
<td>GE Lunar DPX-L</td>
<td>Pencil</td>
<td>1.9%*</td>
</tr>
<tr>
<td>van Marken Lichtenbelt et al.¶¶¶¶¶</td>
<td>34 F</td>
<td>26</td>
<td>8</td>
<td>GE Lunar DPX-L</td>
<td>Pencil</td>
<td>1.7%*</td>
</tr>
<tr>
<td>Williams et al.¶¶¶¶¶¶</td>
<td>27 M</td>
<td>32</td>
<td>6</td>
<td>GE Lunar</td>
<td>Narrow</td>
<td>–0.8</td>
</tr>
<tr>
<td>LaForgia et al.¶¶¶¶¶¶¶</td>
<td>8 M</td>
<td>36</td>
<td>13</td>
<td>GE Lunar Prodigy</td>
<td>Narrow</td>
<td>0.3%</td>
</tr>
</tbody>
</table>

F, female; M, male.
* Statistically significant difference between DXA and the 4C model (P<0.05).
† Includes weight-stable, healthy adults.
‡ Includes competitive and non-competitive athletes.
§ Includes non-athletes.
¶ Includes athletes.
¶¶ Includes non-athletes.
¶¶¶ Includes athletes.
¶¶¶ Includes non-athletes.
¶¶¶¶ Includes athletes.
¶¶¶¶ Includes non-athletes.
¶¶¶¶¶ Includes athletes.
¶¶¶¶¶¶ Includes non-athletes.
¶¶¶¶¶¶¶ Includes athletes.

References

Body composition measures by reference methods

1153


