

TOTAL BODY FAT ESTIMATIONS FROM SUBCUTANEOUS ADIPOSE TISSUE

IN SARCOPENIC AND NON-SARCOPENIC WOMEN: A FOUR-COMPARTMENT MODEL APPROACH



Agreement

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ABSTRA CT

Having accurate body composition measurements is crucial for aging men and women. After 65 years of age significant changes can occur in both fat and muscle mass leading to muscle loss, increased fat storage and even sarcopenia. However, the ability to estimate total body fat (%Fat) using simple subcutaneous adipose tissue (SAT) measurements is still unclear. PUR-POSE: Generate and validate %Fat equations using SAT measurements. METHODS: Using a double-cross validation model, 104 women were separated in two groups maintaining similar sarcopenia numbers as well as age range (Group A: Age 73 +/- 6, 61.95 +/- 6.38 kg, 160.5 +/- 5.5 cm, 40.3 +/- 6.0 %Fat, 61% Sarcopenic; Group B: Age 71 +/- 5, 68.17 +/- 12.88 kg, 163 +/- 6 cm, 40.4 +/- 6.3 %Fat, 64% Sarcopenic). Subjects had their body fat measured using a four-compartment model (4C) including body volume via the BodPod, total body water via deuterium dilution, and bone mineral via dual-energy X-ray absorptiometry (DXA). SAT was measured using a B-Mode ultrasound (US). RESULTS: Using all seven US sites (triceps, biceps, abdominal, suprailiac, quadriceps, hamstrings, and calf) resulted in an r value of 0.72, SEE of 4.25 %Fat and a total error of 4.45 %Fat. Using the three most correlated sites (triceps, hamstrings, and suprailiac) improved prediction errors compared to all sites by producing an r value of 0.73, SEE of 4.18 %Fat and a total error (TE) of 4.36 %Fat. Compared to the 4C model, DXA produced an r value of 0.94, an SEE of 2.11 %Fat and a total error of 2.56 %Fat. **CONCLUSIONS**: SAT measurements in elderly women can be used to predict total body fat. However, due to the large TE and SEE values in comparison to DXA, US does not appear to be a substitution for sophisticated lab methods. Still, US can be used with advanced lab methods to quantify both total body %Fat as well as SAT.

INTRODUCTION

bility of accidental falls in older adults, leading to potential hip fractures and other injuries (1,2). Along with a six-fold increase in government health care costs for the aged by 2040, hip fracture costs alone are projected to be six billion dollars in the year 2040 (3). More importantly, 20% of those with hip fractures will not be able to walk (4), and the average individual at 80 years old lacks the muscle capacity to rise unassisted from a chair (5). These muscular and injury-related limitations not only increase health-related costs but detrimentally affect quality of life, as well as the ability to perform ADLs (6). Therefore, due to the direct association between sarcopenia-related injuries and the subsequent effects on ADLs and quality of life, there exists a need for methods that identify the early onset of sarcopenia. Sarcopenia can be caused by both a decrease in muscle mass and an increase in fat. However, the most accurate methods of measuring body composition are large, bulky devices that require a subject to visit a laboratory. A portable device that can accurately measure subcutaneous fat would be useful for diagnosing sarcopenia in the elderly in their own homes. Though equations have been validated for younger populations, a standard set of equations has not yet been determined for older individuals. Therefore, the purpose of this study is to validate new equations for using ultrasound to calculate %Fat in the elderly.

Changes in body composition can lead to sarcopenia (muscle loss), which increases the possi-

EXPERIMENTAL DESIGN

All body composition assessments were performed on the same day following a twelvehour fast (ad libitum water intake was allowed up to one hour prior to testing). Participants were instructed to avoid exercise for at least twenty-four hours prior to testing. Hydration status was determined using specific gravity via handheld refractometry (Model CLX-1, precision = 0.001 +/- 0.001, VEE GEE Scientific, Inc. Kirkland, Washington) prior to all body composition measurements. Specific gravity values indicated all subjects were properly hydrated during both pre- and post-testing sessions (>1.005, <1.030) (7,8).

One Hundred and four healthy older (>65 yrs), participated in the investigation. All subjects were Caucasians without medical implants or prostheses, considered non-obese (< 30 BMI), and without pacemakers.

METHODS

Sarcopenic Classification

SMI = total body muscle mass from single frequency bioimpedance: (TBMM / BM x100) (6,9)

TBMM = $[(HT^2/R) \times 0.401 + Sex \times 3.825 + Age \times -0.071] + 5.102$

Class I Class II SMI: 37 - 31% men SMI: > 31% men SMI: 28 - 22% women SMI: > 22% women

TBMM = Total body muscle mass, BM = Body mass in kg, HT = Height in cm, R = Bioimpedance Resistance, Sex: Men = 1, Women = 2.

Bioelectrical Impedance Analysis (BIA)

Bioelectrical impedance analysis (BIA) was used to calculate total body muscle mass, as mentioned above (ImpTM DF50, ImpediMed Limited, Queensland, Australia). After resting in a supine position for 5 to 10 minutes, resistance and reactance were measured using a single 50kH frequency while the subjects lay supine on a table with their arms ≥ 30 degrees away from their torso with their legs separated. Each pair of total body electrodes was connected by a non-conductive strip allowing for a distance of 5 cm between electrode centers. After hair removal and cleaning with alcohol, whole body electrodes were placed on the right side of the body. Electrodes were placed at the wrist (dorsal surface at the ulnar styloid process) and ankle (dorsal surface between the malleoli) with the connection strip and connected electrode 5 cm distal from the wrist and ankle. Skeletal muscle index (SMI) was calculated using the equation reported by Janssen et al. (6,9) and used to classify sarcopenia. Previous test retest scans of 11 men and women measured 24-48 hours apart resulted in an SEM = 8.91, ICC = 0.99 for Resistance.





Ultrasound

Ultrasound was used at 3 (biceps, chest and quadriceps) or 7 (triceps, biceps, abdominal, suprailiac, quadriceps, hamstrings, and calf) sites to measure %fat (IntelaMetrix BX-2000, IntelaMetrix Inc., Livermore, CA). A thin layer of water-soluble gel was applied to the transducer, which was then applied to the skin site perpendicular to the skin. Without compressing the tissue, the transducer is slid back and forth along the skin so that the signal may be averaged by the device. The measurements are then included in the new equations listed below for both groups A and B.



New Equations for Ultrasound

New Equations From Group A	New Equations From Group B				
SUM 3 DXA	SUM 3 DXA				
%Fat = 0.349267(Age)+0.325488(kg)+0.469910(SUM3)- 0.076180(cm)-20.977768	%Fat = 0.196538(kg)+0.484680(SUM3)-0.210310(cm) -0.051242(Age)+42.522563				
SUM 3 4C	SUM 3 4C				
%Fat = 0.447745(Age)+0.287871(kg)+0.404046(SUM3)- 0.141163(cm)-10.040965	%Fat = 0.153529(kg)+0.562713(SUM3)-0.150083(cm) +0.154896(Age)+20.013941				
SUM 7 DXA	SUM 7 DXA				
%Fat = 0.418404(Age)+0.385702(kg)+0.195575(SUM7)- 0.115549(cm)-26.220273	%Fat = 0.318056(kg)+0.171981(SUM7)-0.381396(cm) -0.098197(Age)+65.103167				
SUM 7 4C	SUM 7 4C				
%Fat = 0.489505(Age)+0.348660(kg)+0.128872(SUM7)- 0.173685(cm)-11.621047	%Fat = 0.308414(kg)+0.174599(SUM7)-0.343102(cm) +0.118744(Age)+44.467232				

Four-Compartment Model

Fat mass (FM, kg) was estimated using the 4C model described by (2) and converted to %fat using the following equations:

 $4C = 2.748 \times BV - 0.699 \times TBW + 1.129 \times Mo - 2.051 \times BM$ %fat = (FM / BM) x 100

BV = Body volume (L); TBW = Total body water (L); Mo = Total body bone mineral content (kg); BM = Body mass (kg); M = total-body mineral (kg).

Body Volume (BV)

BV was measured using the BOD POD®. Before each test, the BOD POD® (BP) was calibrated according to the manufacturer's instructions with the chamber empty using a cylinder of known volume (49.558L). The subject, in a tight-fitting bathing suit and swimming cap only, then entered and sat in the fiberglass chamber. The BP was sealed, and the subject breathed normally for 20 seconds while body volume (Vb) was measured. After this, the subject was connected to a breathing tube internal to the system to measure thoracic gas volume (VTG). This value was used to correct Vb for VTG. All BP measurements were performed by a BOD POD®-certified investigator who had previously demonstrated a SEM of 0.3447 L on eleven men and women measured 24-48 hours apart.

Siri (6) $2C FM = 4.570 \times BV - 4.50 \times BM$ Brozek (1) $2C FM = 4.570 \times BV - 4.142 \times BM$ BV = Body volume (L); BM = Body mass (kg).



Dual-Energy X-Ray Absorptiometry (DEXA)

A D₂O tracer was used to estimate TBW. Prior to D₂O ingestion, urine samples were

collected from all subjects. Subjects were instructed to void their bladder as much as

possible. After voiding the bladder completely, subjects ingested ≈ 11 grams of D₂O

along with a 100ml rinse of deionized water. The exact amount of deuterium oxide in-

gested for each subject was recorded. After a 4-hour equilibration period restricting defe-

cation, urination, and food and water ingestion, subjects were instructed to provide a post

-urine sample. Within 30 minutes of collection, all urine samples were pipetted into cry-

ogenic vials and stored at -80°C for later analysis. The urine-diluted D₂O was analyzed

in triplicate using an isotope-ratio mass spectrometer, and the isotope abundances in the

urine were calculated following the method of (4). TBW was then calculated from the di-

lution of isotopic water and corrected for the exchange of deuterium with nonaqueous

Bone mineral content (BMC) was estimated using dual-energy X-ray absorptiometry (DXA) (software version 10.50.086, Lunar Prodigy Advance, Madison, WI). BMC was converted to totalbody bone mineral (Mo) using the following equa-

Mo (kg) = total body BMC (kg) x 1.0436

Previous test retest scans of eleven men and women measured 24 - 48 hours apart for Mo produced a standard error of measurement (SEM) of 0.050 kg.

Deuterium Oxide (D,O)

tissue (9).



STUDY PARTICIPANTS

Women (n=104)								
	Group A	Group B						
Age (y)	73 +/- 6	71 +/- 5						
Height (cm)	160.5 +/- 5.5	163 +/- 6						
Weight (kg)	61.95 +/- 6.38	68.17 +/- 12.88						
%Fat	40.3 +/- 6.0	40.4 +/- 6.3						
%Sarcopenic	61	64						

RESULTS

DXA%FAT	(Mean +/- 3D)	Slope	Intercept	r	(%)	(%)	CE (%) / Bias +/- 2SD	Upper Limits	Lower Limits	Trend
Women A (n = 59)										
4C	40.35 +/- 6.03									
SUM 3 4C	38.27 +/- 5.09	0.89	6.16	0.75	3.99	4.47	2.07 +/- 7.82	9.90	-5.75	0.19
SUM 7 4C	38.36 +/- 5.03	0.86	7.44	0.72	4.24	4.67	1.98 +/- 8.36	10.34	-6.38	0.21
DXA % Fat	38.73 +/- 5.37	1.06	-0.54	0.94	2.05	2.61	1.62 +/- 4.03	5.65	-2.41	0.12
SUM 3 DXA	37.26+/- 5.86	0.70	12.51	.77	3.48	4.10	1.47+/- 7.56	9.03	-6.10	-0.10
SUM 7 DXA	37.35 +/- 5.62	0.73	11.62	.76	3.52	4.03	1.37 +/- 7.48	8.86	-6.11	-0.05
Women B (n = 45)										
4C	40.44+/- 6.31									
SUM 3 4C	42.99 +/- 5.81	0.92	0.96	0.84	3.42	4.23	-2.55+/- 6.68	4.14	-9.23	0.09
SUM 7 4C	42.96+/- 5.92	0.92	1.37	0.85	3.33	4.15	-2.52 +/- 6.54	4.03	-9.06	0.07
DXA % Fat	39.14 +/- 6.42	0.93	4.18	0.94	2.14	2.51	1.30 +/- 4.25	5.55	-2.95	-0.02
SUM 3 DXA	41.46 +/- 6.28	.82	4.95	0.79	3.97	4.65	-2.32 +/- 7.98	5.66	-10.31	0.05
SUM 7 DXA	41.50+/- 6.28	0.81	5.50	0.79	3.96	4.68	-2.36 +/- 8.02	5.66	-10.37	0.02
Women A+ B (n = 104)										
4C	40.39+/- 6.12									
SUM 3 4C	40.31+/- 5.87	0.76	9.57	0.73	4.18	3.09	0.07+/- 8.59	8.67	-8.52	0.05
SUM 7 4C	40.35 +/- 5.87	0.75	9.99	0.72	4.25	1.75	0.04+/- 8.77	8.80	-8.73	0.05
DXA %fat	38.90 +/- 5.82	0.99	1.96	0.94	2.11	3.02	1.48+/- 4.12	5.60	-2.64	0.05
SUM 3 DXA	39.08 +/- 6.32	0.69	12.09	0.74	3.91	4.35	-0.17 +/- 0.68	8.38	-8.73	-0.09

GONGLUSIONS

Due to the large TE and SEE values, ultrasound cannot be considered as an alternative to more comprehensive lab methods like DXA in sarcopenic and non-sarcopenic elderly women. However, ultrasound can be used in conjunction with other lab methods to quantify subcutaneous adipose tissue as well as %Fat. Future research should evaluate SAT and the usefulness in overall health and performance. This study was funded by a grant provided by Abbott Nutrition:

39.15 +/- 6.24 | 0.69 | 11.74 | 0.74 | 3.91 | 4.32 | -0.24 +/- 8.50 | 8.26 | -8.74 | -0.08



REFERENCES

1. Census UBot. Special tabulations on aging-extensive data on mobility and self-care. Washington, DC: US Bureau of the Cen-

2. Lipsitz LA, Nakajima I, Gagnon M, Hirayama T, Connelly CM, Izumo H, and Hirayama T. Muscle strength and fall rates among residents of Japanese and American nursing homes: an International Cross-Cultural Study. Journal of the American Geriatrics Society. 1994;42(9):953-9.

3. Schneider EL, and Guralnik JM. The aging of America. Impact on health care costs. JAMA. 1990;263(17):2335-40.

4. Miller CW. Survival and ambulation following hip fracture. *The Journal of Bone and Joint Surgery*. 1978;60(7):930-4.

5. Dawson D, Hendershot G, and Fulton J. Aging in the eighties: functional limitations of individuals age 65 and over. Hyattsville, MD: Advance Data from Vital and Health Statistics; 1987. Available from: Advance Data from Vital and Health Statistics.

6. Janssen I, Heymsfield SB, and Ross R. Low relative skeletal muscle mass (sarcopenia) in older persons is associated with functional impairment and physical disability. *Journal of the American Geriatrics Society*. 2002;50(5):889-96.

Armstrong LE, Maresh CM, Castellani JW, Bergeron MF, Kenefick RW, LaGasse KE, and Riebe D. Urinary indices of hydration status. International Journal of Sport Nutrition. 1994;4(3):265-79.

3. Armstrong LE, Soto JA, Hacker FT, Jr., Casa DJ, Kavouras SA, and Maresh CM. Urinary indices during dehydration, exercise, and rehydration. International Journal of Sport Nutrition. 1998;8(4):345-55.

. Janssen I, Heymsfield SB, Baumgartner RN, and Ross R. Estimation of skeletal muscle mass by bioelectrical impedance analysis. J Appl Physiol. 2000;89(2):465-71.