

The Association between the Low Muscle Mass and Osteoporosis in Elderly Korean People

Sunyoung Kim,^{1,2} Chang Won Won,¹
Byung Sung Kim,¹ Hyun Rim Choi,¹
and Min Young Moon¹

¹Department of Family Medicine, College of Medicine, Kyung Hee University, Seoul; ²Department of Medicine, Graduate School Ewha Womans University, Seoul, Korea

Received: 23 December 2013

Accepted: 24 April 2014

Address for Correspondence:

Chang Won Won, MD

Department of Family Medicine, Kyung Hee University Hospital,
23 Kyunghee dae-ro, Dongdaemun-gu, Seoul 130-872, Korea
Tel: +82-2-958-8697, Fax: +82-2-958-8699
E-mail: chunwon62@dreamwiz.com

Funding: There is no research grant or funding with respect to this manuscript.

The purpose of this study was to predict osteoporosis risk as decreasing muscle mass and to declare the cut-off value of low muscle mass in an elderly Korean population. This study was based on data from the 2008–2010 Korea National Health and Nutritional Examination Surveys (KNHANES). The subjects included 1,308 men and 1,171 women over 65 yr. Bone mineral density (BMD) and appendicular skeletal muscle (ASM) were measured by dual energy X-ray absorptiometry (DXA), and appendicular skeletal muscle was adjusted by height as a marker of sarcopenia. After confirming the correlation between low muscle mass and BMD, the best cut-off value of muscle mass to estimate osteoporosis was suggested through the receiver operating characteristic (ROC) curve. For both men and women, BMD correlated positively with low muscle mass when ASM/Ht² was used as a marker for sarcopenia. The ROC curve showed that ASM/Ht² was the best marker for osteoporosis at a cut-off value of 6.85 kg/m² for men and 5.96 kg/m² for women. When these cut-off values were used to determine sarcopenia, the risk of osteoporosis increased 4.14 times in men and 1.88 times in women. In particular, men (OR 2.12) with sarcopenia were more greatly affected than women (OR 1.15), even after adjusting for osteoporosis risk factors. In elderly Korean people, sarcopenia is positively correlated with BMD and there is a strong correlation between sarcopenia and osteoporosis with risk of bone fracture.

Keywords: Osteoporosis; Sarcopenia; Korea; KNHANES; Elderly Population

INTRODUCTION

According to the 2010 data from the Statistics Korea, the elderly population in Korea increased 24.3% from 2005 to 2010, which was 12.2 times greater than the increase in the total population (2.0%) during the same time period (1, 2). With this significant increase in the elderly population, it is important that increasing attention be given to help maintain their physical and social functioning while minimizing chronic disease and disability to enhance their quality of life.

Decreased muscle mass and bone mineral density are significant changes that occur with aging, and these are often associated with an inability to adapt to external stress resulting in falls, trauma, functional disability, increased hospitalization, decreased quality of life, and increased mortality (3–5). Thus, diagnosing osteoporosis and sarcopenia is not only important clinically, but also has significant public and social ramifications. Determining the relationship between osteoporosis and sarcopenia may help maintain musculoskeletal function and prevent fractures associated with accidental falls.

The diagnostic criteria have already been established for osteoporosis (5), but there is still controversy regarding the diagnostic criteria and markers for sarcopenia. Early studies by Baum-

gartner et al. suggested defining sarcopenia based on appendicular skeletal muscle divided by height squared (Ht²) (7). Janssen et al. used biochemical impedance analysis (BIA) to measure total muscle mass and divided this mass by weight to define sarcopenia (4). In recent studies, sarcopenia is defined as a systemic continuous decrease in skeletal muscle strength and muscle mass so that the diagnosis of sarcopenia must be supported by decreased muscle mass and functional muscle strength (8). Until now, many studies of muscle mass and BMD have been carried out (9–11), and they showed that low muscle mass is correlated with low BMD. Unfortunately, their study populations were inconsistent and their definitions of sarcopenia different. In our study, by presenting a cut-off value of the muscle loss that can predict osteoporosis in Korean elderly individuals, we attempted to identify the relationship between muscle loss and osteoporosis.

MATERIALS AND METHODS

Subjects

This study is based on data obtained in the second and third years (2008–2009) of the Korea National Health and Nutritional Examination Survey (KNHANES) IV and the first year (2010) of

the KNHANES V (2). The KNHANES is a cross-sectional study and has been conducted periodically since 1998 to assess the health and nutritional status of the civilian non-institutionalized population of Korea and has generated nationwide and representative statistical data by self-administered questionnaires on health status, health behaviors, and nutritional status. Data was collected via household units and selected based on a stratified, multistage probability sampling design. The selection was made from sampling units based on the geographical area, gender, and age using household registries. The details of the KNHANES IV and V have been previously described (12, 13). A total of 4,772 individuals aged ≥ 65 yr participated. Of the 4,772 participants, we excluded individuals who did not have DXA results ($n = 1,452$) and those with a history of stroke, coronary artery disease, thyroid disease, lung disease, liver and renal disease, and any cancers and those taking medications likely to affect bone or soft tissue metabolism, such as thyroid hormone, bisphosphonates, SERM, or weight controlling drugs ($n = 841$). The exclusion criteria were aimed to ensure selection of healthy participants and to minimize the confounding effects on muscle mass. The present study included 2,479 elderly participants (1,308 men and 1,171 women).

Measurement of bone density and appendicular skeletal muscle mass

All participants underwent the DXA (DISCOVERY-W fan-beam densitometer; Hologic Inc., MA, USA) for assessment of BMD and body composition. Standardized daily quality control of these DXA instruments was performed using spine phantom provided by the manufacturer before the study commenced. The BMD (g/cm^2 or T-score) of the whole body, femoral neck, and lumbar spine (L1-4) were analyzed. The diagnosis of osteoporosis was made using the WHO T-score criteria (T-score < -2.5), and the maximum BMD value for Japanese patients was used as a reference due to the lack of established diagnostic criteria for Koreans (14, 15). Appendicular skeletal muscle (ASM) mass was calculated as the sum of muscle mass in arms and legs, assuming that all non-fat and non-bone tissue is skeletal muscle (16). According to the definition proposed in previous studies, we used muscle mass indexes, the height adjusted ASM (ASM/Ht^2 , kg/m^2) (5, 7, 17).

Anthropometric and laboratory measurements

Body weight and height were measured to the nearest 0.1 kg and 0.1 cm, respectively. BMI was calculated as $\text{weight}/\text{height}^2$ (kg/m^2). Waist circumference was measured at the midpoint between the lower border of the rib cage and the iliac crest using a non-elastic tapeline. Blood pressure was measured twice at a 5-min interval on the right arm using a standard mercury sphygmomanometer (Baumanometer; Baum, Copiague, NY, USA) and recorded as an average of the two readings. Blood

samples were collected during the health examination survey. These samples, taken from the antecubital veins, were refrigerated immediately and transported on dry ice to the designated central testing facility, and were analyzed within 24 hr of sampling. Serum 25(OH)D levels were measured using a gamma counter (1470 Wizard, Perkin-Elmer, Turku, Finland) with a radioimmunoassay (RIA) kit (DiaSorin, Stillwater, MN, USA) (2).

Statistical analysis

All data are presented as the mean \pm standard error (SE) for the continuous variables or as proportion (SE) for the categorical variables. The characteristics of the participants were compared according to sex using independent sample Student's *t*-tests for the continuous measures and Rao scatt chi-square tests for the categorical measures. Regression analysis was applied to assess the association between various parts of the BMDs and the two indexes of low muscle mass. The receiver operating characteristic (ROC) curve analysis was performed to determine the best cut-off value of low muscle mass for indicating osteoporosis. The areas under the ROC curve (AUC) were investigated for osteoporosis. The ROC curve was a graph of sensitivity plotted against (1-specificity) over all possible diagnostic cut-off values. The optimal cut-off values were obtained from the maximal Youden's index, calculated as (sensitivity+specificity-1) and the best combination of sensitivity and specificity. The odds ratios (ORs) and 95% confidence intervals (CIs) were calculated by logistic analysis to confirm the impact of low muscle mass, which was based on the cut-off value for osteoporosis, after making adjustments for age, fat mass, calcium intake, vitamin D status, smoking, alcohol consumption, and physical activity. Statistical analyses were performed using the survey procedure of SAS software (version 9.2; SAS Institute, Cary, NC, USA) to account for the complex sampling design and to provide nationally representative prevalence estimates. $P < 0.05$ was considered statistically significant.

Ethics statement

This study was approved by the institutional review board of KyungHee University Hospital, in Seoul, Korea (approval ID: KMCIRB 1413-05). Informed consent was waived by the board.

RESULTS

The study population comprised 1,308 men and 1,171 women with an average age of 71.9 yr for men and 74.2 yr for women. Height, weight, waist circumference, and muscle index were higher in men, and body fat index was higher in women. Diabetes was more prevalent in men, and hypertension and metabolic syndrome were more prevalent in women (Table 1). Of the index for sarcopenia, ASM/Ht^2 was positively correlated with BMD in both men and women, and adjusting for age and

Table 1. General characteristics of Korean elderly population

Parameters	Mean \pm SD		P value
	Men (n = 1,308)	Women (n = 1,171)	
Age (yr)	71.9 \pm 0.2	74.2 \pm 0.2	< 0.001
Weight (kg)	62.8 \pm 0.3	54.1 \pm 0.3	< 0.001
Height (cm)	164.8 \pm 0.2	150 \pm 0.2	< 0.001
Body mass index (kg/m ²)	23.1 \pm 0.1	24 \pm 0.1	< 0.001
Waist circumference (cm)	84.5 \pm 0.3	83.1 \pm 0.3	0.004
ASM/Weight	31 \pm 0.1	24.7 \pm 0.1	< 0.001
ASM/Height ² (kg/m ²)	7.1 \pm 0	5.9 \pm 0.07	< 0.001
Total femur BMD (g/cm ³)	0.878 \pm 0.004	0.687 \pm 0.004	< 0.001
Femoral neck BMD (g/cm ³)	0.7 \pm 0.004	0.542 \pm 0.003	< 0.001
Lumbar spine BMD (g/cm ³)	0.928 \pm 0.005	0.732 \pm 0.005	< 0.001
Whole body total BMD (g/cm ³)	1.143 \pm 0.005	0.942 \pm 0.005	< 0.001
Total femur T-score	-0.453 \pm 0.032	-1.427 \pm 0.034	< 0.001
Spine T-score	-0.798 \pm 0.044	-2.38 \pm 0.042	< 0.001
Femoral neck T-score	-1.181 \pm 0.033	-2.442 \pm 0.031	< 0.001
Past history (%)			
Hypertension	56.5 (1.8)	66.6 (1.8)	< 0.001
Diabetes mellitus	21.2 (1.2)	19.1 (1.6)	0.265
Metabolic syndrome*	41.2 (1.8)	63.1 (1.9)	< 0.001
Drinking (%)			< 0.001
Non to moderate drinker	33.4 (1.5)	71.7 (1.7)	
Heavy drinker†	66.6 (1.5)	28.9 (1.7)	
Smoking (%)			< 0.001
Non or ex-smoker	72.8 (1.5)	92.6 (1.1)	
Current smoker	27.2 (1.5)	7.4 (1.1)	
Exercise (%)			< 0.001
No	78.0 (1.5)	84.7 (1.2)	
Moderate to severe‡	22.0 (1.5)	15.2 (1.2)	
25(OH)D (ng/mL)	18.4 \pm 0.3	22.3 \pm 0.4	< 0.001
Calcium intake (mg)	359.6 \pm 21.7	482.0 \pm 12.3	< 0.001

*Metabolic syndrome: three or more of the following five criteria were defined as having metabolic syndrome 1) abdominal obesity by waist circumference cut off values of ≥ 90 cm for men and ≥ 80 cm for women), 2) systolic blood pressure ≥ 130 mmHg or diastolic blood pressure ≥ 85 mmHg or on antihypertensive medication, 3) elevated fasting blood glucose (≥ 5.6 mM/L), 4) hypertriglyceridemia (≥ 1.7 mM/L), and 5) low serum HDL cholesterol (< 1.3 mM/L in men and < 1.29 mM/L in women). †Heavy drinker was defined who drank more than 30 g/day; ‡Moderate exercise was defined as strenuous physical activity performed for at least 20 min at a time at least three times a week. SD, standard deviation; ASM, appendicular skeletal muscle; BMD, bone mineral density.

Table 2. Association of sarcopenia index and bone mineral density, stratified by sex, age and fat mass

Indexes	Men			Women		
	β	SE	P	β	SE	P
Whole body BMD (g/cm ³)						
ASM/Ht ² (kg/m ²)	0.039 [†]	-0.005	< 0.001	0.023*	-0.007	0.001
Model 1 [‡]	0.031 [†]	-0.005	< 0.001	0.016*	-0.006	0.006
Model 2 [§]	0.026 [†]	-0.006	< 0.001	0.007	-0.006	0.24
Femur neck BMD (g/cm ³)						
ASM/Ht ² (kg/m ²)	0.055 [†]	-0.004	< 0.001	0.034 [†]	-0.005	< 0.001
Model 1	0.046 [†]	-0.005	< 0.001	0.025 [†]	-0.004	< 0.001
Model 2	0.039 [†]	-0.005	< 0.001	0.016 [†]	-0.004	< 0.001
Femur total BMD (g/cm ³)						
ASM/Ht ² (kg/m ²)	0.067 [†]	-0.005	< 0.001	0.047 [†]	-0.005	< 0.001
Model 1	0.059 [†]	-0.005	< 0.001	0.036 [†]	-0.004	< 0.001
Model 2	0.051 [†]	-0.005	< 0.001	0.027 [†]	-0.005	< 0.001
L-Spine BMD (g/cm ³)						
ASM/Ht ² (kg/m ²)	0.061 [†]	-0.006	< 0.001	0.034 [†]	-0.007	< 0.001
Model 1	0.06 [†]	-0.007	< 0.001	0.028 [†]	-0.006	< 0.001
Model 2	0.042 [†]	-0.006	< 0.001	0.005	-0.006	0.48

*P < 0.05; †P < 0.001; ‡Model 1: adjusted for age; §Model 2: adjusted for age and fat mass. BMD, bone mineral density; ASM, appendicular skeletal muscle; Ht, height; Wt, weight.

body fat, still resulted in a positive correlation (Table 2).

The ROC curve reflecting osteoporosis from ASM/Ht² is shown in Fig. 1. The AUC of ASM/Ht² for the prediction of osteoporosis was 0.705 (95% CI, 0.679-0.730; P < 0.001) in men, and 0.598 (95% CI, 0.569-0.627; P < 0.001) in women. The best cut-off value of ASM/Ht² for indicating osteoporosis, according to the maximum of the Youden index, was 6.85 kg/m² in men and 5.96 kg/m² in women. Sensitivity and specificity were 66.9% and 66% in men, and 62% and 53.5% in women, respectively.

Odds ratios (ORs) for the low muscle mass in relation to osteoporosis are presented in Table 3. Both men and women subjects in the lower muscle mass group had increased risk for osteoporosis (men: OR, 4.14; 95% CI, 2.67-6.78; P < 0.001, women: OR, 1.88; 95% CI, 1.41-2.5; P < 0.001). After adjusting for osteoporosis risk factors (age, fat mass, calcium intake, vitamin D status, smoking, alcohol consumption, and physical activity), this association remained statistically significant in men (men: OR, 2.21; 95% CI, 1.33-3.37; P = 0.0016, women: OR, 1.15; 95% CI, 0.81-1.65; P = 0.44) (Table 3).

DISCUSSION

In our study, using ASM/Ht² as an index for sarcopenia, muscle mass and BMD showed a positive correlation, verifying that as muscle mass decreases, the BMD of each body part shows a lower value as well. Especially through the ROC curve, ASM/Ht² well reflected the risk of osteoporosis and indicated the best cut-off values (men 6.85 kg/m², women 5.96 kg/m²) for sarcopenia. This result is similar to studies involving postmenopausal women and men concluding that there are correlations among body weight, muscle mass, and osteoporosis (10, 18-20). It can be explained that body weight acts as a stress on the skeletal system to promote bone formation mechanically by stimulat-

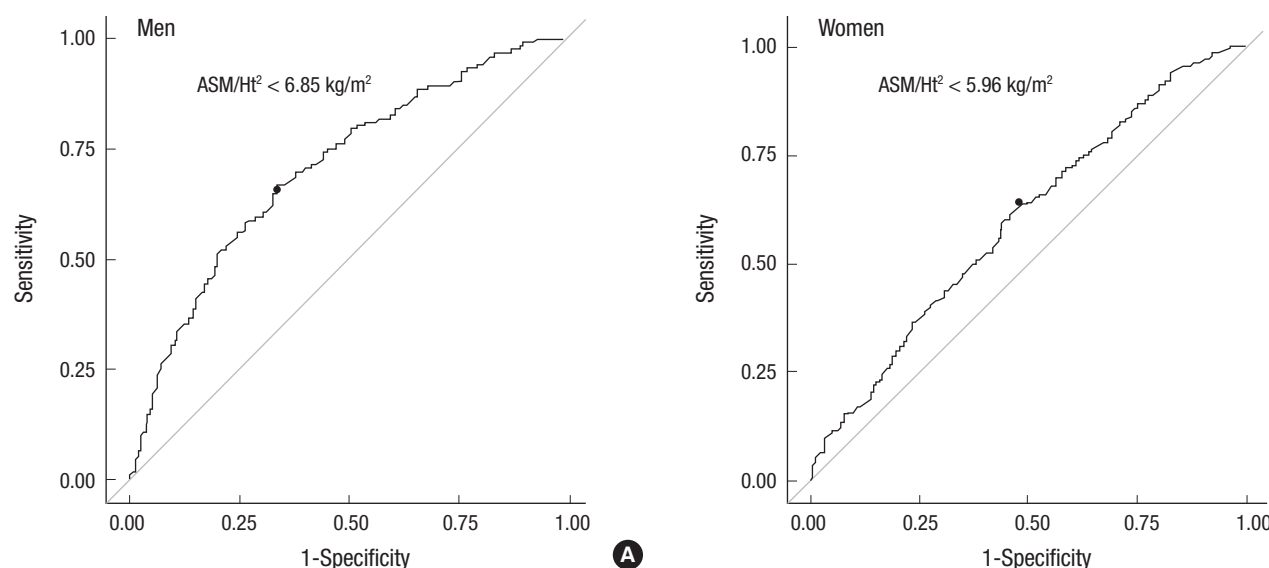


Fig. 1. Receiver operating characteristic (ROC) curve of sarcopenia reflecting osteoporosis in Korean Elderly. The cut-off value of ASM/Ht² was (A) 6.85 kg/m² in men (area under the curve = 0.705; 95% CI, 0.679-0.730; $P < 0.001$ sensitivity 66.9%, specificity 66%), (B) 5.96 kg/m² in women (area under the curve = 0.598; 95% CI, 0.569-0.627; $P < 0.001$ sensitivity 62%, specificity 53.5%).

Table 3. Odds ratios (OR) and 95% confidence intervals (CI) for osteoporosis according to cutoff value of sarcopenia index in Korean Elderly

Sarcopenia index	Osteoporosis*	
	OR (95% CI)	P value
Men		
ASM/Ht ² (kg/m ²) < 6.85	4.14 [†] (2.67-6.78)	< 0.001
Model 1 [‡]	3.85 [†] (2.34-5.47)	< 0.001
Model 2 [‡]	2.63 [†] (1.73-4)	< 0.001
Model 3 [§]	2.12 (1.33-3.37)	0.0016
Women		
ASM/Ht ² (kg/m ²) < 5.96	1.88 [†] (1.41-2.5)	< 0.001
Model 1 [‡]	1.77 [†] (1.3-2.39)	< 0.001
Model 2 [‡]	1.31 (0.96-1.79)	0.08
Model 3 [§]	1.15 (0.81-1.65)	0.44

*Osteoporosis: T-score ≤ -2.5 at femoral neck, total or lumbar spine; [†]Model 1: adjusted for age; [‡]Model 2: adjusted for age and fat mass; [§]Model 3: adjusted for age, fat mass, calcium intake, vitamin D status, smoking, alcohol consumption, physical activity; ^{||} $P < 0.05$; [†] $P < 0.001$. ASM, appendicular skeletal muscle.

ing osteocytes to increase bone density (21), and muscle mass, like fat mass may have a protective effect on bone density through hormones (22). Also, the association between greater muscle mass and greater bone density is likely to be determined by multiple factors including common nutrition, life style and genes regulating size (23-24).

Many studies have focused on sarcopenia since Irwin Rosenberg first identified the condition in 1989 (25) but a precise definition, measurement method, and index of sarcopenia have yet to be established. Baumgartner et al. measured ASM with DXA and defined sarcopenia as less than -2 standard deviations from the mean of ASM/Ht² in a young population (7).

In Korea, interest in sarcopenia has increased, and numerous studies have been carried out. Kim et al. (26) studied sarcopenia in a Korean population and found a lower prevalence of

sarcopenia (men 6.3%, women 4.1%) compared to Western countries when using ASM/Ht² as the sarcopenia index (cut-off values for men and women were 7.40 kg/m² and 5.14 kg/m², respectively). In a KLoSHA study with a sample population of adults older than 65 yr, the prevalence of sarcopenia was 35.3% in men and 13.4% in women when ASM/Ht² was used as the sarcopenia index (cut-off values for men and women were 7.09 kg/m² and 5.27 kg/m², respectively) (17). But a study of the prevalence of sarcopenia in Korea using the KNHANES data showed that weight-adjusted ASM is better than height-adjusted ASM in defining sarcopenia. When using ASM/Ht², the prevalence of sarcopenia was 12.4% in men and 0.1% in women (cut-off values for men and women were 6.58 kg/m² and 4.59 kg/m², respectively) (12). Though these studies performed in Korea seem to suggest different cut-off values, this is because each study enrolled different study populations including various group of ages.

The present study, similar to other Korean studies, found that ASM/Ht² was the best sarcopenia index for indicating osteoporosis, with cut-off values at 6.85 kg/m² in men and 5.96 kg/m² in women. The cut-off value for men ranged between Sarcopenia I (7.50) and II (6.58), and the cut-off value in women was higher than Sarcopenia I (5.38), which was the standard in previous Korean studies (12). The prevalence of sarcopenia in this study was 39.5% in men and 58.1% in women. Previous studies were limited in that the cut-off value for women was too low to determine prevalence, but using ASM/Ht² as an index seems to have addressed this issue. In addition, when sarcopenia was based on the optimal cut-off value, the sarcopenia group had increased risk of osteoporosis both in men (OR, 4.14; $P < 0.001$) and women (OR, 1.88; $P < 0.001$). In particular, men (OR, 2.12; $P = 0.0016$) with sarcopenia were more greatly affected than women (OR,

1.15; $P = 0.44$), even after adjusting for age, fat mass, calcium intake, vitamin D status, smoking, alcohol consumption, and physical activity. This was demonstrated in previous studies showing that bone mass in men was more closely related to muscle mass than in women (27-31). The gender difference in the relationship between bone and muscle are explained by gender specific effects of sex hormones. In men, changes to bone and muscle are controlled by increasing levels of testosterone and IGF-1 resulting in increased muscle mass and strength, whereas in women, the higher level of estrogen results in bone mass tending to increase more rapidly in relation to muscle (27, 29). And in both genders, aging causes loss in bone and muscle; and the bone and muscle relationship is affected by gender differences in the rate of bone and muscle loss (32). Especially, age-related decrease in testosterone and IGF-1 levels may lead to decrease in muscle and bone in men, while, absolute level and degree of decline in testosterone are much lower in women, muscle mass can be relatively preserved. In addition, mechanical strain and estrogen share a common pathway involving activation of estrogen receptor- α (ER- α), and a decline in ER- α with menopause reduces the ability of mechanical loading to induce an osteogenic response (33-35). This mechanostat re-setting due to estrogen deficiency may explain the decoupling of muscle mass and bone density in women, with men tending to show higher correlations between muscle mass and real bone density. Therefore, this study suggests that for preventing osteoporosis, sarcopenia could be more emphasized in men.

A cross-sectional study, we were not able to verify causal relationships. However, participants had minimal radiation exposure during the study as they underwent the DXA study for assessment of both muscle mass and BMD simultaneously. And this showed a positive correlation between muscle mass and BMD in the Korean population. In addition, this study examined the relationship between sarcopenia and osteoporosis separately in each sex. In previous studies the prevalence rate of sarcopenia in women was too low because the cut-off value used for this group was 4.59-5.27. But in this study, the cut-off value for women was 5.96 so that it was possible to predict that the prevalence rate would be higher than previously calculated with the ASM/Ht² index.

As mentioned above, osteoporosis and sarcopenia have common causal factors such as low levels of vitamin D, testosterone, estrogen, and insulin, as well as, reduced calorie intake and malnutrition (9-11). The present study identified that sarcopenia can increase the risk of osteoporosis and may lead to increased risk of bone fracture.

In conclusion, muscle mass is positively correlated with bone density in both men and women using the ASM/Ht² index. The risk of osteoporosis increases when sarcopenia is diagnosed with a cut-off value of 6.85 kg/m² in men and 5.96 kg/m² in women. Concurrently diagnosing osteoporosis with sarcopenia

may help prevent accidental falls and bone fractures.

ACKNOWLEDGMENT

The authors wish to acknowledge the statistical support of KD Han.

DISCLOSURE

The authors have no conflicts of interest to disclose.

ORCID

Sunyoung Kim <http://orcid.org/0000-0003-4115-4455>

Chang Won Won <http://orcid.org/0000-0002-6429-4461>

Byung Sung Kim <http://orcid.org/0000-0002-4984-6918>

Hyun Rim Choi <http://orcid.org/0000-0003-4356-2831>

Min Young Moon <http://orcid.org/0000-0002-7207-8155>

REFERENCES

1. Statistics Korea. *Korean Statistical Information Service*. Available at http://kosis.kr/statisticsList/statisticsList_01List.jsp?vwcd=MT_ZTITLE&parentId=A [accessed on 20 February 2014].
2. Korea Centers for Disease Control and Prevention. *The fourth and fifth Korea National Health and Nutrition Examination Survey (KNHANES IV-2,3 and V-1), 2008-2010*. Available at <http://knhanes.cdc.go.kr> [accessed on 20 February 2014].
3. Kannus P, Parkkari J, Niemi S, Palvanen M. *Fall-induced deaths among elderly people*. *Am J Public Health* 2005; 95: 422-4.
4. Smulders E, Weerdesteyn V, Groen BE, Duysens J, Eijssbouts A, Laan R, van Lankveld W. *Efficacy of a short multidisciplinary falls prevention program for elderly persons with osteoporosis and a fall history: a randomized controlled trial*. *Arch Phys Med Rehabil* 2010; 91: 1705-11.
5. Janssen I, Heymsfield SB, Ross R. *Low relative skeletal muscle mass (sarcopenia) in older persons is associated with functional impairment and physical disability*. *J Am Geriatr Soc* 2002; 50: 889-96.
6. NIH Consensus Development Panel on Osteoporosis Prevention, Diagnosis, and Therapy. *Osteoporosis prevention, diagnosis, and therapy*. *JAMA* 2001; 285: 785-95.
7. Baumgartner RN, Koehler KM, Gallagher D, Romero L, Heymsfield SB, Ross RR, Garry PJ, Lindeman RD. *Epidemiology of sarcopenia among the elderly in New Mexico*. *Am J Epidemiol* 1998; 147: 755-63.
8. Cruz-Jentoft AJ, Baeyens JP, Bauer JM, Boirie Y, Cederholm T, Landi F, Martin FC, Michel JP, Rolland Y, Schneider SM, et al. *Sarcopenia: European consensus on definition and diagnosis: report of the European Working Group on Sarcopenia in Older People*. *Age Ageing* 2010; 39: 412-23.
9. Coin A, Sergi G, Marin S, Vianello A, Perissinotto E, Sarti S, Rinaldi G, Mosele M, Inelmen EM, Enzi G, et al. *Predictors of low bone mineral density in elderly males with chronic obstructive pulmonary disease: the role of body mass index*. *Aging Male* 2010; 13: 142-7.
10. Blain H, Jaussent A, Thomas E, Micallef JP, Dupuy AM, Bernard PL, Mariano-Goulart D, Cristol JP, Sultan C, Rossi M, et al. *Appendicular skeletal*

- tal muscle mass is the strongest independent factor associated with femoral neck bone mineral density in adult and older men. *Exp Gerontol* 2010; 45: 679-84.
11. Frisoli A Jr, Chaves PH, Ingham SJ, Fried LP. Severe osteopenia and osteoporosis, sarcopenia, and frailty status in community-dwelling older women: results from the Women's Health and Aging Study (WHAS) II. *Bone* 2011; 48: 952-7.
 12. Kim YS, Lee Y, Chung YS, Lee DJ, Joo NS, Hong D, Song Ge, Kim HJ, Choi YJ, Kim KM. Prevalence of sarcopenia and sarcopenic obesity in the Korean population based on the Fourth Korean National Health and Nutritional Examination Surveys. *J Gerontol A Biol Sci Med Sci* 2012; 67: 1107-13.
 13. Lim JS, Kim KM, Rhee Y, Lim SK. Gender-dependent skeletal effects of vitamin D deficiency in a younger generation. *J Clin Endocrinol Metab* 2012; 97: 1995-2004.
 14. Orimo H, Hayashi Y, Fukunaga M, Sone T, Fujiwara S, Shiraki M, Kushida K, Miyamoto S, Soen S, Nishimura J, et al. Diagnostic criteria for primary osteoporosis: year 2000 revision. *J Bone Miner Metab* 2001; 19: 331-7.
 15. Kanis JA. Assessment of fracture risk and its application to screening for postmenopausal osteoporosis: synopsis of a WHO report: WHO Study Group. *Osteoporos Int* 1994; 4: 368-81.
 16. Heymsfield SB, Smith R, Aulet M, Bensen B, Lichtman S, Wang J, Pierson RN Jr. Appendicular skeletal muscle mass: measurement by dual-photon absorptiometry. *Am J Clin Nutr* 1990; 52: 214-8.
 17. Kim JH, Hwang BY, Hong ES, Ohn JH, Kim CH, Kim HW, Ahn HY, Yoon JW, Kang SM, Park YJ, et al. Investigation of sarcopenia and its association with cardiometabolic risk factors in elderly subjects. *J Korean Geriatr Soc* 2010; 14: 121-30.
 18. Genaro PS, Pereira GA, Pinheiro MM, Szejnfeld VL, Martini LA. Influence of body composition on bone mass in postmenopausal osteoporotic women. *Arch Gerontol Geriatr* 2010; 51: 295-8.
 19. Van Langendonck L, Claessens AL, Lefevre J, Thomis M, Philippaerts R, Delvaux K, Lysens R, Vanden Eynde B, Beunen G. Association between bone mineral density (DXA), body structure, and body composition in middle-aged men. *Am J Hum Biol* 2002; 14: 735-42.
 20. Pluijm SM, Visser M, Smit JH, Popp-Snijders C, Roos JC, Lips P. Determinants of bone mineral density in older men and women: body composition as mediator. *J Bone Miner Res* 2001; 16: 2142-51.
 21. Rochefort GY, Pallu S, Benhamou CL. Osteocyte: the unrecognized side of bone tissue. *Osteoporos Int* 2010; 21: 1457-69.
 22. Matsumine H, Hirato K, Yanaihara T, Tamada T, Yoshida M. Aromatization by skeletal muscle. *J Clin Endocrinol Metab* 1986; 63: 717-20.
 23. Coin A, Perissinotto E, Enzi G, Zamboni M, Inelmen EM, Frigo AC, Manzato E, Busetto L, Buja A, Sergi G. Predictors of low bone mineral density in the elderly: the role of dietary intake, nutritional status and sarcopenia. *Eur J Clin Nutr* 2008; 62: 802-9.
 24. Seeman E, Hopper JL, Young NR, Formica C, Goss P, Tsalamandris C. Do genetic factors explain associations between muscle strength, lean mass, and bone density? a twin study. *Am J Physiol* 1996; 270: E320-7.
 25. Rosenberg IH. Epidemiologic and methodologic problems in determining nutritional status of older persons. *Am J Clin Nutr* 1989; 50: 1231-3.
 26. Kim TN, Yang SJ, Yoo HJ, Lim KI, Kang HJ, Song W, Seo JA, Kim SG, Kim NH, Baik SH, et al. Prevalence of sarcopenia and sarcopenic obesity in Korean adults: the Korean sarcopenic obesity study. *Int J Obes (Lond)* 2009; 33: 885-92.
 27. Zofková I. Hormonal aspects of the muscle-bone unit. *Physiol Res* 2008; 57: S159-69.
 28. Hasselström H, Karlsson KM, Hansen SE, Grønfeldt V, Froberg K, Andersen LB. Sex differences in bone size and bone mineral density exist before puberty: the Copenhagen School Child Intervention Study (CoSCIS). *Calcif Tissue Int* 2006; 79: 7-14.
 29. Macdonald H, Kontulainen S, Petit M, Janssen P, McKay H. Bone strength and its determinants in pre- and early pubertal boys and girls. *Bone* 2006; 39: 598-608.
 30. Rauch F, Bailey DA, Baxter-Jones A, Mirwald R, Faulkner R. The 'muscle-bone unit' during the pubertal growth spurt. *Bone* 2004; 34: 771-5.
 31. Frost HM. The Utah paradigm of skeletal physiology: an overview of its insights for bone, cartilage and collagenous tissue organs. *J Bone Miner Metab* 2000; 18: 305-16.
 32. Lang TF. The bone-muscle relationship in men and women. *J Osteoporos* 2011; 2011: 702735.
 33. Lee KC, Lanyon LE. Mechanical loading influences bone mass through estrogen receptor alpha. *Exerc Sport Sci Rev* 2004; 32: 64-8.
 34. Fricke O, Schoenau E. The 'Functional Muscle-Bone Unit': probing the relevance of mechanical signals for bone development in children and adolescents. *Growth Horm IGF Res* 2007; 17: 1-9.
 35. Taaffe DR, Cauley JA, Danielson M, Nevitt MC, Lang TF, Bauer DC, Harris TB. Race and sex effects on the association between muscle strength, soft tissue, and bone mineral density in healthy elders: the Health, Aging, and Body Composition Study. *J Bone Miner Res* 2001; 16: 1343-52.