ORIGINAL ARTICLE

Atrophy of the lower limbs in elderly women: is it related to walking ability?

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Abstract This study investigated the relationship between walking ability and age-related muscle atrophy of the lower limbs in elderly women. The subjects comprised 20 young women and 37 elderly women who resided in nursing homes or chronic care institutions. The elderly subjects were divided into three groups according to their walking ability. The muscle thickness of the following ten lower limb muscles were measured by B-mode ultrasound: the gluteus maximus, gluteus medius, gluteus minimus, psoas major, rectus femoris, vastus lateralis, vastus intermedius, biceps femoris, gastrocnemius and soleus. Compared to the young group, muscle thicknesses of all muscles except the soleus muscle were significantly smaller in all the elderly groups. There were no significant differences between the fast- and slow-walking groups in the thickness of any muscle. In the dependent elderly group, noticeable muscle atrophy was observed in the quadriceps femoris muscle. The results of this study suggest that the elderly who are capable of locomotion, regardless of their walking speed, show a moderate degree of age-related atrophy, while those who do not walk exhibit more severe atrophy, especially in the quadriceps femoris muscle.

Keywords Aging · Muscle atrophy · Ultrasonography · Lower limb muscles · Elderly women

Introduction

Skeletal muscle mass decreases as people age, and it decreases by 30–50% between the ages of 40 and 80 years (Akima et al. 2001; Allen et al. 1960; Lexell et al. 1988; Young et al. 1985). Muscle atrophy is reported to be greater in the lower limbs than in the upper limbs (Bemben et al. 1991; Brooks and Faulkner 1994; Janssen et al. 2000; Kubo et al. 2003a). Kubo et al. (2003a) reported that relative muscle thickness of the vastus lateralis and medial gastrocnemius muscles, as measured by B-mode ultrasound, decreased significantly with advancing age, whereas no significant age-related change in relative muscle thickness was observed for the triceps brachii muscle.

In recent years, there have been many studies on agerelated atrophy (sarcopenia) of lower limb muscles using ultrasonographic measurement of muscle thickness. However, in these studies, the quadriceps (Arts et al. 2010; Kubo et al. 2003a, b; Reimers et al. 1998) and triceps surae muscles (Fujiwara et al. 2010; Kubo et al. 2003a, b; Reimers et al. 1998) have been mainly evaluated. There have been no studies that have focused on sarcopenia in the individual muscles of the lower limb, including those around the hip, knee and ankle joints.

Loss of skeletal muscle mass is part of the aging process and is exacerbated by inactivity (Sandler et al. 1991). However, there is evidence that older adults, even frail older adults, can increase muscle mass with exercise (Macaluso and De Vito 2004). In this way, the degree of daily physical activity influences the amount of muscle mass, especially in the elderly.

It has been pointed out that daily physical activity in the elderly is closely associated with their walking ability, such as the amount of walking that they do on a daily basis. For example, Park et al. (2010) demonstrated that muscle mass

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in the lower limbs of older adults, as measured by wholebody dual X-ray absorptiometry, is associated with physical activity such as the walking step count in daily life. Conversely, a previous study that measured thigh muscle volume using magnetic resonance imaging (MRI) showed decreases in muscle mass in the early stage after a 7-day period of strict bed rest that prohibited weight-bearing activity such as locomotor activity (Ferrando et al. 1995). However, it is unclear whether sarcopenia of individual lower limb muscles is related to walking ability in the elderly.

The aim of this study was to investigate the relationship between walking ability and sarcopenia of the lower limbs using ultrasound.

Methods

Subjects

The subject groups were comprised of 20 healthy young women (age 19.8 ± 0.8 years; height 1.58 ± 0.06 m; weight 53.8 ± 5.0 kg) and 37 elderly women (85.5 ± 6.3 years; 1.44 ± 0.07 m; 41.1 ± 9.0 kg). The young subjects were healthy university student volunteers. All subjects were physically active but did not include athletes. All elderly subjects were residents of nursing homes or chronic care institutions in Kyoto, Japan. We excluded elderly subjects who had an unstable physical condition, a history of lower extremity surgery or physical dysfunctions, such as acute neurological impairment (acute stroke, Parkinson's disease, paresis of the lower limbs), or a severe musculoskeletal impairment, such as limited range of motion of the lower limbs.

The elderly subjects were divided into 3 groups according to their walking ability: (1) a fast-walking group of elderly persons who were able to walk independently or with an assistive device at a maximum walking speed of more than 1 m/s—this speed has been considered to be the cut-off value that predicts a risk of falls in older people (Shimada et al. 2009); (2) a slow-walking group of elderly persons who were able to walk independently or with an assistive device at a maximum walking speed of <1 m/s and (3) a dependent group of elderly persons who were able to maintain a sitting position in a wheelchair but were not able to walk independently and had not walked for more than half a year. Table 1 shows the characteristics of the three elderly groups. No significant differences were found between elderly groups in age and height.

The subjects were informed about the study procedures before testing and provided a written informed consent before participating. The study was approved by Kyoto University Graduate School and Faculty of Medicine Ethics Committee.

Table 1 Characteristics of the elderly subjects (mean \pm SD)

	Fast-walking group	Slow-walking group	Dependent group
Number	14	11	12
Age (years)	83.1 ± 5.7	85.5 ± 6.8	88.1 ± 5.5
Height (m)	1.47 ± 0.07	1.44 ± 0.07	1.41 ± 0.07
Weight (kg)	$45.0 \pm 6.8^{**}$	45.1 ± 8.8**	32.8 ± 4.5
BMI (kg/m ²)	$21.0 \pm 3.5^{**}$	$21.8 \pm 3.9^{**}$	16.5 ± 1.9

** p < 0.01 significant difference for the dependent group



Fig. 1 Representative ultrasound image of gastrocnemius and soleus

Muscle thickness measurements

Using B-mode ultrasound imaging (LOGIQ Book Xp; GE Healthcare Japan, Tokyo, Japan) with an 8-MHz transducer, lower limb muscle thicknesses on the longitudinal plane were measured (Fig. 1). Strong correlations have been reported between muscle thickness measured by B-mode ultrasound and site-matched skeletal muscle mass measured by MRI (Dupont et al. 2001; Fukunaga et al. 2001; Miyatani et al. 2004; Sanada et al. 2006; Walton et al. 1997). Therefore, it is plausible to use musclethickness measurements to estimate muscle size and degree of muscle atrophy.

Ten muscles in the lower limbs were examined: the gluteus maximus, gluteus medius, gluteus minimus, psoas major, rectus femoris, vastus lateralis, vastus intermedius,

Muscles	Positions	Measurement sites
Gluteus maximus	Prone	30% proximal between posterior superior iliac spine the greater trochanter
Gluteus medius	Prone	Midway between the proximal end of iliac crest and the greater trochanter
Gluteus minimus	Prone	Midway between the proximal end of iliac crest and the greater trochanter
Psoas major	Prone	7 cm lateral from the L3 spinous process
Rectus femoris	Supine	Midway between the anterior superior iliac spine and the proximal end of the patella
Vastus lateralis	Supine	3 cm lateral of 60% distal between the anterior superior iliac spine and the proximal end of the patella
Vastus intermedius	Supine	Midway between the anterior superior iliac spine and the proximal end of the patella
Biceps femoris	Prone	Midway between the ischial tuberosity and the lateral condyle of the tibia
Gastrocnemius	Prone	Medial head of gastrocnemius at 30% proximal between the lateral malleolus of the fibula and the lateral condyle of the tibia
Soleus	Prone	30% proximal between the lateral malleolus of the fibula and the lateral condyle of the tibia

biceps femoris, gastrocnemius and soleus. We examined the muscles on the right lower limbs of subjects. Measurement positions and measurement sites for each muscle are shown in Table 2. During the examination, care was taken to maintain the same standardized position of the subjects and the exact location of the transducer. To improve acoustic coupling, a water-soluble transmission gel was placed over the scan head. The transducer was held perpendicular to the skin surface using the minimum pressure required to achieve a clear image. Muscle thickness was measured by the examiner who was blinded with respect to the subjects' walking ability, such as walking speed.

A previous study has shown the reliability of the ultrasound technique for measuring muscle thickness of the quadriceps femoris (Blazevich et al. 2006; Thoirs and English 2009), hamstring (Kellis et al. 2009; Thoirs and English 2009) and triceps surae muscles (Maganaris et al. 1998; Narici et al. 1996). However, the reliability of the measurements of the gluteus maximus, gluteus medius, gluteus minimus and psoas major muscles has not been shown before. Therefore, the reliability of the measurements of these muscles was assessed in a pilot study. Sixteen healthy volunteers (mean age 20.3 years; SD 0.49) were recruited for the reliability analysis. The intraclass correlation coefficients (ICC) for the test-retest reliability of the muscle thickness measurements were 0.991 (95% CI 0.974-0.997) for the gluteus maximus, 0.990 (95% CI 0.971-0.996) for the gluteus medius, 0.965 (95% CI 0.900-0.988) for the gluteus minimus and 0.968 (95% CI 0.910-0.989) for the psoas major muscles; these results indicated a high degree of reproducibility in measuring muscle thickness of these muscles.

Statistical analyses

All data are presented as mean \pm SD. We calculated the magnitude of age-related decline compared to the young

reference group using the following formula: Magnitude of age-related decline (%) = (mean thickness in the young – muscle thickness in the elderly) × (mean thickness in the young)⁻¹ × 100.

Furthermore, we calculated the *t* scores to take into account the variability in muscle thickness between muscles in the young reference group. The *t* score was calculated using the following equation: *t* score = (muscle thickness in the elderly – mean thickness in the young) \times (SD of thickness in the young)⁻¹.

Differences in muscle thickness and the magnitude of age-related decline between groups (the young, fast-walking, slow-walking and dependent groups) were examined using the Kruskal–Wallis test. If a significant main effect was found, differences among muscle thickness or magnitude of decline were determined with the Steel–Dwass post hoc test. Differences in the t score between muscles were examined using Friedman's test and a multiple comparison method (Steel–Dwass test).

Results

Comparison of muscle thickness (mm) between the young and elderly groups

Table 3 shows the mean and standard deviation values of muscle thicknesses for the young and the three elderly groups. Statistical analysis revealed a significant main effect for all muscles (p < 0.01). When the muscle thicknesses were compared between the young and elderly groups, all muscles except the soleus muscle, i.e. the gluteus maximus, gluteus medius, gluteus minimus, psoas major, rectus femoris, vastus lateralis, vastus intermedius, biceps femoris and gastrocnemius, were significantly thinner in all the elderly groups. As for the soleus muscle, although there was a significant difference between the

	Young	Fast-walking group	Slow-walking group	Dependent group
Gluteus maximus	25.0 ± 2.98	$14.9 \pm 3.56^{**}$	$14.7 \pm 4.46^{**}$	$8.60 \pm 2.42^{**}$
Gluteus medius	22.9 ± 5.80	$15.0 \pm 4.16^{**}$	$14.6 \pm 3.97^{**}$	9.69 ± 2.23**
Gluteus minimus	19.3 ± 6.47	$12.8 \pm 4.08^{**}$	$12.3 \pm 4.36^{**}$	9.54 ± 1.76**
Psoas major	28.7 ± 4.11	$13.0 \pm 5.58^{**}$	$15.0 \pm 2.93^{**}$	$10.9 \pm 4.07^{**}$
Rectus femoris	22.9 ± 3.39	$16.7 \pm 3.50 **$	$15.4 \pm 5.42^{**}$	$4.08 \pm 1.36^{**}$
Vastus lateralis	22.0 ± 3.25	$14.1 \pm 3.76^{**}$	$12.2 \pm 3.08^{**}$	3.73 ± 1.26**
Vastus intermedius	21.5 ± 3.52	$16.9 \pm 2.99^{**}$	$12.8 \pm 4.68^{**}$	4.69 ± 2.39**
Biceps femoris	36.5 ± 4.87	$17.9 \pm 5.01^{**}$	$17.1 \pm 3.83^{**}$	$11.2 \pm 4.40^{**}$
Gastrocnemius	16.3 ± 2.31	$10.8 \pm 2.63^{**}$	$10.5 \pm 4.01^{**}$	$7.04 \pm 1.67^{**}$
Soleus	34.5 ± 6.11	29.8 ± 8.25	27.1 ± 4.93	$17.6 \pm 6.26^{**}$

Table 3 Comparison of muscle thickness (mm) between the young and elderly groups

Values are expressed as mean \pm the standard deviation (SD)

** p < 0.01 significant difference with fast-walking group

young and dependent elderly groups, no differences were found between the young as well as fast- and slow-walking elderly groups.

Comparison of the magnitude of age-related decline (%) between the elderly groups

Table 4 shows the magnitude of age-related decline (%) in the elderly groups. When the magnitude of age-related decline was compared between the three elderly groups, there were no significant differences in magnitude of decline of any muscle between the fast- and slow-walking groups. Significant differences were found between the dependent group and the two walking groups in the magnitude of decline of gluteus maximus, gluteus medius, rectus femoris, vastus lateralis, vastus intermedius, biceps femoris, gastrocnemius and soleus. There were no

Table 4 Comparison of the magnitude of age-related decline (%)between the elderly groups

	Fast walking	Slow walking	Dependent
Gluteus maximus	40.3 ± 14.8	41.0 ± 17.9	$65.5 \pm 10.2^{**,\dagger\dagger}$
Gluteus medius	34.7 ± 18.9	36.1 ± 17.3	$57.7 \pm 10.2^{**,\dagger\dagger}$
Gluteus minimus	33.9 ± 21.9	36.4 ± 22.6	50.6 ± 9.51
Psoas major	54.8 ± 20.1	45.9 ± 10.2	62.3 ± 14.8
Rectus femoris	26.9 ± 15.9	32.8 ± 23.7	$82.2 \pm 6.19^{**,\dagger\dagger}$
Vastus lateralis	36.0 ± 17.7	44.3 ± 14.0	$83.0 \pm 6.00^{**,\dagger\dagger}$
Vastus intermedius	21.3 ± 14.5	40.3 ± 21.8	$78.2 \pm 11.6^{**,\dagger\dagger}$
Biceps femoris	51.0 ± 14.3	53.1 ± 10.5	$69.4 \pm 12.6^{*,\dagger}$
Gastrocnemius	34.0 ± 16.7	35.4 ± 24.6	$56.8 \pm 10.7^{**,\dagger}$
Soleus	13.5 ± 24.8	21.6 ± 14.3	$49.1 \pm 18.9^{**,\dagger}$

Values are expressed as mean \pm the standard deviation (SD)

** p < 0.01, * p < 0.05 significant difference with fast-walking group ^{††} p < 0.01, [†] p < 0.05 significant difference with slow-walking group significant differences in magnitude of decline between the three elderly groups in the gluteus minimus and psoas major muscles.

Differences in the *t* score between muscles in the elderly groups

Table 5 shows the t score of muscle thickness in the elderly groups. The t score of muscle thickness in the fast-walking group was the highest for the soleus muscle, followed in order by the gluteus minimus, gluteus medius, vastus intermedius, rectus femoris, gastrocnemius, vastus lateralis, gluteus maximus, biceps femoris and psoas major muscles. The Friedman test indicated that there was a significant

Table 5 Differences in the t score between muscles in the elderly
groups

	Fast walking	Slow walking	Dependent
Gluteus maximus	-3.13 ± 1.16^a	-3.19 ± 1.39	-5.09 ± 0.79^{b}
Gluteus medius	-1.37 ± 0.74	-1.42 ± 0.68	-2.28 ± 0.40
Gluteus minimus	-1.01 ± 0.65	-1.08 ± 0.67	-1.51 ± 0.28
Psoas major	-3.84 ± 1.41^{b}	-3.22 ± 0.72^{b}	-4.37 ± 1.04^{c}
Rectus femoris	-1.82 ± 1.07	-2.22 ± 1.60	-5.55 ± 0.42^d
Vastus lateralis	-2.47 ± 1.22	-3.04 ± 0.96	-5.70 ± 0.41^{d}
Vastus intermedius	-1.75 ± 1.64	-2.62 ± 1.42	$-5.10\pm0.76^{\rm b}$
Biceps femoris	-3.80 ± 1.06^{b}	-3.96 ± 0.78^b	-5.17 ± 0.94^{b}
Gastrocnemius	-2.40 ± 1.18	-2.50 ± 1.74	$-4.01\pm0.76^{\rm c}$
Soleus	-0.76 ± 1.40	-1.22 ± 0.81	-2.77 ± 1.07

Values are expressed as means \pm the standard deviation (SD)

^b Significant difference with soleus, gluteus minimus and gluteus medius

^c Significant difference with gluteus minimus and gluteus medius

^d Significant difference with soleus, gluteus minimus, gluteus medius and gastrocnemius

^a Significant difference with soleus and gluteus minimus

main effect on all muscles in the fast walking group (p < 0.01).

The *t* score of muscle thickness in the slow-walking group was the highest for the soleus muscle, followed in order by the gluteus minimus, gluteus medius, rectus femoris, gastrocnemius, vastus intermedius, vastus lateralis, gluteus maximus, psoas major and biceps femoris. The Friedman test revealed a significant main effect for muscle in the slow-walking group (p < 0.05).

The *t* score of muscle thickness in the dependent group was the highest for the gluteus minimus, followed in order by the gluteus medius, soleus muscle, gastrocnemius, psoas major, gluteus maximus, vastus intermedius, biceps femoris, rectus femoris and vastus lateralis. The Friedman test revealed a significant main effect for muscle in the dependent group (p < 0.01). The post hoc test used to compare the *t* score of muscle thickness in the dependent group showed that the quadriceps group of muscles, such as the vastus lateralis and rectus femoris, has significantly smaller values than gluteus minimus, gluteus medius, soleus or gastrocnemius.

Discussion

There were two main findings in this study. First, sarcopenia was lowest for the soleus muscle among the ten muscles of the lower limbs in the elderly who were able to walk independently. Second, a larger magnitude of agerelated decline occurred in the quadriceps group of muscles than in other muscles among the lower limb muscles in the elderly who could not walk independently. To our knowledge, this is the first report showing sarcopenia of individual muscles constituting the lower limb muscles, including those around the hip, knee and ankle joint. In addition, this is the first report showing the relationship between sarcopenia and walking ability in elderly women.

In this study, the measurement of the thickness of ten muscles of the lower limbs showed the smallest degree of atrophy in the soleus muscle in the elderly groups. In addition, no significant difference was seen in the soleus muscle thickness between the young and walking elderly groups. Our findings showed only slight age-related changes of the soleus muscle in elderly people who were able to do ambulatory activity. In general, greater rates of agerelated loss occur among type II fibers, whereas only moderate losses occur among type I fibers (Doherty et al. 1993; Lexell et al. 1988; Roos et al. 1997). Therefore, the postural role of the soleus muscle in the elderly people who were able to do ambulatory activity may protect this muscle from marked reductions in thickness with aging, given its predominant type I composition (Johnson et al. 1973).

Our analyses of the comparison between the elderly groups indicated no difference between the fast- and slowwalking groups in magnitude of decline of all muscles. Previous studies reported that maximum walking speed in the elderly was closely associated with their muscle strength in the lower limbs. However, the relationship between walking speed and muscle strength was non-linear; i.e. there was a threshold value of muscle strength at which walking speed could not improve even if muscle strength increased (Buchner et al. 1996; Ferrucci et al. 1997; Kwon et al. 2001; Rantanen et al. 1998). Our results suggest that elderly women in the slow-walking group had enough muscle mass to give muscle force at the constant standard. We believe that factors other than muscle mass, such as balance control or motor coordination, may contribute to the difference in walking speed between the fastand slow-walking groups.

Although the magnitude of decline of muscle thickness in the dependent elderly group was 49.1-83.0% greater than that in the walking elderly groups, there were no significant differences between the three elderly groups in magnitude of decline of the gluteus minimus and psoas major muscles. These results suggest that despite the marked reduction of overall muscle thickness in the elderly women who did not engage in walking for a long period, a relatively smaller degree of atrophy was found in their deep hip muscles. Studies on rats have frequently reported that experimental cast immobilization results in preferential reduction in slow-twitch muscle mass (Booth 1982; Ohira et al. 1992; Templeton et al. 1988), whereas it was noted that greater decrease of human fast-twitch muscle mass occurred in skeletal muscle as a result of inactivity (Hikida et al. 1989). Because the mass of skeletal muscle is located in a deep part of the muscle and a large proportion of that appears to be slow-type muscle mass, relatively minor atrophy may occur in deep muscles of the hip joint, such as the gluteus minimus and psoas major muscles, even among elderly who do not engage in locomotor activity. Deep muscles in close proximity to a hip joint are believed to provide joint stability (Andersson et al. 1995). Gravity generates muscle loading at the hip joint during sitting activities, which may lead to lesser atrophy in the deep hip muscles even for the elderly who do not engage in locomotor activities. (Hides et al. 2007) also reported that cross-sectional area (CSA) of the multifidus, as measured by MRI, decreased by day 14 of bed rest in healthy male subjects, whereas that of the psoas muscle increased. This may reflect muscle shortening during bed rest. The results of that study suggest that changes in muscle architecture, such as muscle shortening resulting from a long period inactivity may also influence the degree of muscle atrophy.

The t score of muscle thickness in the dependent elderly group was the lowest for quadriceps femoris such as vastus

lateralis and rectus femoris, suggesting particularly marked changes relative to walking disability in this muscle among muscles of the lower limb. Reduced physical activity leads to a loss of skeletal muscle mass. Lower limb muscles, especially antigravity muscles such as the quadriceps femoris that are required for weight-bearing activities (i.e. walking, stair climbing), are likely to be affected most by reduced gravitational loading and inactivity (bed rest). Previous studies have documented that human muscle mass in the quadriceps femoris was reduced by 3% after 7 days (Ferrando et al. 1995), 8% after 20 days (Akima et al. 1997) and 14% after 42 days (Berg et al. 1997) of bed rest. Thus, loss of skeletal muscle mass is exacerbated by periods of inactivity or bed rest. In the present study, a significant impact on muscle atrophy might have been observed in the quadriceps femoris muscle due to inactivity caused by physical disability, because dependent elderly group did not engage in weight-bearing activities for an extended period of time.

This study had some limitations. One limitation arose from measurement of the muscle thickness for determining the muscle mass of lower limb muscles. In general, the muscle cross-sectional area is assumed to reflect muscle strength. In this study, the muscle thickness in the longitudinal axis was measured. However, the obtained measurement values did not reflect the transverse axis.

The other limitation of this study was the small number of subjects. The stringent exclusion criteria, which limited the number of subjects, may also limit the generalisability of the findings.

Another limitation arose from the cross-sectional structure of this study. Thus, for noticeable muscle atrophy of quadriceps femoris in the dependent elderly group, the causality was not clear; i.e. whether muscle atrophy of quadriceps femoris progressed as a result of inactivity or whether muscle atrophy of quadriceps femoris led to disability. In addition, the inclusion criterion for the dependent group selected elderly women who had not walked for more than a half year, which was not standardized as the period when they were not walking.

Further longitudinal studies are required to clarify the association between the degree of sarcopenia in the lower limbs and the locomotor capacity, and to confirm the findings reported in the present study.

Conclusions

We found that the fast- and slow-walking elderly groups showed the same extent of atrophy, while the dependent elderly group showed more severe atrophy. This study suggested that sarcopenia in the lower limbs was least for the soleus muscle in the elderly women who were able to walk independently, and that muscle atrophy in the quadriceps group of muscles was at a maximum in the elderly who had not walked for a long period of time.

Conflict of interest None.

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