

**Objectively-measured sedentary time, habitual physical activity and bone strength in adults aged 62 years: the Newcastle Thousand Families Study**

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## **Abstract**

### **Background**

The influence of sedentary time (ST) and habitual physical activity (PA) on the bone health of middle aged adults is not well known.

### **Methods**

Bone mineral density (BMD) and hip bone geometry were evaluated in 214 men (n=92) and women (n=112) aged  $62.1 \pm 0.5$  years from the Newcastle Thousand Families Study birth cohort. Accelerometry was used to measure PA and ST over four days. Regression models were adjusted for clinical risk factor covariates.

### **Results**

Men were more sedentary than women ( $p < 0.05$ ). ST was negatively associated with spine BMD in men, with 84 minutes more ST corresponding to  $0.268 \text{g.cm}^{-2}$  lower BMD ( $\beta = -0.268$ ;  $p = 0.017$ ). In men, light PA and steps/day were positively associated with bone geometry and BMD. Steps/day was positively associated with bone geometry and femur BMD in women, with a positive difference of 1415 steps/day corresponding to  $0.232 \text{g.cm}^{-2}$  greater BMD ( $\beta = 0.232$ ,  $p = 0.015$ ).

### **Conclusions**

ST was unfavourably associated with bone strength in men born in North East England at age 62 years. Higher volumes of light PA and meeting public health daily step recommendations (reaching 10,000 steps/day) was positively associated with bone health in both sexes.

## **Introduction**

Osteoporosis is defined by the World Health Organisation (WHO) as a 'progressive systemic skeletal disease characterized by low bone mass and microarchitectural deterioration of bone tissue, with a consequent increase in bone fragility and susceptibility to fracture', and is more common with age [1]. In the United Kingdom alone, there are over half a million new fragility fractures annually, and this number is expected to rise by 27% between 2010 and 2025 [2].

There are a number of factors that influence the risk for osteoporosis including glucocorticoids and parental history of osteoporosis. Modifiable factors include physical activity (PA) [3-8]. It is established that bone adapts to its loading environment [3] and that weight-bearing and muscle loading exercises are osteogenic in younger [4] and older populations [5-10]. In the absence of loading, bone strength is reduced [11-13]. As such, habitual sedentary behaviour might have a similar detrimental effect on bone as bed-rest or microgravity environments [11], through reduced muscle forces and skeletal unloading [12, 13].

Older adults are more sedentary than younger individuals [14] and less likely to meet the public health PA recommendations of 10 000 steps per day, or 150 minutes of moderate or 75 minutes of vigorous PA per week [15-17]. However, there is scant evidence on the contribution of objectively-measured habitual PA and sedentary time to bone strength in this population. One study reported significant adverse associations between sedentary time and hip bone mineral density (BMD) in older women, but not men [18], and another reported no associations in women [19]. The aim of this current study was to determine if objectively-measured sedentary time and PA are associated with BMD and bone geometry in men and women at age 62 years. Additionally, sex differences in PA levels and sedentary time, were explored.

## **Methods**

### **Study population**

Two hundred and fourteen adults (men, n = 92; women, n = 122) aged  $62.1 \pm 0.5$  years, from the Newcastle Thousand Families Study birth cohort were included in this analysis. The NTFS was initiated in 1947, when 1142 children born in May and June that year to mothers resident in the city of Newcastle upon Tyne were recruited to a study of the health of children living in the city, in response to the high infant mortality rate and poverty levels in the city at that time [20]. The birth cohort has been described in detail at age 50 years [21]. The current study evaluated participants during the most recent follow up wave in 2011, where 352 participants returned for a clinical examination, having completed health and lifestyle questionnaires. The remainder of the

sample could not be contacted, had moved away from the area, did not wish to take part or had died. Of those who returned for clinical assessments, 214 provided sufficient accelerometry data to be included in the analyses. A favourable ethical opinion was obtained from the Sunderland Research Ethics Committee [Reference 09/H0904/40]. All included study members gave their written informed consent.

### **Accelerometer-measured sedentary time and physical activity**

Sedentary time and PA were assessed using validated Actigraph accelerometry [GT1M, Actigraph Corp, US], which participants were asked to wear for seven consecutive days [22, 23, 24]. Participants were requested to wear the monitor for as much of the day as possible, removing it for washing and bathing, swimming and for bed at night. The Actigraph detects vertical accelerations over a user-specified time interval epoch, and used is used to calculate time spent in activities of different intensity [25]. In this study, an epoch length of 60 seconds was specified. The accelerometer data files were processed using the Actilife Software package. Sedentary time was defined as less than 100 counts per minute [cpm], light PA as 100 - 1951 cpm, moderate intensity PA as 1952 - 5725 cpm and vigorous PA as more than 5725 cpm [25]. As the time spent in vigorous activity was very low, minutes of moderate and vigorous PA [MVPA] were combined. Runs of zero counts lasting more than 60 minutes were excluded, as it was considered the monitor must have been removed for this time. A valid day of recording was defined as one in which more than 500 minutes of monitored on-time were recorded in a 24 hour period [26]; only participants recording at least four valid days of accelerometry were included and only valid days were analysed. The total daily time [in minutes] spent in MVPA were obtained by totalling the duration of all the bouts at each level for each day. The values were then averaged over the number of valid days to derive the percentage of time spent in MVPA per day. Time spent sedentary and time spent in light PA were recorded. The number of daily steps were also obtained, which enabled comparisons with public health recommendations.

### **Bone strength**

Participants wore light-weight clothing with shoes and all jewellery removed. Body weight was measured to the nearest 0.1 kg using calibrated electronic scales and standing height was measured to the nearest 0.1 cm using a stadiometer. BMI was subsequently calculated (weight [kg]/height[m]<sup>2</sup>). Dual energy X-ray absorptiometry (Lunar iDXA, GE Healthcare, Madison, WI) was used to evaluate BMD (g.cm<sup>-2</sup>) of the lumbar spine (L1 - L4), total hip and sub-regions of the hip. Structural geometry of the right proximal femur was estimated from the acquired scans by utilising the GE Lunar Advanced Hip Structural Analysis programme. Cross-sectional area

(CSA in cm<sup>2</sup>), cross-sectional moment of inertia (CSMI, in cm<sup>4</sup>) and femoral strength index (FSI) values were derived. Precision (CV) for the iDXA measurements are 0.4% for lumbar spine BMD and 0.9% for femoral neck BMD [27]. DXA precision error for CSMI and CSA are 3.7% and 3.1% respectively [28]. Scan analysis was performed by the same trained operator using the Lunar enCORE software (version 12.0, GE Healthcare, Madison, WI).

### **Clinical risk factors and falls**

Clinical risk factors for bone fragility were evaluated and treated as covariates in the analysis. In addition to sex and BMI, the covariates included previous fracture, parent hip fracture, current smoking, rheumatoid arthritis, corticosteroid use and alcohol consumption, all ascertained from general health questionnaires. Participants also recorded the number of falls within the last 12 months.

### **Statistical analysis**

All analyses were done using SPSS version 22.0 (IBM Corporation, US). Comparisons of BMI and femoral neck BMD between participants with (n = 214) and without (n = 138) accelerometer data were made by independent T-tests. Normality was tested using the Kolmogorov-Smirnov test. In men, total hip BMD, Wards BMD and FSI were not normally distributed, and in women, total hip BMD was not normally distributed. Therefore, the natural log-transformed values for these variables were created to be used in bivariate analysis and modelling. All descriptive data are expressed as means and their standard deviations (SDs). Associations with bone strength were initially explored using Pearson's correlation coefficients. Covariates (including CRFs) with a p-value < 0.05 (two tailed test) were included in the linear regression models, which were used to examine sex-specific relationships between bone strength, PA levels and sedentary time, with bone strength parameters as the dependent variables. Standardised coefficients enabled the interpretation of differentiation in bone strength parameters according to sedentary time (minutes) or steps per day.

### **Results**

There were fewer men (44% v 51%) and more women (49% v 56%) in the current sample than in the 1947 cohort (p = 0.008). There were no differences in BMI or femoral neck BMD between those who did and did not have accelerometry data (p > 0.05). Over half of women had used hormone replacement therapy (58%). Clinical

risk factors for osteoporosis are given in Table I and Table II presents the anthropometric, PA and bone strength results for the cohort by sex.

Men spent more time sedentary and less time in light PA than women (Table 2). Sedentary time ranged from 305 to 676 minutes per day in men and 283 to 691 minutes per day in women. Time spent in MVPA in women ranged from 1 to 115 minutes per day and in men, 0 to 111 minutes. The average daily steps ranged from 1371 to 19 294 in women and 889 to 19 922 in men, and mean values fell short of the recommended 10 000 steps/day [15 - 17]. Twelve women (10%) and 7 men (8%) met or exceeded 10 000 steps per day. There were no differences in sedentary time or PA levels between participants who had or had not fallen in the last 12 months (Table III).

Lumbar spine BMD, total hip BMD, CSMI and CSA were higher in men than women, but there were no differences in femoral neck BMD and FSI (Table 2). In this cohort, 63% (n = 71) of women and 48% (n = 44) of men were either osteopenic or osteoporotic.

## **Women**

Sex-specific unadjusted correlation coefficients between dependent and independent variables are given in Table IV. BMI was positively associated with all BMD outcomes. There were no associations between self-reported hormone therapy and BMD. Smoking was associated with lower CSA, and use of corticosteroids was associated with lower total hip, femoral neck and Wards area BMD, and lower CSMI. Higher number of steps per day were associated with higher femoral neck and Wards area BMD, and with a greater CSA.

After adjustment for BMI and use of corticosteroids, steps per day remained associated with femoral neck BMD ( $\beta = 0.232 \text{ g.cm}^{-2}$ ; CI 0.228 - 0.236;  $p = 0.015$ ). The standardised coefficient corresponds to a  $0.232 \text{ g.cm}^{-2}$  higher femoral neck BMD for 1415 more steps per day. After adjusting for BMI ( $\beta = 0.174 \text{ mm}^2$ ) and smoking ( $\beta = -0.256 \text{ mm}^2$ ), the regression coefficient for CSA and steps per day was  $\beta = 0.253 \text{ mm}^2$  (CI 0.249 - 0.257;  $p = 0.006$ ).

## **Men**

BMI was associated with lumbar spine BMD but no other measures of bone strength, and smoking was associated with lower CSA (Table IV). Sedentary time was associated with lower lumbar spine, total hip and Wards area BMD, and time spent in light PA was positively associated with lumbar spine, femoral neck, Wards area and total

hip BMD (Table IV). Steps per day were positively associated with total hip BMD and all hip bone geometry outcomes, FSI, CSMI and CSA (Table IV).

Lumbar spine BMD remained associated with sedentary time after adjustment for BMI ( $\beta = -0.268 \text{ g.cm}^{-2}$ ; CI  $-0.347 - -0.189$ ;  $p = 0.017$ ). The standardised coefficient corresponds to  $0.268 \text{ g.cm}^{-2}$  lower lumbar spine BMD for 84 more minutes per day spent sedentary. As TH and Wards area BMD were transformed a similar interpretation was not feasible. After adjustment for BMI, lumbar spine BMD remained positively associated with light PA ( $\beta = 0.322 \text{ g.cm}^{-2}$ ; CI  $0.237 - 0.395$ ;  $p = 0.004$ ), corresponding to a  $0.322 \text{ g.cm}^{-2}$  higher lumbar spine BMD for 82 minutes more per day spent in light PA. The linear regression coefficient between steps per day and CSA, after adjustment for smoking was  $\beta = 0.210 \text{ mm}^2$  (CI  $0.109 - 0.319$ ;  $p = 0.05$ ).

## **Discussion**

### **Main findings of this study**

In this study of men and women aged 62 years and born in the North East of England, there were negative associations between sedentary time and BMD in men. Notably men were also found to be more sedentary than women. Although there were no associations with MVPA, the amount of time men spent in light PA was favourably associated with spine and hip BMD. In both sexes, a higher number of steps/day was associated with higher BMD and positive bone geometrical properties, suggesting that a more active lifestyle was associated with better bone health.

### **What is already known on this topic**

Previous studies have reported negative associations between objectively-measured sedentary time and bone strength only in women [18, 29]. A negative association between self-reported sedentary time and bone strength in men has been reported previously [30]. Self-reported non-study use of the internet in male adolescents was associated with lower total body and femoral neck bone mineral content [30]. In older men, greater self-reported sitting time has been shown to be associated with reduced lean mass and an increased risk of sarcopenia [31], suggesting a similar disuse trajectory.

We observed positive associations between light PA and bone strength in men and steps per day and bone strength in both sexes. This is in agreement with previous cross-sectional and prospective studies measuring PA by self-report [32-35] suggesting the importance of habitual PA for bone health. Most previous studies have

reported positive associations between PA and femur BMD only, and not the lumbar spine [18, 32-35]. Although we found no associations at the lumbar spine in women, BMD at this site was greater in men who engaged in higher levels of light PA and who recorded more steps per day. This might reflect the site-specific mechanics of exercise [36] and it should be considered that bone can become accustomed to repetitive patterns of loading [e.g. walking and running] [37]. It is also possible that earlier life PA might have impacted on later life bone strength [38].

### **What this study adds**

To our knowledge this is the first study to demonstrate an unfavourable relationship between objectively-measured sedentary time and bone strength in men. Our study found that additional 84 minutes per day spent sedentary was associated with a 0.268 g.cm<sup>-2</sup> lower lumbar spine BMD. This is clinically relevant given that in real terms, it could mean the difference between having normal T-score or an osteopenic T-score. It should also be considered that the strength of the association between sedentary time and total hip BMD (R = -0.195) was not dissimilar to that of smoking (R = -0.209), which is a known risk factor for osteoporosis. The findings of the present study together with evidence elsewhere, suggest that sedentary behaviour is a modifiable factor influencing bone health in men, as well as in women [18, 29]. Our findings also support recent calls for interventions aimed at reducing sedentary behaviour including sitting time, in older adults [39], following accumulation of evidence linking sedentary behaviours with cardiometabolic health [40]. We suggest that there may be benefits in promoting a reduction in sedentary time for improving bone health in men in addition to the well documented cardiometabolic benefits.

Interestingly, MVPA was not associated with any bone health outcomes, unlike in several previous accelerometry-based studies, which have reported positive associations between comparable levels of MVPA and femoral neck, total hip and tibial BMD [18, 41]. Steps/day and light PA were more predictive of bone strength outcomes than MVPA. In women, the regression coefficient describing the relationship between steps per day and femoral neck BMD after correction for significant CRFs, indicates that a change of 1415 steps per day corresponds to a 0.232 g.cm<sup>-2</sup> higher femoral neck BMD. This would constitute as a clinically meaningful difference in terms of equivalent diagnostic T-score. The number of steps is equivalent to around 15 minutes extra of moderate intensity walking per day [42]. This figure is also similar to that derived by Chastin et al. of 0.300 g.cm<sup>-2</sup> higher total femur BMD for every 10 minutes of MVPA in men [18]. Walking is an accessible and inexpensive form of PA with multifarious health benefits, and the aforementioned PA levels are comparable to those shown to deliver

improved cardiovascular health, lower cancer and mortality rates, through active commuting in adults [43] and of relevance to policy makers, support the current public health recommendation of 10 000 steps/day [15 - 17].

It is notable that in both sexes, the strength of the associations observed between steps/day and light PA with BMD were greater or comparable to those for smoking, corticosteroid use and BMI. Furthermore, adjustment for CRFs did not change the positive associations between PA and bone strength. These findings suggest that PA interventions for bone health in older adults could include light PA, associated with achievable benefits regardless of BMI, smoking and corticosteroid use. However, there are currently no specific guidelines for light PA or for reducing sedentary time at population level. Such an approach might be particularly useful for older adults who may find MVPA difficult to achieve independently, and who may, following retirement, be more likely to engage in sedentary behaviours such as increased television viewing time [44].

### **Limitations of this study**

The firstly limitation is that although we found no associations between MVPA and bone strength, accelerometry is not able to classify all types of PA including some high impact loading exercises [e.g resistance training], which are known to be especially oestrogenic [12, 45]. Secondly, it has been hypothesised that attaining a 10% higher peak bone mass in young adulthood could delay the development of osteoporosis by around 13 years [46]. Therefore it is possible that individuals who have greater BMD at age 62 years, had previously achieved a higher peak bone mass at a time when the skeleton was especially responsive to exercise [4]. It is not possible to confirm this from the current cross-sectional analysis and there is risk for reverse causation arising from morbidity, none-the-less, elsewhere, bone loss has been clearly evidenced in adults who are sedentary or in bed rest [18]. Finally, caution should be taken when making inferences given that the study was performed among members of a birth cohort born in Newcastle upon Tyne, UK who were aged 61 to 63 years. This age group does however, have particular public health importance because the risk of osteoporosis and fracture is increased compared to younger ages, and effective risk factor modification is still viable.

### **Conclusions**

In conclusion, the results from this present study demonstrate an unfavourable relationship between sedentary time and bone strength in men. Light PA and steps taken per day were associated with better bone strength in older men and women independent of clinical risk factors, and more so than MVPA. These findings provide

impetus for interventions in older adults that seek to reduce sedentary time and increase habitual PA levels regardless of intensity.

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### **Conflict of interest**

None declared.

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**Table I.** Clinical risk factors for osteoporosis and fracture, age 62 years

	<i>Women</i>	<i>Men</i>
Use of corticosteroids [ $>3$ months]	7 (5.8%)	5 (5.7%)
Parental fracture	4 (3.3%)	6 (6.7%)
Previous fracture	21 (17.1%)	16 (18.2%)
Hip fracture	1 (0.8%)	2 (2.2%)
Rheumatoid arthritis	1 (0.8%)	1 (1.1%)
Current smoking	9 (7.4%)	12 (13.6%)
Alcohol consumption $\geq 3$ units/daily	15 (12.4%)	23 (26.1%)
Fallen within last 12 months	20 (16.5%)	16 (17.4%)

**Table II.** Anthropometric, bone strength and physical activity parameters (mean, SD)

	<i>Women</i>	<i>Men</i>	<i>P</i>
Height, cm	161.2 ± 6.8	173.3 ± 7.2	< 0.001
Weight, cm	73.3 ± 14.5	82.7 ± 15.2	< 0.001
Body mass index, kg.m <sup>-2</sup>	28.2 ± 5.4	27.6 ± 4.9	0.346
Sedentary time, minutes/day	493.6 ± 69.7	523.2 ± 83.7	0.008
Light activity time, minutes/day	275.9 ± 59.7	253.2 ± 82.2	0.028
MPVA time, minutes/day	22.1 ± 19.9	23.8 ± 17.9	0.525
Steps per day	8,035.9 ± 1,415.2	8,113.9 ± 1,572.1	0.970
Lumbar spine BMD, g.cm <sup>-2</sup>	1.105 ± 0.175	1.207 ± 0.168	< 0.001
Lumbar spine T-score	- 0.7 ± 1.4	- 0.2 ± 1.4	< 0.001
Total hip BMD, g.m <sup>-2</sup>	0.987 ± 0.138	1.077 ± 0.155	< 0.001
Femoral neck BMD, g.cm <sup>-2</sup>	0.918 ± 0.169	0.961 ± 0.166	0.070
Femoral neck T-score	- 0.5 ± 1.2	- 0.8 ± 1.2	0.070
Wards area BMD, g.cm <sup>-2</sup>	0.739 ± 0.161	0.771 ± 0.176	0.179
Trochanter BMD, g.cm <sup>-2</sup>	0.821 ± 0.135	0.985 ± 0.178	< 0.001
Femoral Strength Index	1.5 ± 0.5	1.5 ± 0.4	0.640
CSMI [mm <sup>4</sup> ]	10,092.6 ± 2,603.5	17,231.5 ± 4,480.2	< 0.001
CSA [mm <sup>2</sup> ]	141.2 ± 29.9	173.9 ± 31.5	< 0.001

MPVA: moderate to vigorous physical activity; BMD: bone mineral density; CSMI: cross-sectional moment of inertia; CSA: cross-sectional area

**Table III.** Sedentary time, physical activity levels and reported falls in the last 12 months (mean, SD)

	<i>Women</i>			<i>Men</i>		
	<i>Fall (n=17)</i>	<i>No fall (n=105)</i>	<i>P</i>	<i>Fall (n=15)</i>	<i>No fall (n=77)</i>	<i>P</i>
Sedentary time, min/day	483.0 ± 74.1	495.3 ± 69.1	0.516	517.1 ± 70.1	526.9 ± 87.9	0.697
Light physical activity, min/day	271.8 ± 62.7	276.6 ± 59.6	0.767	233.4 ± 90.3	260.5 ± 82.2	0.275
MVPA, min/day	20.6 ± 26.6	23.5 ± 20.6	0.614	23.9 ± 13.7	24.1 ± 18.8	0.984
Steps per day	6348.4 ± 3591.3	7017.7 ± 3001.1	0.572	7351.4 ± 3744.7	6545.1 ± 3235.3	0.400

**Table IV.** Unadjusted Pearson correlation coefficients [p values] of physical activity levels and bone strength parameters

	<i>Sedentary time</i>	<i>Light physical activity</i>	<i>MPVA</i>	<i>Steps per day</i>	<i>BMI</i>	<i>Smoking</i>	<i>Corticosteroids</i>
<i>Women</i>							
Lumbar spine BMD	0.017 [0.860]	0.038 [0.699]	-0.145 [0.129]	0.123 [0.183]	0.369 [<0.001]*	-0.051 [0.595]	-0.186 [0.051]
Total hip BMD <sup>†</sup>	-0.018 [0.856]	0.079 [0.415]	-0.024 [0.806]	0.145 [0.116]	0.326 [<0.001]*	-0.116 [0.227]	-0.225 [0.018]*
Femoral neck BMD	-0.077 [0.431]	0.074 [0.447]	-0.100 [0.273]	0.203 [0.037]*	0.255 [0.008]*	-0.141 [0.145]	-0.208 [0.030]*
Wards area BMD	0.051 [0.626]	0.043 [0.658]	-0.030 [0.753]	0.183 [0.057]	0.248 [0.010]*	-0.176 [0.068]	0.211 [0.028]*
Trochanter BMD	-0.024 [0.805]	0.079 [0.417]	-0.072 [0.458]	0.156 [0.109]	0.330 [<0.001]*	-0.039 [0.683]	0.187 [0.051]
Femoral Strength Index	-0.094 [0.334]	0.159 [0.101]	0.126 [0.194]	0.172 [0.079]	n/a	-0.115 [0.233]	0.069 [0.477]
Femur CSMI	-0.014 [0.888]	-0.002 [0.986]	0.088 [0.338]	0.171 [0.080]	0.086 [0.378]	-0.004 [0.964]	0.185 [0.050]*
Femur CSA	-0.045 [0.645]	0.048 [0.627]	0.052 [0.574]	0.243 [0.011]*	0.137 [0.174]	-0.235 [0.014]*	0.162 [0.093]
<i>Men</i>							
Lumbar spine BMD	-0.221 [0.050]*	0.260 [0.021]*	0.036 [0.741]	0.057 [0.614]	0.229 [0.033]*	-0.037 [0.736]	-0.050 [0.653]
Total hip BMD <sup>†</sup>	-0.195 [0.048]*	0.257 [0.021]*	0.041 [0.705]	0.282 [0.038]*	0.160 [0.137]	-0.209 [0.050]*	-0.132 [0.232]
Femoral neck BMD	-0.131 [0.242]	0.230 [0.039]*	0.020 [0.981]	0.145 [0.190]	0.100 [0.353]	-0.021 [0.851]	-0.023 [0.837]
Wards area BMD <sup>†</sup>	-0.264 [0.017]*	0.254 [0.023]*	0.052 [0.626]	0.175 [0.110]	-0.012 [0.909]	-0.058 [0.595]	-0.081 [0.462]
Trochanter BMD	-0.083 [0.467]	0.168 [0.135]	0.075 [0.483]	0.117 [0.292]	0.101 [0.349]	-0.129 [0.239]	-0.209 [0.056]
Femoral Strength Index <sup>†</sup>	-0.093 [0.407]	0.064 [0.571]	0.113 [0.210]	0.320 [0.002]*	n/a	-0.150 [0.169]	0.011 [0.921]
Femur CSMI	-0.174 [0.121]	0.117 [0.297]	0.198 [0.060]	0.261 [0.017]*	0.028 [0.795]	-0.165 [0.129]	-0.045 [0.682]
Femur CSA	-0.166 [0.138]	0.171 [0.126]	0.148 [0.166]	0.308 [0.005]*	0.135 [0.208]	-0.275 [0.010]*	-0.119 [0.280]

BMD: bone mineral density; CSMI: cross-sectional moment of inertia; CSA: cross-sectional area; BMI: body mass index; \* significant association at  $p < 0.05$ ; <sup>†</sup>Log transformed