



Associations of protein source, distribution and healthy dietary pattern with appendicular lean mass in oldest-old men: the Helsinki Businessmen Study (HBS)

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Key summary points

Aim To investigate how food and dietary intakes, protein daily distribution and source were associated with appendicular lean mass (ALM)/m² in the oldest-old community-dwelling men.

Findings ALM/m² was associated with total protein intake, source and distribution as well as fruit and vegetable intakes.

Message Not only protein intake, but also source and distribution as well as healthy overall diet characterized by abundant amounts of fruits and vegetables were important in maintaining muscle mass in the oldest-old men in our study.

Abstract

Purpose We explored how food and dietary intakes, protein daily distribution and source are associated with appendicular lean mass (ALM)/m² of the oldest-old community-dwelling men.

Methods Cross-sectional analyses of Helsinki Businessmen Study (HBS, mean age 87 years) participants who came to clinic visit in 2017/2018. Nutritional status, physical performance and fasting blood samples were measured. Food and dietary intakes were retrieved from 3-day food diaries. Body composition was measured and appendicular lean mass (ALM) per m² was dichotomized as ALM/m² < 7 kg/m² and ≥ 7 kg/m². Differences between lower and higher ALM were analyzed using *t* test or Mann–Whitney *U* test. Analysis of covariance was used to investigate independent associations with ALM/m².

Results Random sample of 130 participants took part in the medical examinations, 126 returned food diaries, and 102 underwent DXA-scan. ALM/m² was associated with total protein (*p* = 0.033), animal protein (*p* = 0.043) and meat protein (*p* = 0.033) intakes. Protein distribution between daily meals differed at lunch; those with higher ALM/m² ate more protein (*p* = .047) at lunch. Consumption of fruits, vegetables (*p* = 0.022) and meat (*p* = 0.006) was associated with ALM/m².

Conclusion Protein intake, source and distribution as well as fruit and vegetable intakes were associated with higher ALM in oldest-old men.

Study registration The study is registered with ClinicalTrials.gov identifier: NCT02526082.

Keywords Appendicular lean mass · Protein intake · Protein distribution · Protein source · Fruits and vegetables · Animal protein · Meat intake

Introduction

Skeletal muscle is highly important for metabolic health and maintenance of physical function in older age [1]. Skeletal muscle mass and strength decline steadily after the fourth decade of life and the rate of decline are accelerated with aging [2]. Loss of skeletal muscle mass is an independent risk factor for osteoporosis, falls and fractures, impaired functioning and mortality [3].

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Skeletal muscle is also a major organ of glucose metabolism and thus low skeletal muscle may impair glucose tolerance and insulin resistance [4]. For these reasons, there has been a great interest to define lifestyle-related risk factors of skeletal muscle loss. Of nutritional factors, especially inadequate protein intake has been associated with accelerated loss of lean mass, and an increased risk of functional impairments, whereas adequate protein intake has been linked to muscle protein balance and slower rate of muscle mass decline [5]. Higher protein intake has also been associated with increase in muscle mass especially in relation to exercise [6]. Of dietary patterns, Mediterranean diet has been positively associated with muscle mass in older adults [7, 8]. However, data on the fastest growing age group of the oldest old (> 85 years) are very limited with respect to nutrition and muscle mass.

To address this limitation, we explored detailed food and nutrition intakes, distribution of daily protein intake and source between lower and higher appendicular lean mass (ALM) groups in oldest old, community-dwelling men.

Methods

In the Helsinki Businessmen Study (HBS) socioeconomically homogenous cohort of men, born between 1919 and 1934, have been followed-up since the 1960s [9]. In the present cross-sectional analysis, we report findings from the most recent clinic visit including a random sub-cohort of home-living survivors of HBS in 2017–2018 (mean age 87 years of age). At the clinic visit, body mass index (BMI) was calculated as weight (kg/height (m) squared), Mini Nutritional Assessment (MNA) [10] and Short Physical Performance Battery (SPPB) [11] were performed as instructed, body composition measured with DXA-scans, and appendicular lean mass (ALM) per m^2 was calculated. ALM/ m^2 was dichotomized as $< 7 \text{ kg}/m^2$ and $\geq 7 \text{ kg}/m^2$ according to the classification of Gould et al. [12]. Blood insulin and glucose levels were analyzed from blood samples after the 12 h fast. Food, energy and nutrient intakes, daily protein distribution and protein source (amounts of vegetable, animal; milk, meat, fish, and egg proteins) were calculated from the 3-day food diaries.

Statistical significance for group differences was evaluated using independent t test for evenly distributed continuous variables and Mann–Whitney U test for unevenly distributed variables. In addition, we used analysis of covariance (ANCOVA) to investigate independent associations with

ALM/ m^2 . Adjustments were made for age, BMI, protein intake, g, insulin levels and tea drinking. Analyses were performed using the SPSS statistical program, version 24 (SPSS IBM, Armonk, NY, USA).

Selection of covariates

Covariates were selected based on the results of our analysis and prior research. Insulin was selected as a covariate because muscle is a major organ for glucose metabolism [4], tea because it was inversely associated with higher ALM in test, and protein intake for its importance to muscle. Age is associated with loss of skeletal muscle mass and BMI with higher muscle mass.

Results

130 men participated in the clinic visit, 126 returned food diaries and 102 underwent the DXA scan. Age, MNA, SPPB or insulin and glucose levels did not differ significantly between the two ALM/ m^2 groups. Higher ALM/ m^2 was associated with higher BMI ($p < 0.001$), total protein intake ($p = 0.033$), consumption of protein of animal origin ($p = 0.043$) and meat protein ($p = 0.033$). Protein distribution between daily meals differed at lunch only; those in the higher ALM/ m^2 group ate 26 g protein at lunch, compared to 20 g in those with low ALM/ m^2 ($p = 0.047$, Table 1). Of foods consumed, total fruits and vegetables intakes differed significantly between the ALM/ m^2 groups; those with higher ALM/ m^2 consumed more fruits and vegetables: 341 g/d compared to 243 g/d ($p = 0.022$, Table 2). Meat intake was higher in those who had higher ALM/ m^2 ($p = 0.006$), whereas tea intake was negatively associated with ALM/ m^2 ($p = 0.027$).

General linear model confirmed the bivariate findings with respect to protein intake and BMI to ALM/ m^2 . Protein intake remained significant to ALM/ m^2 after adjusting for age, and additionally with insulin levels, BMI, and tea intake (Table 3).

Discussion

In this study, total protein intake, protein source and distribution as well as meat, fruit and vegetable intakes were associated with higher ALM/ m^2 in oldest old community-dwelling men, whereas tea drinking was inversely associated with

Table 1 Baseline characteristics, protein distribution between daily meals and protein source by level of appendicular lean mass (ALM)/m²

Baseline characteristics	ALM groups		<i>p</i> value ^a
	ALM < 7 kg/m ² <i>n</i> = 45	ALM ≥ 7 kg/m ² <i>n</i> = 57	
Age, years	87 (3)	87 (3)	0.670
BMI, kg/m ²	24.7 (2)	26.6 (3)	< 0.001
MNA points (range 0–14) (SD)	13 (1)	13 (1)	0.249
SPPB points (range 1–12) (SD)	9 (3)	10 (2)	0.215
Insulin, mmol/L	8.8 (5.1)	7.3 (3.3)	0.112
Glucose, mmol/L	6.2 (1.0)	6.2 (0.8)	0.905
Protein distribution between daily meals, (g)			
Breakfast, g (SD)	16 (8)	16 (7)	0.927
Morning snack, g (SD)	2 (3)	4 (8)	0.071
Lunch, g (SD)	20 (12)	26 (15)	0.047
Dinner, g (SD)	20 (20)	19 (16)	0.692
Afternoon snack, g (SD)	5 (8)	5 (7)	0.615
Evening snack, g (SD)	6 (8)	8 (7)	0.141
Total Protein, g (SD)	69 (24)	79 (21)	0.033
Protein % of total energy	19.4% (3.3)	17.8% (3.3)	0.016
Protein source, g			
Animal protein total, g (SD)	48 (21)	56 (18)	0.043
Meat protein, g (SD)	19 (12)	24 (13)	0.033
Milk protein, g (SD)	16 (10)	18 (10)	0.374
Egg protein, g (SD)	2 (3)	2 (3)	0.943
Fish protein, g (SD)	11 (12)	12 (11)	0.717
Plant protein, g (SD)	21 (7)	22 (7)	0.373
Total protein, g (SD)	69 (24)	79 (21)	0.033

BMI body mass index, *MNA* Mini Nutritional Assessment, *SPPB* short physical performance battery, *SD* standard deviation, *ALM* appendicular lean mass

^aDifference between higher and lower ALM was tested with independent *t* test in even distributed variables and Mann–Whitney *U* test for unevenly distributed variables

ALM/m². Only participants in the higher ALM/m² group reached the amount of protein in a single meal considered to be sufficient for effective protein synthesis.

Proteins of animal origin are high in essential amino acids important for the muscle. Especially the amino acid leucine that is abundant in foods of animal origin has been shown to be important for muscle development and strength [13]. Therefore, it was not surprising that meat protein and meat consumption were associated with higher ALM/m² in our study. Earlier studies have suggested that even protein distribution in daily meals would be most beneficial for older people in relation to muscle health [14]. However, meals with high protein bolus have also been found to be beneficial [15]. In our study protein distribution was relatively even in both ALM groups, as breakfast, lunch,

and dinner were the meals with the highest protein intake, whereas the daily snacks contained relatively low amounts of protein. It has been suggested that ingestion of approximately 25–30 g of protein per meal maximally stimulates muscle protein synthesis in older people [13]. Only the participants classified under the higher ALM group reached this amount in a single meal at lunch, which was the only meal that differed significantly between the ALM groups in our study. The lower ALM group did not reach this threshold in any of their daily meals. These findings underline the importance of educating older people about timing and distribution of their protein intake.

Those with higher ALM ate more fruits and vegetables, which are an essential part of several healthy dietary patterns, including Mediterranean, Nordic, and DASH diets.

Table 2 Food, energy and nutrient intake by level of appendicular lean mass (ALM)/m²

Food intake/day	ALM groups		<i>p</i> values ^a
	ALM/m ² < 7 kg <i>n</i> = 45	ALM/m ² ≥ 7 kg <i>n</i> = 57	
Fruits and berries, g (SD)	112 (119)	170 (183)	0.056
Vegetables, g (SD)	138 (104)	176 (153)	0.163
Total fruits and vegetables, g (SD)	243 (163)	341 (265)	0.022
Whole grain products, g (SD)	98 (60)	102 (65)	0.738
Other grain products, g (SD)	233 (130)	242 (148)	0.749
Legumes, g (SD)	9 (24)	6 (18)	0.448
Nuts, g (SD)	1 (3)	8 (20)	0.060
Fish, g (SD)	56 (56)	66 (61)	0.364
Milk products, g (SD)	280 (222)	347 (220)	0.138
Meat, g (SD)	89 (48)	120 (60)	0.006
Egg, g (SD)	17 (30)	15 (26)	0.761
Alcohol, g (SD)	7 (10)	4 (7)	0.136
Tea, g (SD)	87 (141)	160 (177)	0.027
Coffee, g (SD)	218 (159)	294 (216)	0.051
Energy and nutrient intakes			
Energy, kcal (SD)	1550 (391)	1634 (358)	0.262
Protein, g (SD) g/kg BW/d	69 (24) 0.95 (0.3)	80 (21) 0.99 (0.24)	0.033 0.584
Carbohydrates, g (SD)	164 (44)	170 (43)	0.505
Fat, g (SD)	63 (21)	67 (22)	0.314
Vitamin D, µg (SD)	9 (8)	10 (7)	0.310
Vitamin E, mg (SD)	10 (4)	11 (6)	0.181
Iron, mg, (SD)	10 (3)	11 (3)	0.174

BMI body mass index, *SD* standard deviation, *ALM* appendicular lean mass

^aDifference was between higher and lower ALM was tested with independent *t* test

Fruits and vegetables contain high amounts of vitamins, minerals, antioxidants, and other bioactive compounds and in addition, alkaline salts may also be important in preserving muscle [7, 8, 15]. Tea drinking was associated with lower ALM/m², but as we did further testing, we found that tea drinking was associated with lower BMI which would explain the result.

The strengths of this study include its robust findings—despite the relatively small study sample—and the fact that there are few other studies on oldest-old (> 85 years of age) people. Limitations involve generalization: the survivors of HBS differ in many ways from the general population by being men from upper socioeconomic class; and the cross-sectional design of the

study, which prevents drawing conclusions about causal relationships.

Conclusions and implications

Our study extends previous findings on the importance of protein intake and the threshold in a single meal in the maintenance of muscle mass of the oldest-old. Moreover, fruit and vegetable intakes, emphasized in healthy dietary patterns, were also important for muscle mass. Therefore, the importance of protein intake and distribution, as well as healthy dietary patterns should be highlighted also for the oldest-old.

Table 3 ANCOVA models of factors associated with appendicular lean mass (ALM)/m²

	<i>B</i>	95% confidence interval		<i>p</i> value
		Lower bound	Upper bound	
Model 1				
Intercept	9.724	5.641	13.807	<0.001
Age	− 0.038	− 0.085	0.009	0.111
Protein, g	0.009	0.003	0.015	0.003
Adjusted <i>R</i> ²	0.087			
Model 2				
Intercept	5.205	1.161	9.249	0.012
Age	− 0.018	− 0.060	0.025	0.406
Protein, g	0.005	− 0.026	0.825	0.067
BMI	0.119	0.073	0.165	<0.001
Adjusted <i>R</i> ²	0.273			
Model 3				
Intercept	5.368	1.379	9.357	0.009
Age	− 0.018	− 0.060	0.024	0.392
Protein g	0.005	0.000	0.011	0.063
BMI	0.124	0.078	0.169	<0.001
Insulin	− 0.033	− 0.061	− 0.005	0.020
Adjusted <i>R</i> ²	0.300			
Model 4				
Intercept	5.368	1.437	9.299	0.008
Age	− 0.017	− 0.058	0.024	0.420
BMI	0.120	0.099	0.185	<0.001
Protein, g	0.418	0.076	0.165	0.048
Insulin	− 0.030	− 0.056	0.003	0.032
Tea, g/d	− 0.001	− 0.001	<0.000	0.052
Adjusted <i>R</i> ²	0.320			

Bold values indicate the amount of observed variation that can be explained by the model's inputs

B unstandardized beta, *BMI* body mass index, *BW* body weight, *kg* kilograms, *g* grams, *d* day

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Data availability The data can be enquired from Professor Timo Strandberg, timo.strandberg@oulu.fi.

Compliance with ethical standards

Conflict of interest SKJ: reports no conflict of interest. AU: reports no conflict of interest. MK: was supported by NordForsk, the Academy of Finland (311492), and Helsinki Institute of Life Science. TES: reports having various educational and consultative cooperation with several companies, including Nutricia, Abbott, Amgen, Merck, Pfizer, Novartis, and Novo-Nordisk; a minor amount of stock in Orion Pharma; and is a board member and former president of executive board of European Union Geriatric Medicine Society which has cooperation also with the nutrition industry.

Ethical approval The study protocol was approved by the Ethics Committee of the Helsinki University Hospital, Department of Medicine. The study is registered with ClinicalTrials.gov identifier: NCT02526082.

Informed consent All participants signed an informed consent.

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