



Influence of skeletal muscle mass and fat mass on the metabolic and inflammatory profile in sarcopenic and non-sarcopenic overfat elderly

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Abstract

Background Sarcopenic elderly present low muscle mass and strength, however, it is not clear if the inflammatory and metabolic profile is more related to low lean mass or high fat mass in sarcopenic and non-sarcopenic overfat elderly.

Aim To verify the difference in inflammatory and metabolic responses in sarcopenic and non-sarcopenic overfat elderly and the relationship between these markers, body composition, and strength in this population.

Methods Fifty-seven elderly were divided into two groups: sarcopenic ($n = 30$) and non-sarcopenic ($n = 27$). Body composition was evaluated with octopolar bioimpedance. Total cholesterol, high-density lipoprotein cholesterol, triacylglycerol, glucose, cortisol, leptin, adiponectin, Plasminogen activator inhibitor-1 (PAI-1), TNF- α , IL-6, IL-8, and IL-10 were assessed. The handgrip test was used to evaluate strength.

Results When comparing the inflammatory profile, sarcopenic individuals showed greater adiponectin concentration ($p = 0.019$), adiponectin/fat mass ratio ($p < 0.001$), adiponectin/visceral fat ($p < 0.001$), and higher PAI-1 ($p = 0.019$) than non-sarcopenic overfat elderly. After adjusting the inflammatory profile by skeletal muscle mass the significant differences between groups were maintained ($p < 0.05$) but no significant differences between groups were observed when adjusting by fat mass, despite a tendency to a significant difference for adiponectin concentration ($p = 0.06$). In addition, after adjusting leptin by fat mass there was a statistically significant lower concentration in the sarcopenic compared to non-sarcopenic overfat elderly.

Conclusion Non-sarcopenic overfat elderly presented lower anti-inflammatory and anti-atherogenic responses than sarcopenic elderly. Furthermore, fat mass but not skeletal muscle mass seem to change these responses.

Keywords Sarcopenic · Inflammation · Elderly

Introduction

Aging, associated with high muscle mass loss can result in Sarcopenic syndrome [1], which leads to decreased muscle strength, and impaired locomotion and balance, thus increasing the number of falls due to frailty [2]. Low lean mass plus low muscle strength are predictors of mortality risk among elderly people [3]. Furthermore, muscle mass has an important role in inflammatory and metabolic response, given that cytokines such as IL-6 originate from muscle tissue (in exercise). Furthermore, IL-6 may have an anti-inflammatory effect by inhibiting TNF- α secretion [4], and increasing fat oxidation and glucose uptake via AMPK activation [5]. Not only this, but by activating muscle satellite cells, IL-6 initiates the hypertrophic process [6].

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Several cross-sectional [7] and longitudinal studies [8, 9] have verified the relationship between muscle mass and inflammatory and metabolic markers. Aleman et al. [8] conducted a 5-year longitudinal study with 115 men and women aged 60–84 and concluded that the risk of total and appendicular skeletal muscle mass loss was 1.29 times higher per unit of increase in IL-6 (pg/ml) and 1.28 times higher per unit of increase in high-sensitivity C-reactive protein (CRP) (mg/l).

Recently, studies have used high body fat combined with low skeletal muscle mass to define sarcopenic obesity [10]. Yang et al. [11] performed a cross-sectional study in 844 people aged 65 years old living in the community and compared the inflammatory response among sarcopenic, obese, sarcopenic obese, and non-sarcopenic or non-obese individuals and observed no difference in the serum levels of IL-6 and TNF- α between groups. However, CRP was higher among the obese and sarcopenic obese subjects. In addition, Zoico et al. [12] investigated the relationship between different muscle qualities by conducting muscle biopsies and comparing these with the body composition and inflammation in 16 sarcopenic men (aged between 58 and 80 years old) and demonstrated that the aging of skeletal muscle mass (SMM) is associated not only with muscle atrophy but also with adipose tissue, which worsens the metabolic profile.

Additionally, studies have shown positive correlations between the inflammatory profile, adiposity, metabolic alterations, and arterial properties, such as thickness and stiffness in the elderly [13, 14]; therefore, microvascular damage correlated with stiffer artery may be the link between adiposity, inflammation, and systemic vascular damage. Furthermore, despite sarcopenic people presenting low muscle mass and strength, it is not clear whether the inflammatory and metabolic profile is more related to low lean mass or high fat mass in sarcopenic and non-sarcopenic people.

Thus, the objective of this study was to verify the differences in the inflammatory and metabolic response of sarcopenic and non-sarcopenic overfat elderly and the relationship between these markers, body composition, and strength in this population.

Method

Participants

This cross-sectional study was conducted in May 2015 in the city of Presidente Prudente (approximately 210,000 inhabitants), located in the western region of the state of São Paulo, Brazil. The participants were chosen through convenience sampling. A total of 307 adults of both genders, aged 60 years and older from the abovementioned city, were invited to participate in a previous cohort study,

investigating elderly people and their relationship with physical activity, over 24 months. This study was advertised in the local media, and the individuals voluntarily presented themselves at the institution (Fig. 1).

The exclusion criteria for the current study were: inability to walk, being bedridden, and using pacemakers. The participants were informed regarding the study objectives and data collection methodology. Only individuals who signed the informed consent form were allowed to join the study. All protocols were reviewed and approved by the Research Ethics Committee of the São Paulo State University (Process n°: 15995113.8.0000.5402).

Body composition assessment

Initially, the body composition was assessed using a lunar dual-energy X-ray absorptiometry (DXA) scanner (model DPX-MD, software 4.7, General Electric Healthcare, Lunar DPX-NT; England). The percent body fat as DXA measurements from the 1999 to 2004 National Health and Nutrition Examination Survey (NHANES) [15] with body fat cutoffs from Lohman et al. [16] were used to determine the overfat condition.

For classification of sarcopenic, the appendicular lean soft tissue (ALST) was also determined by DXA. The ALST index was calculated as the ratio of ALST and height to the square (ALSTi), and then the z-score was calculated as the number of SD units from the sample ALSTi mean after normalization of the variable ($Z = [\text{value} - \text{mean}] / \text{SD}$), according to gender. Once the z score was calculated, all the elderly, regardless of gender, were organized from the highest to the lowest value. After this procedure, the elderly with the 40 highest and 40 lowest z score values were invited to participate in an intervention program involving resistance training. Of these, the 60 elderly who agreed to participate in the program were divided into a group of 30 sarcopenic individuals and 30 non-sarcopenic overfat individuals.

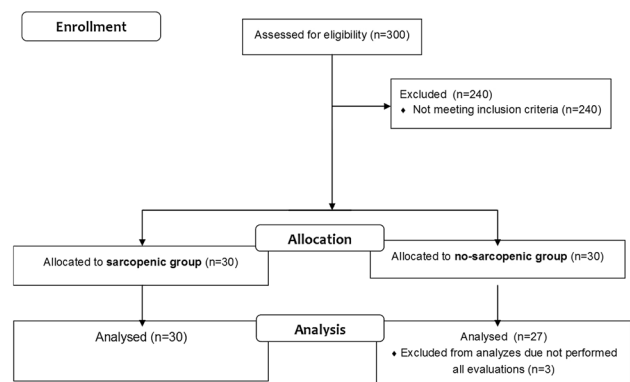


Fig. 1 Flow chart of the study ($n = 57$)

Body mass, intra- and extracellular water, fat mass, musculoskeletal mass and fat and bone free mass from upper and lower limbs and trunk was measured with octopolar bioelectrical impedance equipment InBody 720 (Biospace, Seoul, Korea). Frequencies of 1, 5, 50, 250, 500 and 1000 kHz was emitted and received from the contact of the subject heels and soles and thumbs and palms with the equipment. The minimum clothing was recommended to perform the measurement, according equipment manufacturer's recommendations: not eating and exercising before the test, urinate before the test and standing for 5 min before testing. Tests were performed indoors, at a controlled temperature between 20 and 25 °C.

Strength test procedures

Handgrip was measured on the dominant side, using an electronic hand dynamometer (model: EH101, Camry, China), with a precision of 0.1 kg. To obtain the best performance, the dynamometer was adjusted so that it fitted comfortably to the subject's hand size. Participants, while seated with their elbow bent at an angle of 90°, were instructed to grip the dynamometer with maximum strength with the dynamometer facing outwards from the body. Three trials were performed, with a rest period of at least 1 min between trials. The highest value was considered as handgrip strength and the relative strength was obtained by dividing arm strength by skeletal muscle mass.

Blood sampling and analyses

Blood samples were collected after a 12-h fast, in the morning. The blood samples (15 ml) were immediately allocated into two 5 ml vacutainer tubes (Becton Dickinson, BD, Juiz de Fora, MG, Brazil) containing EDTA for plasma separation and one 5 ml dry vacutainer tube for serum separation. The tubes were centrifuged at 3500g for 15 min at 4 °C, and plasma and serum samples were stored at –20 °C until analysis. PAI-1 (ng/ml), leptin (ng/ml), and adiponectin (µg/ml)

were assessed in serum samples using ELISA commercial kits (R&D Systems, 614 McKinley Place NE, Minneapolis, MN 55413, USA). Glucose (mg/dl), total cholesterol (mg/dl), triacylglycerol (mg/dl), and HDL (mg/dl) were assessed in plasma samples using commercial kits (Labtest®, São Paulo, Brazil). Cortisol was assessed using ELISA commercial kits (Monobind Inc. 100 North Point Drive, Lake Forest, CA 92630 USA). Interleukins (IL-6, IL-10 pg/ml) and TNF-α (pg/ml) cytokines were assessed in serum samples using ELISA commercial kits (affimetrix/eBioscience, Ambriex S/A, São Paulo Brazil). To eliminate inter-assay variance, all samples were analyzed in identical runs, resulting in an intra-assay variance of < 7%.

Data analysis

Initially, data distribution was checked using the Kolmogorov–Smirnov test and, based on the dataset, parametric statistics were performed and the data are described as mean and standard deviation. The independent *t* test was used to verify the differences between groups in the body composition and performance. ANCOVA was performed to compare the metabolic and inflammatory profile, including skeletal muscle mass, fat mass, and the interaction term of both to see whether the role of selected variables differs between lean and overfat subjects with or without sarcopenia. The Pearson correlation (*r*) was used to analyze the relationship between variables. Statistical analysis was performed using the *SPSS* statistical package version 17.0 (*SPSS*, Inc. Chicago, IL, USA) software and the level of statistical significance was set at 5%.

Results

Table 1 presents the comparisons of body composition and performance values for the sarcopenic and non-sarcopenic overfat subjects.

Table 1 Comparison between age, height, weight, total and appendicular body composition and performance in sarcopenic and non-sarcopenic overfat subjects

Variables	Non-sarcopenic (<i>n</i> =27)	Sarcopenic (<i>n</i> =30)	<i>p</i>
Body composition			
Fat mass (kg)	35.6 ± 10.9	20.4 ± 6.2	< 0.001
Visceral fat (cm ²)	139.4 ± 34.9	90.8 ± 24.0	< 0.001
Arm lean mass (kg)	6.0 ± 1.4	3.8 ± 1.0	< 0.001
Leg lean mass (kg)	13.9 ± 2.7	10.6 ± 2.3	< 0.001
SMM (Kg)	27.7 ± 5.3	20.2 ± 3.9	< 0.001
Performance			
Arm strength (kg)	28.5 ± 9.7	23.9 ± 6.2	0.085
Relative strength	4.9 ± 1.4	6.1 ± 1.3	0.014

SMM skeletal muscle mass

It can be observed that the sarcopenic participants presented lower values of fat mass, visceral fat, appendicular muscle mass, and skeletal muscle mass when compared to the non-sarcopenic overfat subjects.

In relation to strength, there was a statistically significant difference between relative strength but no difference was observed in arm strength.

Table 2 presents the comparisons of metabolic and inflammatory profile between sarcopenic and non-sarcopenic overfat subjects.

When comparing the inflammatory profile, the sarcopenic group presented greater adiponectin concentration ($F=5.816$, $p=0.019$), adiponectin/fat mass ratio ($F=20.480$, $p<0.001$), adiponectin/visceral fat ($F=18.054$, $p<0.001$), and higher PAI-1 ($F=5.817$, $p=0.019$) than

non-sarcopenic overfat elderly. After adjusting the metabolic profile by skeletal muscle mass the significant differences between groups were maintained ($p<0.05$) but there were no significant differences between groups when adjusting by fat mass, despite a tendency to a significant difference for adiponectin concentration ($p=0.06$). In addition, after adjusting leptin by fat mass there was a statistically significant lower concentration in the sarcopenic compared to non-sarcopenic overfat elderly.

For TNF, IL-6, IL-8, IL-10, and metabolic profile (glucose, triacylglycerol, total cholesterol, and HDL-c) there were no significant differences between groups.

Table 3 presents the relationship between inflammatory and metabolic response, body composition, and strength in both groups studied.

Table 2 Comparison between sarcopenic and non-sarcopenic overfat subjects in the inflammatory and metabolic profile

Variables	Non-sarcopenic ($n=27$)	Sarcopenic ($n=30$)	p	p Adjusted by SSM	p Adjusted by fat mass	p Adjusted by SSM and fat mass
Glucose (mg/dl)	96.3 ± 21.4	91.8 ± 18.4	0.402	0.717	0.790	0.682
Triacylglycerol (mg/dl)	149.8 ± 86.5	156.1 ± 81.5	0.778	0.934	0.154	0.379
Chol (mg/dl)	187.5 ± 92.0	204.4 ± 79.3	0.501	0.393	0.759	0.997
HDL-c (mg/dl)	70.9 ± 116.3	62.2 ± 61.5	0.746	0.941	0.933	0.735
Cortisol (µg/ml)	15.7 ± 6.3	13.6 ± 6.5	0.213	0.386	0.067	0.228
Leptin (ng/ml)	221.4 ± 149.9	210.4 ± 264.6	0.851	0.219	0.030	0.395
Adiponectin (µg/ml)	12.3 ± 6.9	18.2 ± 10.9	0.019	0.036	0.057	0.145
Adiponectin/fat mass	0.34 ± 0.2	0.99 ± 0.67	<0.001	0.005	0.085	0.241
Adiponectin/visceral fat	0.08 ± 0.05	0.22 ± 0.1	<0.001	0.015	0.115	0.389
PAI-1 (ng/ml)	36.8 ± 20.7	54.5 ± 32.8	0.019	0.036	0.057	0.145
TNF-α (pg/ml)	39.4 ± 38.8	39.1 ± 37.3	0.977	0.697	0.271	0.230
IL-6 (pg/ml)	11.4 ± 30.4	3.5 ± 5.0	0.174	0.352	0.675	0.774
IL-8 (pg/ml)	31.3 ± 65.0	19.5 ± 39.0	0.406	0.121	0.941	0.398
IL-10 (pg/ml)	3.0 ± 5.3	1.8 ± 1.4	0.236	0.478	0.987	0.896

Chol total cholesterol, PAI-1 Plasminogen activator inhibitor-1, TNF-α tumor necrosis factor alpha, IL-6 Interleukin-6, IL-8 Interleukin-8, IL-10 Interleukin-10

Table 3 Correlation between inflammatory and metabolic response, body composition and strength in sarcopenic and non-sarcopenic overfat subjects

Variables	Group	Fat mass (kg)	Visceral fat (cm ²)	Arm lean mass (kg)	Leg lean mass (kg)	SMM (kg)	Arm strength (kg)
Leptin	Non-sarcopenic	0.43*	0.27	-0.33	-0.38	-0.38	-0.06
	sarcopenic	0.61**	0.53**	-0.13	-0.21	-0.21	-0.32
Adiponectin	Non-sarcopenic	-0.21	-0.31	-0.40	-0.33	-0.37	-0.14
	Sarcopenic	-0.07	-0.01	0.01	0.11	0.04	0.26
PAI-1	Non-sarcopenic	-0.21	-0.31	-0.40	-0.33	-0.37	-0.14
	Sarcopenic	-0.07	-0.01	0.01	0.11	0.04	0.26

SMM skeletal muscle mass, PAI-1 Plasminogen activator inhibitor-1

* $p<0.05$; ** $p=0.001$

There was a moderate positive correlation between leptin and fat mass in the non-sarcopenic ($r=0.43$, $p=0.034$) and sarcopenic groups ($r=0.61$, $p=0.001$) and there were moderate positive correlations between leptin and visceral fat in the sarcopenic elderly ($r=0.53$, $p=0.004$). No relationship was observed between adiponectin and PAI-1 concentration with body composition and strength in either group investigated.

Discussion

The main findings of this study were that the sarcopenic elderly presented a higher adiponectin/adiposity tissue ratio and lower PAI-1 than non-sarcopenic overfat elderly. Furthermore, fat mass, but not skeletal muscle mass, seems to change these responses. In addition, there were positive relationships between leptin, and total and visceral fat in both the sarcopenic and non-sarcopenic overfat elderly, although there was no correlation with skeletal muscle mass.

The higher adiponectin concentration and lower PAI-1 in sarcopenic people may be a consequence of the low fat mass in this group. Previous studies on nutritionally induced weight loss [17], gastric surgery [18], and exercise interventions demonstrated an increase in adiponectin [19]. Fatouros et al. [20] conducted a 1-year randomized, controlled trial that analyzed adiponectin responses after different intensities of exercise in 50 inactive men (65–78 years old)—both after exercise training and detraining—and observed that the percentage increase in adiponectin after training was associated with skinfold sum changes ($r=-0.64$; $p=0.001$). Hsieh et al. [21] found that adiponectin levels increased with the reduction in weight, percentage of fat mass, and waist circumference in patients with type 2 diabetes.

Obesity, primarily visceral fat, also contributes significantly to increased plasma PAI-1, IL-6 levels, and low chronic inflammation, resulting in a higher risk of atherosclerosis and is linked to insulin resistance, impaired glucose tolerance, and elevated diabetes risk and metabolic syndrome [22], however, inhibition of PAI-1 in adipocytes protects against insulin resistance by promoting glucose uptake and adipocyte differentiation via increased PPAR γ expression [23]. Studies have demonstrated that PAI-1 is overexpressed in adipose tissue. On the other hand, fat loss is associated with lower PAI-1 [24, 25].

In relation to muscle mass, the aging process is associated with the continuous transactivation of NF- κ B-dependent genes, resulting in many pro-inflammatory markers (TNF- α , IL-6, IL-10) [26]. Furthermore, in advanced age, reactive oxygen species (ROS) can stimulate signaling pathways, resulting in increased NF- κ B activity and chronic inflammation [27]. In sarcopenic people, low muscle mass can occur via induction of the ubiquitin–proteasome (UbP) pathway

and up-regulation of MuRF1 and atrogin-1 gene expression that modulates muscle wasting [28], which, once again increases TNF- α expression and implicates its function as a modifier of MuRF1 and atrogin-1 expression in older muscle [29].

Several studies in the literature demonstrate the relationship between these markers, body composition, and strength [7–9, 30, 31], however, in the present study there was a significant moderate–strong correlation in both sarcopenic and non-sarcopenic elderly only between leptin and total and visceral fat. Our findings are consistent in parts with the results from Lubkowska and colleagues [32] who correlated adiponectin and leptin concentrations with body fat distribution among obese subjects (23–54 years old) and found a positive correlation between leptin, body weight, BMI, and the waist-to-hip ratio. However, in Lubkowska's study, there was a significant negative correlation between adiponectin and total lean mass and skeletal muscle mass. High levels of inflammatory markers were correlated to a fast decline in functionality or a limitation in physical performance [33], given that TNF- α presented the strongest association with the decline in muscle mass and strength [34], however, the study by Legrand et al. [35], as well as our study, did not observe a relationship between inflammatory response and strength. On the other hand, it is known that muscle mass and strength do not decline similarly, thus, longitudinal studies are needed to better elucidate this relationship.

Limitations

The limitations of this study need to be considered when interpreting the findings. The cross-sectional design does not allow any inference of cause and effect and there was no nutritional control. Furthermore, we recommend future longitudinal studies and comparisons with sarcopenic overfat subjects.

Conclusions

In conclusion, the present investigation demonstrated that non-sarcopenic overfat elderly presented lower anti-inflammatory and anti-atherogenic responses than sarcopenic elderly. Furthermore, fat mass, but not skeletal muscle mass, seems to change these responses.

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Compliance with ethical standards

Conflict of interest The authors declare that they have no conflicts of interest.

Ethical approval All procedures performed were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards. The study was reviewed and approved by the Research Ethics Committee of the São Paulo State University (Protocol n°: 15995113.8.0000.5402).

Informed consent The present study complied with ethical standards and informed consent was obtained from all individual participants included in the study.

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