

RESEARCH ARTICLE

The prevalence of probable sarcopenia and intrinsic capacity impairment among a group of community-dwelling older people in an urban area in Cameroon

Marie-Josiane Ntsama Essomba^{1*}, Régine Mylène Mballa Mba², Jean Jacques Noubiap³, Florence Denise Mvondo Lema², Nadine Simo⁴, Maturin Tabue Teguo⁴

1 Department of Internal Medicine and specialties, Faculty of Medicine and Biomedical Sciences, University of Yaounde I, Yaounde, Cameroon, **2** Geriatrics Unit, Yaounde Central Hospital, Yaounde, Cameroon, **3** Division of Cardiology, Department of Medicine, University of California-San Francisco, San Francisco, California, United States of America, **4** Clinical Epidemiology and Aging Team, French West Indies University, French West Indies, France

* ebomaj2012@yahoo.fr



OPEN ACCESS

Citation: Ntsama Essomba M-J, Mballa Mba RM, Noubiap JJ, Mvondo Lema FD, Simo N, Tabue Teguo M (2026) The prevalence of probable sarcopenia and intrinsic capacity impairment among a group of community-dwelling older people in an urban area in Cameroon. *PLoS One* 21(3): e0344528. <https://doi.org/10.1371/journal.pone.0344528>

Editor: Marina De Rui, University Hospital of Padova, ITALY

Received: June 17, 2025

Accepted: February 20, 2026

Published: March 20, 2026

Copyright: © 2026 Ntsama Essomba et al. This is an open access article distributed under the terms of the [Creative Commons Attribution License](https://creativecommons.org/licenses/by/4.0/), which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Data availability statement: The datasets used and/or analysed during the current study are available from the Yaounde Central Hospital

Abstract

Although intrinsic capacity serves as an indirect measure of an individual's functional reserve, whether and in which way it interacts with sarcopenia is still to be addressed in Cameroon. This study aimed to describe the relationship between probable sarcopenia and domains of intrinsic capacity among older people in Cameroon. This cross-sectional study included community-dwelling older aged ≥ 60 years from two senior citizen's association in Cameroon. Probable sarcopenia was assessed using grip strength and Short Physical Performance Battery (SPPB). Screening for intrinsic capacity(IC) impairment was done using the Integrated Care to Older People(ICOPE) approach. Probable sarcopenia was defined by low handgrip strength and low physical performance. We included variables with a p -value $< .1$ in the multivariable analysis model. The significance level was $P < .05$. We included 108 participants [64.8% female, mean age (standard deviation) 70.4 (6.6) years]. The prevalence of probable sarcopenia was 34.3%. All participants had a positive screening for IC impairment and the main impaired domains were vision (88%), locomotion (61.1%) and cognition (50%). The probable sarcopenia group was likely to be older (72.9 ± 7.9 vs 69.6 ± 6.1 $P = .018$), achieved lower education ($P = .012$) and had frequent history of stroke ($P = .038$). After adjusting for age, sex and comorbidities, participants with impaired vitality (OR 3.60, 95%CI 1.08–11.94) were likely to have probable sarcopenia. Participants with preserved locomotion (OR 0.12, 95%CI 0.02–0.66) and a lower number of IC domains impaired (OR 0.52, 95%CI 0.29–0.95) were less likely to have probable sarcopenia. With regard to the components of sarcopenia, preserved locomotion (OR 0.01, 95%CI 0.012–0.07) was associated with a higher physical performance. Lower number of IC domains impaired was associated with higher physical performance

Data Center on reasonable request at the following address dugeronto.fmsb@gmail.com.

Funding: The author(s) received no specific funding for this work.

Competing interests: The authors have declared that no competing interests exist.

(OR 0.29, 95%CI 0.12–0.76) and higher handgrip strength (OR 0.58, 95%CI 0.37–0.92). This study suggests that IC impairment is associated to probable sarcopenia in this group of older adults in Cameroon. Further research on IC trajectories monitoring and incident confirmed sarcopenia as outcome are needed to plan targeted interventions taking into account local resources.

Introduction

Despite the increase in capacity building, manpower development and implementation of geriatric care in many African countries, the rise in older population and longer lifespans are putting pressure on healthcare systems especially in sub-Saharan Africa (SSA) [1–3]. Several evidence support high prevalence of frailty [4] and its impact on poor quality of life and mortality in many African countries [5–7]. Thus, identifying strategies to prevent or delay transition to frailty and dependency are not only relevant but necessary in a context where policies on aging are still scarce. According to a systematic review published in 2024, the weighted prevalence of sarcopenia in Africa was 25.7% [8]. Although it has been recognized as a disease in the International Classification of Diseases (ICD) for several years [9], the lack of a universal and standardized diagnostic criterion remains one of the main issues for reporting sarcopenia in various settings [10].

The decrease of physical abilities and functional decline that can be caused by sarcopenia, can lead to dependency [11], increased risk of falls [12], poor quality of life [13] as well as increased mortality [14]. Furthermore, sarcopenia leads to a high economic burden and healthcare expenditures [15]. In 2019, the World Health Organisation (WHO) has proposed an Integrated Care to Older People (ICOPE) approach, based on the intrinsic capacity (IC) to tackle rising frailty and dependency in the older population [16]. The domains of IC have been suggested to interact with each other and some of them may overlap with clinical manifestations of sarcopenia. Indeed, IC encompasses domains directly related to muscle function including vitality and locomotion, thus it may capture components that overlap with sarcopenia [16]. Furthermore, handgrip strength is commonly used as an indicator of IC [17].

Although IC serves as an indirect measure of an individual's functional reserve, whether and in which way it interacts with sarcopenia is still to be addressed in Cameroon. According to a cross-sectional study conducted in Cameroon in 2020, 26% of people aged 55 and over living in an urban setting had a possible sarcopenia [18]. Higher prevalence was found in another group of community-dwellers [7], reaching 52.3% in a hospital-based study conducted in 2021 [19]. Another crucial epidemiological point is that this high prevalence refers to a population having a mean age of 66 years [7,18,19] which is notably younger than the mean age (78 years) in which sarcopenia is typically observed in other settings [20,21]. Given that background, sarcopenia could become a future public health burden in the country. Thus we aim to examine the relationship between probable sarcopenia and domains of IC in a group of older Cameroonians.

Methods

Study design and setting

This cross-sectional study was carried out from the 30th July to 1st September 2022 in two senior citizens association of Yaounde, the capital of Cameroon. YOMEHCAM (Our contribution to develop Cameroon Association) and APAN (Aide aux personnes âgées de Nkolbisson Association) are two senior citizens association involved in promoting well being and advocacy for the rights of older people in Cameroon.

Study population

Participants were recruited during physical activities sessions. Community-dwelling adults aged ≥ 60 years, who provided an informed written consent were included. Participants with severe disability or severe dementia were not included.

Data collection

Demographic and clinical data. Demographic data included: age (in years), sex, marital status and educational level (illiterate, primary, secondary, university). Clinical data included past medical history including falls history, comorbidities such as hypertension, diabetes, osteoarthritis, obesity and the number of drugs. Polypharmacy was defined as the concomitant use of 5 or more medications.

Domains of intrinsic capacity impairment screening. The IC impairment screening was done as followed:

- Cognition was assessed by the Mini-Cog tool. The Mini-Cog is a quick screening tool and has high sensitivity and specificity to screen for cognitive impairment in older adults across various healthcare settings [22]. A score <3 was considered as impaired cognition.
- Locomotion was assessed by asking the participants to rise from chair five consecutive times without using arms. The incapacity to complete the five-chair rise in less than 14 seconds was considered abnormal [16].
- Vitality was assessed by the Mini-Nutritional Assessment short-form (MNA-SF). The MNA-SF represent a valuable tool for rapid and reliable nutritional screening in the community but also a previously used metric of the vitality domain [17,23]. A score <11 was considered as impaired vitality.
- Psychology was assessed by the 4-item Geriatric Depression Scale (mini-GDS). The mini-GDS 4 yields can be used in screening depression in primary care with a threshold ≥ 1 [24].
- Sensory: Vision was assessed subjectively by asking the participant about difficulties in seeing far, reading, eye diseases or currently under medical treatment for eye disease. If the answer was “yes” to one question, the test was considered abnormal. Audition was assessed by the whispering test for each ear. The examiner stand 1m behind the participant and whisper 2 words at each ear. Failure to repeat one word was considered abnormal.

Any impairment reported for one of the six categories (with vision and hearing considered as two distinct categories) was given a score of 1, otherwise 0. The score was then calculated by adding the number of impaired categories, ranged from 0 to 6. The lower number of IC domains impaired indicates a greater IC.

Probable sarcopenia assessment

Probable sarcopenia was assessed using the muscle strength and the physical performance. Muscle strength was measured by the handgrip strength on the dominant hand in an individual sitted in upright position, using an electronic Jamar Dynamometer. Participants were instructed to exert maximal force. For each individual, the maximum force (kg) after two measurements was recorded as the participant’s handgrip strength. The cut-off for low handgrip strength was <20 kg for

women and <30 kg for men [25]. The Short Physical Performance Battery (SPPB) was used to assess the physical performance, with a score ranging from 0 to 12 [26]. Low physical performance was considered with a SPPB score ≤ 9 . Probable sarcopenia was defined by the presence of low handgrip strength and low physical performance.

Data analysis

Data were analyzed with the Statistical Package for Social Sciences (SPSS 23.0) for Windows (SPSS, Chicago, Illinois, USA). Quantitative variables were presented as mean and standard deviation (SD) or median and interquartile range (IQR). Categorical variables were presented as frequencies and proportions. Quantitative variables were compared by Student's T test or U-Mann Whitney test when needed. Categorical variables were compared using Chi-squared test or Fisher exact T test as needed. Variables that yielded a $P < 0.10$ by univariable analysis were included in the multivariable logistic regression analysis, which estimated odds ratios (OR) and 95% confidence intervals (CI). In this model, the presence of sarcopenia was the dependent variable and the independent variables were demographic and clinical factors, with the no sarcopenia group as the reference. Additionally, IC was considered the main exposure in separate multivariable logistic regression analysis where each defining component of sarcopenia (muscle strength and physical performance) served as a dependent variable. A p -value < 0.05 was statistically significant.

Ethical considerations

This study was approved by the institutional review board of the Yaounde Central Hospital, under the reference number: 17/ACE/CIE/MINSANTE/DHCY/PCE/SG. All participants provided a written consent to participate.

Results

Baseline characteristics of participants

108 participants were included, predominantly female (64.8%) with a mean age (standard deviation SD) of 70.4 (6.6) years. As presented in [Table 1](#), the majority of participants was widowed and achieved secondary education. Sixteen (14.8%) of participants have experienced falls. The prevalence of probable sarcopenia was 34.3%. The probable sarcopenia group was likely to be older (72.9 ± 7.9 vs 69.6 ± 6.1 $P = .018$), achieved lower education ($P = .012$) and demonstrated a higher prevalence of past history of stroke ($P = .038$). Other characteristics are presented in [Table 1](#).

Relationship between probable sarcopenia and intrinsic capacity

All participants had a positive screening for IC impairment and the median (interquartile range) number of IC domains impaired was 3 [2–3]. The IC characteristics are presented in [Table 2](#). The main domains involved were vision (88%), locomotion (65.7%) and cognition (50%). There were no statistical difference in the probable sarcopenia and the non sarcopenia group in terms of cognition, psychological and sensory impairment. However, the probable sarcopenia group had a higher number of impaired domains (4 vs 3, $P = 0.038$) and was likely to have higher locomotion (92.9% vs 56.2%, $P < 0.001$) and vitality impairment (67.9% vs 36.3%, $P = 0.004$). In the multivariable models (see [Table 3](#)), after adjusting for age, sex and comorbidities, participants with preserved locomotion (OR 0.12, 95%CI 0.02–0.66) and a lower number of IC domains impaired (OR 0.52, 95%CI 0.29–0.95) were likely to have a lower prevalence of probable sarcopenia. Impaired vitality (OR 3.60, 95%CI 1.08–11.94) was significantly associated with increased prevalence of probable sarcopenia. With regard to the components of sarcopenia, preserved locomotion (OR 0.01, 95%CI 0.012–0.07) and lower number of IC domains impaired (OR 0.29, 95%CI 0.12–0.76) were associated with higher physical performance. Participants with lower number of IC domains impaired (OR 0.58, 95%CI 0.37–0.92) had a higher handgrip strength.

Table 1. Baseline characteristics of participants.

	Terms	Probable sarcopenia N=28(%)	No sarcopenia N=80(%)	All N=108(%)	P
Sex^a	Male	9 (32.1)	29 (36.3)	38 (35.2)	.695
	Female	19 (67.9)	51 (63.7)	70 (64.8)	–
Age (years)^c	Mean age(SD)	72.9 (7.2)	69.6 (6.1)	70.4 (6.6)	<.001
Marital status^a	Single	2 (7.1)	11 (13.7)	13 (12)	.339
	Married	10 (35.7)	35 (43.8)	45 (41.7)	–
	Widowed	15 (53.6)	28 (35)	43 (39.8)	–
	Divorced	1 (3.6)	6 (7.5)	7 (6.5)	–
Education^a	Illiterate	8 (28.6)	8 (10)	16 (14.8)	.012
	Primary	13 (46.4)	26 (32.5)	39 (36.1)	–
	Secondary	7 (25)	41 (51.2)	48 (44.4)	–
	University	0 (0)	5 (6.3)	5 (4.6)	–
Comorbidities(yes)^a	Hypertension	15 (53.6)	39 (48.8)	54 (50)	.661
	Diabetes	3 (10.7)	19 (23.8)	22 (20.4)	.140
	Osteoarthritis	14 (50)	31 (38.8)	45 (41.7)	.299
	Heart failure	2 (7.1)	3 (3.8)	5 (4.6)	.462
	History of stroke	4 (12.3)	2 (2.5)	6 (5.6)	.038
Falls (yes)^a		6 (21.4)	10 (12.5)	16 (14.8)	.252
Low physical performance	Yes	28 (100)	33 (41.2)	61 (56.5)	<.001
	No	0 (0)	47 (58.8)	47 (43.5)	

SD standard deviation.

^a Categorical variables are shown as number of cases (percentage).

^b Continuous variables are shown as median and interquartile range.

^c Continuous variables are shown as mean and standard deviation.

All variables with a *p*-value at the 0.1 threshold have been included in the final model.

The *p*-value indicated in bold are statistical significant.

<https://doi.org/10.1371/journal.pone.0344528.t001>

Discussion

In this cross-sectional study in a group of community-dwelling older adults, we examined the association between IC, sarcopenia as well as its components. Probable sarcopenia was highly prevalent, affecting approximately one-third of participants. All individuals were screened positive for impairment of at least one domain of IC, with visual, cognitive and locomotor domains being the most frequently affected. Older age, low education and history of previous stroke were more common among participants with probable sarcopenia. After adjustment for age, sex and comorbidities, impairment in the vitality domain was independently associated with a higher likelihood of probable sarcopenia, whereas preserved locomotion and lower number of IC domains impaired were associated with lower odds. With regard to the components of sarcopenia, preserved locomotion and lower number of IC domains impaired were associated with greater physical performance and higher handgrip strength.

Consistent with previous research, sarcopenia was strongly associated with higher number of IC domains impaired and with impairment in specific IC domains, particularly locomotion and vitality. In hospitalized older patients [27] and in community-dwelling octogenarians [28] lower IC composite scores and impairments in locomotion, vitality and cognition are associated with higher odds of sarcopenia. Similarly, studies conducted in disease-specific populations, such as older adults with type 2 diabetes has demonstrated that a greater number of impaired IC domains is associated with increased risk of

Table 2. Relationship between probable sarcopenia and impaired IC.

Impaired IC domains	Terms	Probable sarcopenia N=28(%)	No sarcopenia N=80(%)	All N=108(%)	P
Median IC score (IQR) ^b		4 (3–4)	3 (2–4)	3 (2–3)	.038
Vision ^a	No	1 (3.6)	12 (15)	13 (12)	.177
	Yes	27 (96.4)	68 (85)	95 (88)	–
Audition ^a	No	19 (67.9)	57 (71.3)	76 (70.4)	.735
	Yes	9 (32.1)	23 (28.7)	32 (29.6)	–
Cognition ^a	No	16 (57.1)	38 (47.5)	54 (50)	.380
	Yes	12 (42.9)	42 (52.5)	54 (50)	–
Locomotion ^a	No	2 (7.1)	35 (43.8)	37 (34.3)	<.001
	Yes	26 (92.9)	45 (56.2)	71 (65.7)	–
Vitality ^a	No	9 (32.1)	51 (63.7)	60 (55.6)	.004
	Yes	19 (67.9)	23 (36.3)	48 (44.4)	–
Psychology ^a	No	18 (64.3)	55 (68.8)	73 (67.6)	.664
	Yes	10 (35.7)	25 (31.3)	35 (32.4)	–

IC intrinsic capacity, IQR interquartile range SD standard deviation.

^a Categorical variables are shown as number of cases (percentage).

^b Continuous variables are shown as median and interquartile range.

All variables with a *p*-value at the 0.1 threshold have been included in the final model.

The *p*-value indicated in bold are statistical significant.

<https://doi.org/10.1371/journal.pone.0344528.t002>

Table 3. Multivariable logistic regression of factors associated with probable sarcopenia and its components.

	Probable sarcopenia			Handgrip strength			Physical performance		
	aOR	95%CI	P	aOR	95%CI	P	aOR	95%CI	P
Age	/	/	.152	/	/	.609	/	/	.084
Sex	/	/	.564	2.75	1.07-7.03	.034	/	/	.106
Educational level	/	/	.299	/	/	.836	/	/	.753
Stroke	/	/	.484	/	/	.993	/	/	.999
Vitality	3.60	1.08-11.94	.036	/	/	.271	/	/	.057
Locomotion	0.12	0.02-0.66	.014	/	/	.352	0.01	0.012-0.07	<.001
Number of impaired IC domains	0.52	0.29-0.95	.029	0.58	0.37-0.92	.020	0.29	0.12-0.76	.012

aOR adjusted odds ratio, CI confidence interval, IC intrinsic capacity.

<https://doi.org/10.1371/journal.pone.0344528.t003>

sarcopenia, with locomotion and sensory domains playing a major role [29]. These results are further supported by a systematic review reporting robust interrelation between decline in IC domains, particularly in locomotion and vitality and sarcopenia [30]. Beyond its geographical context, this study contributes by simultaneously evaluating IC domains alongside distinct components of sarcopenia within a single community-based population, thereby extending existing findings beyond selected clinical or disease-specific settings. While our results corroborate this existing body of evidence, they extend prior work in several important ways. First, unlike studies focusing on highly selected populations (e.g., hospitalized patients, octogenarians or individuals with specific chronic diseases) our analysis was conducted in a community-dwelling population, potentially capturing earlier stages of functional decline. Secondly, by examining not only probable sarcopenia as a binary condition but also its components we provide an understanding of how different aspects of muscle function may relate to IC.

With regard to the prevalence of probable sarcopenia, our findings differ from those of previous studies conducted in Cameroon. In 2021, 26% of adults aged ≥ 55 years in an urban setting had possible sarcopenia [18]. In another study involving 403 older community-dwellers, the prevalence of sarcopenia was 47.9% [7]. Our prevalence was also far from the 53% reported in a group of hospitalized patients aged 55 years and over [19]. Our prevalence is higher in comparison to those observed in other African countries. In 2024, Ajuonuma et al reported a prevalence of 21.1% in a group of retirees in Nigeria [31]. In a group of Gambian aged between 40 and 75 years, the prevalence of sarcopenia varied depending on the definition used; in men 20% and 19% and in women 45% and 10% [32]. Although it has been recognized as a disease in the International Classification of Diseases (ICD) for several years [9], the lack of a universal and standardized diagnostic criteria for sarcopenia remains one of the main issues in assessing this condition. As expected, the different criteria used have led to a very heterogeneous prevalence of sarcopenia, even in the same settings. These discrepancies are due to several approaches for the diagnosis of sarcopenia, from those integrating muscle strength, physical performance, and body composition parameters to those using only body composition. Furthermore, several societies and organizations has proposed definitions which are region specific in Western [33] and Asian [34] countries but to date, no operational definition of sarcopenia exists. Another concern when dealing with sarcopenia in SSA, is the lack of specific cut-off values in the muscle mass and muscle strength assessment for the Black Africans sub-populations.

We found that impaired vitality was independently associated with an increased prevalence of probable sarcopenia. This association between vitality and sarcopenia are in line with previous research. In a study conducted in a group of 599 octogenarians in China, those with possible sarcopenia or sarcopenia were more likely to have decline in vitality domain [28]. In another cross-sectional study investigating the relationship between sarcopenia and IC in hospitalized older adults, the sarcopenia group had lower scores in locomotion and vitality domains than the non-sarcopenia group and a strong association was found between impaired vitality and sarcopenia [27]. In a cross-sectional study of a large community-dwelling population, almost all definitions of sarcopenia were associated with a poor nutritional status measured by the MNA [35]. According to a recent systematic review, vitality, locomotion, and cognition are key functional areas of IC linked to sarcopenia [30]. Impaired vitality reflects nutritional and energy-related disorders. Our findings support the notion that impaired vitality may represent a key pathway linking intrinsic capacity impairment to declines in muscle strength and physical performance in older adults. From a pathophysiological point of view, both malnutrition and sarcopenia share many components. With regard to nutritional status, vitality domain is considered as the underlying physiological determinant of IC, resulting from the interaction between multiple physiological systems, reflected in metabolism, neuromuscular function and immune and stress response functions of the body [36]. Beside that, sarcopenia arises from the complex interaction between several factors including chronic inflammation, mitochondrial dysfunction, impaired neuromuscular signaling and reduced protein synthesis [37–40]. Poor nutritional status is common among older people in Cameroon [41] and the rising prevalence of poor physical performance is also a matter of concern [42,43].

We found that preserved locomotion was independently associated with a lower prevalence of probable sarcopenia and higher physical performance. However in multivariable models including probable sarcopenia, locomotion and IC domains, effect estimates showed instability in direction, likely due to collinearity between closely related measures of physical function. These findings should therefore be interpreted with caution and do not suggest true inverse associations. Few studies reported the relationship between IC and physical performance. Individuals with higher IC were less likely to report recent falls, with locomotion being an independently associated domain in a group of octogenarians [44]. Tay et al found that IC was significantly associated with fitness performance, independent of age and gender [45]. Other studies have focused on the relationship between IC and physical activity which although conceptually differs from physical performance, remains closely related. In a study involving a group of healthy older adults, higher moderate-to-vigorous physical activity levels were associated with higher IC mobility, vitality and psychological domain in active versus inactive individuals [46]. Furthermore, the authors found that the inactive category experienced a significant or nearly significant decline in IC, mobility and psychology, while no significant change was observed in the active group [46]. One of the most

prominent change associated to sarcopenia is decline in locomotion [11]. This decline is further exacerbated by reduced physical activity which may accelerate muscle loss. Older adults may therefore exhibit slow gait and poor balance, thereby impacting physical performance and increasing the risk of recurrent falls [44]. Although physical activity was not directly measured in our study, the context of recruitment suggests a predominantly active population. Indeed, participants in our study were involved in physical activity sessions organized by their associations. This behavioural context may have contributed to the presentation of locomotion and physical performance observed in our study. However, a substantial proportion of participants met the criteria for probable sarcopenia. Several hypothesis may explain this apparent paradox. First, physical activity sessions may be insufficient in intensity or resistance loading to counteract impaired muscle function. Activities in these associations usually focus on maintaining mobility and social engagement and may not adequately stimulate muscle strength. In addition, the high prevalence of cerebrovascular disease among participants with probable sarcopenia may compromise muscle strength and physical performance.

In our study, lower number of IC domains impaired was independently associated with a lower prevalence of sarcopenia. It was also independently associated with higher handgrip strength and higher physical performance. Our results are in line with previous studies. Zhu et al found that higher IC composite score was an independent risk factor for sarcopenia and deficits in its associated components namely handgrip strength, skeletal muscle mass and gait speed [27]. Handgrip strength asymmetry and weakness were both significantly associated with an increased risk of IC impairment in older Chinese adults, with the simultaneous presence of both conditions conferring the highest risk. Additionally, a substantial association was observed between handgrip strength status and each domain of IC [47]. In a cross-sectional study conducted in Colombia, participants with optimal handgrip strength had better IC than weak older adults, including both men (OR 0.62, 95%CI 0.53–0.71) and women (OR 0.79, 95% CI 0.68–0.92) [48]. The mechanism for the association between IC and sarcopenia is not yet completely understood. However, sarcopenia is characterized by age-related loss of muscle mass and function which overlap with some domains of IC.

While the IC domains are separate entities and interventions may be domain specific, our findings are in line with previous studies suggesting significant interactions between domains notably vitality and locomotion. Studies on relevant interventions are lacking. Nevertheless, this is an active area of research with promising results. With regard to sarcopenia, current evidence suggests that protein supplementation is effective in improving muscle strength and muscle mass when used as an adjunct to resistance exercise training [49]. Multidomains lifestyle interventions on nutrition, cognition and mobility may be effective in reversing sarcopenia and improving muscle mass and function among community-dwelling older [50]. In another randomized control trial, 6-month active combination of psychological, nutrition and exercise interventions improved sarcopenia indices [51].

Strength and limitations

To the best of our knowledge, this is the first study to examine the relationship between probable sarcopenia and IC in Cameroon, highlighting the impact of vitality, locomotion and number of impaired IC domains in sarcopenia, handgrip strength and physical performance. Our findings suggest that despite their involvement in senior associations our participants did not demonstrate a lower prevalence of locomotion impairment relative to other older populations in Cameroon. This suggests that participation in activity groups alone may not be sufficient to preserve locomotion or that underlying factors such as comorbidities, environmental constraints or early functional decline may still contribute significantly. However, we acknowledge several limitations. First, the cross-sectional design did not allow us to determine the causality between IC impairment and sarcopenia. Second, our sample size is not large enough to generalize our findings. Third, possible selection bias as we choose to recruit among a group of active individuals, thus older people living with frailty are probably underrepresented in this population. Finally, comorbidities were not collected using a standardized index limiting analysis to participants conditions and muscle mass was not measured to confirm the diagnosis of sarcopenia.

Conclusion

This study showed a high prevalence of probable sarcopenia and IC impairment in a group of active older Cameroonians. Our findings enable a nuanced evaluation of the relationship between muscle function components and IC. Further research on IC trajectories monitoring and incident confirmed sarcopenia as outcome are needed to plan targeted interventions taking into account local resources.

Supporting information

S1 File. Inclusivity-in-global-research-questionnaire.
(DOCX)

Acknowledgments

The authors want to acknowledge all the participants and the members of senior citizen's associations of Yaoundé.

Author contributions

Conceptualization: Marie-Josiane Ntsama Essomba, Maturin Tabue Teguo.

Data curation: Marie-Josiane Ntsama Essomba, Regine Mylene Mballa Mba, Maturin Tabue Teguo.

Formal analysis: Marie-Josiane Ntsama Essomba, Jean Jacques Noubiap.

Investigation: Regine Mylene Mballa Mba, Florence Denise Mvondo Lema.

Methodology: Marie-Josiane Ntsama Essomba, Regine Mylene Mballa Mba, Florence Denise Mvondo Lema, Maturin Tabue Teguo.

Supervision: Maturin Tabue Teguo.

Validation: Nadine Simo, Maturin Tabue Teguo.

Writing – original draft: Marie-Josiane Ntsama Essomba, Jean Jacques Noubiap.

Writing – review & editing: Marie-Josiane Ntsama Essomba, Regine Mylene Mballa Mba, Jean Jacques Noubiap, Florence Denise Mvondo Lema, Nadine Simo, Maturin Tabue Teguo.

References

- Dotchin CL, Akinyemi RO, Gray WK, Walker RW. Geriatric medicine: services and training in Africa. *Age Ageing* [Internet]. 2013 [cited 2023 Feb 5];42(1):124–8. Available from: <https://academic.oup.com/ageing/article/42/1/124/25768>
- Cassim B, Tipping B. Healthcare for older people in Africa. *Age Ageing* [Internet]. 2022 [cited 2023 Jul 25];51(11):1–4. Available from: <https://dx.doi.org/10.1093/ageing/afac217>
- Akoria OA. Establishing in-hospital geriatrics services in Africa: insights from the University of Benin Teaching Hospital geriatrics project. *Ann Afr Med*. 2016;15(3):145–53. <https://doi.org/10.4103/1596-3519.188896> PMID: 27549420
- O’Caoimh R, Sezgin D, O’Donovan MR, Molloy DW, Clegg A, Rockwood K, et al. Prevalence of frailty in 62 countries across the world: a systematic review and meta-analysis of population-level studies. *Age Ageing*. 2021;50(1):96–104. <https://doi.org/10.1093/ageing/afaa219> PMID: 33068107
- Ajayi SA, Adebosoye LA, Olowookere OO, Akinyemi RO, Afolayan KO, Akinyemi JO, et al. Prevalence and correlates of frailty syndrome among older adults attending Chief Tony Anenih Geriatric Centre, University College Hospital, Ibadan. *West Afr J Med* [Internet]. 2021;38(3):255–67. Available from: <http://www.ncbi.nlm.nih.gov/pubmed/33765747>
- Adebosoye LA, Ogunbode AM, Olowookere OO, Ajayi SA, Ladipo MM. Factors associated with sarcopenia among older patients attending a geriatric clinic in Nigeria. *Niger J Clin Pract*. 2018;21(4):443–50. https://doi.org/10.4103/njcp.njcp_374_17 PMID: 29607855
- Metanmo S, Kuate-Tegueu C, Gbessemehlan A, Dartigues J-F, Ntsama M-J, Nguegang Yonta L, et al. Self-reported visual impairment and sarcopenia among older people in Cameroon. *Sci Rep*. 2022;12(1):17694. <https://doi.org/10.1038/s41598-022-22563-9> PMID: 36271132
- Veronese N, Smith L, Koyanagi A, Hoffman J, Snoussi M, Prokopidis K, et al. Prevalence of sarcopenia in Africa: a systematic review and meta-analysis of observational studies. *Aging Clin Exp Res*. 2024;36(1):12. <https://doi.org/10.1007/s40520-023-02671-w> PMID: 38281246

9. Anker SD, Morley JE, von Haehling S. Welcome to the ICD-10 code for sarcopenia. *J Cachexia Sarcopenia Muscle*. 2016;7(5):512–4. <https://doi.org/10.1002/jcsm.12147> PMID: [27891296](https://pubmed.ncbi.nlm.nih.gov/27891296/)
10. Kirk B, Cawthon PM, Arai H, Ávila-Funes JA, Barazzoni R, Bhasin S, et al. The conceptual definition of sarcopenia: delphi consensus from the global leadership initiative in sarcopenia (GLIS). *Age Ageing*. 2024;53(3). <https://doi.org/10.1093/ageing/afae052>
11. Beaudart C, Zaaria M, Pasleau F, Reginster J-Y, Bruyère O. Health outcomes of sarcopenia: a systematic review and meta-analysis. Wright JM, editor. *PLoS One* [Internet]. 2017;12(1):e0169548. Available from: <https://dx.plos.org/10.1371/journal.pone.0169548>
12. Yeung SSY, Reijnierse EM, Pham VK, Trappenburg MC, Lim WK, Meskers CGM, et al. Sarcopenia and its association with falls and fractures in older adults: a systematic review and meta-analysis. *J Cachexia Sarcopenia Muscle*. 2019;10(3):485–500. <https://doi.org/10.1002/jcsm.12411> PMID: [30993881](https://pubmed.ncbi.nlm.nih.gov/30993881/)
13. Beaudart C, Demonceau C, Reginster J-Y, Locquet M, Cesari M, Cruz Jentoft AJ, et al. Sarcopenia and health-related quality of life: a systematic review and meta-analysis. *J Cachexia Sarcopenia Muscle*. 2023;14(3):1228–43. <https://doi.org/10.1002/jcsm.13243> PMID: [37139947](https://pubmed.ncbi.nlm.nih.gov/37139947/)
14. Xu J, Wan CS, Ktoris K, Reijnierse EM, Maier AB. Sarcopenia is associated with mortality in adults: a systematic review and meta-analysis. *Gerontology*. 2022;68(4):361–76. <https://doi.org/10.1159/000517099> PMID: [34315158](https://pubmed.ncbi.nlm.nih.gov/34315158/)
15. Bruyère O, Beaudart C, Ethgen O, Reginster J-Y, Locquet M. The health economics burden of sarcopenia: a systematic review. *Maturitas*. 2019;119:61–9. <https://doi.org/10.1016/j.maturitas.2018.11.003> PMID: [30502752](https://pubmed.ncbi.nlm.nih.gov/30502752/)
16. WHO. Integrated care for older people (ICOPE): guidance for person-centred assessment and pathways in primary care. Geneva: World Health Organization; 2019.
17. George PP, Lun P, Ong SP, Lim WS. A rapid review of the measurement of intrinsic capacity in older adults. *J Nutr Health Aging*. 2021;25(6):774–82. <https://doi.org/10.1007/s12603-021-1622-6> PMID: [34179933](https://pubmed.ncbi.nlm.nih.gov/34179933/)
18. Essomba MJN, Atsa D, Noah DZ, Zingui-Ottou M, Paula G, Nkeck JR. Geriatric syndromes in an urban elderly population in Cameroon: a focus on disability, sarcopenia and cognitive impairment. *Pan Afr Med J*. 2020;37:1–14.
19. Marie-josiane NE, Dimitri ZN, Njonou S, Raoul S, Gloria A. Facteurs Associés à la Sarcopénie chez des Patients Âgés de 55 Ans et plus dans un Service de Médecine Interne au Cameroun. *Health Sci Dis*. 2022;23:58–61.
20. Bahat G, Yilmaz O, Kiliç C, Oren MM, Karan MA. Performance of SARC-F in regard to sarcopenia definitions, muscle mass and functional measures. *J Nutr Health Aging* [Internet]. 2018;22(8):898–903. Available from: <http://link.springer.com/10.1007/s12603-018-1067-8>
21. Moreira VG, Perez M, Lourenço RA. Prevalence of sarcopenia and its associated factors: the impact of muscle mass, gait speed, and handgrip strength reference values on reported frequencies. *Clinics (Sao Paulo)*. 2019;74:e477. <https://doi.org/10.6061/clinics/2019/e477> PMID: [30994709](https://pubmed.ncbi.nlm.nih.gov/30994709/)
22. Abayomi SN, Sritharan P, Yan E, Saripella A, Alhamdah Y, Englesakis M, et al. The diagnostic accuracy of the Mini-Cog screening tool for the detection of cognitive impairment-A systematic review and meta-analysis. *PLoS One*. 2024;19(3):e0298686. <https://doi.org/10.1371/journal.pone.0298686> PMID: [38483857](https://pubmed.ncbi.nlm.nih.gov/38483857/)
23. Kaiser MJ, Bauer JM, Ramsch C, Uter W, Guigoz Y, Cederholm T, et al. Validation of the Mini Nutritional Assessment short-form (MNA-SF): a practical tool for identification of nutritional status. *J Nutr Health Aging*. 2009;13(9):782–8. <https://doi.org/10.1007/s12603-009-0214-7> PMID: [19812868](https://pubmed.ncbi.nlm.nih.gov/19812868/)
24. Lafont C, Chah Wakilian A, Lemogne C, Gouraud C, Fossey-Diaz V, Orvoen G, et al. Performance diagnostique de la « Geriatric Depression Scale » pour le repérage de la dépression chez les sujets âgés atteints de cancer: l'étude de cohorte ELCAPA. *Rev Epidemiol Sante Publique* [Internet]. 2018;66:S149. Available from: <https://linkinghub.elsevier.com/retrieve/pii/S0398762018303043>
25. Fernandes S, Rodrigues da Silva E, New York B, Macedo P, Gonçalves R, Camara S, et al. Cutoff points for grip strength in screening for sarcopenia in community-dwelling older-adults: a systematic review. *J Nutr Health Aging*. 2022;26(5):452–60. <https://doi.org/10.1007/s12603-022-1788-6> PMID: [35587757](https://pubmed.ncbi.nlm.nih.gov/35587757/)
26. Guralnik JM, Simonsick EM, Ferrucci L, Glynn RJ, Berkman LF, Blazer DG. A short physical performance battery assessing lower extremity function: association with self-reported disability and prediction of mortality and nursing home admission. *J Gerontol*. 1994;49(2):M85-94. <https://doi.org/10.1093/geronj/49.2.M85>
27. Zhu L, Zong X, Shi X, Ouyang X. Association between intrinsic capacity and sarcopenia in hospitalized older patients. *J Nutr Health Aging*. 2023;27(7):542–9. <https://doi.org/10.1007/s12603-023-1946-5> PMID: [37498101](https://pubmed.ncbi.nlm.nih.gov/37498101/)
28. Hsu P-S, Lee W-J, Peng L-N, Lu W-H, Meng L-C, Hsiao F-Y, et al. Safeguarding vitality and cognition: the role of sarcopenia in intrinsic capacity decline among octogenarians from multiple cohorts. *J Nutr Health Aging*. 2024;28(6):100268. <https://doi.org/10.1016/j.jnha.2024.100268> PMID: [38810513](https://pubmed.ncbi.nlm.nih.gov/38810513/)
29. Lin C-L, Wu H-C, Yu N-C, Liu Y-C, Wu C-L, Chien W-C. Association between intrinsic capacity and sarcopenia in older adults with type 2 diabetes: a cross-sectional study. *Aging Clin Exp Res*. 2025;37(1):252. <https://doi.org/10.1007/s40520-025-03160-y> PMID: [40833515](https://pubmed.ncbi.nlm.nih.gov/40833515/)
30. Sales WB, Silva PV de S, Vital BSB, Câmara M. Sarcopenia and intrinsic capacity in older adults: a systematic review. *Arch Gerontol Geriatr*. 2025;135:105875. <https://doi.org/10.1016/j.archger.2025.105875> PMID: [40318296](https://pubmed.ncbi.nlm.nih.gov/40318296/)
31. Ajuonuma FO, Ibrahim BY, Zubairu HD. Sarcopenia among elderly retired soldiers attending the retiree's clinic in an army reference hospital in Kaduna, northwestern, Nigeria. *PAMJ Clin Med* [Internet]. 2024;16(7). Available from: <https://www.clinical-medicine.panafrican-med-journal.com/content/article/16/7/full>

32. Zengin A, Jarjou LM, Prentice A, Cooper C, Ebeling PR, Ward KA. The prevalence of sarcopenia and relationships between muscle and bone in ageing West-African Gambian men and women. *J Cachexia Sarcopenia Muscle*. 2018;9(5):920–8. <https://doi.org/10.1002/jcsm.12341> PMID: [30221478](https://pubmed.ncbi.nlm.nih.gov/30221478/)
33. Cruz-Jentoft AJ, Bahat G, Bauer J, Boirie Y, Bruyère O, Cederholm T, et al. Sarcopenia: revised European consensus on definition and diagnosis. *Age Ageing*. 2019;48(1):16–31. <https://doi.org/10.1093/ageing/afy169> PMID: [30312372](https://pubmed.ncbi.nlm.nih.gov/30312372/)
34. Chen LK, Woo J, Assantachai P, Auyeung TW, Chou MY, Iijima K, et al. Asian Working Group for Sarcopenia: 2019 consensus update on sarcopenia diagnosis and treatment. *J Am Med Dir Assoc [Internet]*. 2020;21(3):300–307.e2. Available from: <https://linkinghub.elsevier.com/retrieve/pii/S1525861019308722>
35. Calcaterra L, Abellan van Kan G, Steinmeyer Z, Angioni D, Proietti M, Sourdets S. Sarcopenia and poor nutritional status in older adults. *Clin Nutr*. 2024;43(3):701–7. <https://doi.org/10.1016/j.clnu.2024.01.028> PMID: [38320461](https://pubmed.ncbi.nlm.nih.gov/38320461/)
36. Bautmans I, Knoop V, Amuthavalli Thiyagarajan J, Maier AB, Beard JR, Freiburger E, et al. WHO working definition of vitality capacity for healthy longevity monitoring. *Lancet Healthy Longev*. 2022;3(11):e789–96. [https://doi.org/10.1016/S2666-7568\(22\)00200-8](https://doi.org/10.1016/S2666-7568(22)00200-8) PMID: [36356628](https://pubmed.ncbi.nlm.nih.gov/36356628/)
37. Azzolino D, Spolidoro GCI, Saporiti E, Luchetti C, Agostoni C, Cesari M. Musculoskeletal changes across the lifespan: nutrition and the life-course approach to prevention. *Front Med*. 2021;8. <https://doi.org/10.3389/fmed.2021.697954>
38. Granic A, Suetterlin K, Shavlakadze T, Grounds MD, Sayer AA. Hallmarks of ageing in human skeletal muscle and implications for understanding the pathophysiology of sarcopenia in women and men. *Clin Sci [Internet]*. 2023;137(22):1721–51. Available from: <https://portlandpress.com/clinsci/article/137/22/1721/233761/Hallmarks-of-ageing-in-human-skeletal-muscle-and>
39. da Costa Teixeira LA, Avelar NCP, Peixoto MFD, Parentoni AN, Santos JM dos, Pereira FSM. Inflammatory biomarkers at different stages of sarcopenia in older women. *Sci Rep*. 2023;13(1):10367. <https://doi.org/10.1038/s41598-023-37229-3>
40. Sayer AA, Cooper R, Arai H, Cawthon PM, Ntsama Essomba MJ, Fielding RA, et al. Sarcopenia. *Nat Rev Dis Prim*. 2024;10(1):1–16.
41. Mabiama G, Adio D, Preux PM, Desport J-C, Fayemendy P, Jésus P. Nutritional status and associated factors among community-dwelling elderly. *Clin Nutr ESPEN*. 2021;45:220–8. <https://doi.org/10.1016/j.clnesp.2021.08.021> PMID: [34620321](https://pubmed.ncbi.nlm.nih.gov/34620321/)
42. Ndoboe-Koe V, Kemnang Y, Ebode-Ntsama MJ, Nganou-Gnindjio CN, Biatu N, Ba H, et al. Quality of life of a group of Cameroonian patients aged over 65 years in an urban setting: a cross-sectional study. *Heal Sci Dis [Internet]*. 2022;24(1 SE-Medicine and Surgery in the Tropics). Available from: <https://www.hsd-fmsb.org/index.php/hsd/article/view/4123>
43. Metanmo S, Simo-Tabue N, Kuate-Tegoue C, Bonnet M, Gbessemehlan A, Metanmo F, et al. Short physical performance battery and study of osteoporotic fractures index in the exploration of frailty among older people in Cameroon. *Int J Public Health*. 2023;68:1605900. <https://doi.org/10.3389/ijph.2023.1605900> PMID: [37609077](https://pubmed.ncbi.nlm.nih.gov/37609077/)
44. Cacciatore S, Marzetti E, Calvani R, Picca A, Salini S, Russo A, et al. Intrinsic capacity and recent falls in adults 80 years and older living in the community: results from the iSIRENTE Study. *Aging Clin Exp Res*. 2024;36(1):169. <https://doi.org/10.1007/s40520-024-02822-7> PMID: [39126523](https://pubmed.ncbi.nlm.nih.gov/39126523/)
45. Tay L, Tay E-L, Mah SM, Latib A, Koh C, Ng Y-S. Association of intrinsic capacity with frailty, physical fitness and adverse health outcomes in community-dwelling older adults. *J Frailty Aging*. 2023;12(1):7–15. <https://doi.org/10.14283/jfa.2022.28> PMID: [36629078](https://pubmed.ncbi.nlm.nih.gov/36629078/)
46. Raffin J, Fourteau M, Virecoulon Giudici K, Rolland Y, Vellas B, de Souto Barreto P, et al. Cross-sectional and longitudinal associations between physical activity and intrinsic capacity in healthy older adults from the MAPT study. *Arch Gerontol Geriatr*. 2025;130:105724. <https://doi.org/10.1016/j.archger.2024.105724> PMID: [39700710](https://pubmed.ncbi.nlm.nih.gov/39700710/)
47. Li D, Wang Y, Guo S, Ren Z, Su B, Zhang L, et al. Association of handgrip strength asymmetry and weakness with intrinsic capacity impairment among older adults in China. *Exp Gerontol*. 2025;199:112656. <https://doi.org/10.1016/j.exger.2024.112656> PMID: [39672282](https://pubmed.ncbi.nlm.nih.gov/39672282/)
48. Ramírez-Vélez R, Correa-Bautista JE, García-Hermoso A, Cano CA, Izquierdo M. Reference values for handgrip strength and their association with intrinsic capacity domains among older adults. *J Cachexia Sarcopenia Muscle*. 2019;10(2):278–86. <https://doi.org/10.1002/jcsm.12373> PMID: [30843369](https://pubmed.ncbi.nlm.nih.gov/30843369/)
49. Kirwan RP, Mazidi M, Rodríguez García C, Lane KE, Jafari A, Butler T, et al. Protein interventions augment the effect of resistance exercise on appendicular lean mass and handgrip strength in older adults: a systematic review and meta-analysis of randomized controlled trials. *Am J Clin Nutr*. 2022;115(3):897–913. <https://doi.org/10.1093/ajcn/nqab355> PMID: [34673936](https://pubmed.ncbi.nlm.nih.gov/34673936/)
50. Lu Y, Niti M, Yap KB, Tan CTY, Zin Nyunt MS, Feng L, et al. Assessment of sarcopenia among community-dwelling at-risk frail adults aged 65 years and older who received multidomain lifestyle interventions. *JAMA Netw Open [Internet]*. 2019;2(10):e1913346. Available from: <https://jamanetwork.com/journals/jamanetworkopen/fullarticle/2752992>
51. Chan D-CD, Tsou H-H, Chang C-B, Yang R-S, Tsauo J-Y, Chen C-Y, et al. Integrated care for geriatric frailty and sarcopenia: a randomized control trial. *J Cachexia Sarcopenia Muscle*. 2017;8(1):78–88. <https://doi.org/10.1002/jcsm.12132> PMID: [27897406](https://pubmed.ncbi.nlm.nih.gov/27897406/)