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Brief Report

“Skeletal Muscle Function Deficit” in A Nationally Representative British Birth Cohort in Early Old Age

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Abstract

Background. Recommendations for identifying age-related muscle dysfunction have recently been published. We aimed to compare definitions for clinically relevant weakness and low lean mass proposed by the Foundation for the National Institutes of Health (FNIH) Sarcopenia project with the definition of sarcopenia proposed by the European Working Group on Sarcopenia in Older People (EWGSOP).

Methods. A total of 1566 men and women from a British birth cohort had measures of appendicular lean mass, grip strength and timed up, and go speed at ages 60–64. Prevalence of low lean mass, weakness and slowness, identified using the FNIH and EWGSOP recommendations were estimated and compared: using kappa statistics and; by testing cross-sectional associations of both definitions of low lean mass and weakness with slowness and self-reported difficulties walking.

Results. The combined prevalence of low lean mass and weakness ranged from 1.1% in men using FNIH criteria to 6.4% in women using EWGSOP criteria. There was limited overlap between the groups identified using the different criteria, driven by limited agreement between the two definitions of low lean mass. Using FNIH criteria, both low lean mass and weakness were associated with higher odds of slowness and self-reported difficulties walking; whereas low lean mass classified using EWGSOP criteria was not associated with these markers of mobility impairment.

Conclusions. At relatively young ages, signs of skeletal muscle function deficit with potential clinical relevance are already identifiable in the general population. This suggests that implementation of strategies to prevent mobility limitations, related to age-related muscle dysfunction, may need to start at least as early as midlife.

Key words: Epidemiology—Life course—Physical capability.

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across life that drive these changes and; (b) identifying modifiable targets for intervention which may reduce the detrimental impact of these changes on functional and clinical outcomes. If these and other initiatives result in more widely agreed operational definitions for what has recently been termed “skeletal muscle function deficit” (6), this may offer greater opportunities for evidence synthesis.

In direct response to the call made to evaluate the FNIH recommendations in other study populations (1,4,5), we aimed to use data from a nationally representative British birth cohort study (7), to compare the definitions for clinically relevant weakness and low lean mass proposed by the FNIH (2,3) with the definition of sarcopenia proposed by the European Working Group on Sarcopenia in Older People (EWGSOP) (8) which is now being applied in a wide range of settings (9–11). As mobility impairment has been identified as an important primary outcome (1,4) this comparison included tests of the associations of low lean mass and weakness with slow walking speed and self-reported difficulties walking.

Methods

Study Population

The Medical Research Council National Survey of Health and Development (NSHD) is a socially stratified sample of 5362 singleton births that took place in one week of March 1946 in mainland Britain. Between 2006 and 2010 (at 60–64 years), 2856 eligible study members (those known to be alive and living in England, Scotland, or Wales) were invited for assessment at one of six clinical research facilities (CRFs) or to be visited by a research nurse at home; 2229 were assessed (1690 at a CRF) (7). Relevant ethical approval was obtained and participants provided written consent.

Measures At 60–64 Years

As described in detail elsewhere (12,13), measures of body composition (including appendicular lean mass (ALM) [the sum of the fat free mass in the limbs excluding bone mineral content]) were obtained for study participants who attended a CRF, in the supine position using a QDR 4500 Discovery DXA scanner (Hologic Inc, Bedford, MA).

Strength and physical performance were assessed by trained nurses during the CRF or home visit using standardized protocols. Grip strength (kg) was assessed using an electronic handgrip dynamometer (14). Three values were recorded for each hand; then the highest was used in analyses. Walking speed was assessed using the timed-up and go (TUG) test, with participants instructed to rise from a chair, walk 3 m at a normal pace, turn around, return to the chair, and sit down. The time taken to complete the test was recorded and the speed (m/s) calculated. Participants were also asked whether they had difficulty walking for quarter of a mile on the level because of long-term health problems. Height and weight were measured and used to calculate body mass index (BMI;kg/m²).

Analyses

After deriving the variables required (ie ALM adjusted for height², ALM:BMI ratio, and walking speed, by multiplying TUG speed by 1.62, based on comparisons of TUG and walking speeds in other British study populations with both measures (15)), the FNIH and EWGSOP cut-points for low lean mass, weakness and slowness were applied. Those participants in the potential analytic sample, who were unable to complete the grip strength (n = 17) or TUG (n = 8) assessment for health reasons (which included arthritis, other musculoskeletal conditions, and cardiorespiratory problems), were categorised as weak and slow, respectively.

Prevalence estimates of low lean mass, weakness, slowness, and the combinations of these outcomes were calculated. Cohen’s kappa was used to compare the different definitions. Logistic regression was used to estimate the odds ratios of slowness and self-reported difficulty walking associated with the different definitions of low lean mass and weakness. Gender-specific analyses were conducted on the sample with complete data on lean mass, grip strength, and walking speed (n = 1566). Sensitivity analyses, conducted to test the impact of including those people unable to perform the grip strength and TUG assessments in the main analyses and to assess the robustness of findings, are described in Appendix 1.

Results

To aid comparison with similar estimates in other studies, baseline characteristics of the sample included in analyses are presented in Appendix 2. Prevalence estimates of low lean mass, weakness and slowness, and the combinations of these outcomes are shown in Table 1. Estimates of each of these three outcomes alone were much higher than combinations of these outcomes, reflecting limited overlap (see Appendix 3). As the FNIH criteria are more restrictive (5), prevalence estimates using these criteria were slightly lower than those using EWGSOP criteria.

There was limited overlap between the groups identified as having both low lean mass and weakness using the different criteria (K = .23 in men, .09 in women), driven by limited agreement between the two definitions of low lean mass (K = .10 in men, .02 in women); 31.6% of men and 40.5% of women had low lean mass according to at least one definition, but only 4.8% of men and 5.0% of women were identified by both definitions.

Table 1. Prevalence Estimates (%) of Low Lean Mass, Weakness and Slowness Identified Using FNIH (1) and EWGSOP (8) Criteria (n = 747 men, 819 women)

|                | FNIH   | EWGSOP |
|----------------|--------|--------|
|                | Men    | Women  | Men    | Women  |
| Low lean mass  |        |        |        |        |
| Men            | 15.7   | 14.9   | 20.8   | 30.7   |
| Women          | 4.0    | 6.4    | 7.5    | 18.2   |
| Weakness       |        |        |        |        |
| Men            | 9.0    | 9.7    | 9.0    | 9.7    |
| Women          | 3.7    | 2.7    | 3.7    | 7.7    |
| Weakness and low lean mass |        |        |        |        |
| Men            | 1.1    | 2.0    | 2.3    | 6.4    |
| Women          | 3.2    | 2.7    | 4.4    | 7.3    |
| Low lean mass  |        |        |        |        |
| Men            | 0.3    | 0.7    | 0.3    | 0.9    |
| Women          |        |        |        |        |

Cut-points:
Low lean mass (applied to ALM:BMI ratio for FNIH and ALM/ht² for EWGSOP):
FNIH: Men < 0.789; women < 0.512.
EWGSOP: Men ≤ 7.23 kg/m²; women ≤ 5.67 kg/m².
Weakness (applied to grip strength):
FNIH: Men < 26 kg; women < 16 kg.
EWGSOP: Men < 30 kg; women < 20 kg.
Slowness (applied to TUG speed after conversion to walking speed):
FNIH and EWGSOP: ≤0.8 m/s.
ALM, appendicular lean mass; BMI, body mass index; EWGSOP, European Working Group on Sarcopenia in Older People; FNIH, Foundation for the National Institutes of Health; TUG, timed-up and go test.
Using the FNIH criteria, low lean mass and weakness were both associated with higher odds of slowness and self-reported difficulties walking (Table 2). Low lean mass classified using the EWGSOP criteria was not associated with mobility limitation.

Findings from sensitivity analyses suggested that there was no major impact on findings of including those people unable to complete the grip strength and TUG assessments for health reasons (Appendix 1).

Discussion
In a nationally representative British birth cohort study, assessed in early old age, between 2.7% and 7.3% of the population were identified as having both low lean mass and either weakness or slowness according to two sets of recently published criteria (FNIH: 3.1% of men, 2.7% of women; EWGSOP: 4.4% of men, 7.3% of women) (1,8). Both definitions of weakness and the FNIH definition of low lean mass were strongly associated with slowness and self-reported difficulties walking. Despite producing lower prevalence estimates of low lean mass and weakness, the FNIH definitions were more strongly associated with mobility limitations than the EWGSOP definitions.

These analyses respond to the call to evaluate recently proposed criteria for the identification of clinically meaningful age-related changes in muscle mass, strength, and performance in other study populations (1,4,5). While the focus of existing research has been on populations aged 65 years and older (often with a mean age of 70 or older), our analyses of the NSHD at ages 60–64 highlights that at relatively young ages, signs of skeletal muscle function deficit associated with mobility limitations than the EWGSOP definitions. These analyses respond to the call to evaluate recently proposed criteria for the identification of clinically meaningful age-related changes in muscle mass, strength, and performance in other study populations (1,4,5). While the focus of existing research has been on populations aged 65 years and older (often with a mean age of 70 or older), our analyses of the NSHD at ages 60–64 highlights that at relatively young ages, signs of skeletal muscle function deficit associated with mobility limitations than the EWGSOP definitions. The prevalence of low lean mass and weakness classified using these definitions was lower than that observed in existing studies, particularly in men, where the prevalence of low lean mass was 2.7% and the prevalence of weakness was 3.4%.

Table 2. Odds Ratios of Slowness and Self-reported Difficulties Walking by Low Lean Mass and Weakness Identified Using FNIH and EWGSOP Criteria

| Low lean mass | Slowness* | Difficulty Walking† |
|---------------|-----------|---------------------|
| **FNIH**      |           |                     |
| Men           | 2.35 (1.33, 4.17) | 3.44 (1.93, 6.11)  |
| Women         | 1.80 (1.03, 3.18) | 2.59 (1.48, 4.53)  |
| **EWGSOP**    |           |                     |
| Men           | 1.22 (0.68, 2.21) | 0.77 (0.38, 1.55)  |
| Women         | 0.86 (0.51, 1.44) | 0.61 (0.34, 1.10)  |
| Weakness      |           |                     |
| **FNIH**      |           |                     |
| Men           | 4.87 (2.13, 11.12) | 4.73 (2.01, 11.16) |
| Women         | 3.13 (1.57, 6.26) | 3.34 (2.76, 10.33) |
| **EWGSOP**    |           |                     |
| Men           | 3.57 (1.81, 7.04) | 2.84 (1.35, 5.97)  |
| Women         | 2.63 (1.39, 4.35) | 2.10 (1.48, 5.24)  |
| Low lean mass and weakness |           |                     |
| **FNIH**      |           |                     |
| Men           | 3.46 (0.68, 17.47) | 7.30 (1.70, 31.32) |
| Women         | 6.00 (2.12, 16.98) | 15.88 (5.71, 44.13) |
| **EWGSOP**    |           |                     |
| Men           | —         | 0.72 (0.09, 5.54)  |
| Women         | 1.50 (0.65, 3.45) | 1.81 (0.78, 4.18)  |

See Table 1 for definitions of low lean mass, weakness and slowness.

* n = 747 men, 819 women.
† n = 745 men, 817 women.
FNIH, Foundation for the National Institutes of Health; EWGSOP, European Working Group on Sarcopenia in Older People.

Supplementary Material
Supplementary material can be found at: http://biomedgerontology.oxfordjournals.org/

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Conflicts of interest

None.

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