# Associations of Skeletal Muscle Mass and Fat Mass With Incident Cardiovascular Disease and All-Cause Mortality: A Prospective Cohort Study of UK Biobank Participants 

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#### Abstract

BACKGROUND: There is debate whether body mass index is a good predictor of health outcomes because different tissues, namely skeletal muscle mass (SMM) and fat mass (FM), may be differentially associated with risk. We investigated the association of appendicular SMM (aSMM) and FM with fatal and nonfatal cardiovascular disease (CVD) and all-cause mortality. We compared their prognostic value to that of body mass index.


METHODS AND RESULTS: We studied 356590 UK Biobank participants aged 40 to 69 years with bioimpedance analysis data for whole-body FM and predicted limb muscle mass (to calculate aSMM). Associations between aSMM and FM with CVD and all-cause mortality were examined using multivariable Cox proportional hazards models. Over 3749501 person-years of follow-up, there were 27784 CVD events and 15844 all-cause deaths. In men, aSMM was positively associated with CVD incidence (hazard ratio [HR] per 1 SD $1.07 ; 95 \% \mathrm{Cl}, 1.06-1.09$ ) and there was a curvilinear association in women. There were stronger positive associations between FM and CVD with HRs per SD of 1.20 ( $95 \% \mathrm{Cl}, 1.19-1.22$ ) and 1.25 ( $95 \% \mathrm{Cl}, 1.23-1.27$ ) in men and women respectively. Within FM tertiles, the associations between aSMM and CVD risk largely persisted. There were J-shaped associations between aSMM and FM with all-cause mortality in both sexes. Body mass index was modestly better at discriminating CVD risk.

CONCLUSIONS: FM showed a strong positive association with CVD risk. The relationship of aSMM with CVD risk differed between sexes, and potential mechanisms need further investigation. Body fat and SMM bioimpedance measurements were not superior to body mass index in predicting population-level CVD incidence or all-cause mortality.

Key Words: all-cause mortality $\square$ cardiovascular disease ■ cohort study $\square$ fat mass $■$ skeletal muscle mass

The increasing prevalence of obesity is a significant public health concern because it is a known risk factor for several noncommunicable diseases, ${ }^{1-5}$ estimated to account for 56 million deaths globally in 2017. ${ }^{6}$ Evidence from prospective cohort studies ${ }^{7-10}$ and meta-analyses of such studies ${ }^{2,11,12}$ has repeatedly
shown a J- or U-shaped relationship between body mass index (BMI), cardiovascular disease (CVD), and all-cause mortality, even after efforts to account for confounding and reverse causality. ${ }^{12,13}$ A potential explanation for this is that BMI does not distinguish between fat mass (FM) and skeletal muscle mass

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## CLINICAL PERSPECTIVE

## What Is New?

- Reports of the relationship between muscle mass and cardiovascular disease (CVD) are inconsistent and have rarely been considered in the context of adiposity; analysis of body composition measured by bioimpedance in this large cohort of UK adults showed that fat mass showed strong positive associations with CVD events.
- Appendicular skeletal muscle mass had a curvilinear association with CVD events in women and a positive association in men; the associations of appendicular skeletal muscle mass and fat mass with all-cause mortality followed a Jshape in both men and women.
- Measurements of body fat and skeletal muscle mass were not superior to body mass index in predicting CVD events or mortality.


## What Are the Clinical Implications?

- Body mass index has been criticized as an inaccurate measure of health risks, but at a population level, more specific measurements of body composition, namely appendicular skeletal muscle mass and fat mass, were generally not more predictive of CVD events or mortality.
- Although body mass index may be the simplest measurement to assess health risk, which is important from a public health perspective, some of this risk may not be attributable solely to adiposity, particularly if the association observed with appendicular skeletal muscle mass in men is confirmed.
- Further research is needed to better understand the biological mechanisms and impact of different body tissue compartment on health outcomes.


## Nonstandard Abbreviations and Acronyms

| aSMM | appendicular skeletal muscle mass |
| :--- | :--- |
| BIA | bioimpedance analysis |
| FM | fat mass |
| SMM | skeletal muscle mass |

(SMM), ${ }^{12,13}$ yet their contribution to the pathogenesis of disease is likely to be different.

A systematic literature review of the associations between body composition and CVD or mortality (Data S1, Tables S1 and S2) showed that the majority of studies found null or inverse associations with SMM, although a minority of studies reported
positive or curvilinear associations. More studies have investigated the relationship of FM with these outcomes, with the majority of them reporting positive associations. Very few studies have investigated the combined impact of both types of tissues, yet weight change is associated with changes in both these body tissue compartments.

Dual energy X-ray absorptiometry (DEXA) or magnetic resonance imaging techniques are considered to be reference methods for the measurement of body composition because of their precision and reliability. ${ }^{14}$ However, these are often not feasible for large studies given they are expensive and not easily portable. ${ }^{15}$ Bioimpedance analysis (BIA) is a noninvasive and practical method to assess FM and SMM in clinical practice and at scale in population-based studies. ${ }^{16-18}$ The UK Biobank uses a bioimpedance analyzer previously validated against DEXA in a mixed population of children and adults, and body composition estimates were found to be more accurate than those obtained from previous BIA estimates. ${ }^{19}$ A recent validation study comparing BIA to DEXA in a subsample of the UK Biobank participants showed BIA to be a valid method for the assessment of appendicular skeletal muscle mass (aSMM) and FM. ${ }^{20}$

In this study we aimed to use BIA-derived aSMM and FM measurements to look at their associations with incident CVD and all-cause mortality in the UK Biobank population. Furthermore, to investigate the prognostic value of these BIA-derived measurements in comparison to more traditional measures such as BMI, grip strength, and waist circumference.

## METHODS

## Data Availability Statement

Researchers can apply to use the UK Biobank resource and access the data used. No additional data are available.

## Study Design and Participants

The UK Biobank recruited 502664 participants aged 40 to 69 years between 2006 and 2010 (response rate $5.5 \%$ ) via mailed invitations to the general public living within 25 miles of one of the 22 assessment centers in England, Scotland, and Wales. ${ }^{21,22}$ At the baseline assessment clinic, participants completed a touch-screen questionnaire and computer-assisted interview, had physical measurements taken, and biological samples collected. ${ }^{23,24}$ UK Biobank received ethical approval from the North West Multicentre Research Ethics Committee (REC reference: 11/NW/03820). All participants gave written informed consent before enrolment in the study, which was
conducted in accordance with the principles of the Declaration of Helsinki.

Participants were excluded from analyses if they had prior CVD (defined later) or diseases that may affect body composition including fractures in the past year, respiratory diseases, musculoskeletal conditions, and some infectious diseases ( $\mathrm{n}=116$ 679; Figure S1). ${ }^{25}$ Participants were also excluded if they had missing data for the exposures ( $n=6751$ ) or were not of a White race ( $n=22644$ ), because BIA estimates are derived from algorithms in White populations, which represent $\approx 95 \%$ of the UK Biobank sample. ${ }^{26-28}$

## Measurement of Exposures

Measures of body weight and body composition (muscle mass and fat mass) were derived from BIA in bare-footed participants wearing light clothing using a Tanita BC418MA single frequency segmental bodycomposition analyzer (Tanita, Tokyo, Japan) at the baseline assessment center visit. The aSMM (kg) was calculated as the sum of the predicted muscle mass from the 4 limbs. Whole body FM (kg) was also obtained from BIA. Standing height was measured using a Seca 202 scale (Seca, Hamburg, Germany). BMI was calculated by dividing weight (kg) by the squared height in meters.

Waist circumference (cm) was measured at the umbilicus using a tape measure. The mean grip strength (kg) of the left and right hands was taken once using a Jamar J00105 hydraulic hand dynamometer.

## Ascertainment of Outcomes

Participants were followed via linkage to National Health Service hospital in-patient data from hospital episode statistics in England, the Scottish Morbidity Records, and the Patient Episode Database for Wales. Patients were identified if they died of any cause or developed incident CVD, defined using International Statistical Classification of Diseases, Tenth Revision (ICD-10) categories: coronary heart disease (I21-I24, 125.6, I42, I43, K49, K50, K75, K40-K46), congestive heart failure or cardiomyopathy ( $150,150.1,150.9,111.0$, I13.0, I13.2), and total stroke (I60-I64). ${ }^{29}$ Follow-up was available until June 30, 2020, October 31, 2016, and February 29, 2016 for England, Scotland, and Wales respectively; and until July 31, 2020 for all-cause mortality for all regions.

## Statistical Analysis <br> Association of Skeletal Muscle Mass and Fat Mass With CVD and Mortality

First, age-adjusted partial correlation coefficients between aSMM, FM, and height were calculated to examine the relationships between the body composition
measurements and overall body size. As aSMM is highly correlated with FM and height, ${ }^{30}$ aSMM was regressed on height and FM and the residuals from this model were divided into sex-specific quintiles for the main analysis.

Multivariable Cox regression analyses with age as the underlying timescale were used to estimate hazard ratios (HRs) and 95\% Cls for the associations of sex-specific fifths of aSMM and fifths of FM as well as per 1 SD with incident CVD and all-cause mortality. All analyses were sequentially adjusted for height and height ${ }^{2}$ (continuous), Townsend index of deprivation (quintiles), level of education (none, vocational qualifications, any degree, higher degree, other), smoking (never, previous, current), alcohol intake (none, <1 unit/ week, 1-14 units/week, 14+ units/week), physical activity derived from metabolic equivalent of task scores (low, moderate, high), ${ }^{31}$ and dietary factors (oily fish intake, saturated fat intake, fruit and vegetable intake [none, low, medium, high intake]), prior medical history (diabetes mellitus, cancer history $>5$ years ago, and menopausal status in women [binary for each]). We created a category for missing values for each of these covariates. Additionally, FM and aSMM were mutually adjusted for each other to assess the independent effects of each type of tissue. See Table S3 for the details on the derivation of these covariates.

The HRs and $95 \%$ Cls were computed using group-specific variances ${ }^{32}$; these reflect the uncertainty in the estimate of risk in each group (including the reference group), thereby allowing comparisons between any 2 quintiles independently of the reference group. Restricted cubic splines with 5 knots were also computed to visually explore nonlinear associations for continuous exposures, and departures from linearity were tested via the likelihood ratio statistic test used to evaluate if models with linear or categorical exposures were a better fit. ${ }^{33}$ Five knots were specified to be consistent with the quintile analysis but also to provide enough flexibility to the model while also not being too many knots so that the model is oversensitive to the smallest fluctuations. ${ }^{34,35}$ To correct for the measurement error that can arise from using a single baseline measurement to estimate long-term exposure status (ie, regression dilution bias), ${ }^{36}$ mean values of BIA measurements at resurvey (2012-2013) from 15694 participants were used in 2 ways. First, the HR ( $95 \% \mathrm{Cl}$ ) in the baseline-defined groups of aSMM and FM were plotted against the mean resurvey values in those baseline-defined groups (termed the "usual" value). Second, where there was evidence of a log-linear reIationship, regression dilution ratios were calculated using the MacMahon-Peto method. ${ }^{37}$ The log HRs (and theirSEs) per 1 SD of baseline aSMM and FM were then divided by the relevant regression dilution
ratio to obtain HRs (and associated 95\% CI) per 1 SD of usual aSMM and FM. ${ }^{37}$

Sensitivity analyses were conducted to assess potential residual confounding or reverse causality so additional exclusions were made for events that occurred during the first 2 years of follow-up to reduce the impact of reverse causality, for outliers, or for participants with BMI over $35 \mathrm{~kg} / \mathrm{m}^{2}$ for whom BIA measurements may be less accurate. Additional adjustments were made for BMI (instead of FM in the SMM model and instead of SMM in the FM model) as well as for hypertension (diagnosed by doctor, taking medication, or blood pressure measurement), and blood cholesterol (defined as taking medication, plus levels of non-high-density lipoprotein cholesterol and triglycerides) to investigate if these are potential mediators of the associations. Finally, Cox regression models were conducted with BMI as the exposure as a "positive control" to confirm that the specified models would produce the same association that has been documented previously. ${ }^{9}$

## Associations of aSMM Within Tertiles of FM

To better assess the independent association of aSMM with the risk of disease, irrespective of its strong correlation with FM, we examined the sexspecific associations of aSMM within subgroups of FM tertiles (subsequently referred to as "body composition groups," because we looked at low/moderate/ high groups [tertiles] of aSMM within groups of FM). Multivariable Cox models adjusting for all covariates listed previously were used to assess the associations with CVD and all-cause mortality using "moderate" aSMM as the reference category within each tertile of FM.

## Prognostic Comparison of aSMM, FM, and Body Composition Groups With BMI, Waist Circumference, and Grip Strength

The relative importance of the various measures in prediction of CVD or mortality was assessed in several ways. First, where a linear association was present, the HRs associated with 1 usual SD change were compared for each measure. In order to assess the discriminatory ability of each measure with CVD and mortality Harrell's C-statistic from the area under the receiver operating curve was computed. ${ }^{38}$ Third, the Wald test $\chi^{2}$ statistic was used to compare a model with just confounders to a model with confounders plus the exposure of interest to explore how much of the variation in risk is explained by each exposure of interest, given confounders. ${ }^{39}$

All analyses were conducted using Stata 15.0 for analyses and R 3.5.2 for graphs. Analyses used

2 -sided $P$ values ( $a=0.05$ ) without any correction for multiple testing.

## RESULTS

## Study Participants

After exclusions, the final sample included 356590 adults who were followed for a median of 10.5 years during which there were 27784 CVD events and 15844 deaths due to all causes. The mean age at recruitment was 56 (SD 8) years. Men had a higher aSMM (median 27.2 kg in men; and 18.3 kg in women), although the difference between the sexes was smaller for FM (median 21.8 kg in men, 26.3 kg in women; Table 1, Tables S4 and S5). There were strong partial correlations between aSMM and FM (men $r=0.71$, women $r=0.78$ ) and aSMM and height (men $r=0.52$, women $r=0.44$ ) but not between FM and height ( $r=$ men 0.14 , women 0.15; Table S6).

Participants in the highest quintiles of aSMM and FM had similar diets (ie, high saturated fat intake, low oily fish intake, but similar fruit and vegetable intakes), and a higher percentage of participants had low physical activity and a higher prevalence of type 2 diabetes mellitus and hypertension. A higher percentage of participants in the highest FM quintile were taking medication for cholesterol but there were no differences across aSMM quintiles (Table 1, Tables S4 and S5).

## Associations of Skeletal Muscle Mass and Fat Mass With Health Outcomes

There was a potential curvilinear association between aSMM and CVD in women (likelihood ratio test statistic of nonlinearity [df=4], $P<0.001$ ) with the nadir approximately at the median; this curvilinear shape was even more pronounced in the cubic spline analysis (Figure 1, Figure S2, Table S7). There was a positive linear association in men with an HR per 1 usual SD of 1.07 (95\% CI, 1.06-1.09). FM showed much stronger positive log-linear associations with the risk of CVD with HRs per 1 usual SD of 1.20 ( $95 \% \mathrm{CI}$, $1.19-1.22)$ in men and 1.25 ( $95 \% \mathrm{Cl}, 1.23-1.27$ ) in women (Figure 1, Figure S2, Table S7). These associations were similar across CVD subtypes (nonfatal, fatal, coronary heart disease, congestive heart failure, and stroke) for both aSMM and FM (Figure S3). The associations of aSMM and FM with all-cause mortality generally followed a J-shape in both men and women, although the association with FM was more clear (Figure 2, Figure S2, Table S8).

These findings remained robust after sensitivity analyses (Tables S9 and S10). Exclusion of the first 2 years of follow-up to reduce the risk of reverse

Table 1. Baseline Characteristics of the Study Population According to Appendicular Skeletal Muscle Mass and Fat Mass Quintiles in 356590 UK Biobank Participants

|  | Men Appendicular Skeletal Muscle Mass Quintiles, Range (kg) |  |  | Men Fat Mass Quintiles, Range (kg) |  |  | Total |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $\begin{gathered} \text { Q1 } \\ 13.5 \text { to } \leq 24.0 \end{gathered}$ | $\begin{gathered} \text { Q3 } \\ 26.1 \text { to } \leq 27.6 \end{gathered}$ | $\begin{gathered} \text { Q5 } \\ 30.1 \text { to } \leq 54.5 \end{gathered}$ | $\begin{gathered} \text { Q1 } \\ 5.0 \text { to } \leq 15.7 \end{gathered}$ | $\begin{gathered} \text { Q3 } \\ 19.5 \text { to } \leq 22.9 \end{gathered}$ | $\begin{gathered} \text { Q5 } \\ 27.6 \text { to } \leq 98.6 \end{gathered}$ |  |
| Age at recruitment, y , mean (SD) | 61.0 (6.4) | 56.0 (7.8) | 51.7 (7.8) | 54.9 (8.3) | 56.5 (8.1) | 56.9 (7.8) | 56.2 (8.1) |
| aSMM, kg, mean (SD) | 24.6 (2.9) | 26.9 (2.9) | 30.5 (3.6) | 24.6 (2.7) | 26.7 (2.7) | 31.0 (3.6) | 27.2 (3.7) |
| FM, kg, mean (SD) | 22.4 (7.1) | 21.4 (7.4) | 22.2 (9.2) | 12.4 (2.4) | 21.0 (1.0) | 33.6 (6.2) | 21.8 (7.8) |
| BMI, kg/m², mean (SD) | 26.0 (3.5) | 27.4 (3.6) | 29.7 (4.4) | 23.3 (1.8) | 27.1 (1.6) | 33.1 (3.5) | 27.6 (4.0) |
| Higher education, n (\%) | 12740 (39.7\%) | 13140 (40.9\%) | $\begin{aligned} & 12645 \\ & (39.3 \%) \end{aligned}$ | 13909 (43.1\%) | 12987 (40.6\%) | 11735 (36.7\%) | 64706 (40.2\%) |
| Current smokers, n (\%) | 3714 (11.6\%) | 3573 (11.1\%) | 3724 (11.6\%) | 4366 (13.5\%) | 3368 (10.5\%) | 3286 (10.3\%) | 18005 (11.2\%) |
| Low fruit and vegetable intake, n (\%) | 14779 (46.0\%) | 13977 (43.5\%) | 13367 (41.6\%) | 13624 (42.2\%) | 14191 (44.3\%) | 14349 (44.8\%) | 70241 (43.7\%) |
| High saturated fat intake, n (\%) | 11706 (36.4\%) | 11647 (36.2\%) | 11778 (36.6\%) | 10457 (32.4\%) | 11699 (36.6\%) | 12933 (40.4\%) | 58554 (36.4\%) |
| Low oily fish intake, n (\%) | 10919 (34.0\%) | 11342 (35.3\%) | 11988 (37.3\%) | 11197 (34.7\%) | 11518 (36.0\%) | 11661 (36.4\%) | 57348 (35.7\%) |
| Heavy drinkers, n (\%) | 19948 (62.1\%) | 19841 (61.7\%) | 18666 (58.1\%) | 17562 (54.4\%) | 20297 (63.4\%) | 19695 (61.5\%) | 97948 (60.9\%) |
| Low physical activity, n (\%) | 7247 (22.6\%) | 6300 (19.6\%) | 5518 (17.2\%) | 4580 (14.2\%) | 6030 (18.8\%) | 8757 (27.4\%) | 31636 (19.7\%) |
| Hypertension, n (\%) | 20146 (62.7\%) | 18102 (56.3\%) | 17370 (54.0\%) | 12683 (39.3\%) | 18523 (57.9\%) | 23544 (73.5\%) | 91858 (57.1\%) |
| Type 2 diabetes mellitus, n (\%) | 1300 (4.1\%) | 1156 (3.6\%) | 1408 (4.4\%) | 388 (1.2\%) | 906 (2.8\%) | 2883 (9.0\%) | 6305 (3.9\%) |
| Cancer history (>5 y ago), n (\%) | 1176 (3.7\%) | 816 (2.5\%) | 652 (2.0\%) | 780 (2.4\%) | 856 (2.7\%) | 889 (2.8\%) | 4233 (2.6\%) |
| Cholesterol medication, n (\%) | 6204 (19.5\%) | 4646 (14.6\%) | 3842 (12.0\%) | 2175 (6.8\%) | 4664 (14.7\%) | 7440 (23.4\%) | 24008 (15.0\%) |
|  | Women <br> Ma | ndicular Skel uintiles, Rang | Muscle <br> g) | Women F | Mass Quintiles | Range (kg) |  |
|  | $\begin{gathered} \text { Q1 } \\ 10.3 \text { to } \leq 16.5 \end{gathered}$ | $\begin{gathered} \text { Q3 } \\ 17.6 \text { to } \leq 18.6 \end{gathered}$ | $\begin{gathered} \text { Q5 } \\ 20.0 \text { to } \leq 39.2 \end{gathered}$ | $\begin{gathered} \text { Q1 } \\ 5.0 \text { to } \leq 18.5 \end{gathered}$ | $\begin{gathered} \text { Q3 } \\ 22.9 \text { to } \leq 27.1 \end{gathered}$ | $\begin{gathered} \text { Q5 } \\ 33.4 \text { to } \leq 109.8 \end{gathered}$ | Total |
| Age at recruitment, y , mean (SD) | 58.9 (7.0) | 55.8 (7.9) | 53.2 (8.1) | 54.1 (8.1) | 56.6 (7.9) | 56.4 (7.8) | 55.9 (8.0) |
| aSMM, kg, mean (SD) | 17.0 (1.7) | 18.0 (1.8) | 20.3 (2.5) | 16.6 (1.5) | 17.9 (1.4) | 21.1 (2.3) | 18.3 (2.3) |
| FM, kg, mean (SD) | 27.7 (8.5) | 25.5 (8.8) | 27.0 (11.9) | 15.2 (2.6) | 24.9 (1.3) | 41.1 (7.4) | 26.3 (9.6) |
| BMI, mean (SD) | 25.9 (4.2) | 26.3 (4.4) | 28.5 (6.0) | 21.6 (1.7) | 25.9 (1.7) | 33.9 (4.3) | 26.7 (4.9) |
| Higher education, n (\%) | 14330 (36.6\%) | 16033 (40.9\%) | 17056 (43.6\%) | 18112 (45.5\%) | 15834 (39.8\%) | 14209 (36.4\%) | 79178 (40.4\%) |
| Current smokers, n (\%) | 3039 (7.8\%) | 3231 (8.2\%) | 3377 (8.6\%) | 3792 (9.5\%) | 3151 (7.9\%) | 2898 (7.4\%) | 16069 (8.2\%) |
| Low fruit and vegetable intake, n (\%) | 12608 (32.2\%) | 11874 (30.3\%) | 10910 (27.9\%) | 12125 (30.5\%) | 11756 (29.6\%) | 12234 (31.3\%) | 59246 (30.3\%) |
| High saturated fat intake, n (\%) | 11699 (29.9\%) | 11252 (28.7\%) | 11212 (28.6\%) | 10415 (26.2\%) | 11615 (29.2\%) | 12439 (31.9\%) | 57055 (29.1\%) |
| Low oily fish intake, n (\%) | 12200 (31.2\%) | 12619 (32.2\%) | 13137 (33.6\%) | 12971 (32.6\%) | 12669 (31.8\%) | 13082 (33.5\%) | 63668 (32.5\%) |
| Heavy drinkers, n (\%) | 12955 (33.1\%) | 13025 (33.2\%) | 12318 (31.5\%) | 13562 (34.1\%) | 13525 (34.0\%) | 10970 (28.1\%) | 64272 (32.8\%) |
| Low physical activity, n (\%) | 10117 (25.8\%) | 8431 (21.5\%) | 7837 (20.0\%) | 6398 (16.1\%) | 8288 (20.8\%) | 12160 (31.1\%) | 43345 (22.1\%) |
| Hypertension, n (\%) | 19371 (49.5\%) | 16440 (42.0\%) | 16330 (41.7\%) | 11345 (28.5\%) | 17091 (43.0\%) | 23766 (60.9\%) | 85687 (43.8\%) |
| Type 2 diabetes mellitus, n (\%) | 635 (1.6\%) | 669 (1.7\%) | 1250 (3.2\%) | 239 (0.6\%) | 524 (1.3\%) | 2046 (5.3\%) | 4032 (2.1\%) |
| Cancer history (>5 y ago), n (\%) | 2540 (6.5\%) | 2010 (5.1\%) | 1845 (4.7\%) | 1912 (4.8\%) | 2278 (5.7\%) | 2170 (5.6\%) | 10626 (5.4\%) |
| Cholesterol medication, n (\%) | 4294 (11.0\%) | 3163 (8.1\%) | 3080 (7.9\%) | 1536 (3.9\%) | 3349 (8.5\%) | 5721 (14.8\%) | 17157 (8.8\%) |
| Postmenopausal, n (\%) | 28352 (72.4\%) | 23352 (59.6\%) | 17594 (44.9\%) | 21266 (53.5\%) | 24686 (62.1\%) | 22973 (58.8\%) | $\begin{aligned} & 116065 \\ & (59.3 \%) \end{aligned}$ |

$\chi^{2}$ test for trend was performed with $P<0.05$ for all characteristics across the aSMM and FM quintiles. All characteristics were determined at the baseline assessment clinic through touch-screen questionnaires, interviews, and/or physical measurements. Higher education: college or university degree or professional qualifications. Low physical activity: $<600$ metabolic equivalent (MET)-minutes per week. ${ }^{31}$ Heavy alcohol drinker: $>14$ units of alcohol a week). ${ }^{40}$ Hypertension: systolic blood pressure $>140 \mathrm{~mm} \mathrm{Hg}$, diastolic blood pressure $>90 \mathrm{~mm} \mathrm{Hg}$, was diagnosed by a doctor or were taking medication to lower blood pressure. Diabetes mellitus and cholesterol: taking medication for these conditions or diagnosed by a doctor. Cancer history: diagnosed with cancer $>5$ years ago (those with more recent cancer had been excluded). Low fruit and vegetable intake: the lowest consumption tertile (<21 portions per week). High saturated fat: the highest saturated fat tertile, based on portions per week of beef, lamb, pork, and whether they consumed animal- or plant-based spreads. Low oily fish: lowest consumption tertile (<1 portion per week). aSMM indicates appendicular skeletal muscle mass; BMI, body mass index; and FM, fat mass.


Figure 1. Adjusted hazard ratios (HRs) of incident cardiovascular disease associated with appendicular skeletal muscle mass (aSMM) and fat mass (FM).
A, HRs of incident CVD associated with aSMM in men, $1 \mathrm{SD}=5.66 \mathrm{~kg}$. B, HRs of incident CVD associated with aSMM in women, $1 \mathrm{SD}=1.45 \mathrm{~kg}$. C, HRs of incident CVD associated with FM in men, $1 \mathrm{SD}=6.75 \mathrm{~kg}$. D, HRs of incident CVD associated with FM in women, $1 \mathrm{SD}=8.28 \mathrm{~kg}$. For all panels, likelihood ratio tests were used to estimate nonlinearity (aSMM in men, $P=0.04$; aSMM in women, $P<0.001$; FM in men, $P=0.09$; FM in women, $P=0.09$ ). Adjusted HRs and Cls obtained using the floated absolute risk method of Cox proportional hazards regression, number of cases shown above each estimate and HRs shown below. Adjusted for age (underlying timescale variable), height (as a continuous variable in FM and included by regression out of variation due to height for aSMM), Townsend index of deprivation, education, smoking status, alcohol intake, physical activity, oily fish intake, fruit and vegetable intake, saturated fat intake, diabetes mellitus, cancer history, menopause (women), and mutually adjusted for FM (in the aSMM models) and aSMM (in the FM models). HRs are plotted at the mean of the resurvey values for the baseline-defined quintiles ("usual" values) to correct for measurement error. HRs per 1 SD given where there was no evidence of departure from linearity. CVD indicates cardiovascular disease.
causality, outliers, or participants with $\mathrm{BMI}>35 \mathrm{~kg} / \mathrm{m}^{2}$ did not affect the associations.

Adjustment for hypertension and high blood cholesterol as mediators did not affect associations of aSMM
with CVD or all-cause mortality. These mediators explained $\approx 30 \%$ to $40 \%$ of the $\chi^{2}$ statistic in models of FM and CVD, although the association between FM and all-cause mortality was affected less (Tables S11 and


Figure 2. Adjusted hazard ratios (HRs) of all-cause mortality associated with appendicular skeletal muscle mass (ASMM) and fat mass (FM).
A, HRs of all-cause mortality associated with aSMM in men, $1 \mathrm{SD}=5.66 \mathrm{~kg}$. B, HRs of all-cause mortality associated with aSMM in women, $1 \mathrm{SD}=1.45 \mathrm{~kg} . \mathrm{C}$, HRs of all-cause mortality associated with FM in men, $1 \mathrm{SD}=6.75 \mathrm{~kg}$. D, HRs of all-cause mortality associated with FM in women, $1 \mathrm{SD}=8.28 \mathrm{~kg}$. For all panels, likelihood ratio tests were used to estimate nonlinearity $P$ values (aSMM in men, $P=0.002$; aSMM in women, $P=0.008$; FM in men, $P<0.001$; FM in women, $P<0.001$ ). Adjusted HRs and Cls obtained using the floated absolute risk method of Cox proportional hazards regression, number of cases shown above each estimate and HRs shown below. Adjusted for age (underlying timescale variable), height (as a continuous variable in FM and included by regression out of variation due to height for aSMM), Townsend index of deprivation, education, smoking status, alcohol intake, physical activity, oily fish intake, fruit and vegetable intake, saturated fat intake, diabetes mellitus, cancer history, menopause (women), and mutually adjusted for FM (in the aSMM models) and aSMM (in the FM models). HRs are plotted at the mean of the resurvey values for the baseline-defined quintiles ("usual" values) to correct for measurement error.

S12). However, adjustment for BMI (instead of SMM or FM in their respective models) removed the positive association between aSMM and CVD in men but did not change the association observed in women. Associations between FM and CVD and all-cause mortality were largely attenuated after adjustment for BMI, with large \% of the $\chi^{2}$ statistic explained in both men and women.

Analyses of body composition groups (aSMM tertiles within each FM tertile) showed positive linear associations between aSMM and incident CVD risk in men in all FM tertiles, whereas women still had curvilinear associations with incident CVD except those in the highest FM tertile (Figure 3). The associations of aSMM with all-cause mortality within FM tertiles were broadly similar between men and women and to those observed in the main analysis, except for women in the middle tertile of FM, which showed a curvilinear association between aSMM and all-cause mortality.

## Comparing the Prognostic Value of Body Composition Measures

For CVD risk, waist circumference and FM showed the strongest associations whereas aSMM and grip strength showed the weakest associations in both men and women. For all-cause mortality, waist circumference and grip strength showed the strongest associations, whereas aSMM remained the weakest (Figure 4, Table 2). However, the discriminatory performance (Harrell's C-statistics) for total CVD was slightly higher for BMI compared with all the other metrics in men ( $\mathrm{C}=0.63$; $95 \% \mathrm{Cl}, 0.63-0.64$ ) and for BMI and waist circumference in women ( $\mathrm{C}=0.45$; $95 \% \mathrm{Cl}, 0.45-0.45$ ). Similarly, for all-cause mortality, the discriminatory performance was highest for $\mathrm{BMI}(\mathrm{C}=0.62 ; 95 \% \mathrm{Cl}, 0.62-0.63)$ and waist circumference ( $\mathrm{C}=0.61 ; 95 \% \mathrm{Cl}, 0.61-0.62$ ) in men, whereas in women it was FM ( $\mathrm{C}=0.63$; $95 \% \mathrm{Cl}$, $0.62-0.63$ ) and aSMM ( $\mathrm{C}=0.63$; 95\% CI, 0.62-0.63). The $\chi^{2}$ statistic was marginally higher in the BMI and waist circumference model in men and in the BMI and the combined aSMM/FM groups in women.

## DISCUSSION

In this prospective study of 356590 UK adults, FM had a strong positive log-linear association with the risk of CVD in both sexes. There was also a positive log-linear association with aSMM for men and a curvilinear association for women. The associations of aSMM and FM with all-cause mortality followed a J-shape in both men and women. Analysis of the association of aSMM within tertiles of FM supported these associations with CVD and all-cause mortality. The discriminatory ability of BMI was similar to, or better than, more specific measures of body composition (aSMM and FM), waist circumference, or grip strength in relation to CVD events or all-cause mortality.

Few previous studies have specifically examined the association between distinct body compartments with either incident CVD or mortality. In line with previous studies we consistently observed a positive association between FM and CVD. This is consistent with previous analyses from UK Biobank that found significant associations between body fat percentage, waist circumference, and waist-to-hip ratio on CVD outcomes ${ }^{9}$ as well as meta-analyses of prospective cohort studies assessing various adiposity measures. ${ }^{3,41}$

The role of aSMM has been investigated in fewer studies, most of which used older populations with small sample sizes or a proxy for aSMM such as fat-free mass. ${ }^{42-46}$ Our rationale for using aSMM as opposed to whole body muscle or fat-free mass is because this tissue is more likely to be modifiable by lifestyle factors such as physical activity than other components of fat-free mass and it is less likely to be confounded by FM given that higher abdominal FM is often accompanied by greater muscle in the trunk region. ${ }^{30}$ However, aSMM is a large contributor to whole body muscle and it is likely that participants would be classified in the same quintile regardless of the measure used. Although some studies have shown an inverse association between aSMM and CVD risk, our finding of a positive log-linear association among men

Figure 3. Adjusted hazard ratios (HRs) of cardiovascular disease and all-cause mortality associated with appendicular skeletal muscle mass (aSMM) when participants are stratified into fat mass (FM) tertiles.
A, HRs of cardiovascular disease (CVD) associated with aSMM in low fat men. B, HRs of CVD associated with aSMM in moderate fat men. C, HRs of CVD associated with aSMM in high fat men. D, HRs of CVD associated with aSMM in low fat women. E, HRs of CVD associated with aSMM in moderate fat women. F, HRs of CVD associated with aSMM in high fat women. G, HRs of all-cause mortality associated with aSMM in low fat men. H, HRs of all-cause mortality associated with aSMM in moderate fat men. I, HRs of all-cause mortality associated with aSMM in high fat men. J, HRs of all-cause mortality associated with aSMM in low fat women. K, HRs of allcause mortality associated with aSMM in moderate fat women. L, HRs of all-cause mortality associated with aSMM in high fat women. For all panels, adjusted hazard ratios (HR) and Cls obtained using Cox proportional hazards regression, number of cases shown above each estimate and HRs shown below. Adjusted for age (underlying timescale variable), height (included by regression out of variation due to height), Townsend index of deprivation, education, smoking status, alcohol intake, physical activity, oily fish intake, fruit and vegetable intake, saturated fat intake, diabetes mellitus, cancer history, menopause (women), and mutually adjusted for FM (in the aSMM models) and aSMM (in the FM models). HRs are plotted at the mean of the resurvey values for the baseline-defined quintiles ("usual" values) to correct for measurement error.
has been observed previously. The Aerobics Center Longitudinal Study found a similar pattern with fatfree mass index measured by skinfold thicknesses as well as hydrostatic weighing (for which aSMM is
the largest contributor) and had a comparably aged, predominantly male study population. ${ }^{47}$ A plausible physiological mechanism linking higher aSMM with higher CVD risk may be a higher circulating blood

## Cardiovascular disease (CVD)



## All-cause mortality













Figure 4. Independent effects of body mass index (BMI), fat mass (FM), waist circumference, appendicular skeletal muscle mass (aSMM), and grip strength on cardiovascular disease (CVD) subtypes and all-cause mortality. Adjusted hazard ratios (HRs) per SD change.
A, The independent effects of BMI, FM, waist circumference, aSMM and grip strength on CVD subtypes and all-cause mortality in men. B, The independent effects of BMI, FM, waist circumference, aSMM, and grip strength on CVD subtypes and all-cause mortality in women. Range excludes outliers. Adjusted hazard ratios (HR) and Cls obtained using Cox proportional hazard regression. Adjusted for age (underlying timescale variable), height (as a continuous variable in all models except aSMM where it was included by regression out of variation due to height for aSMM), Townsend index of deprivation, education, smoking status, alcohol intake, physical activity, oily fish intake, fruit and vegetable intake, saturated fat intake, diabetes mellitus, cancer history, menopause (women), and mutually adjusted for FM (in the aSMM models) and aSMM (in the FM models). HRs are corrected for regression dilution bias by the MacMahonPeto method.
volume, which increases cardiac output and increases systolic blood pressure and the risk of heart failure, a phenomenon previously described mainly among people with obesity. ${ }^{48-50} \mathrm{~A}$ recent literature review has provided a more counterintuitive view of the role of lean mass on metabolic health, proposing the possibility of publication bias, especially if unexpected results were found. ${ }^{51}$ Our analysis within tertiles of FM confirmed the increased risk of CVD with aSMM even among men with lower FM levels, reducing the possibility of residual confounding by FM although this cannot be completely ruled out. Our exploratory mediation analyses showed that the association between aSMM and CVD was no longer significant in men after adjusting for BMI. This implies that if aSMM increases, FM plus all the other body compartments have to decrease in order to hold BMI constant, such that changes in body composition that increase skeletal muscle while lowering total body fat, as expected with physical training, may not be associated with increased CVD risk. Furthermore, the large changes observed in the $\chi^{2}$ statistic after adjustment for BMI in the aSMM model suggest a
large part of the association may be explained by confounding by BMI, especially among men.

Nevertheless, it is unusual to find a CVD risk factor with such different associations between sexes. ${ }^{52}$ A potential explanation for this disparity could be because of differences in lifestyle factors between men and women classified as high aSMM within each FM tertile. For example, compared with women, a higher percentage of men in the same aSMM and FM tertile, reported poorer diets (low fruit and vegetable intake, high saturated fat intake), heavy drinking (over 14 units/ week, National Health Service guidelines), or presented a higher prevalence of hypertension and cholesterol medication. Our findings are largely consistent with those reported from a recent study of 38000 middleaged men that demonstrated a U-shaped association between predicted lean mass and CVD death and mortality; however, these participants may have been healthier because they recruited health professionals rather than the general population. ${ }^{53}$ Although we adjusted for several potential lifestyle confounders our study may still have residual confounding in relation to lifestyle factors.

Table 2. The Discrimination Ability of Each Body Composition Measure for the Prediction of Cardiovascular Events and All-Cause Mortality, as Calculated by Harrell's C-Statistic From the Area Under the Receiver Operating Curve

|  | Men |  | $\chi_{1}^{2}$ | Women |  | $\chi_{1}^{2}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | HR Per SD (95\% CI) | Harrell's C-Statistic (95\% CI) |  | HR Per SD ( $95 \% \mathrm{CI}$ ) | Harrell's C-Statistic (95\% CI) |  |
| Cardiovascular disease |  |  |  |  |  |  |
| BMI | 1.15 (1.14-1.16) | 0.63 (0.63-0.64) | 597 | 1.18 (1.16-1.19) | 0.45 (0.45-0.45) | 462 |
| FM | 1.20 (1.19-1.22) | 0.56 (0.55-0.56) | 445 | 1.25 (1.23-1.27) | 0.59 (0.58-0.59) | 374 |
| Waist circumference | 1.23 (1.21-1.25) | 0.61 (0.61-0.62) | 492 | 1.31 (1.28-1.33) | 0.45 (0.44-0.45) | 464 |
| aSMM | 1.07 (1.06-1.09) | 0.54 (0.54-0.54) | 73 | 1.00 (0.98-1.02) | 0.57 (0.56-0.57) | 0 |
| Decreasing grip strength | 1.09 (1.07-1.12) | 0.55 (0.54-0.55) | 42 | 1.11 (1.08-1.14) | 0.44 (0.43-0.44) | 51 |
| Body composition groups | $\ldots$ | 0.56 (0.55-0.56) | 504 | ... | 0.45 (0.44-0.45) | 400 |
| All-cause mortality |  |  |  |  |  |  |
| BMI | 1.08 (1.06-1.09) | 0.62 (0.62-0.63) | 79 | 1.06 (1.05-1.08) | 0.61 (0.61-0.62) | 40 |
| FM | 1.08 (1.06-1.1) | 0.60 (0.59-0.60) | 39 | 1.05 (1.02-1.08) | 0.63 (0.62-0.63) | 11 |
| Waist circumference | 1.13 (1.1-1.15) | 0.61 (0.61-0.62) | 84 | 1.09 (1.06-1.12) | 0.61 (0.6-0.62) | 29 |
| aSMM | 1.04 (1.01-1.06) | 0.60 (0.59-0.60) | 9 | 0.98 (0.92-1.04) | 0.63 (0.62-0.63) | 0 |
| Decreasing grip strength | 1.17 (1.13-1.2) | 0.60 (0.60-0.61) | 70 | 1.08 (1.04-1.11) | 0.61 (0.60-0.61) | 18 |
| Body composition groups | $\ldots$ | 0.61 (0.61-0.62) | 83 | $\ldots$ | 0.61 (0.61-0.62) | 41 |

Harrell's C-statistic and hazard ratios (HR) per SD change calculated from the fully-adjusted model, which adjusted for: age (underlying timescale variable), height, Townsend index of deprivation, education, smoking status, alcohol intake, physical activity, oily fish intake, fruit and vegetable intake, saturated fat intake, diabetes mellitus, cancer history, menopause (women), and mutually adjusted for FM (in the aSMM models) and aSMM (in the FM models). HRs are corrected for regression dilution bias using the MacMahon-Peto method. One SD of aSMM is 3.34 kg (men), 1.95 kg (women) and FM is $6.79 \mathrm{~kg}(\mathrm{men}), 8.29 \mathrm{~kg}$ (women). The model for aSMM in men is for increasing aSMM; in women is for decreasing aSMM. Wald test $\chi_{1}^{2}$ statistic was used to compare a model with just confounders to a model with confounders plus the exposure of interest. aSMM indicates appendicular skeletal muscle mass; BMI, body mass index; and FM, fat mass.

## Clinical and Public Health Implications

BMI has been criticized as an inaccurate measure of health risks, ${ }^{54,55}$ but at a population level, more specific measurements of body composition, namely aSMM and FM, were generally not more predictive of CVD events or mortality; an observation that also been reported elsewhere. $47,56,57$ The moderately improved prognostic value of BMI may reflect the combined effects of height, FM, and SMM that are each individually associated with CVD risk. ${ }^{58}$ In addition, BMI has less measurement error than other measures that could contribute to its marginally stronger prognostic ability. ${ }^{50}$ Waist circumference and other measures of central adiposity have been reported to better discriminate CVD risk in some studies, ${ }^{59}$ although not superior to BMI in others ${ }^{60}$ as happened in our study. However, waist circumference is particularly liable to observer error, ${ }^{50}$ whereas measures of central adiposity do not indicate whole-body adiposity, nor is there an equivalent measure for fat-free mass. Grip strength is often used as a functional indicator of SMM; however, it includes a volitional component and the European Working Group of Sarcopenia in Older People recommends measuring the amount of SMM to assess risk. ${ }^{61}$ However, the commonly used measure of the mid-upper arm circumference is vulnerable to overestimation because it cannot distinguish between muscle fibers and intramuscular fat deposits. ${ }^{61}$

However, although BMI may be the simplest measurement to assess health risk, which is important from a public health perspective, some of this risk may not be attributable solely to adiposity, particularly if the association observed with aSMM in men is confirmed, although further research is needed to better understand the biological mechanisms and impact of different body tissue compartment on health outcomes. It could therefore be beneficial to reframe BMI as a composite measure of risk. ${ }^{47,58}$ In addition, at the individual level, additional measurements of CVD risk factors (eg, blood lipids, blood pressure) in addition to BMI or body composition are needed to classify individuals at risk and propose adequate treatments.

## Strengths and Weaknesses of the Study

The strengths of this study include its large sample size, which reduces the risk of chance findings owing to random error, and the detailed measurements of the exposures, potential confounders, and outcomes from the hospital episode statistics follow-up. Some participants had repeated measurements taken at resurvey, which allowed for the correction for random measurement error and consequent regression dilution bias. ${ }^{36}$

Although BIA has many practical strengths in research and clinical settings, it is not as accurate as other methods that use physical properties of the
body to measure composition, such as densitometry or DEXA or imaging methods such magnetic resonance imaging scans, and is vulnerable to estimation errors, especially at the extreme ranges of BMI or in people with conditions that affect water retention. ${ }^{17,62}$ However, validation studies against DEXA show that it performs well in healthy individuals with a stable electrolyte and water balance. ${ }^{20}$ Because algorithms to estimate body composition by BIA vary, it may be that our results can be replicated only using a Tanita BC-418 MA segmental body composition analyzer. However, studies comparing different analyzers from this manufacturer or others have reported only small differences in \% body fat (eg, equivalent to 0.7 kg of difference in FM ), ${ }^{63}$ suggesting that a participant would likely fall into the same quintile regardless of the method used.

Despite the large sample of participants studied here, one of the main limitations is that the UK Biobank presents a low response rate for the United Kingdom (5.5\%); however, the associations in this study should still be valid and not affected by selection bias. ${ }^{64}$ The participants were predominantly people of White race, which limits the generalizability of our findings. Despite the performance of BMI in this high-income population of UK Biobank, there is emerging evidence that suggests that BMI is not an informative measure of risk for mortality in lean populations from low- and middle-income countries. For example, research from a large cohort study of 0.5 million adults in India found little association between BMI and cardiac mortality. ${ }^{65}$ It may be that BMI is a better indicator of risk in populations where fat mass is the dominant type of body tissue, supported in the current study by the largely equivalent associations of BMI and fat mass with the risk of CVD. Research has also suggested that the distribution of fat mass within equivalent levels of BMI may have distinct associations with the risk of cardiometabolic diseases in different ethnic groups. ${ }^{66}$ Thus, although BMI may be the most effective tool for risk assessment within high-income populations further work is needed to compare its prognostic ability with more detailed measures of body composition in diverse populations. Finally, as this is an observational study we cannot eliminate the possibility that residual confounding affected our results.

## CONCLUSIONS

FM showed a strong positive association with CVD risk whereas SMM showed a positive log-linear association with CVD risk in men but curvilinear in women. Although BMI has been criticized as an
inaccurate measure of risk, more specific measurements of body composition did not demonstrate improved prognostic ability to detect the risk of CVD or all-cause mortality.

## ARTICLE INFORMATION

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Author contributions: Carter, Piernas, and Jebb conceived and designed the research question. Knowles, Carter, and Piernas prepared the data for analysis, analyzed the data, and wrote the first draft of the article; and Jebb, Lewington, and Bennett provided input on data analysis and interpretation of results. All authors revised the article critically for important intellectual content and read and approved the final article. Piernas is the guarantor. The corresponding author attests that all listed authors meet authorship criteria and that no others meeting the criteria have been omitted.

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## Disclosures

None.

## Supplementary Material

Data S1
Tables S1-S12
Figures S1-S3
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## REFERENCES

1. Yusuf S, Hawken S, Ounpuu S, Dans T, Avezum A, Lanas F, McQueen M, Budaj A, Pais P, Varigos J, et al. Effect of potentially modifiable risk factors associated with myocardial infarction in 52 countries (the INTERHEART study): case control study. Lancet. 2004;364:937-952. DOI: 10.1016/S0140-6736(04)17018-9.
2. Whitlock G, Lewington S, Sherliker P, Clarke R, Emberson J, Halsey J, Qizilbash N, Collins R, Peto R, MacMahon S, et al. Body-mass index and cause-specific mortality in 900000 adults: collaborative analyses of 57 prospective studies. Lancet. 2009;373:1083-1096. DOI: 10.1016/S0140-6736(09)60318-4.
3. The Emerging Risk Factor Collaboration. Separate and combined associations of body-mass index and abdominal adiposity with cardiovascular disease: collaborative analysis of 58 prospective studies. Lancet. 2011;377:1085-1095. DOI: 10.1016/S0140-6736(11)60105-0.
4. Renehan AG, Tyson M, Egger M, Heller RF, Zwahlen M. Body-mass index and incidence of cancer: a systematic review and meta-analysis of prospective observational studies. Lancet. 2008;371:569-578. DOI: 10.1016/S0140-6736(08)60269-X.
5. Ng M, Fleming T, Robinson M, Thomson B, Graetz N, Margono C Mullany EC, Biryukov S, Abbafati C, Abera SF, et al. Global, regional, and national prevalence of overweight and obesity in children and adults during 1980-2013: a systematic analysis for the Global Burden of Disease Study 2013. Lancet. 2014;384:766-781. DOI: 10.1016/ S0140-6736(14)60460-8.
6. IHME. GBD results tool, global health data exchange. 2019
7. Hubert HB, Feinleib M, McNamara PM, Castelli WP. Obesity as an independent risk factor for cardiovascular disease: a 26-year fol-low-up of participants in the Framingham Heart Study. Circulation. 1983;67:968-977. DOI: 10.1161/01.CIR.67.5.968.
8. Adams KF, Schatzkin A, Harris TB, Kipnis V, Mouw T, Ballard-Barbash R, Hollenbeck A, Leitzmann MF. Overweight, obesity and mortality in a large prospective cohort of persons 50 to 71 years old. N Engl J Med. 2006;355:763-778. DOI: 10.1056/NEJMoa055643.
9. Iliodromiti S, Celis-Morales CA, Lyall DM, Anderson J, Gray SR, Mackay DF, Nelson SM, Welsh P, Pell JP, Gill JMR, et al. The impact of confounding on the associations of different adiposity measures with the incidence of cardiovascular disease: a cohort study of 296535 adults of white European descent. Eur Heart J. 2018;39:1514-1520. DOI: 10.1093/eurheartj/ehy057.
10. Wade KH, Carslake D, Sattar N, Davey Smith G, Timpson NJ. Obesity BMI and Mortality in UK Biobank: revised estimates using Mendelian randomization. Obesity. 2018;26:1796-1806. DOI: 10.1002/oby. 22313.
11. Romero-Corral A, Montori VM, Somers VK, Korinek J, Thomas RJ, Allison TG, Mookadam F, Lopez-Jimenez F. Association of bodyweight with total mortality and with cardiovascular events in coronary artery disease: a systematic review of cohort studies. Lancet. 2006;368:666678. DOI: 10.1016/S0140-6736(06)69251-9.
12. Di Angelantonio E, Bhupathiraju SN, Wormser D, Gao P, Kaptoge S, de Gonzalez AB, Cairns BJ, Huxley R, Jackson CL, Joshy G, et al. Body-mass index and all-cause mortality: individual-participant-data meta-analysis of 239 prospective studies in four continents. Lancet. 2016;388:776-786. DOI: 10.1016/S0140-6736(16)30175-1.
13. Flegal KM, Kit BK, Orpana H, Graubard BI. Association of allcause mortality with overweight and obesity using standard body mass index categories. JAMA. 2013;309:71-82. DOI: 10.1001/ jama.2012.113905.
14. Janssen I, Heymsfield SB, Baumgartner RN, Ross R. Estimation of skeletal muscle mass by bioelectrical impedance analysis. J App/ Physiol. 2000;89:465-471. DOI: 10.1152/jappl.2000.89.2.465.
15. Kuriyan R. Body composition techniques. Indian $J$ Med Res. 2018;148:648-658. DOI: 10.4103/ijmr.IJMR_1777_18.
16. Franssen FME, Rutten EPA, Groenen MTJ, Vanfleteren LE, Wouters EFM, Spruit MA. New reference values for body composition by bioelectrical impedance analysis in the general population: results from the UK Biobank. J Am Med Dir Assoc. 2014;15:448.e1-448.e6. DOI: 10.1016/j.jamda.2014.03.012.
17. Uszko-Lencer NHMK, Bothmer F, Van Pol PEJ, Schols AMWJ. Measuring body composition in chronic heart failure: a comparison of methods. Eur J Heart Fail. 2006;8:208-214. DOI: 10.1016/j.ejhea rt.2005.07.007.
18. Kyle UG, Bosaeus I, De Lorenzo AD, Deurenberg P, Elia M, Gómez JM, Heitmann BL, Kent-Smith L, Melchior JC, Pirlich M, et al. Bioelectrical impedance analysis-part I: review of principles and methods. Clin Nutr. 2004;23:1226-1243. DOI: 10.1016/j.clnu.2004.06.004.
19. Pietrobelli A, Rubiano F, St-Onge M-P, Heymsfield SB. New bioimpedance analysis system: improved phenotyping with wholebody analysis. Eur J Clin Nutr. 2004;58:1479-1484. DOI: 10.1038/ sj.ejen. 1601993.
20. Lee MM, Jebb SA, Oke J, Piernas C. Reference values for skeletal muscle mass and fat mass measured by bioelectrical impedance in 390565 UK adults. J Cachexia Sarcopenia Muscle. 2020;11:487-496. DOI: 10.1002/jcsm. 12523.
21. Collins R. What makes UK Biobank special? Lancet. 2012;379:11731174. DOI: 10.1016/S0140-6736(12)60404-8.
22. Allen N, Sudlow C, Downey P, Peakman T, Danesh J, Elliott P, Gallacher J, Green J, Matthews P, Pell J, et al. UK Biobank: current status and what it means for epidemiology. Health Policy Technol. 2012;1:123126. DOI: 10.1016/j.hlpt.2012.07.003.
23. Biobank UK. Protocol for a large-scale prospective epidemiological resource. 2007.
24. Sudlow C, Gallacher J, Allen N, Beral V, Burton P, Danesh J, Downey P, Elliott P, Green J, Landray M, et al. UK Biobank: an open access
resource for identifying the causes of a wide range of complex diseases of middle and old age. PLoS Med. 2015;12:1-10. DOI: 10.1371/ journal.pmed.1001779.
25. Kalyani RR, Corriere M, Ferrucci L. Age-related and disease-related muscle loss: the effect of diabetes, obesity, and other diseases. Lancet Diabetes Endocrinol. 2014;2:819-829. DOI: 10.1016/S2213 -8587(14)70034-8.
26. Abramowitz MK, Hall CB, Amodu A, Sharma D, Androga L, Hawkins M. Muscle mass, BMI, and mortality among adults in the United States: a population- based cohort study. PLoS One. 2018;13:1-16. DOI: 10.1371/journal.pone.0194697.
27. Baumgartner R, Koehler K, Gallagher D, Romero L, Heymsfield S, Ross R, Garry P, Lindeman R. Epidemiology of sarcopenia among the elderly in New México. Am J Epidemiol. 1998;147:755-763. DOI: 10.1093/oxfordjournals.aje.a009520.
28. Newman AB, Kupelian V, Visser M, Simonsick E, Goodpaster B, Nevitt M, Kritchevsky SB, Tylavsky FA, Rubin SM, Harris TB. Sarcopenia: alternative definitions and associations with lower extremity function. J Am Geriatr Soc. 2003;51:1602-1609. DOI: 10.1046/j.1532-5415.2003.51534.x.
29. WHO. International Statistical Classification of Diseases and Related Health Problems. 10th Revision ed. WHO; 2016.
30. Bosy-Westphal A, Muller MJ. Identification of skeletal muscle mass depletion across age and BMI groups in health and disease-there is need for a unified definition. Int J Obes. 2005;39:379-386. DOI: 10.1038/ijo.2014.161.
31. IPAQ. IPAQ scoring protocol. 2005.
32. Plummer M. Improved estimated of floating absolute risk. Stat Med. 2004;23:93-104. DOI: 10.1002/sim. 1485.
33. Collett D. Modelling Survival Data in Medical Research. Boca Raton, FL: CRC Press; 2015.
34. Durrleman S, Simon R. Flexible regression models with cubic splines. Stat Med. 1989;8:551-561. DOI: 10.1002/sim.4780080504.
35. Govindarajulu US, Spiegelman D, Thurston SW, Ganguli B, Eisen EA. Comparing smoothing techniques in Cox models for exposureresponse relationships. Stat Med. 2007;26:3735-3752. DOI: 10.1002/ sim. 2848.
36. Clarke R, Shipley M, Lewington S, Youngman L, Collins R, Marmot M, Peto R. Underestimation of risk associations due to regression dilution in long-term follow-up of prospective studies. Am J Epidemiol. 1999;150:341-353. DOI: 10.1093/oxfordjournals.aje.a010013.
37. MacMahon S, Peto R, Cutler J, Collins R, Sorlie P, Neaton J, Abbott R, Godwin J, Dyer A, Stamler J. Blood pressure, stroke, and coronary heart disease. Part 1, prolonged differences in blood pressure: prospective observational studies corrected for the regression dilution bias. Lancet. 1990;335:765-774. DOI: 10.1016/0140-6736(90)90878 -9.
38. Altman D. Practical Statistics for Medical Research. London: Chapman \& Hall; 1999.
39. Agresti A. Building and Extending Loglinear/Logit Models. 2nd ed. Hoboken, NJ: John Wiley \& Sons; 2002:357-408.
40. NHS. NHS alcohol units. 2018.
41. Dijk SBV, Takken T, Prinsen EC, Wittink H. Different anthropometric adiposity measures and their association with cardiovascular disease risk factors: a meta-analysis. Neth Heart J. 2012;20:208-218. DOI: 10.1007/s12471-011-0237-7.
42. Brown JC, Harhay MO, Harhay MN. Sarcopenia and mortality among a population-based sample of community-dwelling older adults. J Cachexia Sarcopenia Muscle. 2016;7:290-298. DOI: 10.1002/ jcsm. 12073.
43. Cesari M, Pahor M, Lauretani F, Zamboni V, Bandinelli S, Bernabei R, Guralnik JM, Ferrucci L. Skeletal muscle and mortality results from the InCHIANTI study. J Gerontol A Biol Sci Med Sci. 2009;64:377-384. DOI: 10.1093/gerona/gln031.
44. Batsis JA, Mackenzie TA, Emeny RT, Lopez-Jimenez F, Bartels SJ. Low lean mass with and without obesity, and mortality: results from the 1999-2004 National Health and Nutrition Examination Survey. J Gerontol A Biol Sci Med Sci. 2017;72:1445-1451. DOI: 10.1093/geron a/glx002.
45. Spahillari A, Mukamal KJ, DeFilippi C, Kizer JR, Gottdiener JS, Djoussé L, Lyles MF, Bartz TM, Murthy VL, Shah RV. The association of lean and fat mass with all-cause mortality in older adults: the Cardiovascular Health Study. Nutr Metab Cardiovasc Dis. 2016;26:1039-1047. DOI: 10.1016/j.numecd.2016.06.011.
46. Srikanthan P, Karlamangla AS. Muscle mass index as a predictor of longevity in older adults. Am J Med. 2014;127:547-553. DOI: 10.1016/j. amjmed.2014.02.007.
47. Ortega FB, Sui X, Lavie CJ, Steven N. Body mass index, the most widely used but also widely criticized index: would a gold-standard measure of total body fat be a better predictor of cardiovascular disease mortality? Mayo Clin Proc. 2017;91:443-455. DOI: 10.1016/j. mayocp.2016.01.008.
48. Lavie CJ, Alpert MA, Arena R, Mehra MR, Milani RV, Ventura HO Impact of obesity and the obesity paradox on prevalence and prognosis in heart failure. JACC Heart Fail. 2013;1:93-102. DOI: 10.1016/j. jchf.2013.01.006.
49. Alpert MA, Omran J, Mehra A, Ardhanari S. Impact of obesity and weight loss on cardiac performance and morphology in adults. Prog Cardiovasc Dis. 2015;56:391-400. DOI: 10.1016/j.pcad.2013.09.003.
50. Malden D, Lacey B, Emberson J, Karpe F, Allen N, Bennett D, Lewington S. Body fat distribution and systolic blood pressure in 10,000 adults with whole-body imaging: UK Biobank and Oxford BioBank. Obesity. 2019;27:1200-1206. DOI: 10.1002/oby. 22509.
51. Lagacé JC, Brochu M, Dionne IJ. A counterintuitive perspective for the role of fat-free mass in metabolic health. J Cachexia Sarcopenia Muscle. 2020;11:343-347. DOI: 10.1002/jcsm. 12520.
52. Appelman Y, Rijn BBV, Monique E, Boersma E, Peters SAE. Sex differences in cardiovascular risk factors and disease prevention. Atherosclerosis. 2015;241:211-218. DOI: 10.1016/j.atheroscle rosis.2015.01.027.
53. Lee DH, Keum N, Hu FB, Orav EJ, Rimm EB, Willett WC, Giovannucci EL. Predicted lean body mass, fat mass, and all cause and cause specific mortality in men: prospective US cohort study. BMJ. 2018;362:k2575. DOI: 10.1136/bmj.k2575.
54. Gallagher D, Visser M, Sepulveda D, Pierson RN, Harris T, Heymsfield SB. How useful is body mass index for comparison of body fatness across age, sex, and ethnic groups? Am J Epidemiol. 1996;143:228239. DOI: 10.1093/oxfordjournals.aje.a008733.
55. Pischon T. Commentary: use of the body mass index to assess the risk of health outcomes: time to say goodbye? Int J Epidemiol. 2010;39:528-529. DOI: 10.1093/ije/dyp388.
56. Kuper H, Taylor A, Krishna KVR, Ben-Shlomo Y, Gupta R, Kulkarni B, Prabhakaran D, Davey Smith G, Wells J, Ebrahim S, et al. Is vulnerability to cardiometabolic disease in Indians mediated by abdominal adiposity or higher body adiposity. BMC Public Health. 2014;14:1239. DOI: 10.1186/1471-2458-14-1239.
57. Willett K, Jiang R, Lenart E, Spiegelman D, Willett W, Jiang RUI. Comparison of bioelectrical impedance and BMI in predicting obesityrelated medical conditions. Obesity. 2006;14:480-490. DOI: 10.1038/ oby.2006.63.
58. Wells JCK. Commentary: the paradox of body mass index in obesity assessment: not a good index of adiposity, but not a bad index of cardio-metabolic risk. Int J Epidemiol. 2014;43:672-674. DOI: 10.1093/ ije/dyu060.
59. Lee CMY, Huxley RR, Wildman RP, Woodward M. Indices of abdominal obesity are better discriminators of cardiovascular risk factors than BMI: a meta-analysis. J Clin Epidemiol. 2008;61:646-653. DOI: 10.1016/j.jclinepi.2007.08.012.
60. Taylor AE, Ebrahim S, Ben-Shlomo Y, Martin RM, Whincup PH, Yarnell JW, Wannamethee SG, Lawlor DA. Comparison of the associations of body mass index and measures of central adiposity and fat mass with coronary heart disease, diabetes, and all-cause mortality: a study using data from 4 UK cohorts. Am J Clin Nutr. 2010;91:547-556. DOI: 10.3945/ajcn.2009.28757.
61. Cruz-Jentoft AJ, Baeyens JP, Bauer JM, Boirie Y, Cederholm T, Landi F, Martin FC, Michel JP, Rolland Y, Schneider SM, et al. Sarcopenia: European consensus on definition and diagnosis. Age Ageing. 2010;39:412-423. DOI: 10.1093/ageing/afq034.
62. Kyle UG, Bosaeus I, De Lorenzo AD, Deurenberg P, Elia M, Manuel Gómez J, Lilienthal Heitmann B, Kent-Smith L, Melchior J-C, Pirlich M , et al. Bioelectrical impedance analysis-part II: utilization in clinical practice. Clin Nutr. 2004;23:1430-1453. DOI: 10.1016/j. clnu.2004.09.012.
63. Hemmingsson E, Udden J, Neovius M. No apparent progress in bioelectrical impedance accuracy: validation against metabolic risk and DXA. Obesity (Silver Spring). 2009;17:183-187. DOI: 10.1038/ oby.2008.474.
64. Fry A, Littlejohns TJ, Sudlow C, Doherty N, Adamska L, Sprosen T, Collins R, Allen NE. Comparison of sociodemographic and healthrelated characteristics of UK Biobank participants with those of the general population. Am J Epidemiol. 2017;186:1026-1034. DOI: 10.1093/aje/kwx246.
65. Gajalakshmi V, Lacey B, Kanimozhi V, Sherliker P, Peto R, Lewington S. Body-mass index, blood pressure, and cause-specific mortality in India: a prospective cohort study of 500810 adults. Lancet Glob Health. 2018;6:e787-e794. DOI: 10.1016/S2214-109X(18)30267-5.
66. Abdullah N, Abdul Murad NA, Attia J, Oldmeadow C, Kamaruddin MA, Abd Jalal N, Ismail N, Jamal R, Scott RJ, Holliday EG. Differing contributions of classical risk factors to type 2 diabetes in multi-ethnic Malaysian populations. Int J Environ Res Public Health. 2018;15:2813. DOI: 10.3390/jijerph15122813.
67. Allison DB, Zhu SK, Plankey M, Faith MS, Heo M. Differential associations of body mass index and adiposity with all-cause mortality among men in the first and second National Health and Nutrition Examination Surveys (NHANES I and NHANES II) follow-up studies. Int J Obes. 2002;26:410-416. DOI: 10.1038/sj.ijo.0801925.
68. Warren Andersen S, Shu X-O, Gao Y-T, Zhang X, Cai H, Yang G, Li H-L, Xiang Y-B, Zheng W. Prospective cohort study of central adiposity and risk of death in middle aged and elderly Chinese. PLoS One. 2015;10:e0138429. DOI: 10.1371/journal.pone.0138429.
69. Atkins JL, Whincup PH, Morris RW, Lennon LT, Papacosta O, Wannamethee SG. Sarcopenic obesity and risk of cardiovascular disease and mortality: a population-based cohort study of older men. $J$ Am Geriatr Soc. 2014;62:253-260. DOI: 10.1111/jgs.12652.
70. Bachettini NP, Bielemann RM, Barbosa-Silva TG, Baptista Menezes AM, Tomasi E, Gonzalez MC. Sarcopenia as a mortality predictor in community-dwelling older adults: a comparison of the diagnostic criteria of the European Working Group on Sarcopenia in Older People. Eur J Clin Nutr. 2020;74:573-580. DOI: 10.1038/s41430-019-0508-8.
71. Balogun S, Winzenberg T, Wills K, Scott D, Jones G, Aitken D, Callisaya ML. Prospective associations of low muscle mass and function with 10-year falls risk, incident fracture and mortality in communitydwelling older adults. J Nutr Health Aging. 2017;21:843-848. DOI: 10.1007/s12603-016-0843-6.
72. Batsis JA, Mackenzie TA, Barre LK, Lopez-Jimenez F, Bartels SJ. Sarcopenia, sarcopenic obesity and mortality in older adults: results from the National Health and Nutrition Examination Survey III. Eur J Clin Nutr. 2014;68:1001-1007. DOI: 10.1038/ejcn.2014.117.
73. Bea JW, Thomson CA, Wertheim BC, Nicholas JS, Ernst KC, Hu C, Jackson RD, Cauley JA, Lewis CE, Caan B, et al. Risk of mortality according to body mass index and body composition among postmenopausal women. Am J Epidemiol. 2015;182:585-596. DOI: 10.1093/aje/ kwv103.
74. Bigaard J, Frederiksen K, Tjønneland A, Thomsen BL, Overvad K, Heitmann BL, Sørensen TIA. Body fat and fat-free mass and all-cause mortality. Obes Res. 2004;12:1042-1049. DOI: 10.1038/oby.2004.131.
75. Bigaard J, Frederiksen K, Tjønneland A, Thomsen BL, Overvad K, Heitmann BL, Sørensen TIA. Waist circumference and body composition in relation to all-cause mortality in middle-aged men and women. Int J Obes. 2005;29:778-784. DOI: 10.1038/sj.ijo.0802976.
76. Boloukat RR, Ramezankhani A, Hasheminia M, Tasdighi E, Azizi F, Hadaegh F. Impact of blood pressure, cholesterol and glucose in the association between adiposity measures and coronary heart disease and stroke among Iranian population. Clin Nutr. 2018;37:2060-2067. DOI: 10.1016/j.clnu.2017.09.023.
77. Brown JC, Harhay MO, Harhay MN. Appendicular lean mass and mortality among prefrail and frail older adults. J Nutr Health Aging. 2017;21:5-8. DOI: 10.1007/s12603-016-0753-7.
78. Chen Z, Iona A, Parish S, Chen Y, Guo Y, Bragg F, Yang L, Bian Z, Holmes MV, Lewington S, et al. Adiposity and risk of ischaemic and haemorrhagic stroke in 0.5 million Chinese men and women: a prospective cohort study. Lancet. 2018;6:e630-e640. DOI: 10.1016/ S2214-109X(18)30216-X.
79. Chen G-C, Arthur R, lyengar NM, Kamensky V, Xue X, WassertheilSmoller S, Allison MA, Shadyab AH, Wild RA, Sun Y, et al. Association between regional body fat and cardiovascular disease risk among postmenopausal women with normal body mass index. Eur Heart J. 2019;40:2849-2855. DOI: 10.1093/eurheartj/ehz391.
80. Cl C, Lam KSL, Cheung BMY. Evaluation of cutpoints for low lean mass and slow gait speed in predicting death in the National Health
and Nutrition Examination Survey 1999-2004. J Gerontol A Biol Sci Med Sci. 2016;71:90-95. DOI: 10.1093/gerona/glv112.
81. Chin SO, Rhee SY, Chon S, Hwang Y-C, Jeong I-K, Oh S, Ahn KJ, Chung HY, Woo J-T, Kim S-W, et al. Sarcopenia is independently associated with cardiovascular disease in older Korean adults: the Korea National Health and Nutrition Examination Survey (KNHANES) from 2009. PLoS One. 2013;8:e60119. DOI: 10.1371/journal.pone.0060119.
82. Chuang SY, Chang HY, Lee MS, Chia-Yu Chen R, Pan WH. Skeletal muscle mass and risk of death in an elderly population. Nutr Metab Cardiovasc Dis. 2014;24:784-791. DOI: 10.1016/j.numecd.2013.11.010.
83. Chuang SY, Hsu YY, Chen RCY, Liu WL, Pan WH. Abdominal obesity and low skeletal muscle mass jointly predict total mortality and cardiovascular mortality in an elderly Asian population. J Gerontol A Biol Sci Med Sci. 2016;71:1049-1055. DOI: 10.1093/gerona/glv192.
84. de Almeida Roediger M, de Fátima Nunes Marucci M, Quintiliano Scarpelli Dourado DA, de Oliveira C, Licio Ferreira Santos J, de Oliveira Duarte YA. Body composition changes and 10-year mortality risk in older Brazilian adults: analysis of prospective data from the SABE study. J Nutr Health Aging. 2019;23:51-59. DOI: 10.1007/s1260 3-018-1118-1.
85. de Santana FM, Domiciano DS, Gonçalves MA, Machado LG, Figueiredo CP, Lopes JB, Caparbo VF, Takayama L, Menezes PR, Pereira RM. Association of appendicular lean mass, and subcutaneous and visceral adipose tissue with mortality in older Brazilians: the São Paulo Ageing \& Health Study. J Bone Miner Res. 2019;34:12641274. DOI: 10.1002/jbmr. 3710.
86. Dolan CM, Kraemer H, Browner W, Ensrud K, Kelsey JL. Associations between body composition, anthropometry, and mortality in women aged 65 years and older. Am J Public Health. 2007;97:913-918. DOI: 10.2105/AJPH.2005.084178.
87. Dong B, Peng Y, Wang Z, Adegbija O, Hu J, Ma J, Ma H. Joint association between body fat and its distribution with all-cause mortality: a data linkage cohort study based on NHANES (1988-2011). PLoS One. 2018;13:e0193368. DOI: 10.1371/journal.pone.0193368.
88. Gale CR, Martyn CN, Cooper C, Sayer AA. Grip strength, body composition, and mortality. Int J Epidemiol. 2007;36:228-235. DOI: 10.1093/ije/dyl224.
89. Gillum RF, Mussolino ME, Madans JH. Body fat distribution, obesity, overweight and stroke incidence in women and men: the NHANES I Epidemiologic Follow-up Study. Int J Obes. 2001;25:628-638. DOI: 10.1038/sj.ijo. 0801590.
90. Gnatiuc L, Alegre-Díaz J, Wade R, Ramirez-Reyes R, Tapia-Conyer R, Garcilazo-Ávila A, Chiquete E, Gonzáles-Carballo C, Solano-Sanchez M, Clarke R, et al. General and abdominal adiposity and mortality in Mexico City: prospective study of 150000 adults. Ann Intern Med. 2019;171:397-405. DOI: 10.7326/M18-3502.
91. Graf CE, Karsegard VL, Spoerri A, Makhlouf AM, Ho S, Herrmann FR, Genton L. Body composition and all-cause mortality in subjects older than 65 y. Am J Clin Nutr. 2015;101:760-767. DOI: 10.3945/ ajcn.114.102566.
92. Han SS, Kim KW, Kim K-I, Na KY, Chae D-W, Kim S, Chin HJ. Lean mass index: a better predictor of mortality than body mass index in elderly Asians. J Am Geriatr Soc. 2010;58:312-317. DOI: 10.1111/j.1532-5415.2009.02672.x.
93. Heitmann BL, Erikson H, Bm E, Mikkelsen KL, Larsson B. Mortality associated with body fat, fat-free mass and body mass index among 60 -year-old Swedish men-a 22 -year follow-up. The study of men born in 1913. Int J Obes. 2000;24:33-37. DOI: 10.1038/sj.ijo.0801082.
94. Hirani V, Naganathan V, Blyth F, Le Couteur D, Seibel MJ, Waite LM, Handelsman DJ, Cumming RG. Longitudinal associations between body composition, sarcopenic obesity and outcomes of frailty, disability, institutionalisation and mortality in community-dwelling older men: the Concord Health and Ageing in Men Project. Age Ageing. 2017;46:413-420. DOI: 10.1093/ageing/afw214.
95. Hotchkiss JW, Davies CA, Leyland AH. Adiposity has differing associations with incident coronary heart disease and mortality in the Scottish population: cross-sectional surveys with follow-up. Int J Obes. 2013;37:732-739. DOI: 10.1038/ijo.2012.102.
96. Howell CR, Mehta T, Ejima K, Ness KK, Cherrington A, Fontaine KR. Body composition and mortality in Mexican-American adults: results from the National Health and Nutrition Examination Survey. Obesity. 2018;26:1372-1380. DOI: 10.1002/oby. 22251.
97. Kahn HS, Bullard KM, Barker LE, Imperatore G. Differences between adiposity indicators for predicting all-cause mortality in a
representative sample of United States non-elderly adults. PLoS One. 2012;7:e50428. DOI: 10.1371/journal.pone. 0050428.
98. Katzmarzyk PT, Craig CL, Bouchard C. Adiposity, adipose tissue distribution and mortality rates in the Canada Fitness Survey follow-up study. Int J Obes. 2002;26:1054-1059. DOI: 10.1038/sj.ijo.0802057.
99. Kim Y, Wijndaele K, Lee D-C, Sharp SJ, Wareham N, Brage S. Independent and joint associations of grip strength and adiposity with all-cause and cardiovascular disease mortality in 403,199 adults: the UK Biobank study. Am J Clin Nutr. 2017;106:773-782. DOI: 10.3945/ ajcn.117.156851.
100. Kizer JR, Biggs ML, Ix JH, Mukamal KJ, Zieman SJ, de Boer IH, Mozaffarian D, Barzilay JI, Strotmeyer ES, Luchsinger JA, et al. Original contribution measures of adiposity and future risk of ischemic stroke and coronary heart disease in older men and women. Am J Epidemiol. 2011;173:10-25. DOI: 10.1093/aje/kwq311.
101. Kouvari M, Panagiotakos DB, Chrysohoou C, Notara V, Georgousopoulou EN, Yannakoulia M, Tousoulis D, Pitsavos C, Pitsavos C; Investigators AaGs. A sex-specific evaluation of predicted lean and fat mass composition and cardiovascular disease onset and progression: a combined analysis of the ATTICA and GREECS prospective epidemiological studies. Obes Res Clin Pract. 2019;13:469477. DOI: 10.1016/j.orcp.2019.09.005.
102. Lee JSW, Auyeung TW, Kwok T, Li M, Leung J, Woo J. Survival benefit of abdominal adiposity: a 6-year follow-up study with dual X-ray absorptiometry in 3,978 older adults. Age. 2012;34:597-608. DOI: 10.1007/s11357-011-9272-y.
103. Levitan EB, Yang AZ, Wolk A, Mittleman MA. Adiposity and incidence of heart failure hospitalization and mortality a population-based prospective study. Circ Heart Fail. 2009;2:202-208. DOI: 10.1161/CIRCH EARTFAILURE.108.794099.
104. Li R, Xia J, Zhang X, Gathirua-Mwangi WG, Guo J, Li Y, McKenzie S; Song Y. Associations of muscle mass and strength with all-cause mortality among US older adults. Med Sci Sports Exerc. 2018;50:458467. DOI: 10.1249/MSS.0000000000001448.
105. Myint PK, Kwok CS, Luben RN, Wareham NJ, Khaw KT. Body fat percentage, body mass index and waist-to-hip ratio as predictors of mortality and cardiovascular disease. Heart. 2014;100:1613-1619. DOI: 10.1136/heartjnl-2014-305816.
106. Nalini M, Sharafkhah M, Poustchi H, Sepanlou SG, Pourshams A, Reza A. Comparing anthropometric indicators of visceral and general adiposity as determinants of overall and cardiovascular mortality. Arch Iran Med. 2019;22:301-309.
107. Newman AB, Kupelian V, Visser M, Simonsick EM, Goodpaster BH, Kritchevsky SB, Tylavsky FA, Rubin SM, Harris TB. Strength, but not muscle mass is associated with mortality in the Health, Aging and Body Composition Study cohort. J Gerontol A Biol Sci Med Sci. 2006;61A:72-77. DOI: 10.1093/gerona/61.1.72.
108. Ofstad AP, Sommer C, Birkeland KI, Bjørgaas MR, Gran JM, Gulseth HL, Johansen OE. Comparison of the associations between nontraditional and traditional indices of adiposity and cardiovascular mortality: an observational study of one million person-years of follow-up. Int J Obes. 2019;43:1082-1092. DOI: 10.1038/s4136 6-019-0353-9.
109. Otsuka R, Matsui Y, Tange C, Nishita Y, Tomida M, Ando F, Shimokata H, Arai H. What is the best adjustment of appendicular lean mass for predicting mortality or disability among Japanese community dwellers? BMC Geriatr. 2018;18:8. DOI: 10.1186/s12877-017-0699-6.
110. Padwal R, Leslie WD, Lix LM, Majumdar SR. Relationship among body fat percentage, body mass index, and all-cause mortality: a cohort study. Ann Intern Med. 2016;164:532-541. DOI: 10.7326/M15-1181.
111. Park S, Ham JO, Lee BK. A positive association between stroke risk and sarcopenia in men aged 50 years, but not women: results from the Korean National Health and Nutrition Examination Survey 20082010. J Nutr Health Aging. 2014;18:806-812. DOI: 10.1007/s1260 3-014-0553-x.
112. Park Y, Kim NH, Kwon TY, Kim SG. A novel adiposity index as an integrated predictor of cardiometabolic disease morbidity and mortality. Sci Rep. 2018;8:16753. DOI: 10.1038/s41598-018-35073-4.
113. Reis JP, MacEra CA, Araneta MR, Lindsay SP, Marshall SJ, Wingard DL. Comparison of overall obesity and body fat distribution in predicting risk of mortality. Obesity. 2009;17:1232-1239. DOI: 10.1038/ oby.2008.664.
114. Reis JP, Araneta MR, Wingard DL, Macera CA, Lindsay SP, Marshall SJ. Overall obesity and abdominal adiposity as predictors of mortality
in US white and black adults. Ann Epidemiol. 2009;19:134-142. DOI: 10.1016/j. annepidem.2008.10.008.
115. Rexrode KM, Buring JE, Manson JE. Abdominal and total adiposity and risk of coronary heart disease in men. Int J Obes. 2001;25:10471056. DOI: 10.1038/sj.ijo. 0801615.
116. Sim M, Prince RL, Scott D, Daly RM, Duque G, Inderjeeth CA, Zhu K, Woodman RJ, Hodgson JM, Lewis JR. Sarcopenia definitions and their associations with mortality in older Australian women. J Am Med Dir Assoc. 2019;20:76-82. DOI: 10.1016/j.jamda.2018.10.016.
117. Simpson JA, MacInnis RJ, Peeters A, Hopper JL, Giles GG, English DR, Julie A. A comparison of adiposity measures as predictors of allcause mortality: the Melbourne Collaborative Cohort Study. Obesity. 2007;15:994-1003. DOI: 10.1038/oby.2007.622.
118. Srikanthan P, Horwich TB, Tseng CH. Relation of muscle mass and fat mass to cardiovascular disease mortality. Am J Cardiol. 2016;117:1355-1360. DOI: 10.1016/j.amjcard.2016.01.033.
119. Stefan N, Kantartzis K, Häring H. Cardiorespiratory fitness, adiposity, and mortality. JAMA. 2008;299:1013-1014. DOI: 10.1001/ jama.299.9.1013-b.
120. Sui X, LaMonte MJ, Laditka JN, Hardin JW, Chase N, Jooker SP, Blair SN. Cardiorespiratory fitness and adiposity as mortality predictors in older adults. JAMA. 2007;298:2507-2516. DOI: 10.1001/ jama.298.21.2507.
121. Tanne D, Medalie JH, Goldbourt U. Body fat distribution and long-term risk of stroke mortality. Stroke. 2005;36:1021-1025. DOI: 10.1161/01. STR.0000162584.39366.1c.
122. The Decode Study Group. Does the constellation of risk factors with and without abdominal adiposity associate with different cardiovascular mortality risk? Int J Obes. 2008;32:757-762. DOI: 10.1038/sj.ijo.0803797.
123. Thomson CA, Garcia DO, Wertheim BC, Hingle MD, Bea JW, Zaslavsky O, Caire-Juvera G, Rohan T, Vitolins MZ, Thompson PA, et al. Body shape, adiposity index, and mortality in postmenopausal women: findings from the Women's Health Initiative Cynthia. Obesity. 2016;24:1061-1069. DOI: 10.1002/oby. 21461.
124. Toss F, Wiklund P, Nordstrom P, Nordstrom A. Body composition and mortality risk in later life. Age Ageing. 2012;41:677-681. DOI: 10.1093/ ageing/afs087.
125. Van Aller C, Lara J, Stephan BCM, Maria L, Heyms S, Katzmarzyk PT, Wells JCK, Prado CM, Siervo M. Sarcopenic obesity and overall
mortality: results from the application of novel models of body composition phenotypes to the National Health and Nutrition Examination Survey 1999-2004. Clin Nutr. 2019;38:264-270. DOI: 10.1016/j. clnu.2018.01.022.
126. Wang A, Wu J, Zhou Y, Guo X, Luo Y, Wu S, Zhao X. Measures of adiposity and risk of stroke in China: a result from the Kailuan study. PLoS One. 2013;8:e61665. DOI: 10.1371/journal.pone.0061665.
127. Wannamethee SG, Shaper AG, Lennon L, Whincup PH. Decreased muscle mass and increased central adiposity are independently related to mortality in older men. Am J Clin Nutr. 2007;86:1339-1346. DOI: 10.1093/ajcn/86.5.1339.
128. Wannamethee SG, Shaper AG, Whincup PH, Lennon L, Papacosta O, Sattar N. The obesity paradox in men with coronary heart disease and heart failure: the role of muscle mass and leptin. Int J Cardiol. 2014;171:49-55. DOI: 10.1016/j.ijcard.2013.11.043.
129. Yang M, Jiang J, Zeng Y, Tang H. Sarcopenia for predicting mortality among elderly nursing home residents. Medicine. 2019;98:e14546. DOI: 10.1097/MD. 0000000000014546.
130. Yuki A, Ando F, Otsuka R, Shimokata H. Sarcopenia based on the Asian Working Group for Sarcopenia criteria and all-cause mortality risk in older Japanese adults. Geriatr Gerontol Int. 2017;17:1642-1647. DOI: 10.1111/ggi.12946.
131. Zaslavsky O, Rillamas-Sun E, Li W, Going S, Datta M, Snetselaar L, Zelber-Sagi S. Association of dynamics in lean and fat mass measures with mortality in frail older women. J Nutr Health Aging. 2017;21:112119. DOI: 10.1007/s12603-016-0730-1.
132. Zhang X, Shu XO, Yang G, Li H, Cai H, Gao YT, Zheng W. Abdominal adiposity and mortality in Chinese women. Arch Intern Med. 2007;167:886-892. DOI: 10.1001/archinte.167.9.886.
133. Zhu S, Heo PM, Faith MS, Allison DB. Associations of body mass index and anthropometric indicators of fat mass and fat free mass with all-cause mortality among women in the first and second National Health and Nutrition Examination Surveys follow-up studies. Ann Epidemiol. 2003;13:286-293. DOI: 10.1016/S1047 -2797(02)00417-9.
134. Zong G, Zhang Z, Yang Q, Wu H, Hu FB, Sun Q. Total and regional adiposity measured by dual-energy X-ray absorptiometry and mortality in NHANES 1999-2006. Obesity. 2017;24:2414-2421. DOI: 10.1002/ oby. 21659.

## SUPPLEMENTAL MATERIAL

## Data S1.

## Systematic literature review

Search terms used on PubMed and Embase (up to $18^{\text {th }}$ October 2019):
("muscle mass" or "skeletal muscle" or "sarcopenia" or "fat mass" or "body fat" or "body composition" or "fat free mass" or "lean mass" or "adiposity") AND ("mortality" or "death" or "survival" or "cardiovascular disease" or "CVD" or "stroke" or "coronary heart disease" or "congestive heart failure")
Studies were excluded if they were done in patients or ill people (including but not limited to hospitalised, intensive care or dialysis patients, those with cancer, diabetes, COPD, CKD, dementia, HIV), done in children or adolescents, investigating a diet or exercise intervention, done in animals, molecular, cellular-level studies, or genetic studies. Only findings from prospective cohort studies published after 2000 with more than 600 participants are presented below.

Table S1. Summary of search findings.

\left.| Exposure | Outcome |  | Sex | Number of associations identified in studies |
| :--- | :--- | :--- | :---: | :---: | :---: | :---: |
| U or J- |  |  |  |  |$\right)$

Each association assessed in the studies is counted distinctly (i.e. the number of associations exceeds the number of studies identified since some studies analysed more than one exposure/outcome combination). 'Muscle mass' includes studies that assessed measurements of lean mass from techniques such as bioimpedance analysis (BIA), dual energy x-ray absorptiometry (DEXA), mid upper arm circumference and calf circumference (it does not include studies of muscle quality such as grip strength). 'Fat mass' includes studies that assessed measurement of adiposity such as fat mass as determine by BIA or DEXA, percentage body fat, waist circumference, waist to hip ratio. Studies of BMI or body fat distribution indices are not included.

Table S2. Systematic literature review.

|  |  | Study name (date of recruitment) | Study participants (location, N, age) | Body composition measurement | Exposure definition/coding | Outcome | Shape of association |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 68 | Abramowitz, et al. (2018) | NHANES (19992004) | $\begin{aligned} & \text { USA, n=11687 aged } \\ & \text { >20 yrs } \end{aligned}$ | DEXA (Appendicular skeletal muscle mass) | Low muscle defined as ASMI $<5.45 \mathrm{~kg} / \mathrm{m}^{2}$ (women) or $<7.26 \mathrm{~kg} / \mathrm{m}^{2}$ (men) $[2,3]$ | All-cause mortality | BMI and mortality: U (low muscle group), 0 (preserved muscle group) ASMI and mortality (men and women combined): - |
| 69 | Allison, et al. (2002) | NHANES I \& II | USA, n=10169 men aged 25-75yrs | Skinfolds (indicator of fat mass), upper arm circumference (fat free mass indicator), BMI | Modelled continuously | All-cause mortality | BMI and all-cause mortality: U (men) <br> Fat free mass and mortality: linear - (men) <br> Fat mass and mortality: linear + (men) |
| 70 | Andersen, et al. (2015) | Shanghai Men's Health Study and Shanghai Women's Health Study (1996-2000) | China, $n=116442$ aged 40-74yrs | Waist to hip ratio | Quintiles of WHR | All-cause and CVD mortality | WHR and mortality: linear + (men) <br> WHR and mortality: linear + (women) <br> WHR and CVD mortality: linear + (men) <br> WHR and CVD mortality: linear + (women) |
| 71 | Atkins, et al. (2014) | $\begin{aligned} & \text { NHANES (1999- } \\ & \text { 2004) } \end{aligned}$ | $\begin{aligned} & \text { USA, n=11687 aged } \\ & >20 y r s \end{aligned}$ | Waist circumference, midupper arm circumference, triceps skinfold thickness | 4 groups created: <br> Optimal (MAMC $>25.9 \mathrm{~cm}$ and $\mathrm{WC} \leq 102 \mathrm{~cm}$ ), sarcopenic (MAMC $\leq 25.9 \mathrm{~cm}$ and WC $\leq 102 \mathrm{~cm}$ ), sarcopenic obese (MAMC $\leq 25.9 \mathrm{~cm}$ and $W C>102 \mathrm{~cm}$ ), obese ( $W C>102 \mathrm{~cm}$ and MAMC $>25.9 \mathrm{~cm}$ ) | All-cause mortality, CVD mortality, CVD events | Sarcopenia and mortality: 0 (men) <br> Sarcopenia and mortality: + (women) <br> Sarcopenia and CVD: 0 (men) <br> Sarcopenia and CVD: + (women) |
| 72 | Bachettini et al. (2019) | n/a | Brazil, n=1291 aged $>=60$ years | Using the European Working Group on Sarcopenia in Older People (EWGSOP) definitions of sarcopenia | Sarcopenia, binary | All-cause mortality | Severe sarcopenia and mortality: + (men and women combined) Other sarcopenia and mortality: 0 (men and women combined) |
| 73 | Balogun, et al. (2017) | Tasmanian Older Adult Cohort study | Tasmania, n=1099 aged $>50 \mathrm{yrs}$ | DEXA (Appendicular lean mass), lower limb strength | Calculated ALM/height² [2], ALM/BMI [1], ALM/weight*100 [12] from DEXA. Measured grip, lower-limb muscle, and upper-limb muscle strength and/or quality. Low muscle mass or function defined as participants in the lowest $20 \%$ of the sexspecific distribution for each measurement | Mortality, falls, fracture | Muscle strength or quality and mortality (men and women combined): 0 Low appendicular lean mass/BMI and mortality (men and women combined): + <br> Low appendicular lean mass/height ${ }^{2}$ and mortality (men and women combined): 0 <br> Low appendicular lean mass/weight and mortality (men and women combined): 0 |
| 74 | Batsis, et al. (2014) | NHANES III (1988- <br> 1994) | USA, $n=4652$ aged >60years | BIA (skeletal muscle index, \% body fat) | 4 definitions used: <br> Normal if SMI $\geqslant 10.76 \mathrm{~kg} / \mathrm{m}^{2}$ (men) <br> $>6.76 \mathrm{~kg} / \mathrm{m}^{2}$ (women); <br> Class I sarcopenia if SMI 8.51-10.75 (men) <br> 5.76-6.75 (women); <br> Class II sarcopenia: $\leqslant 8.50 \mathrm{~kg} / \mathrm{m} 2 \leqslant 5.75$ [17]. <br> Obesity defined as body fat >27\% (men) <br> >38\% (women) [3] | All-cause mortality | Sarcopenia only and mortality: 0 (men) <br> Sarcopenia only and mortality: + (women) <br> Sarcopenic obese and mortality: 0 (men) <br> Sarcopenic obese and mortality: 0 (women) <br> Obese only and mortality: 0 (men) <br> Obese only and mortality: 0 (women) |
| 45 | Batsis, et al. (2017) | NHAHES (1999- 2004) | USA, n=4984 aged $>60 \mathrm{yrs}$ | DEXA (Appendicular lean mass), waist circumference, BMI | Low lean mass defined as: ALM $<19.75 \mathrm{~kg}$ (men) <15.02kg (women) [1] or ALM:BMI $<0.789$ (men), <0.512 (women). <br> Obesity defined as body fat >25\% (men) >35\% (women) [9]. | All-cause and CVD mortality | Low lean mass and mortality: + (men) <br> Low lean mass and mortality: 0 (women) <br> Low lean mass and CVD: + (men) <br> Low lean mass and CVD: 0 (women) <br> Low lean mass with obesity and mortality: + (men) <br> Low lean mass with obesity and mortality: 0 (women) <br> Low lean mass with obesity and CVD : 0 (men) <br> Low lean mass with obesity and CVD : 0 (women) <br> Obesity (body fat \%) and mortality: - (men) |


|  |  |  |  |  |  |  | Obesity (body fat \%) and mortality: - (women) Obesity (body fat \%) and CVD: 0 (men) Obesity (body fat \%) and CVD: 0 (women) |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 75 | Bea, et al. <br> (2015) | Women's Health Initiative (19931998) | USA, $n=10525$ <br> postmenopausal <br> women aged 50- <br> 79yrs | DEXA (total body fat \%, lean body mass \%), BMI | Quintiles of each exposure | All-cause mortality | Fat mass and mortality (women aged 50-59): linear + Fat mass and mortality (women aged $>70$ ): linear Lean mass and mortality (women aged 50-59): linear Lean mass and mortality (women aged >70): linear + |
| 76 | Bigaard, et al (2004) | Diet, Cancer and Health study (1993-1997) | Denmark, n=51053 aged 50-64yrs | BIA (fat free mass index, body fat mass index) | Modelled continuously using linear splines, and relative risks estimated per 10\% increase in waist circumference and per $\mathrm{kg} / \mathrm{m}^{2}$ increase in body fat mass index and fat free mass index Results stratified into high and low range corresponding to cut-offs at the $40^{\text {th }}$ and $60^{\text {th }}$ percentiles | All-cause mortality | Fat mass and mortality: $J$ (men) <br> Fat mass and mortality: J (women) <br> Fat free mass and mortality: reverse J (men) <br> Fat free mass and mortality: reverse J (women) <br> WC and mortality: linear + (men) <br> WC and mortality: linear + (women) |
| 77 | Bigaard, et al. (2005) | Diet, Cancer and Health (19931997) | Denmark, n=57058 adults | BIA (body fat mass index, fat free mass index), waist circumference | Modelled continuously using linear splines | All-cause mortality | Body fat mass index and all-cause mortality: $J$ (men) Body fat mass index and all-cause mortality: J (women) Fat free mass index and mortality: reverse J (men) Fat free mass index and mortality: reverse J (women) |
| 78 | Boloukat, et al. (2018) | Tehran Lipid and Glucose Study | Iran, n=4631 aged >40yrs | BMI, waist circumference, waist-to-height ratio, WHR | Modelled continuously per 1 unit increase in each exposure | CHD or stroke incidence | WC and CVD: 0 (men) <br> WC and CVD: 0 (women) <br> WHR and CVD: 0 (men) <br> WHR and CVD: 0 (women) <br> Waist-to-height ratio and CVD: 0 (men) <br> Waist-to-height ratio and CVD: 0 (women) |
| 43 | Brown, et al. (2016) | $\begin{aligned} & \text { NHANES (1988- } \\ & \text { 1994) } \end{aligned}$ | USA, $\mathrm{n}=4425$ aged $>60 \mathrm{yrs}$ | BIA (skeletal muscle index), gait speed | Sarcopenia defined as low gait speed ( $<0.8 \mathrm{~m} / \mathrm{s}$ ) and SMI <10.76 (men), $<6.75$ (women) | All-cause and CVD mortality | ```Sarcopenia and mortality: + (men) Sarcopenia and mortality: + (women) Sarcopenia and CVD: 0 (men) Sarcopenia and CVD: + (women)``` |
| 79 | Brown, et al. (2017) | NHANES III (1988- <br> 1994) | $\begin{aligned} & \text { USA, n=1487 aged } \\ & >65 \mathrm{yrs} \end{aligned}$ | BIA (Appendicular lean mass) | Modelled continuously per 1 SD of ALM | All-cause mortality | Appendicular lean mass and mortality: - linear (men) Appendicular lean mass and mortality: - linear (women) |
| 44 | Cesari, et al (2009) | InChianti study <br> (1998-2000) | Italy, $\mathrm{n}=934$ aged $>65 y r s$ | Peripheral quantitative computerised tomography (muscle density, muscle area, fat area) | Modelled continuously per 1 SD increase of each exposure | All-cause mortality | Muscle area and mortality (men and women combined): 0 Fat area and mortality (men and women combined): 0 |
| 80 | Chen, et al. (2018) | China Kadoorie Biobank (20042008) | China, $\mathrm{n}=489301$ aged 30-79yrs | BMI, body fat \%, waist circumference | 7 groups for each exposure, and BMI per $5 \mathrm{~kg} / \mathrm{m}^{2}, \%$ body fat per $10 \%$, waist circumference per 1 SD | Ischaemic and haemorrhagic stroke incidence | BMI and stroke/haemorrhage (men and women combined): + linear WC and stroke/haemorrhage (men and women combined): + linear Body fat \% and stroke/haemorrhage (men and women combined): + linear |
| 81 | Chen et al. (2019) | Women's Health Initiative | USA, $n=2,683$ <br> postmenopausal women with normal BMI | DEXA - \% whole body and regional fat quartiles, fat mass index (dividing total or regional fat mass in kg by height squared) | Quartiles | CVD | Whole body fat mass and mortality: 0 (women) <br> Leg fat and mortality: - (women) <br> trunk fat and mortality: + (women) |
| 82 | Cheung, et al. (2016) | $\begin{aligned} & \text { NHANES (1999- } \\ & \text { 2004) } \end{aligned}$ | USA, n=2841 aged >65yrs | DEXA (Appendicular lean mass) | Calculated ALM, ALM/BMI, ALM/ $\mathrm{H}^{2}$ from DEXA. Sarcopenia defined using cut-points for each measure (given in their supplementary index). | All-cause mortality | Low appendicular lean mass and mortality: + (men and women combined) |
| 83 | Chin, et al (2013) | $\begin{aligned} & \text { KNHANES (2008- } \\ & \text { 2012) } \end{aligned}$ | South Korea, $\mathrm{n}=1578$ aged $>65 \mathrm{yrs}$ | DEXA (Appendicular skeletal muscle mass) | Sarcopenia defined as ASM/weight <1 SD below the gender-specific mean of a reference group aged 20-39 years. Equates to $<32.2 \%$ men, $<25.6 \%$ women. | CVD incidence | Sarcopenic and CVD (men and women combined): + |


| 84 | Chuang, et al. (2014) | Elderly Nutrition and Health Survey (1999-2000) | Taiwan, n=1512 aged >65yrs | BIA (skeletal muscle mass index), BMI, waist circumference | Low/high risk groups defined as ASMI $<11.45 \mathrm{~kg} / \mathrm{m}^{2}$ (men) $8.51 \mathrm{~kg} / \mathrm{m}^{2}$ (women) based off quartiles | All-cause mortality | Low skeletal muscle index and mortality: 0 (men) Low skeletal muscle index and mortality: + (women) Low skeletal muscle index and CVD: 0 (men) Low skeletal muscle index and CVD: + (women) |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 85 | Chuang, et al. (2016) | Elderly Nutrition and Health Survey | Taiwan, n=1485 aged >65 yrs | BIA (skeletal muscle mass index) | Sarcopenia defined as SMMI men <11.45, women <8.51 [14] <br> Sarcopenic obesity defined as being sarcopenic and having a high triglyceride ( $>150 \mathrm{mg} / \mathrm{dL}$ ) and high waist circumference ( $>90 \mathrm{~cm}$ men, $>80 \mathrm{~cm}$ women) | All-cause and CVD mortality | Abdominal obesity and mortality: 0 (men and women combined) <br> Abdominal obesity and CVD: 0 (men and women combined) <br> Sarcopenic and mortality: 0 (men and women combined) <br> Sarcopenic and CVD: 0 (men and women combined) <br> Sarcopenic obesity and mortality: + (men and women combined) <br> Sarcopenic obesity and CVD: + (men and women combined) |
| 86 | de Almeida Roediger, et al. (2019) | SABE (2000-2010) | Brazil, n=1504 aged >60yrs | BMI, waist circumference, wait-to-hip ratio skinfold, mid-upper arm circumference, calf circumference, arm muscle area | All exposures analysed as binary variables as either high vs low or adequate vs inadequate. | All-cause mortality | WC and mortality: 0 (men and women combined) WHR and mortality: 0 (men and women combined) MAMC (muscle) and mortality: - (men and women combined) Calf circumference (muscle) and mortality: - (men and women combined) |
| 87 | de Santana et <br> al. (2019) | n/a | Brazil, $n=839$ aged $>=65 y r s$ | DEXA (total fat, appendicular lean mass) | Low lean mass, presence of visceral adipose issue | CVD mortality, all cause morality | Low muscle and CVD mortality: + (men and women combined) <br> Low muscle and mortality: + (men and women combined) <br> Fat mass and CVD mortality: + (men and women combined) <br> Fat mass and mortality: + (men and women combined) |
| 88 | Dolan, et al. (2007) | Study of Osteoporotic Fractures | USA, $n=8029$ women aged >65 | BIA (lean mass, fat mass, \% body fat), BMI, waist girth | Quintiles of each exposure | All-cause mortality | Fat mass and mortality (women): U Lean mass and mortality (women): 0 BMI and mortality (women): U |
| 89 | Dong et al (2018) | NHANES III (19881994) | USA, $\mathrm{n}=16415$ ) aged 18-89yrs | BIA (body fat \%), WHR | Body fat \% and WHR modelled continuously using cubic splines at 4 knots. Also created 9 groups to assess joint associations | All-cause mortality | Body fat \% and mortality: $U$ (men) <br> Body fat \% and mortality : U (women) <br> WHR and mortality: $U$ (men) <br> WHR and mortality: + linear (women) |
| 90 | Gale, et al. (2007) | n/a | UK, $\mathrm{n}=800$ aged >65yrs | Skinfolds (fat mass, fat free mass), BMI, grip strength | Modelled continuously per 1 SD increase in each exposure | All-cause and CVD mortality | Body fat $\%$ and CVD: 0 (men) <br> Body fat \% and CVD: 0 (women) <br> Body fat \% and mortality: 0 (men) <br> Body fat \% and mortality: 0 (women) <br> MAMC (muscle) and CVD: 0 (men) <br> MAMC (muscle) and CVD: 0 (women) <br> Fat free mass and CVD: 0 (men) <br> Fat free mass and CVD: 0 (women) <br> MAMC (muscle) and mortality: 0 (men) <br> MAMC (muscle) and mortality: 0 (women) <br> Fat free mass and mortality: 0 (men) <br> Fat free mass and mortality: 0 (women) <br> Grip strength and CVD: + (men) <br> Grip strength and CVD: 0 (women) <br> Grip strength and mortality: + (men) <br> Grip strength and mortality: 0 (women) |
| 91 | Gillum, et al. (2001) | NHANES I (1992) | USA, n=6936 aged 45-74yrs | BMI, Subscapular skinfold (SSF), subscapular to triceps skinfold thickness ratio (SFR) | Each exposure modelled in quartiles and results stratified by race, sex and smoking status | Stroke incidence | Trunk obesity and stroke: weak U (white men ex-smokers) Trunk obesity and stroke: 0 (women) Overall obesity and stroke: $U$ (white men ex-smokers) Overall obesity and stroke: 0 (women) |
| 92 | Gnatiuc et al. (2019) | Mexico City cohort study | Mexico, $\mathrm{n}=159,755$ aged 35 to $<75$ | BMI, waist-to-hip ratio, waist circumference | Continuously | all-cause mortality | BMI and mortality: $J$ (men and women cobined) WHR and mortality: + (men and women combined) |


|  |  |  |  |  |  |  | WC and mortality: + (men and women combined) |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 93 | Graf, et al. (2015) | n/a | Switzerland, n=3181 <br> aged $>65 \mathrm{yrs}$ | BIA (fat mass index, fat free mass index) | Quartiles of FMI and FFMI | All-cause mortality | Fat free mass index and mortality: 0 (women) <br> Fat free mass index and mortality: - (men) <br> Fat mass index and mortality: 0 (men) <br> Fat mass index and mortality: 0 (women) |
| 94 | Han, et al. <br> (2010) | Korean <br> Longitudinal <br> Study on Health and Aging | South Korea, n=877 aged $>65 \mathrm{yrs}$ | BIA (lean mass, fat mass, fat percentage, lean mass index), BMI, waist circumference | 3 groups for each exposure: $<25^{\text {th }}$ percentile, $25-75^{\text {th }}$ percentile, $>75^{\text {th }}$ percentile | All-cause mortality | Lean mass and mortality: - (men and women combined) Fat mass and mortality: 0 (men and women combined) |
| 95 | Heitmann, et al. (2000) | n/a | Sweden, n=787 men aged 60yrs | Whole body potassium counter (lean body mass and fat mass calculated) | Quintiles, fractional polynomials | All-cause mortality | Fat free mass and mortality: reverse J (men) Fat mass and mortality: J (men) BMI and mortality: $U$ (men) |
| 96 | Hirani, et al. (2017) | Concord Health and Aging in Men Project (20052013) | Australia, $\mathrm{n}=1666$ men aged >70yrs | DEXA (Appendicular lean mass, fat \%) | Low lean mass defined as ALM:BMI ratio <0.789 [6], <br> Obesity defined as fat $\%>30 \%$ [7]. <br> 4 groups created: neither obese nor low muscle, obese only, low muscle only, sarcopenic obesity | Frailty, disability, institutionalisati on and mortality | Obesity and mortality (men): 0 <br> Low muscle and mortality (men): 0 <br> Sarcopenic obesity and mortality (men): 0 |
| 97 | Hotchkiss, et al. (2013) | Scottish Health Survey | Scotland, n=9329 <br> aged 18-86yrs | BMI, Waist circumference, waist to hip ratio | Quartiles of WC and WHR. <br> BMI groups (WHO classification) | All-cause <br> mortality, CVD <br> incidence and mortality | Adiposity and incident CHD: + (men) <br> Adiposity and incident CHD: + (women) <br> Adiposity and mortality: + (men) <br> Adiposity and mortality: + (women) <br> Adiposity and CHD mortality: 0 (men) <br> Adiposity and CHD mortality: 0 (women) |
| 98 | Howell, et al. (2018) | NHANES III and NHANES (19992010) | USA, n=5849 <br> Mexican Americans aged >20yrs | BIA (lean mass, fat mass, body fat \%), waist circumference, waist-toheight ratio, skinfolds | Modelled continuously: BMI per $5 \mathrm{~kg} / \mathrm{m}^{2}$, WC per 5 cm , Waist to height ratioper 0.5 , skinfolds per 5 mm , \% body fat per $5 \%$, lean mass per 5 kg , fat mass per 5 kg | All-cause and CVD mortality | Fat mass and CVD mortality: - (men) <br> Fat mass and CVD mortality: + (women) <br> Fat mass and mortality: 0 (men) <br> Fat mass and mortality: + (women) <br> Lean mass and CVD mortality: 0 (men) <br> Lean mass and CVD mortality: + (women) <br> Lean mass and mortality: 0 (men) <br> Lean mass and mortality: - (women) |
| 9 | Iliodromiti, et <br> al. (2018) | UK Biobank (2006-2010) | UK, n=296535 aged 40-69yrs | BMI, waist-circumference, waist-to-hip ratio, BIA (body fat \%) | All exposures treated as continuous variables (BMI $22 \mathrm{~kg} / \mathrm{m}^{2}$ was referent value) | CVD incidence | BMI and CVD: J (men and women combined) <br> WC and CVD: + linear (men and women combined) <br> WHR and CVD: + linear (men and women combined) <br> Body fat \% and CVD: + linear (men and women combined) |
| 99 | Kahn, et al. (2012) | NHANES III (1988- 1994) | USA, $\mathrm{n}=11437$ aged 18 -64 yrs | BMI, waist circumference, waist-to-hip ratio, waist-to-height ratio | Modelled continuously per 1 SD increase in each exposure and also in quartiles | All-cause mortality | WC and all-cause mortality: + (men) <br> WC and all-cause mortality: + (women) <br> WHR and all-cause mortality: + (men) <br> WHR and all-cause mortality: + (women) <br> Waist-to-height ratio and all-cause mortality: + (men) <br> Waist-to-height ratio and all-cause mortality: + (women) |
| 100 | Katzmarzyk, et <br> al. (2002) | Canada Fitness Survey | Canada, n=10323 aged 20-69yrs | Waist circumference, skinfolds principal component of skinfold residuals (indicates subcutaneous adipose tissue distribution) | Modelled each exposure continuously using polynomial models | All-cause mortality | WC and all-cause mortality: $J$ (men) <br> WC and all-cause mortality: linear + (women) <br> Skinfolds (adiposity) and mortality: J (men), <br> Skinfolds (adiposity) and mortality: linear + (women) <br> BMI and mortality: J (men) <br> BMI and mortality: linear + (women) |
| 101 | Kim, et al. (2017) | UK Biobank (2006-2010) | UK, n=403199 aged $40-69 \mathrm{yrs}$ | Grip strength, BMI, waist circumference, \% body fat | Per 5 kg increase in GS, quintiles of GS within BMI/\%body fat/waist circumference categories | All-cause and CVD mortality | Grip strength and all-cause mortality: linear - (men) <br> Grip strength and all-cause mortality: linear - (women) <br> Grip strength and CVD mortality: linear - (men) <br> Grip strength and CVD mortality: linear - (women) |


| 102 | Kizer, et al. (2011) | Cardiovascular Health Study | USA, n=3754 aged 65-100yrs | BIA (fat free mass, fat mass) BMI, waist circumference, waist to hip ratio, waist to height ratio | Quintiles | Ischemic stroke incidence, CHD | Fat mass and stroke or CHD: 0 (men) <br> Fat mass and stroke or CHD: 0 (women) <br> Fat free mass and stroke or CHD: 0 (men) <br> Fat free mass and stroke or CHD: 0 (women) <br> WC and stroke: 0 (men) <br> WC and stroke: 0 (women) <br> WHR and stroke: 0 <br> WC and CHD: + <br> WHR and CHD: + |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 103 | Kouvari et al. (2019) | ATTICA and GREECS | Greece, n=10428 | Lean mass index and fat mass index created through total body lean and fat mass (indirectly calculated through population formals based on body weight, height, waist circumference) divided by height squared | Tertiles | CVD | Fat mass and CVD: + (men) <br> Fat mass and CVD: + (women) <br> Lean mass and CVD: U (men) <br> Lean mass and CVD: $U$ (women) |
| 54 | Lee et al (2018) | Health <br> Professionals <br> Follow-up Study <br> (1987-2012) | USA, n=38006 men aged 40-75 years | Derived predicted lean body mass and fat mass using equations developed by NHANES and based on age, race, height, weight, waist circumference | Quintiles of predicted lean body mass and fat mass | All-cause and cause-specific (CVD, cancer, respiratory, other) mortality | BMI \& all-cause mortality: J (men) Fat mass \& all-cause mortality: linear (men) Lean mass \& all-cause mortality: U (men) Fat \& CVD: linear (men) Lean body mass \& CVD: weak U (men) BMI \& CVD: U (men) |
| 104 | Lee, et al. (2012) | n/a | China, n=3978 aged $>65 y r s$ | DEXA (body fat \%), waist circumference, waist to hip ratio, relative abdominal fat (abdominal fat/whole body fat) | Quintiles | All-cause and CVD mortality | All-cause mortality and \% body fat: - (men) <br> All-cause mortality and \% body fat: 0 (women) <br> All-cause mortality and waist circumference: 0 (men) <br> All-cause mortality and waist circumference: 0 (women) <br> All-cause mortality and WHR: 0 (men) <br> All-cause mortality and WHR: 0 (women) <br> CVD mortality and WC: 0 (men) <br> CVD mortality and WC: 0 (women) <br> CVD mortality and \% body fat: 0 (men) <br> CVD mortality and \% body fat: 0 (women) <br> CVD mortality and WHR: 0 (men) <br> CVD mortality and WHR: 0 (women) |
| 105 | Levitan, et al. (2009) | Swedish <br> Mammography <br> Cohort and the <br> Cohort of Swedish <br> Men | Sweden, $\mathrm{n}=80360$ aged 45-79yrs | BMI, waist circumference, waist to hip ratio, waist to height ratio | Continuously (per 1 IQR increase in each exposure) | Heart failure hospitalisation or mortality | WC and CVD: linear + (men) <br> WC and CVD: linear + (women) <br> WHR and CVD: + (men) <br> WHR and CVD: 0 (women) <br> Waist-to-height ratio and CVD: + linear (men) <br> Waist-to-height ratio and CVD: + linear (women) |
| 106 | Li, et al. (2018) | $\begin{aligned} & \text { NHANES (1999- } \\ & 2002) \end{aligned}$ | USA, n=4449 aged $>50 \mathrm{yrs}$ | DEXA (Appendicular lean mass) | Low muscle mass defined as appendicular lean mass (ALM) <19.75kg (M) or <15.02kg (F) and ALM/BMI <0.512 (M) and $<0.789 \mathrm{~kg}$ (F). Based on FNISH Sarcopenia Project definition [1]. Low muscle strength | All-cause mortality | Low muscle alone and mortality: 0 (men and women combined) Low muscle with low strength and mortality: + (men and women combined) |
| 107 | Myint, et al. <br> (2014) | EPIC-Norfolk (1997-2000) | UK, $\mathrm{n}=15062$ aged 40-79yrs | BIA (body fat \%) | Quartiles of body fat \% | All-cause mortality and CVD incidence | Body fat \% and mortality: 0 (men) Body fat \% and mortality: 0 (women) Body fat \% and CVD: 0 (men) Body fat \% and CVD: 0 (women) |


|  |  |  |  |  |  |  | BMI and all-cause mortality: 0 (men) |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  |  |  |  | BMI and all-cause mortality: 0 (women) |
|  |  |  |  |  |  |  | BMI and CVD mortality: + (men) |
|  |  |  |  |  |  |  | BMI and CVD mortality: + (women) |
|  |  |  |  |  |  |  | Waist to height ratioand mortality: + (men) |
|  |  |  |  |  |  |  | Waist to height ratioand mortality: + (women) |
|  |  |  |  |  |  |  | Waist to height ratioand CVD mortality: + (men) |
| 108 | Nalini et al. | Golestan Cohort | Iran, n=50,045 aged | BMI, waist circumference, waist to hip ratio, waist to | Quintiles | All cause and | Waist to height ratioand CVD mortality: + (women) |
|  |  |  |  | height ratio | Quinties | CVD mortality | WC and mortality: 0 (men) |
|  |  |  |  |  |  |  | WC and mortality 0 (women) |
|  |  |  |  |  |  |  | WC and CVD mortality: + (men) |
|  |  |  |  |  |  |  | WC and CVD mortality: + (women) |
|  |  |  |  |  |  |  | WHR and mortality: + (men) |
|  |  |  |  |  |  |  | WHR And mortality: + (women) |
|  |  |  |  |  |  |  | WHR and CVD mortality: + (men) |
|  |  |  |  |  |  |  | WHR and CVD mortality: + (women) |
| 109 | Newman, et al. (2006) | Health, Aging and Body Composition study | USA, n=29292 aged 70-79yrs | DEXA (leg and arm lean mass), CT scan (thigh muscle area), knee extension strength | Per 1 SD increase | All-cause mortality | Arm or leg lean mass and mortality: 0 (men and women combined) Grip strength and mortality: + (men and women combined) |
| 110 | Ofstad et al (2019) | HUNT 2 | Norway, $\mathrm{n}=61,016$ | BMI, WC, WHR, estimated total body fat (based on the YMCA's gender specific formulas), ABSI | Quartiles | CVD mortality | Body fat (estimated) and CVD mortality: + (men) <br> Body fat (estimated) and CVD mortality: + (women) <br> WC and CVD mortality: + (men) <br> WC and CVD mortality: + (women) <br> BMI and CVD mortality: + |
| 48 | Ortega, et al. (2016) | Aerobics Centre Longitudinal <br> Study (1979-2003) | USA, $\mathrm{n}=60335$ aged >20 years | BMI, body fat \% determined by hydrostatic weighing or skinfold measurements, fat mass index, fat free mass index, fat free mass | For each exposure: very low <5th percentile, low 5-15th percentile, middle 15th-85th percentile, high 85-95th percentile, very high >95th | CVD mortality | BMI and CVD: + linear (men and women combined) <br> Body fat \% and CVD: J (men and women combined) <br> Fat mass index and CVD: $J$ (men and women combined) <br> Fat free mass and CVD: 0 (men and women combined) <br> Fat free mass index and CVD: + (men and women combined) |
| 111 | Otsuka, et al (2018) | Longitudinal Study of Aging (1997-2013) | $\begin{aligned} & \text { Japan, n=1978 aged } \\ & 40-79 \text { yrs } \end{aligned}$ | DEXA (Appendicular lean mass excluding bones [4]) | ALM/leg length, ALM/height, ALM/height ${ }^{2}$, ALM/weight*100, ALM/BMI*10 | All-cause mortality or disability | Lean mass and mortality: 0 (men) Lean mass and mortality: 0 (women) |
| 112 | Padwal, et al. (2016) | Population Health Research Data Repository | Canada, $\mathrm{n}=54420$ aged >40yrs (91\% women) | DEXA (body fat \%), BMI | Quintiles of BMI and body fat \%. Fully adjusted models included both BMI and body fat \% | All-cause mortality | BMI and mortality: U (men) <br> BMI and mortality: $U$ (women) <br> Body fat \% and mortality: J (men) <br> Body fat \% and mortality: reverse J (women) |
| 113 | Park, et al. (2014) | KNHANES (20082012) | South Korea, $\mathrm{n}=7208$ aged >50yrs | DEXA (ASM, SMI) | SMI normal >32\% (men) >25.4\% (women), class I sarcopenia 29-32\% (men) 22.8-25.4\% (women), class II sarcopenia <29\% (men) <22.8\% (women) - corresponded to 1 and 2 SDs below gender-specific means of younger population [18] | CVD incidence | Sarcopenia and CVD (men): + <br> Sarcopenia and CVD (women): 0 |
| 114 | Park, et al. (2018) | Korean National Health Insurance Cohort (20082013) | South Korea, $\mathrm{n}=465,629$ adults | Weight-adjusted-waist index, waist circumference (WC), BMI, waist-to-height | 10 groups of each index | All-cause and CVD mortality | BMI and all-cause mortality: U (men and women combined) <br> WC and all-cause mortality: U (men and women combined) <br> WHR and all-cause mortality: $U$ (men and women combined) |

ratio (WHR), a body shape
index (ABSI)

|  |  |  |  | ratio (WHR), a body shape index (ABSI) |  |  | Weight-adjusted-waist index and all-cause mortality: U (men and women combined) <br> ABSI and all-cause mortality: $U$ (men and women combined) <br> Weight-adjusted-waist index and CVD mortality: linear + (men and women combined) <br> A Body Shape Index and CVD mortality: linear + (men and women combined) <br> BMI and CVD: inverse J (men and women combined) <br> WC and CVD: inverse J (men and women combined) <br> WHR and CVD: inverse J (men and women combined) |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 115 | Reis, et al. (2009) | NHANES III (1988- 1994) | $\begin{aligned} & \text { USA, } n=12,228 \text { aged } \\ & 30-102 y s \end{aligned}$ | Waist-to-thigh ratio, waist-to-hip ratio, waist circumference, BMI | BMI categories, quintiles of WC, WHR and waist-to-thigh ratio | All-cause mortality | WHR and mortality: linear + (men) <br> WHR and mortality: 0 (women) <br> Waist-to-height ratio and mortality: + linear (men) <br> Waist-to-height ratio and mortality: + (women) <br> BMI and mortality: U (men) <br> BMI and mortality: U (women) <br> WC and mortality: J (women) <br> WC and mortality: J (men) |
| 116 | Reis, et al. (2009) | NHANES III (1988- 1994) | USA, $n=5,780$ aged 30-64yrs | BMI, waist circumference, waist to hip ratio, waist to thigh ratio | Quartiles of each exposure, results stratified by sex and race (white/black) | All-cause mortality | BMI and all-cause mortality: 0 (men) <br> BMI and all-cause mortality: 0 (women) <br> WHR and all-cause mortality: linear + (women) <br> WHR and all-cause mortality: 0 (men) <br> Waist-to-thigh ratio and all-cause mortality: linear + (men) <br> Waist-to-thigh ratio and all-cause mortality: linear + (women) <br> WC and all-cause mortality: 0 (men) <br> WC and all-cause mortality: 0 (women) |
| 117 | Rexrode, et al. (2001) | Physicians Health Study | USA, $n=16,164$ men aged 40-84yrs | Waist circumference, waist to hip ratio, BMI | Quintiles of each exposure | CHD | WC and CVD: linear + (men) WHR and CVD: + linear (men) BMI and CVD: + linear (men) |
| 118 | Sim et al (2019) | Perth Longitudinal Study in Aging women | Australia, n=903 older women | DEXA (appendicular lean mass), BMI, grip strength, timed up and go | Various definitions of sarcopenia. Foundation of National Institues of Health (FNIH), European Working Group on Sarcopenia in Older People (EWGSOP), and the adapted FNIH (AUS-POPF) using Australian population-specific cut points (<2 standard deviation below mean of young healthy) | All cause mortality | Lean mass and mortality: 0 (women) |
| 119 | Simpson, et al. (2007) | Melbourne Collaborative Cohort Study | Australia, $\mathrm{n}=41313$ aged 27-75yrs | BIA (fat mass, body fat \%), BMI, waist circumference, waist to hip ratio | Quintiles | All-cause mortality | BMI and all-cause mortality: $U$ (men) <br> BMI and all-cause mortality: $U$ (women) WC and all-cause mortality: linear + (men) WC and all-cause mortality: linear + (women) WHR and all-cause mortality: linear + (men) WHR and all-cause mortality: linear + (women) Fat mass and all-cause mortality: linear + (men) Fat mass and all-cause mortality: 0 (women) Body fat \% and all-cause mortality: linear + (men) Body fat \% and all-cause mortality: 0 (women) |
| 46 | Spahilari, et al. (2016) | Cardiovascular Health Study | USA, $\mathrm{n}=1355$ aged >65yrs | DEXA (Total, appendicular lean and fat mass) | ALM, fat mass and lean principal components modelled linearly. Fat principal components modelled in quartiles | All-cause, CVD and non-CVD mortality | Appendicular lean mass and CVD: linear - (men and women combined) Appendicular lean mass and mortality: linear - (men and women combined) <br> Fat mass and CVD: linear - (men and women combined) <br> Fat mass and mortality: linear - (men and women combined) |


| 47 | Srikanthan, et al. (2014) | NHANES III (19881994) | USA, n=3659 men aged >55 and women aged $>65 \mathrm{yrs}$ | BIA (SMI), non-muscle index (BMI - SMI) | Quartiles of SMI | All-cause mortality | Skeletal muscle index and mortality: weak - (men and women combined) |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 120 | Srikanthan, et al. (2016) | NHANES (19992004) | $\begin{aligned} & \text { USA, n=6541 aged } \\ & >20 y r s \end{aligned}$ | DEXA (Appendicular skeletal muscle mass index, trunk fat mass index) | 4 groups: Low muscle/low fat (ASMI<median, TRFI<median), Low muscle/high fat (ASMI<median, TRFI>=median), High muscle/low fat (ASMI>=median, TRFI<median), high muscle. High fat (ASMI>=median, TRFI>=median) | All-cause and CVD mortality | High muscle/low fat and CVD: - (men and women combined) High muscle/low fat and mortality: - (men and women combined) All other body composition groups and mortality: 0 (men and women combined) <br> All other body composition group and CVD: 0 (men and women combined) |
| 121 | Stefan, et al. (2008) | Cooper Centre Longitudinal study (1970-2005) | USA, $n=11335$ women | BMI, skinfolds (\% body fat), waist circumference, waist to height ratio, waist to hip ratio | Categorised as high or normal for each exposure. Cut-offs: Body fat $30 \%$, Waist to hip ratio 0.75 , waist to height ratio 0.5 | All-cause mortality | Body fat \% and all-cause mortality (women): 0 <br> WC and all-cause mortality (women): 0 <br> WHR and all-cause mortality (women): 0 <br> Waist-to-height ratio and all-cause mortality (women): 0 <br> BMI and all-cause mortality (women): 0 |
| 122 | Sui, et al. (2007) | Aerobics Centre Longitudinal Study (1979-2001) | USA, $n=2603$ aged >60yrs | Hydrostatic weighing or skinfolds (\% body fat, fat mass, fat free mass), waist circumference, BMI | BMI categories, WC categorised as abdominal obese if $>88 \mathrm{~cm}$ (men) or $>102 \mathrm{~cm}$ (women), \% Body fat \% categorised as obese if $>25 \%$ (men), $>30 \%$ (women), quintiles of FFM | All-cause mortality | WC and all-cause mortality: 0 (men and women combined) Body fat \% and all-cause mortality: 0 (men and women combined) Fat free mass and all-cause mortality: + (men and women combined) |
| 123 | Tanne, et al. (2005) | Israeli Ischemic Heart Disease project | Israel, n=9151 men aged $>23$ ys | BMI, Subscapular skinfold (SSF), subscapular to triceps skinfold thickness ratio (SFR) | Exposures analysed per 1 SD increase | Stroke and CHD mortality | Trunk obesity and CVD mortality (men): 0 Body fat distribution and CHD (men): + linear Body fat distribution and stroke (men): 0 |
| 124 | The DECODE Study Group (2008) | DECODE study | Europe, n=15521 <br> aged $30-89 y r s$ | Waist circumference | Abdominal obesity defined as waist circumference $>94 \mathrm{~cm}$ (men) or $>80 \mathrm{~cm}$ (women) | CVD mortality | WC and CVD mortality (men and women combined): 0 |
| 125 | Thomson, et al. (2016) | Women's Health Initiative Observational Study | USA, $n=77505$ women | BMI, a body shape index, body adiposity index | Quintiles of ABSI and WC, BMI groups | All-cause mortality | A Body Shape Index and CVD: linear + (women) BMI and CVD: U (women) Body adiposity index and CVD: U (women) |
| 126 | Toss, et al. (2012) | $\mathrm{n} / \mathrm{a}$ | Sweden, n=921 <br> aged $>65 \mathrm{yrs}$ | DEXA (fat and lean mass) | Assessed continuously per kg | All-cause mortality | Lean mass and mortality: linear - (men) <br> Lean mass and mortality: linear - (women) <br> Fat mass and mortality: linear - (women) <br> Fat mass and mortality: 0 (men) |
| 127 | Van Aller et al (2018) | NHANES (19992004) | $\begin{aligned} & \text { USA, n=3577 aged } \\ & >=50 y r s \end{aligned}$ | DEXA | Body composition phenotype (4 categories based on having low adiposity or high adiposity and low muscle mass or high muscle mass), Trunk FM/ASM ratio, Fat mass/fat free mass ratio. Then defined sarcopenic obese from cut-offs at various percentiles for each of these measures | All cause mortality | Sarcopenic obesity and mortality: + (men) aged 50-70 yr <br> Sarcopenic obesity and mortality: + (women) aged 50-70 yr <br> Sarcopenic obesity and mortality: 0 (men) aged $>70 \mathrm{yr}$ <br> Sarcopenic obesity and mortality: 0 (women) aged $>70 \mathrm{yr}$ |
| 128 | Wang, et al. (2013) | Kailuan study | China, $\mathrm{n}=94733$ aged 18-98yrs | BMI, waist circumference, waist-to-hip ratio, waist-to-height ratio | Quintiles | Total stroke, ischemic stroke, haemorrhagic stroke | WC and total stroke: + linear (men and women combined) WHR and total stroke: + linear (men and women combined) Waist-to-height ratio and total stroke: + linear (men and women combined) |
| 129 | Wannamethee, et al. (2007) | British Regional Heart Study (1998-2000) | UK, n=4107 men aged 60-79 | Mid-upper arm circumference, mid arm muscle circumference, BMI, BIA (Fat mass index, fat free mass index) | Quartiles of each exposure | All-cause mortality | Fat mass and mortality: 0 (men) <br> Fat free mass and mortality: 0 (men) <br> MAMC (muscle) and mortality: - linear (men) |


| 130 | Wannamethee, et al. (2014) | British Regional Heart Study (1998-2000) | UK, $n=4046$ men aged 60-79yrs | Mid-upper arm circumference, mid arm muscle circumference, BMI | Tertiles of MAMC, and results stratified by prior CVD condition (none, coronary heart disease, heart failure) | All-cause mortality | MAMC (muscle) and CHD (men): linear MAMC (muscle) and heart failure (men): 0 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 131 | Yang, et al. (2019) | n/a | China, n=329 aged $>=70 y r s$ from 4 nursing homes | SARC-F and SARC-CaIF questionnaires to screen for sarcopenia. | Score of $>=4$ (SARC-F) or $>=11$ (SARC-CaIF) indicate sarcopenia | All-cause mortality | Sarcopenia and mortality: + (men and women combined) |
| 132 | Yuki, et al. (2017) | Longitudinal Study of Aging (1997-2013) | Japan, n=700 aged 65-79yrs | DEXA (skeletal muscle mass index), grip strength, gait speed | Low muscle mass defined as $\mathrm{SMI}<7.0$ (men), <5.4 (women) [15]. Low grip strength defined as $<26 \mathrm{~kg}$ (men) <18kg (women) [15]. Low gait speed defined as $<0.8 \mathrm{~m} / \mathrm{s}$ [15]. Sarcopenia if low SMI and one of low grip strength or low gait speed | All-cause mortality | Sarcopenia and mortality: + (men) <br> Sarcopenia and mortality: 0 (women) |
| 133 | Zaslavsky, et al. (2017) | Women's Health Initiative Observational Study (1993-1998) | USA, $n=876$ frail women aged >65 | DEXA (Appendicular, central and total lean and fat mass) | Quartiles of each measurement | All-cause mortality | Lean mass \& mortality: 0 (women) Fat mass \& mortality: linear - (women) |
| 134 | Zhang, et al. (2007) | Shanghai Women's Health Study (1996-2000) | China, $\mathrm{n}=72773$ women aged 4070yrs | Waist to hip ratio | Quintiles | All-cause and CVD mortality | WHR and mortality: linear + (women) WHR and CVD mortality: linear + (women) |
| 135 | Zhu, et al. (2003) | NHANES I \& II | USA, $n=13369$ women aged 2575yrs | Skinfolds (fat mass, fat free mass) | All exposures analysed as continuous | All-cause mortality | BMI and mortality (women): U <br> Fat free mass and mortality (women): linear - <br> Fat mass and mortality (women): linear - |
| 136 | Zong, et al. (2016) | $\begin{aligned} & \text { NHANES (1996- } \\ & \text { 2006) } \end{aligned}$ | $\begin{aligned} & \text { USA, n=9471 aged } \\ & >20 y r s \end{aligned}$ | DEXA (body fat \% for whole-body, trunk, leg) | Quartiles of each measure of FM\% | All-cause and CVD mortality | Body fat \% and CVD: U (men and women combined) Body fat \% and all-cause mortality: U (men and women combined) |
|  <br>  Korea National Health and Nutrition Examination Survey; NHANES: National Health and Nutrition Examination Survey; SMI: Skeletal muscle mass index (Skeletal muscle mass of whole body, kg/height ${ }^{2}$, m ${ }^{2}$ ) |  |  |  |  |  |  |  |

## Table S3. Derivation of variables used in analysis from the UK Biobank questionnaire and interviews.

| Covariate | Categories used in analysis | UK Biobank variable used (question ID) and source |
| :---: | :---: | :---: |
| Body composition variables (exposures) |  |  |
| Appendicular skeletal muscle mass (aSMM) | Quintiles of the residuals from the model of aSMM regressed on height | Arm predicted mass (left) (UKBBID: 23126) ${ }^{\ddagger}$, arm predicted mass (right) (UKBBID: 23122) ${ }^{\ddagger}$, leg predicted mass (left) (UKBBID: 23118) ${ }^{\ddagger}$, leg predicted mass (right) (UKBBID: 23114). <br> Height (UKBBID: 12144) ${ }^{\ddagger}$. |
| Fat mass (FM) | Quintiles | Whole body fat mass (UKBBID: 23100) ${ }^{\ddagger}$ |
| Body composition groups | 1. Low aSMM / low FM <br> 2. Low aSMM / moderate FM <br> 3. Low aSMM / high FM <br> 4. Moderate aSMM / low FM <br> 5. Moderate aSMM / moderate FM <br> 6. Moderate aSMM / high FM <br> 7. High aSMM / low FM <br> 8. High aSMM / moderate FM <br> 9. High aSMM / high FM | Tertiles of aSMM within tertiles of FM derived from: arm predicted mass (left) (UKBBID: 23126) ${ }^{\ddagger}$, arm predicted mass (right) (UKBBID: 23122) ${ }^{\ddagger}$, leg predicted mass (left) (UKBBID: 23118) ${ }^{\ddagger}$, leg predicted mass (right) (UKBBID: 23114), Whole body fat mass (UKBBID: 23100) ${ }^{\ddagger}$, Height (UKBBID: 12144) ${ }^{\ddagger}$. <br> Low aSMM, residuals of aSMM regressed on height: -10.82- $\leq-0.97$ (men), $-5.88-\leq-0.57$ (women) <br> Moderate aSMM: residuals of aSMM regressed on height: -0.97- $\leq 0.85$ (men), $-0.57-\leq 0.45$ (women) <br> High aSMM: residuals of aSMM regressed on height: $0.85-\leq 14.50$ (men), $0.46-\leq 11.99$ (women) <br> Low FM, kg: $5.0-\leq 18.1$ (men), $5.0-\leq 21.4$ (women) <br> Moderate FM: 18.2- $\leq 24.1$ (men), 21.5- $\leq 28.8$ (women) <br> High FM: $24.2-\leq 98.6$ (men), $28.9-\leq 109.8$ (women) |
| BMI | Continuous ( $\mathrm{kg} / \mathrm{m}^{2}$ ) | Body mass index (UKBBID: 21001) ${ }^{\ddagger}$ |
| Waist circumference | Continuous (cm) | Waist circumference (UKBBID: 48) ${ }^{\ddagger}$ |
| Grip strength | Continuous (kg) | Hand grip strength (left) (UKBBID: 46) ${ }^{\ddagger}$, Hand grip strength (right) (UKBBID: 47) ${ }^{\ddagger}$ |
| Sociodemographic characteristics |  |  |
| Sex | Men <br> Women | Sex (UKBBID: 31)* |
| Education | Higher degree (college or university degree, or professional qualifications) <br> Any school degree (A levels, AS levels, O levels, GCSEs or CSEs) <br> Vocational qualifications (NVQ, HND or HNC) <br> Other (none of the above qualifications) | Qualifications (UKBBID: 6138) ${ }^{\dagger}$ |
| Townsend index | Quintiles (high index indicates most deprivation) | Townsend index (UKBBID: 189)* |
| Lifestyle factors |  |  |
| Smoking status | Never <br> Current <br> Previous | Smoking status (UKBBID: 20116) ${ }^{\dagger}$ |
| Alcohol intake | None <br> Occasional (<1 unit/week) <br> Moderate (1-14 units/week) <br> Heavy (>14 units/week) | Alcohol intake frequency (UKBBID: 1558) ${ }^{\dagger}$ <br> Depending on the participants' response, they were asked how much they consumed per week or month of the following alcoholic drinks: |


|  |  | Red wine (UKBBID: 4407, 1568) ${ }^{\dagger}$; champagne and white wine (UKBBID: 4418, 1578) ${ }^{\dagger}$; beer and cider (UKBBID: 4429, $1588)^{\dagger}$; spirits (UKBBID: 4440, 1598) ${ }^{\dagger}$; fortified wine (UKBBID: 4451, 1608) ${ }^{\dagger}$; other (UKBBID: 4462, 5364) ${ }^{\dagger}$. |
| :---: | :---: | :---: |
| Saturated fat score | None <br> Low <br> Medium <br> High <br> Unknown | Beef intake (UKBBID: 1369) ${ }^{\dagger}$, cheese intake (UKBBID: 1408$)^{\dagger}$, pork intake (UKBBID: 1389) ${ }^{\dagger}$, lamb intake (UKBBID: 1379) ${ }^{\dagger}$, processed meat intake (UKBBID: 1349), spread type (UKBBID: 1428) ${ }^{\dagger}$ <br> The amount of each food eaten per week was totalled, then split into groups based off tertiles (with 'none' subsequently separated from the low category). |
| Fruit and vegetable score | None <br> Low <br> Medium <br> High <br> Unknown | Fresh fruit intake (UKBBID: 1309) ${ }^{\dagger}$, raw vegetable intake (UKBBID: 1299$)^{\dagger}{ }^{\dagger}$, cooked vegetable intake (UKBBID: 1289). The amount of each food eaten per week was totalled, then split into groups based off tertiles (with 'none' subsequently separated from the low category). |
| Oily fish score | Unknown <br> None <br> Low <br> Medium <br> High | Oily fish intake (UKBBID: 1329) ${ }^{\dagger}$ <br> The amount of oily fish per week was split into groups based off tertiles (with 'none' subsequently separated from the low category). |
| Physical activity (IPAQ MET scores) | Low <br> Moderate High | Number of days/week of vigorous physical activity $10+$ minutes (UKBBID: 904) ${ }^{\dagger}$; Duration of vigorous activity (UKBBID: 914) ${ }^{+}$ <br> Number of days/week of moderate physical activity 10+ minutes (UKBBID: 884) ${ }^{\dagger}$; Duration of moderate activity (UKBBID: 894) ${ }^{+}$ <br> Number of days/week walked 10+ minutes (UKBBID: 864$)^{\dagger}$; Duration of walks (UKBBID: 874$)^{\dagger}$ |
| Medical history |  |  |
| Type 2 diabetes | No <br> Yes (if diagnosed by doctor or taking insulin for diabetes) | Diabetes diagnosed by doctor (UKBBID: 2443) ${ }^{\dagger}$ <br> Medication for cholesterol, blood pressure or diabetes (men) (UKBBID: 6177) ${ }^{\dagger}$; Medication for cholesterol, blood pressure, diabetes, or take exogenous hormones (women) (UKBBID: 6153) ${ }^{\dagger}$ |
| Hypertension | No <br> Yes (if diagnosed by doctor, had an SBP $>140 \mathrm{mmHg}$, DBP $>90 \mathrm{mmHg}$ or taking medication for blood pressure) | Vascular/heart problems diagnosed by doctor (high blood pressure is one response) (UKBBID: 6150) ${ }^{+}$ <br> Systolic blood pressure, automated reading / manual reading (UKBBID: 4080 / 93) ${ }^{\ddagger}$ <br> Diastolic blood pressure, automated reading / manual reading (UKBBID: 4079 / 94) ${ }^{\ddagger}$ <br> Medication for cholesterol, blood pressure or diabetes (men) (UKBBID: 6177) ${ }^{\dagger}$; Medication for cholesterol, blood pressure, diabetes, or take exogenous hormones (women) (UKBBID: 6153) ${ }^{\dagger}$ |
| Cholesterol | No <br> Yes (if taking cholesterol-lowering medication) | Medication for cholesterol, blood pressure or diabetes (men) (UKBBID: 6177) ${ }^{\dagger}$; Medication for cholesterol, blood pressure, diabetes, or take exogenous hormones (women) (UKBBID: 6153) ${ }^{\dagger}$; blood lipids (UKBBID: 30690,30760,30870) |
| Prior cancer | No <br> Yes (if cancer had been previously diagnosed by a doctor) | Cancer diagnosed by doctor (UKBBID: 2453) ${ }^{+}$ |
| Menopause | No <br> Yes (if responded to questions saying they have experienced the menopause) | Had menopause (women only) (UKBBID 2724) ${ }^{+}$ |
| *Recruitment questions <br> ${ }^{\dagger}$ Touchscreen questions <br> ${ }^{\ddagger}$ Physical measurements |  |  |

Table S4. Baseline characteristics of the study population according to appendicular skeletal muscle mass and fat mass quintiles.

|  | Men appendicular skeletal muscle mass (aSMM) quintiles, range (kg) |  |  |  |  | Men fat mass (FM) quintiles, range (kg) |  |  |  |  | Total |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 13.5- 24.0 | 24.1- $\mathbf{2 6 . 0}$ | 26.1- 27.6 | 27.7- 30.0 | $30.1-\leq 54.5$ | $5.0-\leq 15.7$ | $15.8-\leq 19.4$ | 19.5- 22.9 | 23.0- 27.5 | 27.6- 598.6 |  |
| Age at recruitment, mean (SD) | 61.0 (6.4) | 58.2 (7.4) | 56.0 (7.8) | 54.0 (7.9) | 51.7 (7.8) | 54.9 (8.3) | 56.0 (8.2) | 56.5 (8.1) | 56.7 (8.0) | 56.9 (7.8) | 56.2 (8.1) |
| aSMM (kg), mean (SD) | 24.6 (2.9) | 25.9 (2.8) | 26.9 (2.9) | 28.1 (3.0) | 30.5 (3.6) | 24.6 (2.7) | 25.7 (2.7) | 26.7 (2.7) | 28.0 (2.8) | 31.0 (3.6) | 27.2 (3.7) |
| FM (kg), mean (SD) | 22.4 (7.1) | 21.6 (7.2) | 21.4 (7.4) | 21.5 (7.8) | 22.2 (9.2) | 12.4 (2.4) | 17.5 (1.1) | 21.0 (1.0) | 24.9 (1.3) | 33.6 (6.2) | 21.8 (7.8) |
| Body mass index (BMI), mean (SD) | 26.0 (3.5) | 26.8 (3.5) | 27.4 (3.6) | 28.2 (3.8) | 29.7 (4.4) | 23.3 (1.8) | 25.6 (1.6) | 27.1 (1.6) | 29.0 (1.8) | 33.1 (3.5) | 27.6 (4.0) |
| Weight (kg), mean (SD) | 81.1 (12.1) | 83.0 (12.1) | 84.9 (12.4) | 87.3 (13.0) | 92.6 (15.1) | 71.7 (7.0) | 78.7 (6.3) | 84.0 (6.2) | 90.3 (6.5) | 104.3 (11.8) | 85.8 (13.6) |
| Height (cm), mean (SD) | 176.6 (6.7) | 175.9 (6.7) | 175.8 (6.7) | 176.0 (6.7) | 176.6 (6.7) | 175.3 (6.8) | 175.5 (6.6) | 176.0 (6.5) | 176.6 (6.6) | 177.6 (6.7) | 176.2 (6.7) |
| Higher education, n (\%) | 12740 (39.7) | 13085 (40.7) | 13140 (40.9) | 13096 (40.7) | 12645 (39.3) | 13909 (43.1) | 13669 (42.1) | 12987 (40.6) | 12406 (38.8) | 11735 (36.7) | 64706 (40.2) |
| Current smokers, n (\%) | 3714 (11.6) | 3431 (10.7) | 3573 (11.1) | 3563 (11.1) | 3724 (11.6) | 4366 (13.5) | 3600 (11.1) | 3368 (10.5) | 3385 (10.6) | 3286 (10.3) | 18005 (11.2) |
| Low fruit and vegetable intake, n (\%) | 14779 (46.0) | 14122 (43.9) | 13977 (43.5) | 13996 (43.5) | 13367 (41.6) | 13624 (42.2) | 13909 (42.8) | 14191 (44.3) | 14168 (44.3) | 14349 (44.8) | 70241 (43.7) |
| High saturated intake, n (\%) | 11706 (36.4) | 11786 (36.6) | 11647 (36.2) | 11637 (36.2) | 11778 (36.6) | 10457 (32.4) | 11354 (34.9) | 11699 (36.6) | 12111 (37.9) | 12933 (40.4) | 58554 (36.4) |
| Low oily fish intake, n (\%) | 10919 (34.0) | 11292 (35.1) | 11342 (35.3) | 11807 (36.7) | 11988 (37.3) | 11197 (34.7) | 11452 (35.2) | 11518 (36.0) | 11520 (36.0) | 11661 (36.4) | 57348 (35.7) |
| Heavy drinkers, n (\%) | 19948 (62.1) | 19885 (61.8) | 19841 (61.7) | 19608 (61.0) | 18666 (58.1) | 17562 (54.4) | 19859 (61.1) | 20297 (63.4) | 20535 (64.2) | 19695 (61.5) | 97948 (60.9) |
| Low physical activity, n (\%) | 7247 (22.6) | 6670 (20.7) | 6300 (19.6) | 5901 (18.3) | 5518 (17.2) | 4580 (14.2) | 5351 (16.5) | 6030 (18.8) | 6918 (21.6) | 8757 (27.4) | 31636 (19.7) |
| Hypertension, n (\%) | 20146 (62.7) | 18878 (58.7) | 18102 (56.3) | 17362 (54.0) | 17370 (54.0) | 12683 (39.3) | 16508 (50.8) | 18523 (57.9) | 20600 (64.4) | 23544 (73.5) | 91858 (57.1) |
| Type 2 diabetes, n (\%) | 1300 (4.1) | 1250 (3.9) | 1156 (3.6) | 1191 (3.7) | 1408 (4.4) | 388 (1.2) | 666 (2.1) | 906 (2.8) | 1462 (4.6) | 2883 (9.0) | 6305 (3.9) |
| Cancer history (>5 years ago), n (\%) | 1176 (3.7) | 927 (2.9) | 816 (2.5) | 662 (2.1) | 652 (2.0) | 780 (2.4) | 836 (2.6) | 856 (2.7) | 872 (2.7) | 889 (2.8) | 4233 (2.6) |
| Cholesterol medication, n (\%) | 6204 (19.5) | 5171 (16.2) | 4646 (14.6) | 4145 (13.0) | 3842 (12.0) | 2175 (6.8) | 3807 (11.8) | 4664 (14.7) | 5922 (18.6) | 7440 (23.4) | 24008 (15.0) |
|  | Women appendicular skeletal muscle mass (aSMM) quintiles, range (kg) |  |  |  |  |  | Women fat mass (FM) quintiles, range (kg) |  |  |  | Total |
|  | $10.3-\leq 16.5$ | 16.6- 17.5 | $17.6-\leq 18.6$ | 18.6- 19.9 | 20.0- 39.2 | 5.0- $\leq 18.5$ | 18.6-522.8 | 22.9- 27.1 | 27.2- 33.3 | 33.4- 109.8 |  |
| Age at recruitment, mean (SD) | 58.9 (7.0) | 57.1 (7.6) | 55.8 (7.9) | 54.6 (8.0) | 53.2 (8.1) | 54.1 (8.1) | 55.7 (8.0) | 56.6 (7.9) | 57.0 (7.7) | 56.4 (7.8) | 55.9 (8.0) |
| aSMM (kg), mean (SD) | 17.0 (1.7) | 17.5 (1.7) | 18.0 (1.8) | 18.7 (1.9) | 20.3 (2.5) | 16.6 (1.5) | 17.3 (1.4) | 17.9 (1.4) | 18.8 (1.5) | 21.1 (2.3) | 18.3 (2.3) |
| FM (kg), mean (SD) | 27.7 (8.5) | 26.0 (8.4) | 25.5 (8.8) | 25.3 (9.5) | 27.0 (11.9) | 15.2 (2.6) | 20.7 (1.2) | 24.9 (1.3) | 29.9 (1.7) | 41.1 (7.4) | 26.3 (9.6) |
| Body mass index (BMI), mean (SD) | 25.9 (4.2) | 26.0 (4.2) | 26.3 (4.4) | 26.8 (4.8) | 28.5 (6.0) | 21.6 (1.7) | 24.0 (1.5) | 25.9 (1.7) | 28.3 (2.0) | 33.9 (4.3) | 26.7 (4.9) |
| Weight (kg), mean (SD) | 69.1 (11.6) | 68.8 (11.6) | 69.4 (12.1) | 70.8 (12.9) | 75.9 (16.3) | 56.4 (4.7) | 63.2 (3.7) | 68.5 (3.7) | 75.3 (4.2) | 90.9 (11.1) | 70.8 (13.3) |
| Height (cm), mean (SD) | 163.4 (6.2) | 162.6 (6.1) | 162.5 (6.2) | 162.6 (6.2) | 163.3 (6.2) | 161.6 (6.1) | 162.5 (6.1) | 162.9 (6.1) | 163.3 (6.1) | 164.0 (6.2) | 162.9 (6.2) |
| Higher education, n (\%) | 14330 (36.6) | 15246 (38.9) | 16033 (40.9) | 16513 (42.2) | 17056 (43.6) | 18112 (45.5) | 16503 (42.6) | 15834 (39.8) | 14520 (37.7) | 14209 (36.4) | 79178 (40.4) |
| Current smokers, n (\%) | 3039 (7.8) | 3164 (8.1) | 3231 (8.2) | 3258 (8.3) | 3377 (8.6) | 3792 (9.5) | 3166 (8.2) | 3151 (7.9) | 3062 (8.0) | 2898 (7.4) | 16069 (8.2) |
| Low fruit and vegetable intake, n (\%) | 12608 (32.2) | 12333 (31.5) | 11874 (30.3) | 11521 (29.4) | 10910 (27.9) | 12125 (30.5) | 11518 (29.7) | 11756 (29.6) | 11613 (30.2) | 12234 (31.3) | 59246 (30.3) |
| High saturated intake, n (\%) | 11699 (29.9) | 11407 (29.1) | 11252 (28.7) | 11485 (29.3) | 11212 (28.6) | 10415 (26.2) | 10824 (27.9) | 11615 (29.2) | 11762 (30.6) | 12439 (31.9) | 57055 (29.1) |
| Low oily fish intake, n (\%) | 12200 (31.2) | 12678 (32.4) | 12619 (32.2) | 13034 (33.3) | 13137 (33.6) | 12971 (32.6) | 12521 (32.3) | 12669 (31.8) | 12425 (32.3) | 13082 (33.5) | 63668 (32.5) |
| Heavy drinkers, n (\%) | 12955 (33.1) | 12840 (32.8) | 13025 (33.2) | 13134 (33.5) | 12318 (31.5) | 13562 (34.1) | 13654 (35.2) | 13525 (34.0) | 12561 (32.6) | 10970 (28.1) | 64272 (32.8) |
| Low physical activity, n (\%) | 10117 (25.8) | 8974 (22.9) | 8431 (21.5) | 7986 (20.4) | 7837 (20.0) | 6398 (16.1) | 7114 (18.4) | 8288 (20.8) | 9385 (24.4) | 12160 (31.1) | 43345 (22.1) |
| Hypertension, n (\%) | 19371 (49.5) | 17669 (45.1) | 16440 (42.0) | 15877 (40.5) | 16330 (41.7) | 11345 (28.5) | 14192 (36.6) | 17091 (43.0) | 19293 (50.1) | 23766 (60.9) | 85687 (43.8) |
| Type 2 diabetes, n (\%) | 635 (1.6) | 673 (1.7) | 669 (1.7) | 805 (2.1) | 1250 (3.2) | 239 (0.6) | 363 (0.9) | 524 (1.3) | 860 (2.2) | 2046 (5.3) | 4032 (2.1) |
| Cancer history (>5 years ago), n (\%) | 2540 (6.5) | 2298 (5.9) | 2010 (5.1) | 1933 (4.9) | 1845 (4.7) | 1912 (4.8) | 2052 (5.3) | 2278 (5.7) | 2214 (5.8) | 2170 (5.6) | 10626 (5.4) |
| Cholesterol medication, n (\%) | 4294 (11.0) | 3605 (9.3) | 3163 (8.1) | 3015 (7.8) | 3080 (7.9) | 1536 (3.9) | 2350 (6.1) | 3349 (8.5) | 4201 (11.0) | 5721 (14.8) | 17157 (8.8) |
| Post-menopausal, n (\%) | 28352 (72.4) | 25700 (65.6) | 23352 (59.6) | 21067 (53.8) | 17594 (44.9) | 21266 (53.5) | 22970 (59.3) | 24686 (62.1) | 24170 (62.8) | 22973 (58.8) | 116065 (59.3) |

$\chi^{2}$ test for trend: $\mathrm{p}<0.05$ for all characteristics across the aSMM and FM quintiles. All characteristics were determined at the baseline assessment clinic through touch-screen questionnaires, interviews and/or physical


 saturated fat: the highest saturated fat tertile, based off portions per week of beef, lamb, pork and whether they consumed animal or plant-based spreads. Low oily fish: lowest consumption tertile (< 1 portion per week).

Table S5. Baseline characteristics of the study population according to body composition groups.

| Men | Low Muscle / <br> Low Fat | Med Muscle / <br> Low Fat | High Muscle / <br> Low Fat | Low Muscle / <br> Med Fat | Med Muscle / <br> Med Fat | High Muscle / <br> Med Fat | Low Muscle / High Fat | Med Muscle / High Fat | High Muscle / High Fat |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Age at recruitment, mean (SD) | 59.0 (7.3) | 55.1 (8.0) | 51.7 (8.0) | 60.3 (6.7) | 56.3 (7.7) | 52.6 (8.0) | 60.7 (6.5) | 56.7 (7.6) | 53.2 (7.8) |
| Appendicular skeletal muscle (kg), mean (SD) | 22.9 (2.1) | 24.9 (1.9) | 27.1 (2.3) | 24.5 (1.9) | 26.6 (1.8) | 29.1 (2.2) | 27.3 (2.6) | 29.6 (2.5) | 32.7 (3.3) |
| Fat mass (kg), mean (SD) | 14.6 (2.8) | 14.3 (2.8) | 13.8 (3.1) | 21.0 (1.7) | 21.0 (1.7) | 21.0 (1.7) | 29.8 (5.5) | 30.0 (5.8) | 31.3 (7.2) |
| Body mass index (BMI), mean (SD) | 22.8 (1.6) | 24.1 (1.6) | 25.4 (1.8) | 25.7 (1.3) | 27.2 (1.2) | 28.6 (1.4) | 29.7 (2.8) | 31.4 (2.9) | 33.5 (3.6) |
| Weight, mean (SD) | 70.3 (6.3) | 74.0 (6.3) | 78.3 (7.0) | 79.7 (5.3) | 83.8 (5.1) | 88.8 (5.9) | 93.6 (9.7) | 98.4 (10.0) | 105.4 (12.3) |
| Height, mean (SD) | 175.4 (6.8) | 175.1 (6.7) | 175.5 (6.7) | 176.1 (6.6) | 175.7 (6.5) | 176.2 (6.6) | 177.4 (6.6) | 177.0 (6.6) | 177.3 (6.7) |
| Higher education, n (\%) | 7753 (42.8) | 7814 (43.2) | 7626 (42.1) | 7240 (41.0) | 7309 (41.3) | 6949 (39.3) | 6488 (36.4) | 6760 (37.9) | 6767 (38.0) |
| Current smokers, n (\%) | 2406 (13.3) | 2139 (11.8) | 2285 (12.6) | 1850 (10.5) | 1882 (10.6) | 1951 (11.0) | 1768 (9.9) | 1823 (10.2) | 1901 (10.7) |
| Low fruit and vegetable intake, n (\%) | 8143 (45.0) | 7708 (42.6) | 7199 (39.8) | 7903 (44.7) | 7827 (44.3) | 7593 (42.9) | 8095 (45.5) | 7900 (44.3) | 7873 (44.2) |
| High saturated intake, n (\%) | 6078 (33.6) | 6014 (33.2) | 6006 (33.2) | 6423 (36.4) | 6406 (36.2) | 6510 (36.8) | 6978 (39.2) | 7064 (39.6) | 7075 (39.7) |
| Low oily fish intake, n (\%) | 6141 (33.9) | 6302 (34.8) | 6510 (36.0) | 5948 (33.7) | 6296 (35.6) | 6698 (37.9) | 6280 (35.3) | 6539 (36.7) | 6634 (37.2) |
| Heavy drinkers, n (\%) | 10276 (56.8) | 10373 (57.3) | 10233 (56.5) | 11242 (63.6) | 11271 (63.7) | 10997 (62.2) | 11508 (64.7) | 11370 (63.8) | 10498 (58.9) |
| Low physical activity, n (\%) | 3250 (18.0) | 2669 (14.7) | 2204 (12.2) | 3657 (20.7) | 3413 (19.3) | 2993 (16.9) | 4695 (26.4) | 4475 (25.1) | 4280 (24.0) |
| Hypertension, n (\%) | 8641 (47.7) | 7727 (42.7) | 7208 (39.8) | 10898 (61.7) | 10187 (57.6) | 9600 (54.3) | 12947 (72.7) | 12437 (69.8) | 12213 (68.6) |
| Diabetes, n (\%) | 298 (1.6) | 245 (1.4) | 274 (1.5) | 558 (3.2) | 505 (2.9) | 500 (2.8) | 1233 (6.9) | 1245 (7.0) | 1447 (8.2) |
| Cholesterol medication, n (\%) | 1930 (10.7) | 1505 (8.4) | 1195 (6.7) | 3179 (18.1) | 2527 (14.4) | 2093 (11.9) | 4515 (25.5) | 3723 (21.0) | 3341 (18.9) |
| Cancer diagnosed by doctor (>5 years ago), n (\%) | 573 (3.2) | 407 (2.3) | 373 (2.1) | 591 (3.3) | 480 (2.7) | 340 (1.9) | 651 (3.7) | 468 (2.6) | 350 (2.0) |
| Women | Low Muscle / <br> Low Fat | Med Muscle / Low Fat | High Muscle / Low Fat | Low Muscle / <br> Med Fat | Med Muscle / Med Fat | High Muscle / Med Fat | Low Muscle / High Fat | Med Muscle / High Fat | High Muscle / High Fat |
| Age at recruitment, mean (SD) | 57.1 (7.7) | 54.6 (8.0) | 52.2 (7.9) | 58.7 (7.1) | 56.6 (7.8) | 54.4 (8.1) | 58.7 (7.1) | 56.5 (7.7) | 54.7 (8.0) |
| Appendicular skeletal muscle (kg), mean (SD) | 15.7 (1.0) | 16.7 (1.0) | 18.0 (1.3) | 16.7 (1.0) | 17.8 (0.9) | 19.2 (1.2) | 18.7 (1.5) | 20.0 (1.5) | 22.1 (2.3) |
| Fat mass (kg), mean (SD) | 17.8 (2.8) | 17.1 (3.1) | 16.4 (3.4) | 25.1 (2.1) | 24.9 (2.1) | 24.9 (2.1) | 36.1 (6.7) | 36.3 (6.9) | 38.8 (8.8) |
| Body mass index (BMI), mean (SD) | 21.8 (1.7) | 22.5 (1.8) | 23.1 (1.9) | 24.9 (1.6) | 25.9 (1.6) | 26.9 (1.8) | 30.0 (3.5) | 31.5 (3.7) | 34.0 (4.7) |
| Weight, mean (SD) | 56.8 (4.4) | 58.6 (4.8) | 60.9 (5.3) | 66.1 (3.6) | 68.4 (3.6) | 71.5 (4.0) | 80.9 (9.2) | 84.0 (9.5) | 90.9 (12.6) |
| Height, mean (SD) | 161.6 (6.2) | 161.6 (6.1) | 162.5 (6.1) | 163.1 (6.1) | 162.6 (6.1) | 163.1 (6.2) | 164.2 (6.2) | 163.5 (6.1) | 163.6 (6.2) |
| Higher education, n (\%) | 9086 (41.3) | 9855 (44.8) | 10384 (47.2) | 8069 (37.2) | 8681 (39.9) | 9277 (42.7) | 7548 (35.0) | 7976 (37.0) | 8302 (38.5) |
| Current smokers, n (\%) | 1977 (9.0) | 1946 (8.8) | 2027 (9.2) | 1652 (7.6) | 1721 (7.9) | 1795 (8.3) | 1596 (7.4) | 1618 (7.5) | 1737 (8.1) |
| Low fruit and vegetable intake, n (\%) | 7248 (33.0) | 6664 (30.3) | 6047 (27.5) | 6700 (30.9) | 6491 (29.9) | 6072 (28.0) | 6863 (31.8) | 6737 (31.2) | 6424 (29.8) |
| High saturated intake, n (\%) | 5875 (26.7) | 5960 (27.1) | 5867 (26.7) | 6416 (29.5) | 6303 (29.0) | 6340 (29.2) | 6864 (31.8) | 6720 (31.2) | 6710 (31.1) |
| Low oily fish intake, n (\%) | 6984 (31.8) | 7159 (32.5) | 7342 (33.4) | 6680 (30.8) | 6845 (31.5) | 7228 (33.3) | 6969 (32.3) | 7150 (33.1) | 7311 (33.9) |
| Heavy drinkers, n (\%) | 7167 (32.6) | 7649 (34.8) | 7893 (35.9) | 7415 (34.1) | 7400 (34.0) | 7428 (34.2) | 6840 (31.7) | 6542 (30.3) | 5752 (26.7) |
| Low physical activity, n (\%) | 4468 (20.3) | 3679 (16.7) | 2965 (13.5) | 4922 (22.7) | 4536 (20.9) | 4116 (19.0) | 6478 (30.0) | 6130 (28.4) | 6051 (28.1) |
| Hypertension, n (\%) | 7938 (36.1) | 6817 (31.0) | 5918 (26.9) | 9983 (46.0) | 9342 (43.0) | 8681 (40.0) | 12572 (58.3) | 11981 (55.5) | 12455 (57.8) |
| Diabetes, n (\%) | 155 (0.7) | 138 (0.6) | 175 (0.8) | 247 (1.1) | 285 (1.3) | 362 (1.7) | 626 (2.9) | 786 (3.7) | 1258 (5.9) |
| Cholesterol medication, n (\%) | 1357 (6.2) | 979 (4.5) | 695 (3.2) | 2057 (9.5) | 1829 (8.5) | 1607 (7.5) | 3079 (14.4) | 2720 (12.7) | 2834 (13.3) |
| Cancer diagnosed by doctor (>5 years ago) , n (\%) | 1270 (5.8) | 1077 (4.9) | 932 (4.2) | 1433 (6.6) | 1173 (5.4) | 1092 (5.0) | 1358 (6.3) | 1175 (5.5) | 1116 (5.2) |
| Menopause, n (\%) | 14788 (67.3) | 12378 (56.3) | 9432 (42.9) | 15674 (72.2) | 13585 (62.5) | 11126 (51.2) | 15109 (70.1) | 13077 (60.6) | 10896 (50.6) |

All characteristics were determined at the baseline assessment clinic through touch-screen questionnaires, interviews and/or physical measurements. Higher education: has a college or university degree or professional


 highest saturated fat tertile, based off portions per week of beef, lamb, pork and whether they consumed animal or plant-based spreads. Low oily fish: lowest consumption tertile (< 1 portion per week).

Table S6. Partial correlation coefficients between appendicular skeletal muscle mass, fat mass and height, adjusted for age at recruitment.

| Men | aSMM | FM | Height |
| :---: | :--- | :--- | :--- |
| aSMM | - | 0.7132 | 0.5185 |
| FM | 0.7132 | - | 0.1375 |
| Height | 0.5185 | 0.1375 | - |
| Women | aSMM | FM | Height |
| aSMM | - | 0.7771 | 0.4403 |
| FM | 0.7771 | - | 0.1544 |
| Height | 0.4403 | 0.1544 | - |

Table S7. CVD sequential model adjustment.
Hazard ratios of CVD associated with appendicular skeletal muscle mass (aSMM) and fat mass (FM)

|  | Quintile | No. events | Age and height adjusted | + Sociodemographics* | + Lifestyle factors ${ }^{+}$ | + Medical history ${ }^{\ddagger}$ | + Mutual adjustment for ASM or FM |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Appendicular skeletal muscle mass (aSMM) |  |  |  |  |  |  |  |
| Men | 1 | 4373 | 1 (reference) | 1 (reference) | 1 (reference) | 1 (reference) | 1 (reference) |
|  | 2 | 3715 | 0.99 (0.95-1.03) | 1.00 (0.96-1.04) | 1.01 (0.97-1.06) | 1.01 (0.97-1.06) | 1.02 (0.97-1.06) |
|  | 3 | 3385 | 1.03 (0.98-1.08) | 1.04 (1.00-1.09) | 1.06 (1.01-1.11) | 1.06 (1.01-1.11) | 1.07 (1.02-1.12) |
|  | 4 | 3055 | 1.07 (1.02-1.12) | 1.08 (1.03-1.14) | 1.11 (1.06-1.16) | 1.10 (1.05-1.15) | 1.10 (1.05-1.16) |
|  | 5 | 2971 | 1.25 (1.19-1.31) | 1.26 (1.20-1.32) | 1.29 (1.23-1.35) | 1.27 (1.21-1.33) | 1.25 (1.19-1.31) |
|  | Non-linearity p-value |  | p<0.001 | p<0.001 | p<0.001 | p<0.001 | 0.014 |
|  |  |  | 103 | 106 | 120 | 106 | 87 |
| Women | 1 | 2617 | 1 (reference) | 1 (reference) | 1 (reference) | 1 (reference) | 1 (reference) |
|  | 2 | 2182 | 0.92 (0.87-0.97) | 0.93 (0.88-0.98) | 0.93 (0.88-0.99) | 0.93 (0.88-0.98) | 0.95 (0.90-1.01) |
|  | 3 | 1848 | 0.85 (0.80-0.91) | 0.87 (0.82-0.92) | 0.87 (0.82-0.93) | 0.87 (0.82-0.92) | 0.89 (0.84-0.95) |
|  | 4 | 1753 | 0.89 (0.83-0.94) | 0.90 (0.85-0.96) | 0.91 (0.86-0.97) | 0.90 (0.85-0.96) | 0.92 (0.87-0.98) |
|  | 5 | 1885 | 1.08(1.02-1.15) | 1.09 (1.03-1.16) | 1.09 (1.03-1.16) | 1.06 (1.00-1.13) | 1.03 (0.97-1.09) |
|  | Non-linearity p -value |  | p<0.001 | p<0.001 | p<0.001 | p<0.001 | p<0.001 |
|  |  |  | 69 | 63 | 59 | 51 | 26 |
| Fat mass (FM) |  |  |  |  |  |  |  |
| Men | 1 | 2565 | 1 (reference) | 1 (reference) | 1 (reference) | 1 (reference) | 1 (reference) |
|  | 2 | 3097 | 1.14 (1.08-1.20) | 1.14 (1.08-1.20) | 1.15 (1.09-1.21) | 1.14 (1.09-1.21) | 1.16 (1.10-1.22) |
|  | 3 | 3393 | 1.25 (1.19-1.31) | 1.24 (1.18-1.31) | 1.25 (1.19-1.32) | 1.24 (1.18-1.31) | 1.26 (1.20-1.33) |
|  | 4 | 3835 | 1.42 (1.35-1.49) | 1.40 (1.33-1.47) | 1.40 (1.33-1.48) | 1.38 (1.31-1.45) | 1.40 (1.33-1.47) |
|  | 5 | 4609 | 1.76 (1.68-1.85) | 1.71 (1.63-1.79) | 1.71 (1.62-1.79) | 1.65 (1.57-1.73) | 1.65 (1.57-1.73) |
|  | Non-linearity p-value |  | p<0.001 | 0.002 | 0.005 | 0.024 | 0.081 |
|  |  |  | 654 | 571 | 638 | 455 | 453 |
| Women | 1 | 1313 | 1 (reference) | 1 (reference) | 1 (reference) | 1 (reference) | 1 (reference) |
|  | 2 | 1653 | 1.16 (1.08-1.25) | 1.15 (1.07-1.24) | 1.16 (1.08-1.25) | 1.16 (1.08-1.24) | 1.16 (1.08-1.25) |
|  | 3 | 2037 | 1.32 (1.23-1.42) | 1.30 (1.21-1.39) | 1.31 (1.22-1.40) | 1.30 (1.21-1.39) | 1.31 (1.22-1.40) |
|  | 4 | 2357 | 1.56 (1.46-1.67) | 1.51 (1.41-1.62) | 1.51 (1.41-1.62) | 1.49 (1.39-1.59) | 1.50 (1.40-1.61) |
|  | 5 | 2925 | 2.07 (1.94-2.21) | 1.97 (1.84-2.10) | 1.95 (1.82-2.08) | 1.88 (1.76-2.01) | 1.88 (1.76-2.01) |
|  | Non-linearity p-value |  | p<0.001 | p<0.001 | 0.003 | 0.016 | 0.038 |
|  |  |  | 639 | 540 | 491 | 430 | 429 |

Test for non-linearity across quintiles conducted using LRTs with 4df (p<0.05 indicates significant departure from linearity). Adjusted hazard ratios (HR) and $95 \%$ confidence intervals (CI) obtained using Cox proportional hazard regression. Age at risk adjusted for by using age during study as the underlying timescale for Cox regression. Height adjusted for by inclusion as continuous variable for FM and by regression out of variation due to height for ASM. *Sociodemographic characteristics: Townsend index of deprivation, education. †Lifestyle factors: smoking status, alcohol intake, physical activity, oily fish intake, fruit and vegetable intake, saturated fat intake. $\ddagger$ Medical history: diabetes, cancer history, menopause (women). $\chi^{2}$ values were calculated from likelihood ratio tests to estimate the improvement in model fit.

Table S8. Mortality sequential model adjustment.

## Hazard ratios of all-cause mortality associated with appendicular skeletal muscle mass (aSMM) and fat mass (FM)

|  | Quintile | No. events | Age and height adjusted | + Sociodemographics* | + Lifestyle factors ${ }^{\dagger}$ | + Medical history ${ }^{\ddagger}$ | + Mutual adjustment for ASM or FM |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Appendicular skeletal muscle mass (aSMM) |  |  |  |  |  |  |  |
| Men | 1 | 2706 | 1 (reference) | 1 (reference) | 1 (reference) | 1 (reference) | 1 (reference) |
|  | 2 | 2013 | 0.94 (0.88-0.99) | 0.95 (0.90-1.01) | 0.97 (0.92-1.03) | 0.97 (0.92-1.03) | 0.98 (0.92-1.04) |
|  | 3 | 1673 | 0.94 (0.88-1.00) | 0.96 (0.90-1.02) | 0.99 (0.93-1.05) | 0.99 (0.93-1.05) | 0.99 (0.93-1.06) |
|  | 4 | 1378 | 0.94 (0.88-1.00) | 0.96 (0.89-1.02) | 1.00 (0.93-1.07) | 0.99 (0.93-1.06) | 1.00 (0.93-1.06) |
|  | 5 | 1312 | 1.12 (1.05-1.20) | 1.13 (1.06-1.21) | 1.19 (1.11-1.27) | 1.17 (1.09-1.25) | 1.16 (1.08-1.24) |
|  | Non-linearity p-value |  | $\mathrm{p}<0.001$ | $\mathrm{p}<0.001$ | p<0.001 | p<0.001 | 0.002 |
|  |  | $\chi^{2}$ | 33 | 30 | 34 | 29 | 25 |
| Women | 1 | 1732 | 1 (reference) | 1 (reference) | 1 (reference) | 1 (reference) | 1 (reference) |
|  | 2 | 1391 | 0.91 (0.85-0.98) | 0.92 (0.86-0.99) | 0.93 (0.86-0.99) | 0.92 (0.86-0.99) | 0.93 (0.87-1.00) |
|  | 3 | 1255 | 0.92 (0.86-0.99) | 0.93 (0.87-1.00) | 0.94 (0.87-1.01) | 0.94 (0.88-1.01) | 0.95 (0.89-1.03) |
|  | 4 | 1176 | 0.96 (0.89-1.03) | 0.97 (0.90-1.04) | 0.98 (0.91-1.06) | 0.98 (0.91-1.06) | 0.99 (0.92-1.07) |
|  | 5 | 1208 | 1.11 (1.03-1.20) | 1.11 (1.03-1.20) | 1.13 (1.05-1.22) | 1.12 (1.03-1.20) | 1.11 (1.03-1.19) |
|  | Non-linearity p-value |  | p<0.001 | $\mathrm{p}<0.001$ | $\mathrm{p}<0.001$ | $\mathrm{p}<0.001$ | 0.003 |
|  |  | $\chi^{2}$ | 31 | 28 | 29 | 26 | 21 |
| Fat mass (FM) |  |  |  |  |  |  |  |
| Men | 1 | 1573 | 1 (reference) | 1 (reference) | 1 (reference) | 1 (reference) | 1 (reference) |
|  | 2 | 1542 | 0.89 (0.83-0.95) | 0.89 (0.83-0.95) | 0.90 (0.84-0.96) | 0.89 (0.83-0.96) | 0.90 (0.84-0.96) |
|  | 3 | 1727 | 0.97 (0.91-1.04) | 0.96 (0.90-1.03) | 0.97 (0.91-1.04) | 0.96 (0.90-1.03) | 0.97 (0.90-1.04) |
|  | 4 | 1870 | 1.05 (0.98-1.12) | 1.02 (0.96-1.1) | 1.02 (0.95-1.09) | 1.00 (0.94-1.07) | 1.01 (0.94-1.08) |
|  | 5 | 2370 | 1.35 (1.27-1.44) | 1.28 (1.2-1.37) | 1.27 (1.19-1.35) | 1.22 (1.14-1.31) | 1.22 (1.14-1.31) |
|  | Non-linearity p-value |  | $\mathrm{p}<0.001$ | $\mathrm{p}<0.001$ | $\mathrm{p}<0.001$ | $\mathrm{p}<0.001$ | $\mathrm{p}<0.001$ |
|  |  | $\chi^{2}$ | 198 | 145 | 126 | 101 | 98 |
| Women | 1 | 1126 | 1 (reference) | 1 (reference) | 1 (reference) | 1 (reference) | 1 (reference) |
|  | 2 | 1228 | 0.97 (0.90-1.06) | 0.97 (0.89-1.05) | 0.98 (0.90-1.06) | 0.98 (0.90-1.06) | 0.99 (0.91-1.07) |
|  | 3 | 1321 | 0.95 (0.87-1.03) | 0.94 (0.86-1.01) | 0.94 (0.86-1.02) | 0.93 (0.86-1.01) | 0.94 (0.87-1.02) |
|  | 4 | 1414 | 1.02 (0.94-1.10) | 0.99 (0.92-1.08) | 0.98 (0.91-1.07) | 0.98 (0.90-1.06) | 0.99 (0.92-1.08) |
|  | 5 | 1673 | 1.27 (1.18-1.37) | 1.21 (1.12-1.31) | 1.18 (1.09-1.28) | 1.16 (1.07-1.26) | 1.17 (1.08-1.26) |
|  | Non-linearity p-value |  | p<0.001 | $\mathrm{p}<0.001$ | $p<0.001$ | $\mathrm{p}<0.001$ | p<0.001 |
|  |  | $\chi^{2}$ | 82 | 62 | 47 | 41 | 37 |

Test for non-linearity across quintiles conducted using LRTs with 4 df ( $\mathrm{p}<0.05$ indicates significant departure from linearity). Adjusted hazard ratios $(H R)$ and $95 \%$ confidence intervals (CI) obtained using Cox proportional hazard regression. Age at risk adjusted for by using age during study as the underlying timescale for Cox regression. Height adjusted for by inclusion as continuous variable for FM and by regression out of variation due to height for ASM. *Sociodemographic characteristics: Townsend index of deprivation, education. †Lifestyle factors: smoking status, alcohol intake, physical activity, oily fish intake, fruit and vegetable intake, saturated fat intake. $\ddagger$ Medical history: diabetes, cancer history, menopause (women). $\chi^{2}$ values were calculated from likelihood ratio tests to estimate the improvement in model fit.

Table S9. CVD sensitivity analyses.
Hazard ratios of CVD associated with appendicular skeletal muscle mass (aSMM) and fat mass
(FM)

| Quintile | Remove first 2 years of follow up |  | Remove outliers |  | Remove participants with$\text { BMI > } 35$ |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | No. events | HR (95\% CI) | No. events | HR (95\% CI) | No. events | HR (95\% CI) |
| Appendicular skeletal muscle mass (aSMM) |  |  |  |  |  |  |
| Men 1 | 3804 | 1 (reference) | 4316 | 1 (reference) | 4283 | 1 (reference) |
| 2 | 3250 | 1.06 (1.01-1.11) | 3715 | 1.02 (0.97-1.06) | 3614 | 1.03 (0.98-1.08) |
| 3 | 2939 | 1.12 (1.06-1.17) | 3385 | 1.07 (1.02-1.12) | 3198 | 1.07 (1.02-1.12) |
| 4 | 2666 | 1.19 (1.13-1.25) | 3055 | $1.10(1.05-1.16)$ | 2800 | 1.11 (1.06-1.17) |
| 5 | 2597 | 1.37 (1.30-1.44) | 2819 | 1.23 (1.17-1.29) | 2448 | 1.25 (1.18-1.31) |
| Non-linearity p-value |  | 0.038 |  | 0.057 |  | 0.046 |
| Women 1 | 2327 | 1 (reference) | 2582 | 1 (reference) | 2505 | 1 (reference) |
| 2 | 1947 | 0.97 (0.91-1.03) | 2182 | 0.95 (0.90-1.01) | 2075 | 0.96 (0.90-1.01) |
| 3 | 1667 | 0.93 (0.87-0.99) | 1848 | 0.89 (0.84-0.95) | 1699 | 0.88 (0.83-0.94) |
| 4 | 1570 | 0.96 (0.90-1.02) | 1753 | 0.92 (0.87-0.98) | 1555 | 0.92 (0.86-0.98) |
| 5 | 1681 | 1.08 (1.01-1.15) | 1682 | 1.01 (0.95-1.07) | 1375 | 1.01 (0.94-1.08) |
| Non-linearity p-value |  | $\mathrm{p}<0.001$ |  | $\mathrm{p}<0.001$ |  | $\mathrm{p}<0.001$ |
| Fat Mass (FM) |  |  |  |  |  |  |
| Men $\begin{array}{ll}1 \\ & 2 \\ & 3 \\ & 4 \\ & 5\end{array}$ | 2293 | 1 (reference) | 2565 | 1 (reference) | 2565 | 1 (reference) |
|  | 2701 | 1.13 (1.07-1.19) | 3097 | 1.16 (1.1-1.22) | 3097 | 1.16 (1.10-1.22) |
|  | 2907 | 1.20 (1.14-1.27) | 3393 | 1.26 (1.19-1.32) | 3393 | 1.26 (1.19-1.32) |
|  | 3341 | 1.36 (1.28-1.43) | 3835 | 1.40 (1.33-1.47) | 3828 | 1.40 (1.33-1.47) |
|  | 4014 | 1.60 (1.51-1.68) | 3994 | 1.62 (1.54-1.71) | 3460 | 1.61 (1.53-1.70) |
| $\begin{array}{ll}\text { Women } & \\ & 1 \\ & 2 \\ & 3 \\ & 4 \\ & 5\end{array}$ |  | 0.038 |  | 0.211 |  | 0.286 |
|  | 1173 | 1 (reference) | 1313 | 1 (reference) | 1313 | 1 (reference) |
|  | 1487 | 1.16 (1.07-1.25) | 1653 | 1.16 (1.08-1.25) | 1652 | 1.15 (1.07-1.24) |
|  | 1806 | 1.27 (1.18-1.37) | 2037 | 1.30 (1.21-1.40) | 2037 | 1.29 (1.21-1.39) |
|  | 2118 | 1.48 (1.37-1.59) | 2357 | 1.50 (1.40-1.60) | 2351 | 1.49 (1.39-1.59) |
|  | 2608 | 1.84 (1.71-1.98) | 2493 | 1.82 (1.70-1.96) | 1856 | 1.76 (1.63-1.89) |
| Non-linearity p-value |  | 0.026 |  | 0.230 |  | 0.681 |

Test for non-linearity across quintiles conducted using LRTs with 4 df ( $\mathrm{p}<0.05$ indicates significant departure from linearity). Adjusted hazard ratios $(\mathrm{HR})$ and $95 \%$ confidence intervals (CI) obtained using Cox proportional hazard regression. Adjusted for age at risk, Townsend index of deprivation, education, smoking status, alcohol intake, physical activity, oily fish intake, fruit and vegetable intake, saturated fat intake, diabetes, cancer history, menopause (women), and mutually adjusted for FM and ASM. Height is adjusted for by inclusion as continuous variable for FM and by regression out of variation due to height for ASM. For the analysis without outliers, the top and bottom $25 \%$ of values were removed.

## Table S10. CVD mediation analyses.

## Hazard ratios of CVD associated with appendicular skeletal muscle mass (aSMM) and fat mass (FM)

|  | Quintile | No. events | Adjust for BMI | Adjust for hypertension | Adjust for cholesterol |
| :---: | :---: | :---: | :---: | :---: | :---: |
| Appendicular skeletal muscle mass (aSMM) |  |  |  |  |  |
| Men | 1 | 2861 | 1 (reference) | 1 (reference) | 1 (reference) |
|  | 2 | 2457 | 0.96 (0.92-1.01) | 1.02 (0.98-1.07) | 1.03 (0.99-1.08) |
|  | 3 | 2209 | 0.97 (0.93-1.02) | 1.06 (1.02-1.11) | 1.08 (1.03-1.14) |
|  | 4 | 2004 | 0.98 (0.93-1.03) | 1.10 (1.05-1.16) | 1.14 (1.09-1.20) |
|  | 5 | 1907 | 1.04 (0.99-1.10) | 1.23 (1.17-1.29) | 1.28 (1.21-1.35) |
|  | Non-linea | ity p-value | 0.015 | 0.031 | 0.123 |
|  | Main mod |  | 106 | 87 | 74 |
|  | Adjusted | model $\chi^{2}$ | 12 | 79 | 92 |
|  | \% Change |  | 87\% | 9\% | 20\% |
| Women | 1 | 2617 | 1 (reference) | 1 (reference) | 1 (reference) |
|  | 2 | 2182 | 0.92 (0.87-0.97) | 0.95 (0.90-1.01) | 0.93 (0.88-0.99) |
|  | 3 | 1848 | 0.84 (0.79-0.89) | 0.90 (0.84-0.95) | 0.87 (0.82-0.93) |
|  | 4 | 1753 | 0.84 (0.79-0.90) | 0.92 (0.87-0.98) | 0.94 (0.88-1.00) |
|  | 5 | 1885 | 0.90 (0.85-0.96) | 1.02 (0.96-1.09) | 1.04 (0.98-1.11) |
|  | Non-linea | ity p-value | $\mathrm{p}<0.001$ | $\mathrm{p}<0.001$ | $\mathrm{p}<0.001$ |
|  | Main mod |  | 51 | 26 | 29 |
|  | Adjusted | model $\chi^{2}$ | 43 | 23 | 30 |
|  | \% Change |  | 17\% | 12\% | 3\% |
| Fat mass (FM) |  |  |  |  |  |
| Men | 1 | 2565 | 1 (reference) | 1 (reference) | 1 (reference) |
|  | 2 | 3097 | 1.05 (0.99-1.10) | 1.12 (1.06-1.18) | 1.09 (1.03-1.15) |
|  | 3 | 3393 | 1.06 (1.00-1.12) | 1.19 (1.13-1.25) | 1.15 (1.09-1.22) |
|  | 4 | 3835 | 1.09 (1.03-1.16) | 1.29 (1.22-1.36) | 1.27 (1.20-1.34) |
|  | 5 | 4609 | 1.10 (1.01-1.19) | 1.48 (1.41-1.56) | 1.49 (1.41-1.58) |
|  | Non-linea | ity p -value | 0.726 | 0.102 | 0.009 |
|  | Main mo | $\text { el } \chi^{2}$ | 455 | 453 | 398 |
|  | Adjusted | model $\chi^{2}$ | 8 | 274 | 252 |
|  | \% Change |  | 98\% | 40\% | 37\% |
| Women | 1 | 1313 | 1 (reference) | 1 (reference) | 1 (reference) |
|  | 2 | 1653 | 1.07 (0.99-1.16) | 1.12 (1.05-1.21) | 1.13 (1.05-1.23) |
|  | 3 | 2037 | 1.13 (1.05-1.22) | 1.23 (1.15-1.32) | 1.25 (1.16-1.35) |
|  | 4 | 2357 | 1.21 (1.11-1.32) | 1.37 (1.28-1.47) | 1.40 (1.30-1.52) |
|  | 5 | 2925 | 1.29 (1.15-1.46) | 1.64 (1.54-1.76) | 1.73 (1.60-1.87) |
|  | Non-linea | ity p-value | 0.991 | 0.119 | 0.047 |
|  | Main mo | el $\chi^{2}$ | 430 | 429 | 398 |
|  | Adjusted | model $\chi^{2}$ | 20 | 258 | 259 |
|  | \% Change |  | 95\% | 40\% | 35\% |

Adjusted hazard ratios (HR) and 95\% confidence intervals (CI) obtained using Cox proportional hazard regression. Adjusted for age at risk, Townsend index of deprivation, education, smoking status, alcohol intake, physical activity, oily fish intake, fruit and vegetable intake, saturated fat intake, diabetes, cancer history, menopause (women), and mutually adjusted for FM and ASM ASM (except for the model adjusted for BMI). Height is adjusted for by inclusion as continuous variable for FM and by regression out of variation due to height for ASM. Test for non-linearity across quintiles conducted using LRTs with 4 df ( $\mathrm{p}<0.05$ indicates significant departure from linearity). The $\chi^{2}$ values were calculated from likelihood ratio tests to estimate the improvement in model fit.

Table S11. Mortality sensitivity analyses.

## Hazard ratios of all-cause mortality associated with appendicular skeletal muscle mass (aSMM) and fat mass (FM)



Adjusted hazard ratios (HR) and 95\% confidence intervals (CI) obtained using Cox proportional hazard regression. Adjusted for age at risk, Townsend index of deprivation, education, smoking status, alcohol intake, physical activity, oily fish intake, fruit and vegetable intake, saturated fat intake, diabetes, cancer history, menopause (women), and mutually adjusted for FM and ASM. Height is adjusted for by inclusion as continuous variable for FM and by regression out of variation due to height for ASM. Test for non-linearity across quintiles conducted using LRTs with 4 df ( $\mathrm{p}<0.05$ indicates significant departure from linearity). For the analysis without outliers, the top and bottom $25 \%$ of values were removed.

Table S12. Mortality mediation analyses.

## Hazard ratios of all-cause mortality associated with appendicular skeletal muscle mass (aSMM) and fat mass (FM)

|  | Quintile | No. events | Adjust for BMI | Adjust for hypertension | Adjust for cholesterol |
| :---: | :---: | :---: | :---: | :---: | :---: |
| Appendicular skeletal muscle mass (aSMM) |  |  |  |  |  |
| Men | 1 | 2706 | 1 (reference) | 1 (reference) | 1 (reference) |
|  | 2 | 2013 | 0.95 (0.89-1.01) | 0.98 (0.92-1.04) | 0.98 (0.92-1.05) |
|  | 3 | 1673 | 0.95 (0.89-1.01) | 0.99 (0.93-1.05) | 0.97 (0.91-1.04) |
|  | 4 | 1378 | 0.93 (0.87-1.00) | 0.99 (0.93-1.06) | 0.96 (0.89-1.03) |
|  | 5 | 1312 | 1.06 (0.98-1.14) | 1.15 (1.07-1.23) | 1.13 (1.05-1.22) |
|  | Non-linea | ity p -value | 0.001 | 0.002 | 0.001 |
|  | Main mo | el $\chi^{2}$ | 29 | 25 | 21 |
|  | Adjusted | model $\chi^{2}$ | 16 | 24 | 20 |
|  | \% Change |  | 45\% | 4\% | 5\% |
| Women | 1 | 1732 | 1 (reference) | 1 (reference) | 1 (reference) |
|  | 2 | 1391 | 0.92 (0.86-0.99) | 0.93 (0.87-1.00) | 0.93 (0.86-1.00) |
|  | 3 | 1255 | 0.93 (0.87-1.00) | 0.95 (0.89-1.03) | 0.95 (0.88-1.03) |
|  | 4 | 1176 | 0.96 (0.89-1.03) | 0.99 (0.92-1.07) | 0.99 (0.91-1.07) |
|  | 5 | 1208 | 1.06 (0.98-1.14) | 1.10 (1.02-1.19) | 1.10 (1.02-1.20) |
|  | Non-linea | ity p -value | 0.003 | 0.003 | 0.004 |
|  | Main mo |  | 26 | 21 | 21 |
|  | Adjusted | model $\chi^{2}$ | 16 | 20 | 19 |
|  | \% Change |  | 38\% | 5\% | 9\% |
| Fat mass (FM) |  |  |  |  |  |
| Men | 1 | 1573 | 1 (reference) | 1 (reference) | 1 (reference) |
|  | 2 | 1542 | 0.86 (0.79-0.92) | 0.88 (0.82-0.95) | 0.90 (0.84-0.97) |
|  | 3 | 1727 | 0.90 (0.83-0.97) | 0.94 (0.88-1.01) | 0.98 (0.91-1.06) |
|  | 4 | 1870 | 0.90 (0.82-0.98) | 0.97 (0.91-1.04) | 1.01 (0.94-1.09) |
|  | 5 | 2370 | 1.01 (0.90-1.13) | 1.17 (1.09-1.25) | 1.24 (1.15-1.34) |
|  | Non-linea | ity p -value | p<0.001 | p<0.001 | p<0.001 |
|  | Main mo | $\chi^{2}$ | 101 | 98 | $88$ |
|  | Adjusted | model $\chi^{2}$ | 36 | 81 | 87 |
|  | \% Change |  | 64\% | 17\% | 1\% |
| Women | 1 | 1126 | 1 (reference) | 1 (reference) | 1 (reference) |
|  | 2 | 1228 | 0.91 (0.84-0.99) | 0.98 (0.90-1.06) | 0.97 (0.89-1.06) |
|  | 3 | 1321 | 0.82 (0.75-0.90) | 0.93 (0.86-1.01) | 0.97 (0.89-1.06) |
|  | 4 | 1414 | 0.81 (0.73-0.90) | 0.97 (0.90-1.06) | 1.00 (0.91-1.09) |
|  | 5 | 1673 | 0.83 (0.71-0.96) | 1.13 (1.05-1.23) | 1.16 (1.06-1.27) |
|  | Non-linea | ity p-value | 0.033 | p<0.001 | 0.003 |
|  | Main mo | el $\chi^{2}$ | 41 | 37 | 31 |
|  | Adjusted | model $\chi^{2}$ | 22 | 31 | 26 |
|  | \% Change |  | 46\% | 16\% | 16\% |

Adjusted hazard ratios (HR) and 95\% confidence intervals (CI) obtained using Cox proportional hazard regression. Adjusted for age at risk, Townsend index of deprivation, education, smoking status, alcohol intake, physical activity, oily fish intake, fruit and vegetable intake, saturated fat intake, diabetes, cancer history, menopause (women), and mutually adjusted for FM and ASM (except for the model adjusted for BMI). Height is adjusted for by inclusion as continuous variable for FM and by regression out of variation due to height for ASM. Test for non-linearity across quintiles conducted using LRTs with 4 df ( $p<0.05$ indicates significant departure from linearity). The $\chi^{2}$ values were calculated from likelihood ratio tests to estimate the improvement in model fi

Figure S1. Exclusions and selection of the study population included in the analysis from the UK Biobank.

*Prior CVD: follows the same definition and codes from HES reported in the manuscript as well as self-reported from verbal interviews and touchscreen questionnaire including instances of heart attack/myocardial infarction, heart failure, angina, heart failure, stroke, transient ischaemic attack, subdural haemorrhage, aortic aneurysm rupture, cerebral aneurysm, peripheral vascular disease, leg claudication, arterial embolism, .
${ }^{\dagger}$ Respiratory diseases: chronic obstructive pulmonary disease, emphysema, chronic bronchitis, bronchiectasis (occurred anytime)
${ }^{\ddagger}$ Endocrine diseases: Cushing’s syndrome, hyperthyroidism (occurred anytime)
${ }^{\S}$ Musculoskeletal diseases: motor neurone disease, osteoporosis, rheumatoid arthritis, fractures (occurred anytime) "HIV, cirrhosis or other disease: pancreatitis, encephalitis, meningitis, intracranial abscess, empyema, Stevens Johnson Syndrome and carcinoid syndrome/tumour (occurred in the last year); liver cirrhosis, HIV/AIDS (occurred anytime)

Figure S2. Splines.
Adjusted hazard ratios (HRs) of incident cardiovascular disease (CVD) and all-cause mortality associated with appendicular skeletal muscle mass (aSMM), whole body fat mass (FM) and body mass index (BMI) estimated using restricted cubic splines










Figure S3. Associations with CVD subtypes.
Adjusted hazard ratios for the association between cardiovascular disease (CVD) subtypes (coronary heart disease, stroke and congestive heart failure) with i) appendicular skeletal muscle mass (aSMM) and ii) fat mass (FM)
i) Appendicular skeletal muscle mass
ii) Fat mass

i A, adjusted hazard ratios for the association between CVD subtypes with aSSM in men. i B, adjusted hazard ratios for the association between CVD subtypes with aSSM in women. ii $A$, adjusted hazard ratios for the association between CVD subtypes with FM in men. ii B, adjusted hazard ratios for the association between CVD subtypes with FM in women
Adjusted hazard ratios (HR) and 95\% group-specific confidence intervals (CI) obtained using floated absolute risk method of Cox proportional hazard regression. Adjusted for age at risk, Townsend index of deprivation, education, smoking status, alcohol intake, physical activity, oily fish intake, fruit and vegetable intake, saturated fat intake, diabetes, cancer history, menopause (women), and mutually adjusted for FM and ASM. Height is adjusted for by inclusion as continuous variable for FM and by regression out of variation due to height for ASM. HRs are plotted at the mean of the resurvey values for the baseline-defined quintiles ("usual" values) to correct for measurement error.


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