The association of grip strength from midlife onwards with all-cause and cause-specific mortality over 17 years of follow-up in the Tromsø Study

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ABSTRACT

Background Grip strength has consistently been found to predict all-cause mortality rates. However, few studies have examined cause-specific mortality or tested age differences in these associations.

Methods In 1994, grip strength was measured in the population-based Tromsø Study, covering the ages 50–80 years (N=6850). Grip strength was categorized into fifths, and as z-scores. In this cohort study, models with all-cause mortality and deaths from specific causes as the outcome were performed, stratified by sex and age using Cox regression, adjusting for lifestyle-related and health-related factors.

Results During 17 years of follow-up, 2338 participants died. A 1 SD reduction in grip strength was associated with HR=1.17 (95% CI 1.12 to 1.22) for all-cause mortality in a model adjusted for age, gender and length of follow-up, there was some evidence to suggest that associations between grip strength and all-cause mortality may be higher in older age groups, stronger in males compared with females, and in studies with follow-up of <10 years, however, there was limited statistical power to formally test these differences. Since the publication of this review, subsequent studies have continued to find that weaker grip strength is associated with increased all-cause mortality rates in a range of settings.10–13

Most studies of grip strength in middle and old age have investigated associations with all-cause mortality, and studies that have examined cause-specific mortality are not as numerous.14–19 Furthermore, often such studies are only powered to investigate broad groups of causes of death such as cardiovascular disease (CVD) and cancer, and have not been able to look at additional causes such as ischaemic heart disease (IHD), respiratory diseases and external causes. Recently, findings from a study which had measures of grip strength in almost 140 000 people aged 35–70 years from 17 countries were reported. This study found that weak grip strength was related to increased rates of all-cause mortality, cardiovascular mortality, myocardial infarction and stroke, while no association was found for cancer mortality.14 These findings of an association between grip strength and cardiovascular mortality and no association with cancer mortality are consistent with those from a study of 1 million Swedish male adolescents aged 16–19 years followed over a period of 24 years.15 These results, of an association between grip strength and cardiovascular mortality and no association for cancer, suggest a potential mechanistic explanation related to the cardiovascular system,7,14 which should be further explored.

Larger-scale studies including objectively measured physical performance and strength are lacking in Norway, with one exception; the Tromsø Study, a longitudinal study with six follow-ups in the period 1974–2008. Grip strength was objectively measured in 1994, in a sample of men and women covering a wide range of ages.20 Our aim was to investigate the associations of grip strength with all-cause and cause-specific mortality (CVD, stroke, IHD, cancer, respiratory diseases and external causes) in men and women in this large general Norwegian population. Low grip strength in adolescents has been found to be associated with increased suicide risk in young adulthood,19 but to the best of our knowledge, our paper is the first to investigate the association between grip strength...
and the broader group of external causes of death. Another novel aspect of our study was our aim to formally test whether any associations varied by age and sex at strength assessment.

METHODS

Study population

The Tromsø Study is a longitudinal population-based multipurpose study focusing on lifestyle-related diseases, initiated in 1974 with surveys, inviting earlier participants as well as recruiting new participants, repeated in 1979, 1986, 1994, 2001 and 2008. This study uses baseline data from the survey initiated in 1994. At this time, all those men and women in the municipality aged 25 years and above (n = 37,558) were invited to participate in phase I of the survey. All men aged 55–74 years, all women aged 50–74 years and 5–10% random samples of other birth cohorts aged 25–84 years were preselected to undergo a more extensive phase II examination (10,542 participants) which included grip strength measurements. The attendance rate was 74% in men and 77% in women. In the current analysis, 6,850 participants (58% women) aged 50–80 years at the time of screening and with valid grip strength were included. Participants outside this age range were excluded as there were too few deaths among the 925 individuals below 50 years (n = 41), and there were few participants above 80 years (n = 14). The Regional Committee of Research Ethics recommended and the Norwegian Data Inspectorate approved the study.

Assessment of grip strength

Grip strength of the non-dominant hand was measured in kilopascal (kPa) in the Tromsø Study, wave four (1994–1995) using a Martin vigorimeter. Each participant was allowed two attempts, and the highest score registered was recorded and used in analyses.

Ascertainment of mortality

Data on each participant were linked to death certificate data from the Norwegian Cause of Death Registry, and participants were followed from 1 January 1996 to 31 December 2012, death or emigration, whichever occurred first. Thus, follow-up started ~2 years after the time point of the health examination, which was done to minimise the effect of reverse causation from disease at baseline. Only 51 participants (0.7%) left the study (died or emigrated) during the first 2 years, before 1 January 1996. Participants had mean follow-up time of 14.4 years, and maximum 17.0 years, and mean age at the end of follow-up was 77.5 years (minimum 51.7 and maximum 96.4 years). Causes of death were based on underlying cause of death at the death recorded by a medical doctor on the certificate and categorised using the International Classification of Diseases (ICD) V.10 and grouped as: cancer (C00–C97), CVD (I00–I99), IHD (I20–I25), stroke (I60–I69), respiratory causes (J00–J99) and external causes (V01–Y89) (such as accidents, falls, injuries, poisoning, assault and other adverse events).

Covariates

Covariates known to be associated with grip strength and mortality were selected a priori for inclusion as possible confounders. All these covariates were taken from the baseline study in Tromsø 4 in 1994–1995. Height and weight were measured in light clothing without shoes. Body mass index (BMI) was calculated as weight in kilograms divided by the square of height in metres. Blood pressure was assessed three times after 2 min seated rest with 1 min interval with an oscillometric digital automated device (Dinamap Vital Signs Monitor; Critikon Inc, Tampa, Florida, USA). The mean systolic blood pressure of reading two and three was used. From non-fasting blood samples, total cholesterol level (mmol/L) and triglycerides (mmol/L) were analysed at the Department of Clinical Chemistry, University Hospital of North Norway. The participants completed two self-administered questionnaires, one before entering the study, and the other during the study. From these questionnaires, data on health status were derived (poor, not so good, good, very good), a history of heart attack, angina, stroke, asthma or diabetes. Participants also provided information on dietary physical activity (not sweating or out of breath) and hard physical activity (sweating and/or out of breath) during a typical week in the last year, and for each of the two variables participants were categorised in four groups based on hours per week in activity (none, less than one, one to two, three or more). Participants were asked if they used blood pressure-lowering drugs (currently, previously, but not now, never). Participants were dichotomised as current cigarette smokers or non-current cigarette smokers. Education was stratified into five groups: primary (7–10 years), technical school/middle school, high school diploma (3–4 years), college/university <4 years and college/university 4 years or more.

Statistical methods

To allow investigation of variation in the association between grip strength and mortality by age, participants were assigned to one of the three age groups based on age at the time of grip strength assessment: 50–59, 60–69 and 70–80 years. Sex-specific quintiles for grip strength in each of these age groups were created. Age-specific and gender-specific grip strength z-scores were also created, which had a mean of 0 and SD of 1.

In a first set of analyses, Cox regression was used to assess the associations between fifths of grip strength and hazards of mortality, with the strongest fifth used as the reference category. In a second set of analyses, the associations were assessed using age-specific and gender-specific grip strength z-scores. In a third set of analyses, the associations were modelled using grip strength z-scores as natural cubic splines with three knots to allow for a more flexible analysis of grip strength on the continuous scale. This approach allowed us to investigate deviation from linearity, and test for evidence of thresholds. All three sets of analyses examined associations of grip strength with all-cause and cause-specific mortality rates. In all models, attained age was the time scale, and thereby models were adjusted for age. In addition, models were first adjusted for sex and body size (height and BMI) and then for all health-related and lifestyle-related factors. To investigate differences in associations by age group, an age group by grip strength interaction term was included in a subsequent stage of analyses. In the gender-adjusted model, gender was treated as a covariate. The proportional hazards assumption was checked using Schoenfeld residuals, and on the basis of visual inspection of the estimated hazard function. No severe violations of proportionality were observed. Stata SE V.14.0 was used for the analyses.

RESULTS

Mean value for the absolute grip strength for the total study population was 0.78 kPa with SD 0.21, and 0.84 kPa (SD 0.21) in men and 0.73 kPa (SD 0.20) in women. By age group, the mean values in men were: 0.94 kPa (SD 0.21) (50–59 years), 0.82 kPa (SD 0.19) (60–69 years) and 0.70 kPa (SD 0.17) (70–80 years). Corresponding values in women were: 0.78 kPa (0.20), 0.70 kPa (0.18) and 0.63 kPa (0.15).
Table 1 displays the baseline characteristics of the participants stratified by fifths of grip strength by gender. Those in the higher grip strength categories had higher BMI, were younger, reported better overall health, had less history of heart disease and asthma, smoked less, were more physically active and had higher education than those in the lower grip strength categories (all p<0.05). Overall, there were 2338 deaths among the 6850 participants during follow-up, of which 35% of deaths were caused by CVD (8% of which were stroke and 17% IHD), 35% by cancer, 9% by respiratory diseases, 3% by external causes (table 2) and 18% other causes. Regression models in tables 3 and 4 and in figure 1 were run on a slightly smaller sample with no missing data on covariates (N=6601, with 2225 deaths). Of these deaths, 776 were due to CVD (of which 183 were due to stroke and 366 were due to IHD), 774 were due to cancer, 200 were due to respiratory diseases, 68 were due to external causes and 407 were due to other causes. Minimally adjusted results on the maximum available samples produced very similar results (results not shown).

All-cause mortality
Grip strength was inversely associated with all-cause mortality rates in men and women (table 3), with no grip strength by gender interaction found when grip strength was modelled linearly (p=0.91); 1 SD lower grip strength was associated with an HR of all-cause mortality of 1.17 (95% CI 1.12 to 1.22) in a model adjusted for age, gender and body size, which was only partially attenuated in the fully adjusted model (HR=1.12, 95% CI 1.07 to 1.17). The spline models suggested a linear association between grip and all-cause mortality rates in men, while in women, there was some evidence that only grip strength below the mean was associated with increased mortality rates (figure 1). Weaker grip strength was associated with increased hazards of mortality in all age groups (overall p<0.05; table 4), with no evidence of a grip strength by age interaction (p=0.69; table 4). Results were similar using grip in fifths rather than z-scores (see online supplementary appendix table A1).

Cardiovascular mortality
Weaker grip strength was associated with increased CVD mortality rates (per 1 SD lower grip strength HR=1.21, 95% CI 1.12 to 1.30) in a model adjusted for age, sex and body size, and this was not completely attenuated in the fully adjusted model (HR=1.14, 95% CI 1.06 to 1.22) (table 4). The association between grip strength and CVD mortality was similar in men and women (p=0.98). Furthermore, the association between grip strength and CVD mortality was similar across age groups (grip strength by age interaction, p=0.84) (table 4). Results were similar using grip in fifths rather than z-scores (see online supplementary appendix table A1). The spline models indicated linear associations between grip strength and CVD mortality (figure 1). Similar associations of grip strength with stroke and IHD mortality were found (table 3).

Cancer mortality
Weaker grip strength was associated with increased cancer mortality in the minimally adjusted model (per 1 SD lower grip strength HR=1.07, 95% CI 1.00 to 1.16), but this was fully attenuated in the fully adjusted model (HR=1.05, 95% CI 0.97, 1.13). While there was some evidence in basic models that grip strength was associated with cancer mortality in men (HR=1.11, 95% CI 1.00 to 1.29) but not in women (HR=1.04, 95% CI 0.93 to 1.15), the grip strength by gender interaction term was not significant (p=0.46). However, after adjustment for health-related and lifestyle-related factors, the grip strength–cancer association was not significant in either sex (table 3, figure 1). Furthermore, there was evidence that the association between grip strength and cancer mortality was only significant in the youngest age group (ie, 50–59 years; HR=1.15, 95% CI 1.00 to 1.32), while in the older groups, the association did not reach statistical significance (table 4). However, the interaction between grip strength and age group was non-significant (p=0.29; table 4). Results were similar using grip in fifths rather than z-scores (see online supplementary appendix table A1).

Respiratory and external causes of mortality
Weaker grip strength was significantly associated with increased respiratory disease mortality in the minimally adjusted model (per 1 SD lower grip strength HR=1.35, 95% CI 1.16 to 1.57) (table 3), and the HR did not differ between men and women (p=0.69). The association was robust to adjustment for a range of health-related factors (HR=1.23, 95% CI 1.06 to 1.44). For external causes of death, a 1 SD lower grip strength was associated with HR=1.44 (95% CI 1.13 to 1.84) in the minimally adjusted model, and there was no significant grip strength by gender interaction (p=0.79). The association was robust to further adjustment (HR=1.37, 95% CI 1.06 to 1.76). Numbers of deaths due to respiratory diseases and external factors were too small for age-specific analyses of these causes of death. See figure 1 for a graphical illustration of the associations of grip strength with deaths from respiratory diseases and external causes.

Results for all-cause mortality, CVD mortality and cancer mortality using grip in fifths are reported in online supplementary appendix table A1.

There were 162 participants reporting a history of stroke (2.4%), and as a sensitivity analysis, we performed an additional set of analyses in which these participants were excluded. This sensitivity analysis provided results similar to those described above.

DISCUSSION
In this large population-based study of home-dwelling Norwegians aged 50–80 years followed over 17 years including 2338 deaths, we found weaker grip strength to be associated with significantly higher rates of all-cause mortality in a stepwise fashion. This association was similar across age groups and in men and women. These associations were not fully explained by adjustment for a range of lifestyle-related and health-related factors. Results for deaths due to CVD, respiratory disease and external causes resembled those for all-cause mortality, while cancer mortality was only weakly associated with grip strength in men, and this association was not significant in fully adjusted analyses.

The finding of an association between weaker grip strength and higher rates of all-cause mortality is in line with findings from previous reports.9–11 Our finding of similar associations between grip strength and all-cause mortality across age groups is in line with the few previous reports which have been able to stratify analyses by age.14–16 Our finding of an equally strong inverse association between grip strength and all-cause mortality in men and women is also consistent with findings in other studies.14 17

Analyses of grip strength and cause-specific mortality are less numerous.14–19 Consistent with our findings, those six studies which have explored cause-specific mortality previously have found stronger grip to be associated with lower CVD mortality rates,14–19 and those four studies which included men and...
The two studies that had investigated stroke mortality found similarly to us that this association was similar in both genders. The consistency across studies of the inverse association between grip strength and CVD mortality suggests potential mechanism acting on the cardiovascular pathway. Increased grip strength has been found to be associated with a more favourable cardiovascular risk factor profile, but the strong association found in our study was only weakly attenuated after adjustment for a wide range of CVD risk factors, such as smoking, blood pressure, and body mass index. The consistency across studies of the inverse association between grip strength and CVD mortality suggests potential mechanism acting on the cardiovascular pathway. Increased grip strength has been found to be associated with a more favourable cardiovascular risk factor profile, but the strong association found in our study was only weakly attenuated after adjustment for a wide range of CVD risk factors, such as smoking, blood pressure, and body mass index.
risk factors. Another important finding in support of the cardiovascular pathway is that grip strength has been found to be an equally important prognostic factor as systolic blood pressure regarding CVD mortality risk. In our study, the association between grip strength and cancer mortality was weaker than for CVD, and restricted to men. Results for grip strength and cancer mortality are not consistent in the literature. In line with our findings, Sasaki et al.15 and Gale et al.17 found stronger grip to be associated with lower cancer mortality in men only. Another study of US men also found stronger grip to be associated with lower cancer mortality risk.18 The other two studies reported no significant association,16 19 and Leong et al.14 reported the unexpected finding of a protective effect of low grip strength on cancer mortality. Potential explanations for the inconsistent results regarding cancer mortality might be due to power issues and differences in study populations.

### Table 2 Number of deaths and mortality rate (per 1000 person-years) by cause of death, gender and age group

<table>
<thead>
<tr>
<th>Cause of death (number of deaths)</th>
<th>Men and women (N=6601)</th>
<th>Men (N=2744)</th>
<th>Women (N=3857)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total mortality</td>
<td>2225</td>
<td>1189</td>
<td>1036</td>
</tr>
<tr>
<td>Cardiovascular</td>
<td>776</td>
<td>423</td>
<td>353</td>
</tr>
<tr>
<td>Stroke</td>
<td>183</td>
<td>81</td>
<td>102</td>
</tr>
<tr>
<td>IHD</td>
<td>366</td>
<td>228</td>
<td>138</td>
</tr>
<tr>
<td>Cancer</td>
<td>774</td>
<td>412</td>
<td>326</td>
</tr>
<tr>
<td>Respiratory</td>
<td>200</td>
<td>115</td>
<td>75</td>
</tr>
<tr>
<td>External</td>
<td>68</td>
<td>39</td>
<td>29</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Mortality rate</th>
<th>Total mortality</th>
<th>Cardiovascular</th>
<th>Stroke</th>
<th>IHD</th>
<th>Cancer</th>
<th>Respiratory</th>
<th>External</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total mortality</td>
<td>23.3</td>
<td>8.1</td>
<td>1.9</td>
<td>3.8</td>
<td>8.1</td>
<td>2.1</td>
<td>0.7</td>
</tr>
<tr>
<td>Cardiovascular</td>
<td>1.23</td>
<td>1.11</td>
<td>1.08</td>
<td>1.06</td>
<td>1.11</td>
<td>1.13</td>
<td>1.0</td>
</tr>
<tr>
<td>Stroke</td>
<td>1.26</td>
<td>1.22</td>
<td>1.2</td>
<td>1.8</td>
<td>1.35</td>
<td>1.35</td>
<td>1.44</td>
</tr>
<tr>
<td>IHD</td>
<td>1.35</td>
<td>1.22</td>
<td>1.07</td>
<td>1.0</td>
<td>1.24</td>
<td>1.19</td>
<td>1.44</td>
</tr>
<tr>
<td>Cancer</td>
<td>1.35</td>
<td>1.22</td>
<td>1.11</td>
<td>1.1</td>
<td>1.35</td>
<td>1.35</td>
<td>1.35</td>
</tr>
<tr>
<td>Respiratory</td>
<td>1.35</td>
<td>1.22</td>
<td>1.11</td>
<td>1.1</td>
<td>1.35</td>
<td>1.35</td>
<td>1.35</td>
</tr>
<tr>
<td>External</td>
<td>1.35</td>
<td>1.22</td>
<td>1.11</td>
<td>1.1</td>
<td>1.35</td>
<td>1.35</td>
<td>1.35</td>
</tr>
</tbody>
</table>

Deaths due to external causes are only presented for 50–80 years and not age specific due to small numbers. Study sample has non-missing values for all included covariates (age, sex, BMI, systolic blood pressure, total cholesterol, triglycerides, self-reported general health status, self-reported history of heart attack, stroke, angina, asthma and diabetes, self-reported blood pressure treatment, smoking, leisure-time physical activity and education).

BMI, body mass index; IHD, ischaemic heart disease.

### Table 3 HR (95% CI) for all-cause and cause-specific mortality per 1 SD reduction in grip strength

<table>
<thead>
<tr>
<th>Gender interaction</th>
<th>Men and women (N=6601)</th>
<th>Men (N=2744)</th>
<th>Women (N=3857)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total mortality</td>
<td>1.17 (1.12 to 1.22)</td>
<td>1.22 (1.12 to 1.30)</td>
<td>1.16 (0.99 to 1.22)</td>
</tr>
<tr>
<td>Cardiovascular</td>
<td>1.26 (1.08 to 1.47)</td>
<td>1.26 (1.08 to 1.47)</td>
<td>1.22 (0.99 to 1.49)</td>
</tr>
<tr>
<td>Stroke</td>
<td>1.35 (1.06 to 1.32)</td>
<td>1.22 (1.02 to 1.35)</td>
<td>1.20 (1.01 to 1.43)</td>
</tr>
<tr>
<td>IHD</td>
<td>1.13 (1.10 to 1.57)</td>
<td>1.34 (1.10 to 1.63)</td>
<td>1.39 (1.11 to 1.74)</td>
</tr>
<tr>
<td>Cancer</td>
<td>1.44 (1.13 to 1.84)</td>
<td>1.37 (0.98 to 1.91)</td>
<td>1.51 (1.05 to 2.17)</td>
</tr>
</tbody>
</table>

Age 50–80 years at baseline.

*Model 1. Adjusted for age, sex and body size (BMI and height).
†Model 2. Adjusted for age, sex and body size (BMI and height)+systolic blood pressure, total cholesterol, triglycerides, self-reported general health status, self-reported history of heart attack, stroke, angina, asthma and diabetes, self-reported blood pressure treatment, smoking, leisure-time physical activity and education.

BMI, body mass index; IHD, ischaemic heart disease.
in study designs. Further studies should investigate the association by cancer subtype.

We had sufficient power to study the association between grip strength and mortality from respiratory diseases, and our results suggest that weak muscle strength is associated with deaths from respiratory diseases in men and women. To our knowledge, only two other studies have investigated this, and their results are in line with ours. In contrast with these findings, no association between grip strength and hospital admission with respiratory illness was reported in a recent study. However, this could be due to differences in study outcome, that is, hospital admission versus mortality. Another study from Japan reported an association between grip strength and death due to pneumonia, one important cause of respiratory death.19

Our finding of an association between weak grip strength and increased rates of death due to external causes is novel, and we are not aware of other studies investigating this. Suicide, a sub-category of external causes, was studied in male adolescents in a Swedish study, and here those in the lowest tenth of grip strength were found to have higher rates of mortality due to suicide. It was suggested that physically weaker men might be more psychologically vulnerable, but the mechanisms are not

Table 4  HR for all-cause and cause-specific mortality per 1 SD reduction in grip strength, by age at grip assessment

<table>
<thead>
<tr>
<th>Cause of death (number of deaths in ages: 50–59, 60–69, 70–80 years)</th>
<th>Age 50–59</th>
<th>Age 60–69</th>
<th>Age 70–80</th>
<th>Age interaction</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total mortality (422, 1053, 750)</td>
<td>1.21 (1.10 to 1.34)</td>
<td>1.18 (1.10 to 1.25)</td>
<td>1.13 (1.05 to 1.22)</td>
<td>0.69</td>
</tr>
<tr>
<td>Cardiovascular (115, 349, 312)</td>
<td>1.19 (0.98 to 1.45)</td>
<td>1.22 (1.09 to 1.36)</td>
<td>1.24 (1.10 to 1.39)</td>
<td>0.84</td>
</tr>
<tr>
<td>Cancer (211, 387, 176)</td>
<td>1.15 (1.00 to 1.32)</td>
<td>1.06 (0.96 to 1.18)</td>
<td>0.98 (0.84 to 1.16)</td>
<td>0.29</td>
</tr>
<tr>
<td>Total mortality (422, 1053, 750)</td>
<td>1.14 (1.03 to 1.26)</td>
<td>1.13 (1.06 to 1.20)</td>
<td>1.07 (0.99 to 1.16)</td>
<td>0.61</td>
</tr>
<tr>
<td>Cardiovascular (115, 349, 312)</td>
<td>1.14 (0.94 to 1.39)</td>
<td>1.14 (1.02 to 1.28)</td>
<td>1.16 (1.02 to 1.31)</td>
<td>0.87</td>
</tr>
<tr>
<td>Cancer (211, 387, 176)</td>
<td>1.11 (0.96 to 1.27)</td>
<td>1.04 (0.93 to 1.15)</td>
<td>0.97 (0.82 to 1.14)</td>
<td>0.32</td>
</tr>
</tbody>
</table>

Number of participants: 2847 in ages 50–69 years, 2664 in ages 60–69 years and 1090 in ages 70–80 years.

*Model 1. Adjusted for age, sex and body size (BMI and height).
†Model 2. Adjusted for age, sex and body size (BMI and height)+systolic blood pressure, total cholesterol, triglycerides, self-reported general health status, self-reported history of heart attack, stroke, angina, asthma and diabetes, self-reported blood pressure treatment, smoking, leisure-time physical activity and education.

BMI, body mass index.

Figure 1  Grip z-scores and mortality HRs with 95% CIs (shaded areas) for men (blue solid lines) and women (red dashed lines). Estimated in Cox regression with grip z-scores modelled as natural cubic splines with three knots. Z-score 0 is reference. *A is minimally adjusted for age and body size (BMI and height); *B is fully adjusted for age and body size (BMI and height)+systolic blood pressure, total cholesterol, triglycerides, self-reported general health status, self-reported history of heart attack, stroke, angina, asthma and diabetes, self-reported blood pressure treatment, smoking, leisure-time physical activity and education. BMI, body mass index; CVD, cardiovascular disease.

fully understood. In our study, suicide constituted only 8% of the deaths due to external causes, and results were similar when deaths due to external causes were examined having excluded those due to suicide (results not shown). The absence of an association between grip strength and the risk of injury from falls (another subcategory of deaths due to external causes in our study) was surprising to the authors in another study.13 A large share of deaths due to external causes in the age group we have studied are falls related, about one in three for ages above 75 years and one in four in the age group 70–74 years.23 Thus, our findings contrast those of the other study,14 and this divergence could be due to the relatively young age in the other study (39–74 years at the end of follow-up) compared with our participants (52–96 years at the end of follow-up). It could also be due to differences in study outcome, that is, hospital admission versus mortality.

Most studies on grip strength and cause-specific mortality have been performed among older populations, but associations have also been shown for grip strength assessed at younger ages19,26 and middle age.9,13,16 Our results of similar associations between grip strength and all-cause mortality and CVD mortality in midlife and old age are in line with results from Japan16 and the USA.18 Studies restricted to all-cause mortality also show similar association with mortality, whether grip was measured in midlife or old age.9

Our finding that grip strength is linked to deaths from diseases that are strongly associated with ageing in our study, such as CVD, stroke and respiratory diseases, is consistent with the explanation that grip strength is a marker of underlying ageing and disease processes. The strong association with deaths due to external causes, such as falls, is also likely to be connected to the ageing process, for example, decreasing reaction time and reduced postural control. Thus, these findings suggest that weak grip strength is a marker for advanced biologic age and poor overall health. However, our results were robust and not fully explained by adjustment for a range of health-related factors, which suggests that poor overall health does not fully explain our findings. Furthermore, our consistent findings across all age groups, including the youngest age group, suggest that clinically manifest conditions are unlikely to provide the whole explanation. It suggests that weak grip strength may indicate underlying ageing and disease processes even prior to their clinical manifestation and thus may be a useful early sentinel of later life risk.

The main strengths of the present population-based study are the long follow-up time and the large sample size and high number of deaths, with sufficient statistical power to perform cause-specific analyses stratified by gender and age groups. The Cause of Death Registry has a near-complete coverage and the data quality is good.27 Another strength is our novel finding of an association between grip strength and mortality from external causes, which we were able to show was not driven by the association between grip strength and suicide, which has previously been reported.19 A limitation of our study is also related to study size, as it is underpowered to break down the rather broad group of ‘external causes’ into finer subgroups. Another limitation is our use of a single measurement of grip strength, whereby we were unable to study change in grip strength and its association with mortality, which has been reported previously.28 As in most observational studies, there is also the possibility of unmeasured confounding, which may account for the relationship between grip strength and mortality. In our study, we report age-specific and gender-specific grip strength z-scores; therefore, the age and gender by grip interactions are based on mortality HRs associated with a 1 unit z-score change within each combination of age group and sex. If the grip strength distribution varied markedly between groups, for example, between age groups, a 1 z-score unit change would mean a quite different absolute change in grip strength for young and old, which could affect interpretation of results. While the SD was somewhat smaller in the older age group, it was similar in both genders, and when an additional set of analyses were run using absolute values of grip measured in kPa, these gave results similar to the analyses using z-scores (results not shown).

In conclusion, weaker grip strength was significantly associated with increased rates of all-cause mortality and deaths due to a range of causes, including CVD, stroke, IHD and respiratory diseases. A novel finding was the association between weaker grip strength and deaths due to external causes. These associations were similar in both genders and across age groups, which supports the hypothesis that grip strength might be a useful biomarker of ageing over the lifespan.
Contributors BHS had the idea of the paper, and drafted it together with RC, AB, Lj, HS, VS and NE. BHS did all the statistical analyses. All authors have discussed the paper and seen and approved the final version.

Funding RC is supported by the UK Medical Research Council (Programme code MC_UU_12019/4).

Competing interests None declared.

Patient consent Obtained.

Ethics approval This study was conducted with the approval of the Data Inspectorate, and written informed consent was obtained from the participants.

Provenance and peer review Not commissioned; externally peer reviewed.

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