Obesity in young men, and individual and combined risks of type 2 diabetes, cardiovascular morbidity and death before 55 years of age: a Danish 33-year follow-up study

Morten Schmidt, Aarhus University Hospital
Sigrun A Johannesdottir, Aarhus University Hospital
Stanley Lemeshow, Aarhus University Hospital
Timothy L Lash, Emory University
Sinna P Ulrichsen, Aarhus University Hospital
Hans Erik Botker, Aarhus University Hospital
Henrik Toft Sorensen, Aarhus University Hospital

Journal Title: BMJ Open
Volume: Volume 3, Number 4
Publisher: BMJ Publishing Group | 2013-01-01, Pages e002698-e002698
Type of Work: Article | Final Publisher PDF
Publisher DOI: 10.1136/bmjopen-2013-002698
Permanent URL: https://pid.emory.edu/ark:/25593/s7b4v

Final published version: http://dx.doi.org/10.1136/bmjopen-2013-002698

Copyright information:
© 2013, British Medical Journal

This is an Open Access work distributed under the terms of the Creative Commons Attribution-Noncommercial 3.0 Unported License (http://creativecommons.org/licenses/by-nc/3.0/).

Accessed June 9, 2024 4:14 AM EDT
Obesity in young men, and individual and combined risks of type 2 diabetes, cardiovascular morbidity and death before 55 years of age: a Danish 33-year follow-up study

Morten Schmidt,1,2,3 Sigrun A Johannesdottir,1,2 Stanley Lemeshow,1,2 Timothy L Lash,1,4 Sinna P Ulrichsen,1 Hans Erik Bøtker,3 Henrik Toft Sørensen1

ABSTRACT

Objectives: To examine the association between body mass index (BMI) in young adulthood and cardiovascular risks, including venous thromboembolism, before 55 years of age.

Design: Cohort study using population-based medical databases.

Setting: Outcomes registered from all hospitals in Denmark from 1977 onwards.

Participants: 6502 men born in 1955 and eligible for conscription in Northern Denmark.

Main outcome measures: Follow-up began at participants’ 22nd birthday and continued until death, emigration or 55 years of age, whichever came first. Using regression analyses, we calculated the risks and HRs, adjusting for cognitive test score and years of education.

Results: 48% of all obese young men (BMI ≥30 kg/m²) were either diagnosed with type 2 diabetes, hypertension, myocardial infarction, stroke or venous thromboembolism or died before reaching 55 years of age. Comparing obese men with normal weight men (BMI 18.5 to <25.0 kg/m²), the risk difference for any outcome was 28% (95% CI 19% to 38%) and the HR was 3.0 (95% CI 2.3 to 4.0). Compared with normal weight, obesity was associated with an event rate that was increased more than eightfold for type 2 diabetes, cardiovascular morbidity or premature death of almost 30%.

Conclusions: In this cohort of young men, obesity was strongly associated with adverse cardiometabolic events before 55 years of age, including venous thromboembolism. Compared with those of normal weight, young obese men had an absolute risk increase for type 2 diabetes, cardiovascular morbidity or premature death of almost 30%.

INTRODUCTION

Tripling in prevalence over the last three decades and now exceeding 30% in the USA,1–3 obesity in young adulthood is as a major public health concern worldwide, leading to a shorter lifespan for today’s children.4 Despite the obesity epidemic, cardiovascular morbidity and mortality have continued to decrease in the Western world.5–6

Although obesity in adulthood is a risk factor for type 2 diabetes,1–3 7 cardiovascular disease4 8 and mortality,5 6 9 it remains unclear...
whether an elongated history of overweight, starting early in life, poses additional risks. Previous reports indicate that age modifies the effect of obesity on cardiovascular death with greater impact in younger age groups, including childhood and young adulthood. However, while obese children reduce their additional cardiovascular risks (compared with non-obese children) by ceasing to be obese children, new studies indicate that the risks among obese young adults persist independently of weight changes in later adulthood. These findings emphasise the need to study the adverse outcomes associated with obesity within the age group of young adults specifically.

Several studies have examined the association between body mass index (BMI) in young adults and premature death. Fewer studies have examined long-term risks of type 2 diabetes and ischaemic heart disease. However, despite the link between atherosclerosis, metabolic syndrome and venous thromboembolism, none of the previous studies have included venous thromboembolism as an outcome. Moreover, estimates of the absolute risk, in addition to the relative risk, of adverse events are important for clinical decision making, particularly when counselling patients on risk modification associated with weight, diet and exercise. Nonetheless, no study has examined the combined risk of type 2 diabetes, numerous cardiovascular outcomes and premature death in one study, thereby assessing the complete cardiometabolic risks associated with young adulthood obesity.

We therefore followed a cohort of 22-year-old men for 33 years to examine the association between BMI in young adulthood and the risk of type 2 diabetes, hypertension, myocardial infarction, stroke, venous thromboembolism and death before 55 years of age.

**METHODS**

**Setting**

We conducted this population-based cohort study in the Fifth Military Conscription District in Denmark, populated by approximately 700,000 inhabitants. The Danish National Health Service provides universal tax-supported healthcare, guaranteeing unfettered access to general practitioners and hospitals and partial reimbursement for prescribed medications. Accurate and unambiguous individual-level linkage of all Danish registries is possible using the unique central personal registration number assigned to all residents at birth or upon immigration.

**Study cohort**

Nearly all Danish men have had to register with the military board for an examination of fitness to serve when reaching 18 years of age or shortly thereafter (median age 19 years). In connection with the registration, all examinees complete a health questionnaire, in which they report chronic diseases that could preclude military service, for example, asthma, epilepsy or spinal osteochondrosis, but not obesity. The Draft Board verifies such reports with health care providers, and men deemed ineligible for military service are excused from the draft at this point (approximately 15%).

Information obtained at the examination is registered and stored in regional and national databases. Using a conscription research database, we identified all persons from the 1955 birth cohort who later appeared before the draft board in Northern Denmark (n=6502).

**Body mass index**

All potential conscripts undergo a physical and mental examination. The physical examination includes measurements of height (without shoes) and body weight (wearing trunks only and using sliding scales and calibrated balances). Using these height and weight measurements, we calculated the BMI. Values of weight <45 kg (n=6) were considered to be data entry errors and recoded as missing. We categorised BMI as underweight (<18.5 kg/m²), normal (18.5 to <25 kg/m²), overweight (25 to <30 kg/m²) or obese (≥30 kg/m²).

**Covariates**

From the conscription database, we obtained data on cognitive function as measured by the validated Boerge Prie test. The Boerge Prie test is a 78-item group intelligence test with four subscales (letter matrices, verbal analogies, number sequences and geometric figures) and a single final score, recorded as the number of correctly answered items (range 0–78). The scores are strongly correlated with conventional intelligence-test scores (eg, correlation of 0.82 with the Wechsler Adult Intelligence Scale). Based on quartiles, we categorised the test scores as low (0–31), moderate (32–39), high (40–46) and very high (47–78).

From the conscription database, we also obtained information on years of education at the time of examination. Based on quartiles, we categorised the education length as short, moderate, long and very long.

**Outcomes**

The Danish National Registry of Patients records information on patients discharged from all Danish non-psychiatric hospitals since 1 January 1977 and from all emergency room and outpatient specialty clinic visits since 1995. Each hospital discharge or outpatient visit is recorded in the registry with one primary diagnosis and one or more secondary diagnoses classified according to the *International Classification of Diseases*, 8th revision (ICD-8) until the end of 1993 and the 10th revision (ICD-10) thereafter.

Using the Danish National Registry of Patients, we identified all examinees with a first diagnosis of type 2 diabetes, hypertension, myocardial infarction, stroke (ischaemic stroke, intracerebral haemorrhage or subarachnoid haemorrhage) or venous thromboembolism (deep venous thromboembolism or pulmonary
embolism). To ensure more complete identification of examinees with type 2 diabetes, we also searched the Aarhus University Prescription Database, covering the study region, for any use of antidiabetic drugs from 1 January 1989 through 2010. ICD and ATC codes are provided as online supplementary material.

We obtained information on all-cause mortality from the Danish Civil Registration System. This registry has recorded all changes in vital status and migration for the entire Danish population since 1968, with daily electronic updates. Finally, a combined outcome was defined as the first-time occurrence of any of the individual outcomes.

Statistical analyses
Initially, we used descriptive statistics to characterise the study population by categories of BMI, cognitive test score and years of education. To ensure that the Danish National Registry of Patients (initiated in 1977) would capture events, follow-up started at the examinee’s 22nd birthday. We excluded all men who were censored between their date of examination and their 22nd birthday (17 died and 8 emigrated). Follow-up continued until the first occurrence of an outcome, emigration or 33 years of follow-up (ie, their 55th birthday), whichever came first. First, we illustrated graphically the association between BMI and the predicted cumulative incidence function for each outcome using the proportional sub-hazards model by Fine and Gray. We used the pseudo-value method to calculate the 33-year risk for each BMI group, as well as the risk differences compared with normal weight. We treated death as a competing risk in all analyses of non-fatal outcomes.

We calculated rates for each BMI group and used Cox proportional hazards regression to compute hazard ratios (HRs) associating BMI with all outcomes. BMI was analysed both as a categorical and continuous variable. For the categorical BMI variable, the proportional hazard assumption was assessed by log–log plots and Schoenfeld’s test and found valid. We assessed the scale of the continuous BMI variable using fractional polynomials and found no evidence of non-linearity in the log hazard. We repeated all regression analyses adjusting for cognitive test score and years of education. As there was no recent validation study available for hypertension, we estimated the proportion of patients registered with hypertension in the Danish National Registry of Patients that also redeemed at least one prescription for antihypertensive medication (registered in the prescription database).

The study was approved by the Danish Data Protection Agency (2011-41-5807). All analyses were conducted in STATA software V.12.1 (STATA, College Station, Texas, USA).

Results
Characteristics
The characteristics of the study population are presented in table 1. We identified 6502 male examinees from the 1955 birth cohort in Northern Denmark. Among these, 5407 (83%) were of normal weight, 353 (5%) were underweight, 639 (10%) were overweight and 97 (1.5%) were obese. BMI ranged from a minimum of 14.4 kg/m² to a maximum of 42.7 kg/m². The median BMI was 21.7 kg/m² (IQR 20.3–23.4 kg/m²).

Compared with men of normal weight (25%), a higher proportion of those who were overweight (32%) or obese (41%) scored low on the cognitive test. Also, compared with subjects of normal weight (24%), a higher proportion of overweight (32%) or obese (33%) persons belonged to the lowest quartile of years of education. The validation analysis revealed that 88% of patients with a diagnosis of hypertension redeemed one or more prescriptions for antihypertensive medication.

Combined outcome
The cohort contributed a total of 199 430 person-years of follow-up, providing a mean follow-up time of
31 years. Compared with 20% of men of normal weight, 48% of all obese men were diagnosed with type 2 diabetes, hypertension, myocardial infarction, stroke or venous thromboembolism or died before their 55th birthday (table 2 and figure 1). After adjusting for cognitive test score and years of education, the absolute risk difference between the groups was 28% (95% CI 19% to 38%) and the HR was 3.0 (95% CI 2.3 to 4.0).

**Individual outcomes**

Obesity was strongly associated with all individual outcomes except stroke (table 3 and figure 2). With a risk difference of 22% (95% CI 13% to 31%) and HR of 8.2 (95% CI 5.4 to 12.3) the absolute and relative associations for obesity, compared with normal weight, were the strongest for type 2 diabetes. However, there was also a more than fourfold increased rate of venous thromboembolism and more than twofold increased rate of hypertension, myocardial infarction and death. For stroke, there was a 40% increased rate among overweight persons. The wide CIs made the association between obesity and stroke inconclusive. In the analysis of BMI as a continuous variable, a unit increase in BMI corresponded to an increased rate of approximately 5% for myocardial infarction, 10% for hypertension and venous thromboembolism and 20% for type 2 diabetes (table 4).

**DISCUSSION**

In this population-based 33-year follow-up study, we found that nearly half of all obese young men either were diagnosed with type 2 diabetes, hypertension, myocardial infarction, stroke or venous thromboembolism or died before reaching 55 years of age. Obesity was associated with an event rate that, compared with normal weight, was increased more than eightfold for type 2 diabetes, fourfold for venous thromboembolism and twofold for hypertension, myocardial infarction, and premature death. The strong association between young adulthood obesity and venous thromboembolism was of particular importance because it, to our knowledge, has not previously been reported. Obese subjects had a threefold increased event rate of any of these diseases compared with persons of normal weight, yielding a notable absolute risk increase of almost 30%. The magnitude of the relative and absolute risk estimates emphasise the major clinical and public health implications associated with obesity in young adulthood.

**Strengths and limitations of study**

Several issues should be considered when interpreting our results. With military examination in anticipation of conscription being mandatory and participation enforced by law, the population-based design reduced potential selection bias. We used measured instead of self-reported height and weight. Doing so, we eliminated differential misclassification of BMI, which may have biased other studies because underweight people tend to overestimate their BMI and overweight people tend to underestimate their BMI. The Danish Civil Registration System ensured complete follow-up of all examinees with accurate mortality data. The tax-supported universal healthcare system allowed for

---

**Table 2** Body mass index in young adulthood and the combined risk of type 2 diabetes, hypertension, myocardial infarction, stroke, venous thromboembolism and death before age 55 years*

<table>
<thead>
<tr>
<th>BMI categories</th>
<th>Number</th>
<th>Risk, % (95% CI)</th>
<th>Risk difference†, % (95% CI)</th>
<th>Rate‡ (95% CI)</th>
<th>HR† (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Normal</td>
<td>1083</td>
<td>20 (16 to 23)</td>
<td>0</td>
<td>649 (612 to 689)</td>
<td>1</td>
</tr>
<tr>
<td>Underweight</td>
<td>63</td>
<td>18 (12 to 23)</td>
<td>−2 (−6 to 2)</td>
<td>574 (448 to 734)</td>
<td>0.9 (0.7 to 1.1)</td>
</tr>
<tr>
<td>Overweight</td>
<td>205</td>
<td>31 (26 to 36)</td>
<td>11 (7 to 15)</td>
<td>1086 (947 to 1245)</td>
<td>1.7 (1.4 to 1.9)</td>
</tr>
<tr>
<td>Obesity</td>
<td>48</td>
<td>48 (38 to 59)</td>
<td>28 (19 to 38)</td>
<td>1814 (1367 to 2407)</td>
<td>3.0 (2.3 to 4.0)</td>
</tr>
</tbody>
</table>

*Among the 1955 birth cohort that appeared for military examination in Northern Denmark and who survived until their 22nd birthday.
†In all regression analyses, adjustments were made for cognitive test score and years of education.
‡Rates per 100 000 person-years.
BMI, body mass index.
Table 3  Body mass index in young adulthood and the risks of type 2 diabetes, cardiovascular morbidity or death before age 55 years*  

<table>
<thead>
<tr>
<th>BMI categories</th>
<th>Number</th>
<th>Risk, % (95% CI)</th>
<th>Risk difference†, % (95% CI)</th>
<th>Rate‡ (95% CI)</th>
<th>HR† (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Type 2 diabetes</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Normal</td>
<td>207</td>
<td>5 (3 to 7)</td>
<td>0</td>
<td>122 (106 to 139)</td>
<td>1</td>
</tr>
<tr>
<td>Underweight</td>
<td>7</td>
<td>4 (1 to 6)</td>
<td>-2 (-3 to 0)</td>
<td>63 (30 to 132)</td>
<td>0.5 (0.2 to 1.1)</td>
</tr>
<tr>
<td>Overweight</td>
<td>76</td>
<td>13 (10 to 16)</td>
<td>8 (5 to 10)</td>
<td>387 (309 to 485)</td>
<td>3.1 (2.4 to 4.0)</td>
</tr>
<tr>
<td>Obesity</td>
<td>26</td>
<td>27 (18 to 36)</td>
<td>22 (13 to 31)</td>
<td>946 (644 to 1390)</td>
<td>8.2 (5.4 to 12.3)</td>
</tr>
<tr>
<td><strong>Hypertension</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Normal</td>
<td>394</td>
<td>7 (4 to 9)</td>
<td>0</td>
<td>233 (211 to 257)</td>
<td>1</td>
</tr>
<tr>
<td>Underweight</td>
<td>24</td>
<td>6 (3 to 10)</td>
<td>0 (-3 to 3)</td>
<td>215 (144 to 321)</td>
<td>0.9 (0.6 to 1.4)</td>
</tr>
<tr>
<td>Overweight</td>
<td>92</td>
<td>13 (10 to 17)</td>
<td>7 (4 to 10)</td>
<td>469 (382 to 575)</td>
<td>2.0 (1.6 to 2.5)</td>
</tr>
<tr>
<td>Obesity</td>
<td>14</td>
<td>14 (7 to 21)</td>
<td>8 (1 to 16)</td>
<td>494 (292 to 833)</td>
<td>2.1 (1.2 to 3.6)</td>
</tr>
<tr>
<td><strong>Myocardial infarction</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Normal</td>
<td>134</td>
<td>3 (1 to 5)</td>
<td>0</td>
<td>79 (66 to 93)</td>
<td>1</td>
</tr>
<tr>
<td>Underweight</td>
<td>12</td>
<td>4 (1 to 6)</td>
<td>1 (-1 to 3)</td>
<td>107 (61 to 189)</td>
<td>1.4 (0.8 to 2.5)</td>
</tr>
<tr>
<td>Overweight</td>
<td>18</td>
<td>3 (1 to 5)</td>
<td>0 (-1 to 2)</td>
<td>90 (56 to 142)</td>
<td>1.1 (0.7 to 1.8)</td>
</tr>
<tr>
<td>Obesity</td>
<td>6</td>
<td>7 (2 to 11)</td>
<td>4 (-1 to 9)</td>
<td>208 (93 to 462)</td>
<td>2.5 (1.1 to 5.6)</td>
</tr>
<tr>
<td><strong>Stroke</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Normal</td>
<td>128</td>
<td>2 (1 to 4)</td>
<td>0</td>
<td>75 (63 to 89)</td>
<td>1</td>
</tr>
<tr>
<td>Underweight</td>
<td>8</td>
<td>2 (0 to 4)</td>
<td>0 (-2 to 2)</td>
<td>71 (36 to 143)</td>
<td>1.0 (0.5 to 2.0)</td>
</tr>
<tr>
<td>Overweight</td>
<td>22</td>
<td>3 (1 to 5)</td>
<td>1 (-1 to 2)</td>
<td>109 (72 to 166)</td>
<td>1.4 (0.9 to 2.2)</td>
</tr>
<tr>
<td>Obesity</td>
<td>2</td>
<td>2 (-1 to 5)</td>
<td>0 (-3 to 3)</td>
<td>69 (17 to 276)</td>
<td>0.9 (0.2 to 3.6)</td>
</tr>
<tr>
<td><strong>Venous thromboembolism</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Normal</td>
<td>57</td>
<td>2 (1 to 3)</td>
<td>0</td>
<td>33 (26 to 43)</td>
<td>1</td>
</tr>
<tr>
<td>Underweight</td>
<td>2</td>
<td>1 (0 to 3)</td>
<td>0 (-1 to 0)</td>
<td>18 (4 to 71)</td>
<td>0.6 (0.1 to 2.3)</td>
</tr>
<tr>
<td>Overweight</td>
<td>7</td>
<td>2 (0 to 3)</td>
<td>0 (-1 to 1)</td>
<td>35 (17 to 73)</td>
<td>0.9 (0.4 to 2.1)</td>
</tr>
<tr>
<td>Obesity</td>
<td>5</td>
<td>6 (1 to 10)</td>
<td>4 (0 to 8)</td>
<td>173 (72 to 415)</td>
<td>4.7 (1.9 to 11.9)</td>
</tr>
<tr>
<td><strong>Death</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Normal</td>
<td>416</td>
<td>7 (5 to 10)</td>
<td>0</td>
<td>243 (220 to 267)</td>
<td>1</td>
</tr>
<tr>
<td>Underweight</td>
<td>28</td>
<td>7 (3 to 11)</td>
<td>0 (-3 to 2)</td>
<td>249 (172 to 361)</td>
<td>1.0 (0.7 to 1.5)</td>
</tr>
<tr>
<td>Overweight</td>
<td>47</td>
<td>6 (3 to 9)</td>
<td>-1 (-3 to 1)</td>
<td>232 (174 to 309)</td>
<td>0.9 (0.7 to 1.2)</td>
</tr>
<tr>
<td>Obesity</td>
<td>16</td>
<td>16 (8 to 23)</td>
<td>8 (1 to 16)</td>
<td>547 (335 to 893)</td>
<td>2.1 (1.3 to 3.5)</td>
</tr>
</tbody>
</table>

*Among the 1955 birth cohort that appeared for military examination in Northern Denmark and who survived until their 22nd birthday.
†In all regression analyses, adjustments were made for cognitive test score and education level. In all non-fatal outcomes, death was treated as a competing risk.
‡Rates per 100 000 person-years.
BMI, body mass index.

Figure 2  Body mass index in young adulthood and cumulative incidence (risk) of type 2 diabetes, cardiovascular morbidity or death before age 55 years (note that follow-up starts at age 22 and that the maximum risk is not constant on the y-axis of the various panels).
Table 4 The association between a one-unit increase in body mass index in young adulthood and type 2 diabetes, cardiovascular morbidity or death before 55 years of age*

<table>
<thead>
<tr>
<th>Outcomes</th>
<th>HR (95% CI)†</th>
</tr>
</thead>
<tbody>
<tr>
<td>Type 2 diabetes</td>
<td>1.19 (1.16 to 1.22)</td>
</tr>
<tr>
<td>Hypertension</td>
<td>1.11 (1.08 to 1.13)</td>
</tr>
<tr>
<td>Myocardial infarction</td>
<td>1.05 (1.00 to 1.10)</td>
</tr>
<tr>
<td>Stroke</td>
<td>1.02 (0.97 to 1.08)</td>
</tr>
<tr>
<td>Venous thromboembolism</td>
<td>1.10 (1.03 to 1.18)</td>
</tr>
<tr>
<td>Death</td>
<td>1.01 (0.98 to 1.04)</td>
</tr>
</tbody>
</table>

*Among the 1955 birth cohort that appeared for military examination in Northern Denmark and who survived until their 22nd birthday.†BMI analysed as a continuous variable in a linear regression model adjusted for cognitive test score and education level. BMI, body mass index.

There were, however, some limitations. Examinees had to survive from their examination date until the start of follow-up at 22 years of age. Although BMI is a widely used proxy for adiposity, it does not distinguish adipose from muscle tissue nor does it take the anthropometric distribution of fat into account. Some degree of misclassification is therefore inevitable (for all studies on this topic). Nevertheless, BMI is a valuable tool to provide a standardised definition of obesity, allowing for a comparison of its associated risks across decades. The Aarhus University Prescription Database did not cover the entire study period. However, any potential underreporting of diabetes and hypertension in the Danish National Registry of Patients would provide underestimates of the absolute risks, and thus cannot explain the increased risks.

We cannot exclude confounding from exercise, diet, weight changes and smoking. An advantage of studying death rates among young subjects is that adjustments for other conditions are largely unnecessary. Moreover, analyses of myocardial infarction and death should not adjust for hypercholesterolaemia, hypertension, diabetes or other factors associated with the metabolic syndrome because all of these risk factors may be on the causal pathway. We were unable to consider the impact of weight changes later in life. However, weight loss among obese or weight gain among normal weight conscripts would bias the results towards unity, and thus cannot explain our findings. Furthermore, other studies have found the associations between young adulthood BMI and coronary artery disease and death, independent of later weight changes. Smoking is associated with low and not high BMI. Thus, any unmeasured confounding from smoking would also tend to bias the results towards unity. Supporting the robustness of our results, previous studies on young obese adults have found similar death rates among smokers and non-smokers or even higher death rates among non-smokers. In addition, as a surrogate measure of lifestyle, we did adjust for cognitive test score and years of education. Finally, although the results derive from a study that included only young men, previous reports suggest the associations are likely also to hold for young women.

Comparison with other studies

Our results are in line with previous reports on the individual outcomes. The increased premature mortality risk is consistent. Among prior studies, a similar Swedish study on male conscripts with mean age of 18.7 years found an almost identical association between obesity and premature mortality (HR 2.14, 95% CI 1.61 to 2.85). In contrast to the reports of a U-shaped relationship between BMI and mortality in young adults, our results supported the absence of any association between underweight and premature mortality.

Our results for type 2 diabetes and myocardial infarction are also supported by previous reports. Reporting on the magnitude of the association between BMI and diabetes, a recent meta-analysis of 18 cohort studies reported relative risk estimates almost identical to ours for both overweight (2.92, 95% CI 2.57 to 3.32) and obesity (7.28, 95% CI 6.47 to 8.28). Tirosh et al also examined young adults and found an HR for each unit increase in BMI of 1.10 (95% CI 1.08 to 1.12) for diabetes and of 1.12 (95% CI 1.07 to 1.18) for coronary heart disease. In addition, a recent meta-analysis that pooled the results from seven studies of subjects between 18 and 30 years found a relative risk of 1.08 (95% CI 1.05 to 1.11) associating 1 kg/m² higher BMI with coronary heart disease. Another recent meta-analysis found that obesity overall was associated with a twofold increased risk of venous thromboembolism. However, none of the included studies examined the risk among young adults specifically. Our finding of a more than fourfold increased risk in this age group is thus important.

Pathophysiological explanations and implications

There may be several pathophysiological explanations for our findings. First, the apparent higher cardiovascular risks associated with BMI in young adulthood compared with older adulthood may be caused by early development and clustering of cardiovascular risk factors, in particular the metabolic syndrome. Thus, overweight and obesity are both associated with insulin resistance, higher blood pressure and adverse blood lipid profiles. These risk factors increase short-term and long-term incidence of type 2 diabetes, venous thromboembolism, premature coronary artery disease (through enhanced progression of atherosclerosis)
and finally, as a result, the cardiovascular death rate. Second, physical effects of body fat may limit venous return and create a proinflammatory, prothrombotic and hypofibrinolytic milieu that enhance venous thromboembolic risks.47

The current and projected continued increase in obesity prevalence1 48 may affect negatively the current trends of declining cardiovascular death rates. 3–6 49 Thus, obesity-related morbidity and mortality will, in decades to come, place an unprecedented burden on healthcare systems worldwide. 58 This forecast reinforces the importance of understanding the effects of adiposity in young adults to plan future strategies for weight management and primary prevention.

CONCLUSIONS
In this cohort of young men, obesity was strongly associated with adverse cardiometabolic events before 55 years of age, including venous thromboembolism. Compared with those of normal weight, young obese men had an absolute risk increase for type 2 diabetes, cardiovascular morbidity or premature death of almost 30%.

Author affiliations
1Department of Clinical Epidemiology, Aarhus University Hospital, Aarhus N, Denmark
2Division of Biostatistics, College of Public Health, The Ohio State University, Columbus, OH, USA
3Department of Cardiology, Aarhus University Hospital, Aarhus N, Denmark
4Department of Epidemiology, Rollins School of Public Health, Emory University, Atlanta, GA, USA

Contributors MS and HTS conceived the study idea. MS designed the study. SPU and HTS collected the data. MS and HTS reviewed the literature. MS, SAJ, SL, TLL, HEB and HTS directed the analyses, which were carried out by MS. All authors participated in the discussion and interpretation of the results. MS organised the writing and wrote the initial drafts. All authors critically revised the manuscript for intellectual content and approved the final version. HTS is the guarantor.

Funding The study was supported by Aarhus University Research Foundation, Department of Clinical Epidemiology’s Research Foundation, ‘Augustinusfonden’, ‘Snedkermesters Sophus Jacobsen & Hustru Astrid Jacobsens Fond’ and ‘Fabrikant Karl G. Andersens Fond’. The Department of Clinical Epidemiology is a member of the Danish Center for Strategic Research in Type 2 Diabetes (Danish Research Council, grants 09-075724 and 10-079102). None of the funding sources had a role in the design, conduct, analysis or reporting of the study.

Competing interests None.

Ethics approval As this study did not involve any contact with patients or any intervention, it was not necessary to obtain permission from the Danish Scientific Ethical Committee.

Provenance and peer review Not commissioned; externally peer reviewed.

Data sharing statement No additional data are available.

REFERENCES


