Sex differences in body fat distribution and carotid intima media thickness: cross sectional survey using data from the British regional heart study

Debbie A Lawlor, Shah Ebrahim, Peter Whincup, Jonathan Sterne, Olia Papacosta, Goya Wannamethee, Surinder Dhanjil, Maura Griffin, Andrew N Nicolaides, George Davey Smith

Objective: To determine the role of central adiposity in explaining sex differences in carotid intima media thickness (IMT).

Design: Cross sectional survey.

Setting: Two British towns.

Participants: 800 men and women aged 56–75 years.

Main outcome measures: Carotid IMT.

Results: There was a continuous linear association between waist-hip ratio and IMT in both men and women. The magnitude of the association between waist to hip ratio and IMT was identical in both sexes. In age adjusted analyses IMT was 1.4% greater in men compared with women (age adjusted male to female ratio of geometric means 1.14; 95% confidence interval 1.07 to 1.21) with adjustment for waist to hip ratio this attenuated to no difference (1.00; 0.92 to 1.09). Adjustment for body mass index and for lifestyle risk factors had very little effect on the sex difference in mean intima media thickness.

Conclusions: Sex differences in body fat distribution may explain sex differences in arterial atherosclerosis.
Sex differences in body fat distribution

PREVIOUSLY DESCRIBED METHODS. The within and between ultrasonographer coefficients of variation for IMT measurements were assessed in pilot studies and were between 3% and 5%. The mean difference between ultrasonographers showed no evidence of bias, and was $-0.005$ mm, with 95% limits of agreement of $-0.12$ to $+0.11$. In a previous study using data from this cohort it was found that mean IMT at the bulb origin (carotid bifurcation) was more directly associated with clinical CHD than other sites of IMT measurement for both women and men. This measure of IMT is used as the outcome in our study. Mean IMT of the bulb origin was defined as the point at which the arterial wall diverges to form the bulb.

Other variables

Height was measured without shoes using a Harpenden Stadiometer that recorded to the nearest millimetre. Weight was measured in light clothing without shoes using Soehnle portable scales that measured to the nearest 0.1 kg. Waist measurements were taken using the midpoint between the lowest rib and iliac crest and hip measurements using the largest circumference below the waist. A flexible metal tape was used and two measurements taken to the nearest millimetre. A Dinamap 1846SX vital signs monitor was used to measure blood pressure using standard procedures. The Dinamax 1846SX monitor systematically overestimates systolic blood pressure by $8$ mm Hg and all measurements were corrected for this error. A sample of non-fasting venous blood was taken for analysis of lipids and glycated haemoglobin.

A self administered questionnaire was used to obtain information on smoking behaviour, alcohol consumption, physical activity, diet, and social class. Subjects were classified into never, former, and current smokers. Alcohol consumption was classified as none, occasional (once or twice a month), weekends only, or daily drinking. Sporting or exercise activity was classified as none, occasional (less than once a month), or frequent (at least once a month). Subjects completed a detailed food item questionnaire and “low fat diet” was defined as mainly using either semi-skimmed, skimmed or no milk and using either low calorie margarine or no spread on bread. Social class was derived from the longest held occupation of each man or of each husband in the case of married women and her own occupation in the case of single women. Occupation was coded in accordance with the registrar general’s occupational classification (social classes I, II, III non-manual, III manual, IV, V, where I is professional/managerial and V unskilled manual labour).

Statistical analysis

Multiple linear regression was used to assess the effect of adjustment for waist-hip ratio on sex differences in IMT. The outcome in these models was IMT the main predictor variable was sex. Age, waist-hip ratio, body mass index, systolic and diastolic blood pressure, high density lipoprotein cholesterol, triglycerides (logged), and glycated haemoglobin were entered as continuous variables. Smoking, alcohol consumption, exercise, low fat diet, and social class (I, II, III non-manual, III manual, IV, V) were entered as categorical variables, together with dummy variables representing those for whom data on smoking status ($n = 59$), alcohol ($n = 46$), exercise ($n = 56$), low fat diet ($n = 43$), and social class ($n = 55$) were not available. For continuous variables missing values were not imputed. Sensitivity analyses were undertaken, without these dummy variables, including only subjects with complete data for all categorical as well as continuous variables. The point estimates and $p$ values of these sensitivity analysis were unaltered from those presented here; the confidence intervals were slightly widened with changes at the third decimal place. In all models adjustment was made for age, the clustering effect of each town (to obtain robust standard errors for the calculation of 95% confidence intervals), and for social class because there was a slight sex difference in the distribution of social class in the cohort.

IMT had a skewed distribution but was log normal and therefore geometric means are presented and logged values were used in the linear regression models; with these transformations residuals in the linear regression models were normally distributed. The resulting regression coefficients of these logged variables were back transformed to provide the ratio of geometric means that were then used to provide the proportionate (expressed as a percentage) difference in IMT level between men and women. All analyses were undertaken using Stata version 7 (Stata Corporation, Texas).

RESULTS

A total of 375 women and 425 men attended (response rates of 69% and 83% respectively) and of these adequate ultrasonic images were obtained on 367 women and 418 men. Women who attended and had ultrasound images were younger than non-responders (mean age 65.2 versus 67.7 years, $p = 0.001$). There was no substantive age difference between responders and non-responders among men (65.8 compared with 65.9 years, $p = 0.9$). The mean (SD) age of female study participants was 65.2 (6.0) and of male study participants was 65.8 (5.6), $p = 0.18$ for difference between female and male participants.

Figure 1 shows the distribution of waist-hip ratio in women and men; there is little overlap between the sexes. Table 1 shows age adjusted distributions of IMT, waist-hip ratio, and other CHD risk factors in women and men. IMT was thinner in women compared with men (geometric mean 1.36 mm versus 1.54 mm) and women had smaller waist-hip ratios (mean 0.80 versus 0.93) than men. Compared with women, men had higher systolic and diastolic blood pressures, higher glycated haemoglobin, and triglyceride levels and lower high density lipoprotein levels and low density lipoprotein levels. The proportion of daily drinkers among men was twice that among women and men were less

Key points

- The reason for the sex difference in the occurrence of CHD is unclear.
- Previous studies have suggested that sex differences in fat distribution may be important, but since the distributions in central adiposity are so different in women and men simple adjustment of the odds ratio of CHD comparing women with men for waist to hip ratio is uninformative.
- It needs to be shown both that there is a continuous linear association across the two sexes in the association between waist to hip ratio and CHD risk, and that adjustment for waist to hip ratio attenuates the association between sex and CHD.
- In this study we have shown that there is a linear continuous association between waist to hip ratio and carotid intima media thickness (an indicator of atherosclerosis) across both sexes and in addition that adjustment for waist to hip ratio removes the sex difference in carotid intima media thickness.

www.jech.com
likely to have a low fat diet. There was very little difference between women and men in the proportion undertaking sport or exercise at least monthly. Similar proportions of women and men were current smokers but the proportion of men who were ex-smokers was nearly twice that of women.

Figure 2 shows the unadjusted linear relation between waist-hip ratio and IMT in women and men. This suggests a modest continuous relation across both sexes with similar gradients in the linear relation between waist-hip ratio and IMT in both women and men. After adjustment for age, town, and social class the regression coefficients (95% confidence intervals) of log IMT on waist-hip ratio were identical in women and men: IMT increased by 8% (95% confidence interval: 3% to 14%), \( p < 0.001 \) for a one standard deviation increase in waist-hip ratio in both sexes.

In both sexes waist-hip ratio was positively associated with triglyceride level, glycated haemoglobin level, and was inversely associated with high density lipoprotein cholesterol (all \( p \) values < 0.01). There was no strong evidence of statistical interaction with sex in the associations between waist-hip ratio and any of the metabolic risk factors (all \( p \) values were equal to or greater than 0.1). In women, age was positively associated with waist-hip ratio but in men there appeared to be no association between age and waist-hip ratio (\( p \) for interaction = 0.01). The age, social class, and town adjusted ratio of male to female IMT geometric means (95% CI) was 1.14 (1.07 to 1.21) and when further adjustment for waist to hip ratio was made this attenuated to 1.00 (0.92 to 1.09). These results have been presented as percentage changes in the first row of table 2—that is, 14% (7% to 21%) and 0% (−8% to 9%). Adjustment for metabolic risk factors attenuated the association from 15% to 8%. Adjustment for body mass index and for lifestyle risk factors had very little effect on the sex difference in IMT.

### DISCUSSION

#### Main findings
In this study we have shown that the association between waist-hip ratio and carotid IMT is linear and continuous across both sexes and that the magnitude of the association, although modest, is the same in both sexes. Furthermore, adjustment for waist-hip ratio attenuated the sex difference in IMT to the null value.

#### Study limitations
We have used carotid IMT rather than CHD as our outcome. Carotid IMT has been shown to be strongly associated with prevalent and incident CHD.\(^{13,18}\) It has been suggested that the difference in IMT between women and men may be

### Table 1

<table>
<thead>
<tr>
<th></th>
<th>Women (n = 367)</th>
<th>Men (n = 418)</th>
<th>( p ) for difference</th>
</tr>
</thead>
<tbody>
<tr>
<td>Intima media thickness (mm)</td>
<td>1.36 (1.31 to 1.42)</td>
<td>1.54 (1.49 to 1.61)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Waist-hip ratio</td>
<td>0.80 (0.79 to 0.80)</td>
<td>0.93 (0.92, 0.94)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Waist circumference (cm)</td>
<td>84.90 (83.73 to 86.06)</td>
<td>97.33 (96.25 to 98.42)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Hip circumference (cm)</td>
<td>106.66 (105.73 to 107.58)</td>
<td>104.41 (103.55 to 105.27)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Body mass index (kg/m(^2))</td>
<td>27.72 (27.27 to 28.16)</td>
<td>26.91 (26.49 to 27.32)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>HDLc (mmol/l)</td>
<td>1.55 (1.51 to 1.59)</td>
<td>1.26 (1.23 to 1.29)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>LDLc (mmol/l)</td>
<td>4.14 (4.04 to 4.24)</td>
<td>3.65 (3.56 to 3.74)</td>
<td>0.03</td>
</tr>
<tr>
<td>Triglycerides (mmol/l)</td>
<td>1.87 (1.77 to 1.97)</td>
<td>2.01 (1.92 to 2.12)</td>
<td>0.005</td>
</tr>
<tr>
<td>Glycated haemoglobin (%)</td>
<td>4.69 (4.59 to 4.79)</td>
<td>4.85 (4.75 to 4.95)</td>
<td>0.01</td>
</tr>
<tr>
<td>Systolic blood pressure (mm Hg)</td>
<td>144.3 (142.0 to 146.7)</td>
<td>149.0 (146.8 to 151.2)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Diastolic blood pressure (mm Hg)</td>
<td>84.3 (83.3 to 85.3)</td>
<td>84.3 (83.3 to 85.3)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Current smoker (%)</td>
<td>19.5 (15.5 to 24.3)</td>
<td>20.9 (16.7 to 24.4)</td>
<td>0.13</td>
</tr>
<tr>
<td>Former smoker (%)</td>
<td>28.9 (24.1 to 31.2)</td>
<td>51.8 (46.9 to 56.5)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Daily alcohol (%)</td>
<td>15.1 (11.7 to 19.4)</td>
<td>36.4 (31.9 to 41.2)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Exercise at least once a month (%)</td>
<td>21.8 (17.6 to 26.6)</td>
<td>26.2 (22.1 to 30.7)</td>
<td>0.13</td>
</tr>
<tr>
<td>Low fat diet (%)</td>
<td>36.1 (31.3 to 41.2)</td>
<td>27.0 (23.0 to 31.5)</td>
<td>0.01</td>
</tr>
<tr>
<td>Manual social class (%)</td>
<td>50.9 (45.3 to 56.4)</td>
<td>58.6 (53.9 to 63.3)</td>
<td>0.001</td>
</tr>
</tbody>
</table>

*Geometric means; HDLc, high density lipoprotein cholesterol; LDLc, low density lipoprotein cholesterol.*
Sex differences in body fat distribution

explained by differences in luminal diameter. However, this has been shown not to be the case in the large Tromso study. There was a differential in the response between women and men in our study and a difference in the nature of the sampling frame for women and men. Men were survivors of a prospective cohort, whereas women were a newly selected random sample. While among the men responders did not appear to differ from non-responders with respect to age, women who responded were younger than those who did not and women were on average slightly younger than men in the final study sample. All results presented in this study are adjusted for age. Mean blood pressure, lipid levels, and waist-hip ratios for both women and men from this study are similar to those for similar aged women and men from the health survey for England, and the distributions of IMT for women and men in this study were similar to those of similar aged women and men in other studies. However, the difference in social class distributions between women and men and the lower low density lipoprotein cholesterol level among men compared with women in this study may indicate that there was some selection bias in the study sample. Such selection bias would only importantly bias our results if men with small waist-hip ratios but increased IMT had selectively died during the 20 years follow up of the British regional heart study. Such survivorship bias would be unlikely to affect women who have much lower mortality at these ages. While adjustment for lifestyle factors did not importantly reduce the sex difference in IMT it is possible that our assessment of some of these exposures, for example physical activity and low fat diet, was too crude to allow for full adjustment. Our results are consistent with other prospective studies with CHD outcomes that suggest that sex differences in these lifestyle factors do not explain sex differences in CHD occurrence. Most (98%) of the cohort are white. While this means that the results are not necessarily generalisable to other ethnic groups it does mean that the sample is homogenous with respect to ethnicity and therefore not confounded by differences between ethnic groups.

Our findings suggest an important role for abdominal obesity in explaining sex differences in CHD risk and are consistent with an earlier study showing that adjustment for waist-hip ratio removes the sex difference in CHD occurrence. However, our findings are rather more convincing of such a role than those of the ARIC study, in which adjustment for waist-hip ratio attenuated the sex difference in IMT though did not remove it. The differences between the two studies may reflect the play of chance, or may be related to the comparatively younger age (45–64 years) of the ARIC population. In addition to waist-hip ratio, diabetes, hyperglycaemia, and other components of the metabolic syndrome seem to account for at least some of the sex difference in CHD risk. However, in the ARIC study adjustment for hyperglycaemia and diabetes had no effect on the sex difference, which is surprising.

Implications

There is a pronounced sex difference in body fat distribution and our results suggest that this has a role in explaining sex differences in arterial atherosclerosis and by implication may be important in explaining some of the sex difference in CHD occurrence. Why are men more prone to central adiposity? It has been suggested that oestrogen protects younger women from CHD via an influence on body fat distribution. In our study we found that waist-hip ratio increased in women with increasing age but we found no association between age and waist-hip ratio in men. However, all women in our study were post-menopausal and age specific trends show that the decrease in the sex difference in CHD mortality with increasing age is attributable to a deceleration in the male rate rather than an increase in the female rate after the menopause. The role of male sex hormones in explaining differences in body fat distribution and CHD occurrence is also unclear. Secular trends in CHD mortality suggest that environmental factors are the main drivers of the sex difference in CHD occurrence. Alcohol consumption, smoking, and physical inactivity have all been found to increase waist-hip ratio. In our study adjustment for these factors had only a small effect on the sex difference in IMT. Within each sex there is considerable variation in waist-hip ratio and while these factors may be important predictors of abdominal obesity, in general they may not fully explain the sex difference in body fat distribution. A recent study of trends in childhood obesity in Britain found that increases in waist circumference among children over the past 30 years greatly exceeded those in body mass index. Furthermore, it was found that increases in waist circumference among girls over the time period were greater than those seen among boys. If these trends in childhood central obesity are reflected in similar later adult trends then they could result not only in a slowing in the declines in CHD seen over the past three decades in most industrialised countries, but also a reduction in the sex difference, not because of a positive reduction in men but because of an increase in women.

ACKNOWLEDGEMENTS

We thank the Stroke Association for funding this study and the Department of Health and the British Heart Foundation for programme grant funding of the British Regional Heart Study. We also thank the CDER Trust for funding the ultrasound equipment. We are grateful to the two general practices that collaborated in the field work, to our field team (Stella Barlow RGN, Annalise Hamilton RGN, Lucy Lennon BSc) who carried out the field work and all the participants of the British regional heart study.

Table 2 Effect of adjustment for central adiposity and other risk factors on the sex difference in carotid intima media thickness. Results show the percentage difference between men and women in carotid IMT (95% CI). *These proportionate differences are obtained from the linear regression of log IMT (IMT measurements were skewed but log normal) on sex. The resulting exponent of the regression coefficient is the ratio of geometric means of male:female IMT, which is expressed as a proportionate (%) difference. †Interaction term with age included in first model (waist to hip ratio). ‡Metabolic risk factors: systolic blood pressure, diastolic blood pressure, glycated haemoglobin, high density lipoprotein cholesterol, triglycerides. §Lifestyle risk factors: smoking, alcohol consumption, physical activity, low fat diet.
CONTRIBUTORS
DAL, SE, and GDS developed the idea for this study and SE, PW, GW, and OP set up and organised data collection for the study. SD and MG undertook the arterial ultrasound scans and ANN interpreted the arterial ultrasound videos. DAL and JS undertook the statistical analysis. DAL wrote the first draft of the paper and all authors have contributed to the final version. DAL and PW will act as guarantor.

Authors’ affiliations
D A Lawlor, S Ebrahim, J Sterne, G Davey Smith, Department of Social Medicine, University of Bristol, Bristol, UK
P Whincup, Department of Public Health Sciences, St George’s Hospital Medical School, London, UK
O Papacosta, G Wannamethee, Department of Primary Care and Population Sciences, Royal Free Hospital School of Medicine, London, UK
S Dhanjil, M Griffin, A N Nicolaides, Division of Surgery, Anaesthetics and Intensive Care, Irvine Laboratory, Imperial College School of Medicine, St Mary’s Hospital, London, UK
A N Nicolaides, Department of Neurovascular Sciences, The Cyprus Institute of Neurology and Genetics, Nicosia, Cyprus

Funding: this study was funded by the Stroke Association. The Department of Health and British Heart Foundation have provided funding for the British regional heart study. The CDER Trust provided funding for the ultrasound equipment used in this study. DAL was funded by a Medical Research Council/Department of Health (UK) research training fellowship at the time that this work was undertaken and is now funded by a Department of Health (UK) Career Scientist Award. The opinions expressed are those of the authors.

Conflicts of interest: none declared.

REFERENCES