Skinfold thickness, body mass index and ischaemic heart disease

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ABSTRACT
Study objective: To determine the relationship between obesity and subsequent incidence of ischaemic heart disease (IHD).
Design: Prospective cohort survey.
Setting: Study of three occupational groups, with follow up examinations.
Subjects: 3500 people recruited between 1972 and 1978 (80% response rate), and followed up between 1978 and 1984. This report is based on subgroup of 1511 white men aged 40–64 at entry.
Measurements and main results: Information was obtained on smoking and family history of IHD. Blood pressure, weight, height, skinfold thickness at four sites, fibrinogen, factor VII activity and cholesterol were measured during follow up. Body mass index (BMI) was used as an index of obesity. BMI was found to be more strongly correlated with IHD than any of the skinfold measurements, none of which was significantly associated with IHD when BMI was allowed for. Increase in BMI by 1 SD (~8 kg) was associated with a 44% increase in the risk of IHD. Of the four skinfolds, subcapular was the most closely associated with risk, confirming the relevance of central obesity. The association between obesity and IHD remained when possible mechanisms for its effects were taken into account, and its strength may increase with time: for 1 SD increase in BMI, risk of events within 5 years was increased by 28%, while risk of events after longer than 5 years was increased by 65%.
Conclusions: Preventive strategies for IHD should include avoidance of obesity.

Many studies show an association between obesity and the risk of ischaemic heart disease (IHD), although some have not found a relationship. Most studies which have found an association have concluded that the adverse effect of obesity is mediated through mechanisms such as blood pressure or cholesterol. Another pathway may be through haemostatic variables.

There has recently been considerable interest in the suggestion that centrally located subcutaneous fat is particularly implicated in the causation of IHD. Waist/hip ratio has been shown to be associated with IHD in prospective studies, when body mass index was taken into account. A recent study has suggested that skinfold thickness, and in particular subcapular skinfold, is more closely associated with risk of subsequent heart disease than body mass index.

We have compared the relationships of four separate skinfold thicknesses and body mass index with the risk of IHD. We have also assessed the extent to which the association between obesity and IHD appears to act through various mechanisms.

Methods
The main aim of the Northwick Park Heart Study is to investigate the thrombotic component of ischaemic heart disease by including measures of haemostatic function along with other indices of risk. The design, conduct and principal results have been reported elsewhere. In brief, 3500 people were recruited between 1972 and 1978 from three occupational groups in north west London. The response rate was about 80%. Follow up took place between 1978 and 1984. This report is based on the 1511 white men aged 40–64 at entry.

Participants were interviewed by trained nurses in a standard fashion. Information on smoking habit was obtained and blood pressure was measured using a standard sphygmomanometer with a 12 × 23 cm bladder. A blood sample was taken, if possible without venecompression. Fibrinogen, factor VII activity and cholesterol were assayed using previously described techniques. Weight and height were measured on a Steven’s beam balance with height attachment, the
subject wearing light clothing. Body mass index (weight divided by height squared, in kg/m²) was used as an index of obesity.

Participants were interviewed by a doctor in a standard way to determine whether there was any history of IHD. Skinfold thickness was measured at four sites—forearm, triceps, subcapular and suprailliac—using Holtain skinfold calipers as previously described. Three doctors took the majority of the measurements.

Full details of follow up procedures for mortality and IHD events have already been described. The term "all ischaemic heart disease" refers to the sum of fatal and non-fatal events in persons without a history of myocardial infarction at entry, ie, it excludes 52 men who had had an MI at recruitment.

Survival analysis using Cox’s proportional hazards model were carried out with the skinfold thicknesses and body mass index as covariates, along with age and smoking status. These were entered into the regression first separately and then together. In further analyses, cholesterol, systolic blood pressure, fibrinogen and factor VII activity were also included as covariates. The proportionality assumption was checked by including a quadratic term as a covariate and a time dependent term. There was no significant departure from this assumption. The distributions of skinfold thickness and body mass index are skewed. However, since the use of log transformed data in the analyses made very little difference to the results, the results using untransformed data are presented.

The skinfold thickness results were adjusted for age and observer effects prior to survival analysis by using coefficients derived from multiple regression analysis. The regression coefficients from the survival analysis have been standardised so that the standardised regression effect is equivalent to the change in relative risk associated with a 1 SD change in the variable under consideration.

Results

The mean levels and standard deviations of the unadjusted skinfold thicknesses and body mass index (BMI) at entry to the study by categories of IHD event are shown in table 1.

A comparison of the relative strength of association between the skinfold thicknesses and risk of IHD, and between BMI and risk of IHD is made in table 2. The relative risk of IHD by low, middle and high thirds of the distributions of these measures of obesity is shown. The gradient of risk was steepest for BMI, and then for subscapular skinfold, both for fatal IHD and all IHD.

The standardised regression effects (table 3) also show that BMI was most strongly associated with IHD, followed by subscapular skinfold thickness, after allowing for effects of age and smoking. For a 1 SD increase in BMI (on average 8 kg weight), the risk of an IHD event was 44% greater, while for a 1 SD increase in subscapular skinfold it was 25% greater.

There was a suggestion that the relationship between BMI and all IHD became stronger with time (p = 0.07 for the interaction term). The standardised regression effect for all IHD events within 5 years was 1.28, while for events beyond 5 years it was 1.65. For all IHD events occurring within 5 years of follow up, mean BMI was 26.2, while for events occurring later it was 27.0. Similar results hold for fatal IHD alone.

There was no significant variation with age in the association between BMI and IHD (fatal or non-fatal). The mean BMI in men aged 40–59 having IHD events was 26.8 (25.4 in survivors), while for men aged 60–64 it was 26.1 (25.5 in survivors). Similar results held for fatal IHD alone.

Only BMI was significantly associated with the risk of death from any cause. Mean levels in those dying of cancer (n = 48) or from other causes (n = 27) were 25.4 and 24.6 respectively, compared with 26.9 for fatal IHD.

Multiple regression analyses were performed including all the skinfold measures together. For all IHD, only the subscapular measurement approached significance (p = 0.07). When BMI was also included none of the skinfolds were significantly associated

Table 1  Values for skinfold thickness and body mass index at entry, by categories of IHD event. Results are means (SD)

<table>
<thead>
<tr>
<th></th>
<th>Skinfolds (mm)</th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Forearm</td>
<td>Triceps</td>
<td>Subscapular</td>
<td>Suprailliac</td>
</tr>
<tr>
<td>Still alive and no IHD at follow up (n=1280)</td>
<td>5.9 (2.2)</td>
<td>9.8 (3.8)</td>
<td>16.6 (6.4)</td>
<td>13.1 (6.5)</td>
</tr>
<tr>
<td>Fatal IHD* (n=59)</td>
<td>6.0 (2.4)</td>
<td>10.5 (4.1)</td>
<td>19.1 (8.0)</td>
<td>13.0 (7.1)</td>
</tr>
<tr>
<td>Non-fatal IHD* (n=60)</td>
<td>5.9 (1.8)</td>
<td>10.1 (3.7)</td>
<td>17.2 (7.1)</td>
<td>13.5 (7.0)</td>
</tr>
<tr>
<td>All IHD (n=109)</td>
<td>5.9 (2.1)</td>
<td>10.2 (3.8)</td>
<td>17.8 (7.1)</td>
<td>13.0 (6.6)</td>
</tr>
</tbody>
</table>

* Ten men had an episode of non-fatal IHD followed by a fatal episode.
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Table 2  Relative risks with (95% confidence intervals) of fatal IHD and all IHD by tertiles of skinfold thickness and body mass index

<table>
<thead>
<tr>
<th>Tertile</th>
<th>Forearm</th>
<th>Triceps</th>
<th>Subscapular</th>
<th>Suprailliac</th>
<th>Body mass index</th>
</tr>
</thead>
<tbody>
<tr>
<td>Low</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>Middle</td>
<td>(0.71-2.34)</td>
<td>(0.71-2.63)</td>
<td>(0.97-3.71)</td>
<td>(0.72-2.30)</td>
<td>(0.96-4.29)</td>
</tr>
<tr>
<td>High</td>
<td>(0.57-2.06)</td>
<td>(0.91-3.24)</td>
<td>(1.02-3.88)</td>
<td>(0.46-1.72)</td>
<td>(1.49-6.13)</td>
</tr>
</tbody>
</table>

Fatal IHD (n=59)

<table>
<thead>
<tr>
<th>Tertile</th>
<th>Forearm</th>
<th>Triceps</th>
<th>Subscapular</th>
<th>Suprailliac</th>
<th>Body mass index</th>
</tr>
</thead>
<tbody>
<tr>
<td>Low</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>Middle</td>
<td>(0.91-2.20)</td>
<td>(0.77-1.95)</td>
<td>(0.97-2.46)</td>
<td>(0.72-1.69)</td>
<td>(1.24-3.60)</td>
</tr>
<tr>
<td>High</td>
<td>(0.84-2.11)</td>
<td>(0.90-2.22)</td>
<td>(1.04-2.63)</td>
<td>(0.64-1.57)</td>
<td>(1.70-4.74)</td>
</tr>
</tbody>
</table>

All IHD (n=109)

| Tertile cutting points (mm): Forearm 4.7, 6.5; Triceps 7.9, 10.7; Subscapular 13.3, 18.5; Suprailliac 9.5, 15.1; Body mass index (kg/m²) 24.3, 26.6

Table 3  Standardised regression effects (SRE),* with (95% confidence intervals) and p values for fatal IHD, all IHD and total mortality of separate skinfold thicknesses and body mass index

<table>
<thead>
<tr>
<th>Skinfolds (mm)</th>
<th>Forearm</th>
<th>Triceps</th>
<th>Subscapular</th>
<th>Suprailliac</th>
<th>Body mass index (kg/m²)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fatal IHD SRE</td>
<td>1.18</td>
<td>1.31</td>
<td>1.44</td>
<td>1.16</td>
<td>1.62</td>
</tr>
<tr>
<td>p value</td>
<td>0.02</td>
<td>0.03</td>
<td>0.002</td>
<td>0.3</td>
<td>0.0001</td>
</tr>
<tr>
<td>All IHD SRE</td>
<td>1.14</td>
<td>1.20</td>
<td>1.25</td>
<td>1.14</td>
<td>1.44</td>
</tr>
<tr>
<td>p value</td>
<td>0.2</td>
<td>0.05</td>
<td>0.01</td>
<td>0.2</td>
<td>0.0001</td>
</tr>
<tr>
<td>All deaths SRE</td>
<td>0.96</td>
<td>1.13</td>
<td>1.14</td>
<td>0.96</td>
<td>1.21</td>
</tr>
<tr>
<td>p value</td>
<td>0.7</td>
<td>0.2</td>
<td>0.6</td>
<td>0.6</td>
<td>0.02</td>
</tr>
</tbody>
</table>

* For definition see text.

with IHD. Conversely, body mass index remained significantly associated with IHD after all the skinfolds were added (p=0.0006).

For fatal IHD, only subscapular skinfold thickness was significantly associated (p=0.002). When BMI was added there was a weak negative association with suprailliac skinfold (p=0.04), but there was no association with the subscapular measurement. BMI remained significantly associated after all the skinfolds were added (p=0.003).

To assess the independent effect of obesity on the risk of IHD the survival analyses were repeated, including factor VII, cholesterol, systolic blood pressure and fibrinogen as covariates. For all IHD, BMI remained significantly associated (p=0.002) with a slightly reduced SRE 1.36 (cf 1.44). The SRE was reduced by a similar small amount (between 1 and 3%) as each covariate entered the regression. Systolic blood pressure was associated with the largest reduction. For subscapular skinfold the association was no longer significant (p=0.1) and the SRE was reduced to 1.16 (cf 1.25).

In the case of fatal IHD, the association of body mass index with IHD risk was still significant (p=0.001) with an SRE of 1.57 (cf 1.62). In the regression, cholesterol was associated with the largest reduction in the SRE. Subscapular skinfold (p=0.02) was also independently associated, with an SRE 1.34 (cf 1.44).

For events of IHD within 5 years, after inclusion of
these risk factors the SRE for BMI was reduced to 1.19 (p=0.2). For events beyond 5 years the SRE was reduced to 1.60 (p=0.001).

Discussion

An association between obesity and the risk of IHD was originally found in life insurance data. The results of epidemiological studies carried out more recently have been equivocal. In some, no significant association was found, in others the association was no longer evident after taking risk factors such as blood pressure or cholesterol into account. Some have shown a link independent of the risk factors measured. Our results show a clear link between obesity and IHD which is independent of its association with established risk factors.

The apparently conflicting findings of different studies may arise from differences in the range and degree of obesity, the ages of the cohorts, or the length of follow up. For example, many of the populations in the Seven Countries Study were, on average, leaner than the Northwick Park Heart Study population, which is comparable to the rest of the UK in terms of obesity. As others have shown, the Northwick Park Study found the contribution of obesity to IHD risk is strongest in younger men and over greater lengths of follow up although these associations were not statistically significant. Results of the Whitehall Study showed that initially there was no relationship between obesity and IHD in men aged 40–49 but that after 10 years of follow up there was a significant association.

Obesity and cholesterol, factor VII, fibrinogen and blood pressure are associated in cross sectional data and changes in obesity result in changes in these variables, with the possible exception of fibrinogen. Our results suggest that although obesity may predispose to IHD partly through these mechanisms these do not appear to be solely responsible for its association with IHD.

We have shown that body mass index is more closely associated with subsequent IHD (both fatal and non-fatal) than skinfold thickness. Skinfold thickness is subject to greater measurement error than weight and height, but we have attempted to adjust for systematic variation between observers. Subscapular skinfold is the most closely related of the four skinfolds to IHD. This may be further evidence for the role of centrally located subcutaneous fat in the causation of IHD.

Our results, from a contemporary study of a UK population, suggest that the reduction of obesity should be given more prominence in cardiovascular disease prevention.

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