Diabetes, glaucoma, sex, and cataract: analysis of combined data from two case control studies

John J Harding, Muriel Egerton, Ruth van Heyningen, Ruth S Harding

Abstract

Data from two case control studies in Oxfordshire were combined and analysed. The combined study covered 1940 subjects, 723 cases, and 1217 controls, between the ages of 50 and 79 with a response rate of 97% for cases and 94% for controls. Diabetes was shown to be a powerful and highly significant risk factor for cataract with a relative risk of 5-04. More than 11% of cataracts in Oxfordshire are attributable to diabetes. The relative risk did not increase significantly with age within the range 50 to 79 years but was higher in females than in males. For females with diabetes the relative risk was 7-85 with 95% confidence interval from 4-30 to 14-3 compared with 3-42 with confidence interval from 2-05 to 5-71 for males with diabetes. Diabetes remained a powerful risk factor when other identified risk factors had been controlled for. No known mechanism for the development of diabetic complications provides an explanation for the excess risk in females. Combination of the two studies led to better estimates of the relative risk of glaucoma as a risk factor for cataract (3-96 with 95% confidence interval from 2-35 to 6-68). The relative risk appeared to be greater in women than in men but this difference was not statistically significant. There was no significant change in risk with age. Glaucoma is a powerful and independent risk factor for cataract in both sexes and may be responsible for 5% of all cataracts in our area.

(Clinical science)

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Cataract is an important complication of diabetes, and diabetes is the most important risk factor for cataract identified in Western countries. Diabetes has been reported to increase the risk of cataract up to 12-fold in different populations.1-4 Further evidence that diabetes can cause cataract is derived from observations of cataract development in experimental diabetes and from in vitro studies of the effects of high sugar concentrations on incubated lenses and lens proteins.5-9

In some of the epidemiological studies the subjects were subdivided by sex for analysis and this indicated an enhanced risk of cataract in females with diabetes than in males with diabetes in the HANES study from the United States,7 in Denmark,9 and in England.9-11 The confidence intervals for each sex, where given, however, were wide and the difference between the sexes was not statistically significant. For example in our own studies the relative risks (and confidence intervals) for diabetic males and females were 3-2 (1-3 to 7-6) and 10-5 (4-5 to 24-6) respectively in the first study,9,11 and 3-4 (1-8 to 6-4) and 6-0 (2-6 to 14-2) in the second study.9

Individually each of these results identified diabetes as a highly significant and powerful risk factor in males and females separately, but the difference between the sexes was not significant, although there were 300 cataract patients in the first study and 423 in the second.

The possibility that glaucoma or treatment of glaucoma leads to cataract has been investigated for many years with the more recent emphasis on glaucoma surgery as a cause of cataract.12-16 It has been estimated that glaucoma increases the risk of cataract by up to sixfold in different populations over the age of 50.14-16 No difference between the sexes was reported.14

A claimed decrease in relative risk with age1 was not entirely convincing above the age of 50 and very few glaucoma patients were identified below that age.

Glaucoma was identified as a risk factor for cataract in the two case control studies in Oxfordshire with relative risks of 5-95 with a 95% confidence interval from 2-6 to 13-5 in the first,9 and relative risk of 2-9 with a 95% confidence interval from 1-5 to 5-7 in the second.9 The relative risks seem different but the confidence intervals overlap.

These two studies covered almost 1000 patients each. Both were based on interviews of all the patients and a larger group of age and sex-matched controls. Some questions were the same in both studies and so it has been possible to pool the data on diabetes, glaucoma, and on several other major risk factors from the two studies, giving information on a total of 1940 subjects. The large numbers should permit a more accurate assessment of the relative risk for diabetes and glaucoma, an exploration of differences in risk between the sexes and between age groups, and the assessment of attributable risk. This paper reports the analysis of data from the combined studies in relation to the association between diabetes, glaucoma, and cataract. It has been possible to establish clearly for the first time that the risk of cataract associated with diabetes is greater in females than in males.
Subjects and methods

The subjects, 723 cases and 1217 controls, were pooled from the two original case control studies of cataract in Oxfordshire. Cases were those aged 50 to 79 years admitted to the Oxford Eye Hospital for cataract extraction. Controls were recruited both from other hospital departments and from the age-sex registers of general practitioners. As the Oxford Eye Hospital is a National Health Service hospital with a catchment population of about 560 000 and the vast majority of the population is registered with general practitioners, the cases and controls in the two studies were thought to be representative of all cataract patients, and of the general population respectively. The controls were chosen to give an age-sex distribution that matched the cases. The overall response rates were 97% for cases and 94% for controls with the non-responders including those too ill to be interviewed and those who spoke no English.

The cases and controls were interviewed in the same way by the same interviewers, using the same forms. The questionnaires included questions on medical history and ocular history, and on drug therapy. In the second study self-reported glaucoma was confirmed by examination of the hospital notes after completion of interviewing. This supported the reliability of the self-reporting technique.

In relation to this paper information on diabetes was ascertained when patients were asked if they had ever suffered a serious illness and if they had been admitted to hospital. Positive responses were followed up and diabetes was coded on the questionnaire. In the second study those reporting diabetes were asked its duration. Confirmation of diabetes was not sought. Blood analysis would have been difficult when subjects were interviewed at home up to 30 miles (50 km) from Oxford and would have decreased our response rate. Confirmation via clinical notes would have been invalid as different sources, hospital and general practitioners, would have had to be used for different subjects leading to ascertainment bias. The similarity in the results between the two studies and between the different control groups in each study indicates that we were getting unbiased data for known diabetes.

Data were entered into the university’s Vax computer. In the present study data files from the two original studies were combined before analysis initially by the χ2 test for categorical variables using the SAS statistical package (SAS Institute Inc, Cary, NC, USA). Odds ratios were calculated from contingency tables and taken as valid estimates of relative risk. Confidence intervals and attributable risks were calculated as described.1 Interactions between risk factors were explored using generalised linear modelling (GLIM: Royal Statistical Society, London).

Results

The control subjects were partly hospital controls and partly community controls. Each control group, four groups in the first study and three in the second, was matched to the cases for age and sex distribution so the overall match is very close (Table 1). Results for the different control groups were similar in the two original studies and so all were pooled for this combined analysis.

Diabetes appeared as a powerful risk factor in

Table 1 Number (%) of male and female cases and controls in each age group

<table>
<thead>
<tr>
<th>Age groups</th>
<th>50–54</th>
<th>55–59</th>
<th>60–64</th>
<th>65–69</th>
<th>70–74</th>
<th>75–79</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Males:</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Controls</td>
<td>13 (2-8)</td>
<td>32 (9-8)</td>
<td>53 (14-7)</td>
<td>80 (17-2)</td>
<td>151 (27-4)</td>
<td>148 (26-8)</td>
</tr>
<tr>
<td>Cases</td>
<td>16 (2-8)</td>
<td>47 (14-7)</td>
<td>60 (16-7)</td>
<td>89 (22-5)</td>
<td>133 (27-2)</td>
<td>129 (26-5)</td>
</tr>
<tr>
<td><strong>Females:</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Controls</td>
<td>26 (3-8)</td>
<td>49 (7-4)</td>
<td>93 (14-6)</td>
<td>305 (17-3)</td>
<td>172 (25-9)</td>
<td>196 (29-5)</td>
</tr>
<tr>
<td>Cases</td>
<td>26 (6-8)</td>
<td>50 (13-1)</td>
<td>63 (16-5)</td>
<td>108 (27-0)</td>
<td>124 (32-3)</td>
<td>121 (32-5)</td>
</tr>
</tbody>
</table>

Table 2 Diabetes as a risk factor for cataract

<table>
<thead>
<tr>
<th></th>
<th>Controls</th>
<th>Cases</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>No diabetes</td>
<td>1179</td>
<td>622</td>
<td>1801</td>
</tr>
<tr>
<td>Diabetes</td>
<td>38</td>
<td>101</td>
<td>139</td>
</tr>
<tr>
<td>Total</td>
<td>1217</td>
<td>723</td>
<td>1940</td>
</tr>
<tr>
<td>Percent positive</td>
<td>3-1</td>
<td>1-3</td>
<td>4-5</td>
</tr>
</tbody>
</table>

χ²=80-2; p<0.001. Relative risk=5-04; 95% confidence interval 3-43 to 7-41.

Table 3 Diabetes as a risk factor in males

<table>
<thead>
<tr>
<th></th>
<th>Controls</th>
<th>Cases</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>No diabetes</td>
<td>528</td>
<td>296</td>
<td>824</td>
</tr>
<tr>
<td>Diabetes</td>
<td>24</td>
<td>46</td>
<td>70</td>
</tr>
<tr>
<td>Total</td>
<td>552</td>
<td>342</td>
<td>894</td>
</tr>
<tr>
<td>Percent positive</td>
<td>4-4</td>
<td>3-4</td>
<td>3-8</td>
</tr>
</tbody>
</table>

χ²=24-2; p<0.001. Relative risk=3-42; 95% confidence interval 2-05 to 5-71.

Table 4 Diabetes as a risk factor in females

<table>
<thead>
<tr>
<th></th>
<th>Controls</th>
<th>Cases</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>No diabetes</td>
<td>651</td>
<td>326</td>
<td>977</td>
</tr>
<tr>
<td>Diabetes</td>
<td>60</td>
<td>55</td>
<td>99</td>
</tr>
<tr>
<td>Total</td>
<td>665</td>
<td>381</td>
<td>1046</td>
</tr>
<tr>
<td>Percent positive</td>
<td>2-1</td>
<td>1-4</td>
<td>1-8</td>
</tr>
</tbody>
</table>

χ²=59-8; p<0.001. Relative risk=7-85; 95% confidence interval 4-30 to 14-3.

Table 5 Diabetes as a risk factor for cataract in subjects aged 50 to 69 years

<table>
<thead>
<tr>
<th></th>
<th>Controls</th>
<th>Cases</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>No diabetes</td>
<td>537</td>
<td>267</td>
<td>804</td>
</tr>
<tr>
<td>Diabetes</td>
<td>13</td>
<td>43</td>
<td>56</td>
</tr>
<tr>
<td>Total</td>
<td>550</td>
<td>310</td>
<td>860</td>
</tr>
<tr>
<td>Percent positive</td>
<td>2-4</td>
<td>3-9</td>
<td>6-3</td>
</tr>
</tbody>
</table>

χ²=43-1; p<0.001. Relative risk=6-65; 95% confidence interval 3-52 to 12-6.

Table 6 Diabetes as a risk factor in subjects aged 70 to 79 years

<table>
<thead>
<tr>
<th></th>
<th>Controls</th>
<th>Cases</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>No diabetes</td>
<td>642</td>
<td>355</td>
<td>994</td>
</tr>
<tr>
<td>Diabetes</td>
<td>25</td>
<td>58</td>
<td>83</td>
</tr>
<tr>
<td>Total</td>
<td>667</td>
<td>413</td>
<td>1080</td>
</tr>
<tr>
<td>Percent positive</td>
<td>3-8</td>
<td>14-0</td>
<td>8-0</td>
</tr>
</tbody>
</table>

χ²=38-1; p<0.001. Relative risk=4-20; 95% confidence interval 2-58 to 6-83.

Table 7 Diabetes as a risk factor for cataract in three age groups

<table>
<thead>
<tr>
<th>Age</th>
<th>Relative risk</th>
<th>Confidence interval p</th>
</tr>
</thead>
<tbody>
<tr>
<td>50–59</td>
<td>12-6</td>
<td>2-76–57-9</td>
</tr>
<tr>
<td>60–69</td>
<td>5-56</td>
<td>2-74–11-3</td>
</tr>
<tr>
<td>70–79</td>
<td>4-20</td>
<td>2-58–6-83</td>
</tr>
</tbody>
</table>

Breslow-Day homogeneity test: χ²=2-1; p=0-35.
the combined study with $\chi^2$ higher than before and the confidence interval somewhat narrower (Table 2). The relative risk of 5.04 compares with 6.2 found in the first Oxford study and 4.2 found in the second. In the separate studies the relative risks for female diabetics were two to three times those for male diabetics. Analysing the combined data diabetics are a powerful risk factor for cataract in both males (Table 3) and females (Table 4) separately. Both confidence intervals are well above unity showing a clear risk association. The difference between the sexes was tested by the Breslow-Day test for homogeneity which gave $\chi^2 = 4.31$ and $p = 0.038$ showing that the relative risks are significantly different for males and females with diabetes. The interaction between sex and diabetes determined in a GLIM analysis was also statistically significant ($\chi^2 = 4.11, p < 0.05$). The excess risk for females with diabetes over that for males with diabetes was 2.294 (95% confidence interval = 1.023 to 5.143). In other words diabetes in females is a more powerful risk factor for cataract than diabetes in males. Nevertheless diabetes is a risk factor in both sexes.

Some earlier studies have determined the relative risk or odds ratio for diabetes as a risk factor for cataract at different ages and claimed a decline with age though without showing the age-related decline in risk was statistically significant. The numbers of cataract patients and controls in this combined analysis have enabled us to look again at the age relationship. First we divided the subjects into two groups of similar size: ages 50 to 69 and ages 70 to 79. Diabetes was a highly significant risk factor for cataract in each of these age groups (Tables 5 and 6) with a relative risk of 6.7 in the younger group falling to 4.2 in the older group. The Breslow-Day test of homogeneity shows that the difference between the groups is not statistically significant ($\chi^2 = 1.28; p = 0.26$). Subdividing by decades gives an impression of an age-related fall in risk but the numbers become small, the confidence intervals wide, and the homogeneity test showed no difference between the three relative risks (Table 7). With 723 cataract patients and 1217 controls there is an apparent age-related decline in risk of cataract associated with diabetes but the decline is not significant. The risk in each age group separately is highly significant.

Although diabetes emerges as a powerful risk factor in the approach described above it was necessary to examine the effect of other risk factors on the risk associated with diabetes using a linear modelling program. The risk factors that remained significant and were incorporated into the final model were diabetes, glaucoma, severe diarrhoea, and myopia. Analysis for glaucoma follows in this paper and that for the other factors will be published separately. The four factors did not interact significantly with each other and the presence of the other three in the model made little difference to the relative risk associated with diabetes - final value 5.10 with a 95% confidence interval from 3.43 to 7.58 (compare Table 2). The self-reporting of glaucoma was checked and confirmed by examination of hospital notes. Glaucoma emerged as a powerful risk factor in the combined study increasing the risk of cataract fourfold (Table 8). As earlier studies had not established a possible excess risk in females with glaucoma, we analysed the data for the two sexes separately. Glaucoma emerged as a powerful risk factor for cataract in males alone (Table 9) and in females alone (Table 10) with relative risks of 2.6 and 6.6 respectively. The relative risks indicated that females with glaucoma might experience a greater risk of cataract than males with glaucoma, but when the data were analysed for homogeneity between the sexes it just failed to reach statistical significance (Table 10). In a similar way we subdivided the data by age to identify any age-related change in risk. This is not to investigate aging as a risk factor which is not possible in an age-matched study such as this. Glaucoma appeared as a powerful risk factor for cataract in those aged less than 70 (Table 11; relative risk = 2.9) and in the older group (Table 12; relative risk = 4.5). The Breslow-Day test for homogeneity gave $p = 0.45$.
indicating that the apparent increased risk in
glaucoma patients with age was not significant.

The possibility that glaucoma appears as a risk
factor only because it is associated with other
factors was examined in the two original
studies.\textsuperscript{18,19} It has been re-examined using
the combined data. Four major risk factors remained
in the final model: glaucoma, myopia, diabetes,
diarrhoea. There was no confounding with

glaucoma and indeed the relative risk was close to
the uncorrected value given in Table 8: 3·90
with a confidence interval from 2·26 to 6·71.

Glaucoma is a powerful, independent, and highly
significant risk factor for cataract.

It is clear that diabetes and glaucoma are
powerful risk factors increasing the risk of
cataract fivefold and fourfold in the population as
a whole, but it is also of interest to know what
proportion of cataract is attributable to these
factors. We have examined this by calculations of
attributable risk.\textsuperscript{19} This has seldom been calculated.
The figure for diabetes from the combined
data in Table 2 is 11·2%, and values of population
attributable risk for males, females, older, and
younger subjects are between 10·7% and 12·9% with
little difference between them for the
groups in Tables 3 to 6. This indicates that about
11% of all cataracts would be prevented if
diabetes, or at least the cataractogenic effect of
diabetes, were eliminated. This makes diabetes
the most important risk factor for cataract in
Western populations. Finally we calculated the
proportion of cataract in Oxfordshire attributable
to glaucoma. This worked out to 4·9%. In
septuagenarians, in whom glaucoma is more
common and the relative risk appears slightly
higher, the attributable risk is 6·8%.

\section*{Discussion}

The choice of controls and possible sources of
bias were discussed in the papers on the two
separate studies.\textsuperscript{14} Combining those studies has
provided data on a greater number of patients with
severe cataract, sufficiently impairing to
require surgery, than previous studies. This has
provided a more accurate figure for the relative
risk of cataract associated with diabetes, and has
for the first time demonstrated a significant
excess risk in female diabetics over male diabetics.

Cataract in subjects with diabetes in the age
group studied is not the pure diabetic cataract
seen in years gone by but is simply a component
of age-related cataract. Diabetes in experimental
animals produces cataract and many in vitro
experiments bear witness to the damaging effects
of glucose and other sugars on the lens.\textsuperscript{10} The
combination of epidemiological and laboratory
evidence establishes without doubt the causal
relationship from diabetes to cataract. How
diabetes causes cataract is a more open question.
Several routes have been discussed with the most
interest having followed the osmotic effect of
sorbitol and non-enzymic glycosylation (glucos,
\textsuperscript{a}) The sorbitol mechanism presupposes
the accumulation of sorbitol in the lens to levels high
enough to exert an osmotic effect and damage

less than 1 mM) would scarcely balance the
excess glucose in the aqueous humour.\textsuperscript{20} No
significant accumulation of sorbitol is found in
human lens incubated in high glucose media.\textsuperscript{21-23}

Glycation of lens proteins is increased in human
cataracts from diabetic subjects relative to those
from non-diabetics and relative to normal lenses\textsuperscript{24-29} and may therefore appear to be the
more plausible mechanism. There are probably
additional mechanisms, however, because
neither mechanism has been able to explain all the
biochemical changes in lenses of diabetic
animals notably the early loss of glutathione.\textsuperscript{10}

Now we have the additional information that
diabetic females have a greater risk of cataract
than diabetic males. None of the currently
postulated mechanisms for diabetic complications
to help explain this discrepancy between
the sexes. Other diabetic complications do not
appear to affect females more than males; for
example the risk of retinopathy is no greater in
females than in male diabetics.\textsuperscript{30}

One potential problem relevant to this paper is
the use of interviews to identify subjects with

glaucoma, but in one of the original studies this
was confirmed by subsequent examination of the
hospital notes.\textsuperscript{4} All subjects reporting glaucoma
had indeed been treated for glaucoma in Oxford.

As more studies provide evidence associating

glaucoma and cataract it becomes more convinc-
ing that glaucoma in some way causes cataract.
The present analysis shows a highly significant
association indicating a fourfold increase in the
risk of visually impairing cataract in those with

glaucoma. Glaucoma is a powerful risk factor in
both men and women with a possible increase in
risk in females although that excess is not statistically
significant. It remains a powerful risk factor
throughout the age range 50 to 79 years.

Drainage surgery frequently leads to catarac-
t\textsuperscript{11-15,16} and in Oxford most of the risk
associated with glaucoma has been attributed to
surgical procedures.\textsuperscript{3} As 5% of total cataract


can be attributed to glaucoma it seems that almost 5%

of cataract surgery would be avoided if the
trauma of glaucoma surgery could be avoided.

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Hospital, and to local general practitioners for access to their
patients. Useful discussions with European colleagues were made
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