ORIGINAL ARTICLE

The association between tobacco and the risk of asthma, rhinoconjunctivitis and eczema in children and adolescents: analyses from Phase Three of the ISAAC programme

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ABSTRACT

Background Exposure to parental smoking is associated with wheeze in early childhood, but in 2006 the US Surgeon General stated that the evidence is insufficient to infer a causal relationship between exposure and asthma in childhood and adolescents.

Aims To examine the association between maternal and paternal smoking and symptoms of asthma, eczema and rhinoconjunctivitis.

Methods Parents or guardians of children aged 6–7 years completed written questionnaires about symptoms of asthma, rhinoconjunctivitis and eczema, and several risk factors, including maternal smoking in the child’s first year of life, current maternal smoking (and amount) and paternal smoking. Adolescents aged 13–14 years self completed the questionnaires on these symptoms and whether their parents currently smoked.

Results In the 6–7-year age group there were 220 407 children from 75 centres in 32 countries. In the 13–14-year age group there were 350 654 adolescents from 118 centres in 53 countries. Maternal and paternal smoking was associated with an increased risk of symptoms of asthma, eczema and rhinoconjunctivitis in both age groups, although the magnitude of the OR is higher for symptoms of asthma than the other outcomes. Maternal smoking is associated with higher ORs than paternal smoking. For asthma symptoms there is a clear dose relationship (1–9 cigarettes/day, OR 1.27; 10–19 cigarettes/day, OR 1.35; and 20+ cigarettes/day, OR 1.56). When maternal smoking in the child’s first year of life and current maternal smoking are considered, the main effect is due to maternal smoking in the child’s first year of life. There was no interaction between maternal and paternal smoking.

Conclusions This study has confirmed the importance of maternal smoking, and the separate and additional effect of paternal smoking. The presence of a dose–response effect relationship with asthma symptoms suggests that the relationship is causal, however for eczema and rhinoconjunctivitis causality is less certain.

Key messages

What is the key question?

What is the association between maternal and paternal smoking and symptoms of asthma, eczema and rhinoconjunctivitis?

What is the bottom line?

This study has confirmed the association of maternal smoking, and identified a separate and additive association between paternal smoking and the risk of symptoms of asthma, eczema and rhinoconjunctivitis.

Why read on?

The results of this large study (over 570 000 children and adolescents in 53 countries) were consistent throughout the world and were independent of the income of the country.

The effect of environmental tobacco smoke exposure (ETS) on respiratory health has been of interest for many years. In 2006 the US Surgeon General concluded that ‘the evidence is sufficient to infer a causal relationship between secondhand smoke exposure from parental smoking and the onset of wheeze illnesses in early childhood’ and ‘that the evidence is suggestive but not sufficient to infer a causal relationship between secondhand smoke exposure from parental smoking and the onset of childhood asthma’. While a recent systematic review and meta-analysis has quantified the strength of the associations, the data come predominantly from developed countries. The importance of the age of exposure to ETS is not well established, nor is the relative importance of maternal and paternal smoking.

In contrast with the number of studies examining the association of ETS with asthma, there are fewer studies examining the association between ETS and either eczema or rhinoconjunctivitis, and these have provided conflicting results.

The aims of this study were to examine the association between maternal and paternal smoking and symptoms of asthma, rhinoconjunctivitis and eczema, to assess the relative importance of maternal and paternal smoking, to look for consistency in these findings from many different countries, including developing countries, and to determine if there is a dose–response effect, which if present would support a causal relationship.
Asthma

METHODS
International Study of Asthma and Allergies in Childhood (ISAAC) Phase Three is a multicentre, multicountry, cross-sectional study of children (aged 6–7 years) and adolescents (aged 13–14 years) chosen from a random sample of schools in a defined geographical area. The parent or guardian completed the questionnaire for the children and for the adolescents the questionnaire was self-completed. The methods have been described in detail previously.5

Outcomes
Current wheeze was identified by a positive response to the question: ‘Has your child/have you had wheezing or whistling in the chest in the past 12 months?’

Severe asthma symptoms were identified by fulfilling one or more of the following criteria:
1. Four or more attacks of wheezing in the past 12 months.
2. Sleep disturbance due to wheezing occurring one or more nights per week in the past 12 months.
3. Wheezing limiting speech to one or two words at a time.

Asthma (the diagnosis) was defined as a positive response to the question: ‘Has your child/have you ever had asthma?’

Current rhinoconjunctivitis was identified by a positive response to each of the following questions:
1. Does your child/have you ever had an itchy skin rash, which was coming and going for at least 6 months?
2. Does your father (or male guardian) smoke cigarettes?
3. Has your child/have you ever had this itchy rash at any time affected any of the following places—the folds of the elbows, behind the knees, in front of the ankles, under the buttocks, or around the neck, ears, or eyes?

Current eczema was identified by a positive answer to each of the following questions:
1. Has your child/have you had an itchy skin rash in the past 12 months?
2. If yes, has your child/have you had this itchy rash at any time in the past 12 months?
3. If yes, has this itchy rash at any time affected any of the following places—the folds of the elbows, behind the knees, in front of the ankles, under the buttocks, or around the neck, ears, or eyes?

The prevalence of wheeze and asthma is higher in children with rhinoconjunctivitis and eczema,4 and this might account for the association between parental smoking and eczema and rhinoconjunctivitis; therefore, the risk of symptoms of eczema and rhinoconjunctivitis in the absence of current asthma symptoms was also examined.

In addition, a video questionnaire of asthma symptoms was administered to the adolescents. The video question used was the sequence on current wheeze at rest defined as a positive response to the question: ‘Has your breathing been like this in the past year?’7 The video asthma questionnaire was validated against the written questionnaire.5

Explanatory variables
The following questions were used to assess smoking by the parents of the subjects. For children aged 6–7 years:
1. Does your child’s mother (or female guardian) smoke cigarettes?
2. Does your father (or male guardian) smoke cigarettes?
3. If yes, has this itchy rash at any time affected any of the following places—the folds of the elbows, behind the knees, in front of the ankles, under the buttocks, or around the neck, ears, or eyes?

For adolescents aged 13–14 years:
1. Does your mother (or female guardian) smoke cigarettes?
2. Does your father (or male guardian) smoke cigarettes?

Table 1 Association between tobacco smoking by the parents and symptoms of asthma, eczema and rhinoconjunctivitis at 6–7 and 13–14 years of age

<table>
<thead>
<tr>
<th>Symptom</th>
<th>Tobacco question</th>
<th>Age group 6–7 years</th>
<th>Age group 13–14 years</th>
</tr>
</thead>
<tbody>
<tr>
<td>Current wheeze</td>
<td>Current paternal smoking</td>
<td>1.17 (1.12 to 1.21)</td>
<td>1.20 (1.15 to 1.24)</td>
</tr>
<tr>
<td></td>
<td>Current maternal smoking</td>
<td>1.29 (1.22 to 1.34)</td>
<td>1.32 (1.26 to 1.37)</td>
</tr>
<tr>
<td></td>
<td>Mother smoked first year of life</td>
<td>1.36 (1.29 to 1.43)</td>
<td>*</td>
</tr>
<tr>
<td>Current wheeze (video)</td>
<td>Current paternal smoking</td>
<td>*</td>
<td>1.22 (1.15 to 1.29)</td>
</tr>
<tr>
<td></td>
<td>Current maternal smoking</td>
<td>*</td>
<td>1.41 (1.32 to 1.49)</td>
</tr>
<tr>
<td>Severe asthma symptoms</td>
<td>Current paternal smoking</td>
<td>1.23 (1.16 to 1.30)</td>
<td>1.26 (1.20 to 1.32)</td>
</tr>
<tr>
<td></td>
<td>Current maternal smoking</td>
<td>1.32 (1.24 to 1.41)</td>
<td>1.38 (1.31 to 1.45)</td>
</tr>
<tr>
<td></td>
<td>Mother smoked first year of life</td>
<td>1.44 (1.33 to 1.55)</td>
<td>*</td>
</tr>
<tr>
<td>Asthma</td>
<td>Current paternal smoking</td>
<td>1.11 (1.06 to 1.15)</td>
<td>1.08 (1.04 to 1.12)</td>
</tr>
<tr>
<td></td>
<td>Current maternal smoking</td>
<td>1.26 (1.20 to 1.32)</td>
<td>1.20 (1.16 to 1.25)</td>
</tr>
<tr>
<td></td>
<td>Mother smoked first year of life</td>
<td>1.34 (1.27 to 1.42)</td>
<td>*</td>
</tr>
<tr>
<td>Current eczema symptoms</td>
<td>Current paternal smoking</td>
<td>1.09 (1.04 to 1.13)</td>
<td>1.19 (1.14 to 1.25)</td>
</tr>
<tr>
<td></td>
<td>Current maternal smoking</td>
<td>1.15 (1.09 to 1.21)</td>
<td>1.22 (1.16 to 1.28)</td>
</tr>
<tr>
<td></td>
<td>Mother smoked first year of life</td>
<td>1.20 (1.13 to 1.27)</td>
<td>*</td>
</tr>
<tr>
<td>Current rhinoconjunctivitis</td>
<td>Current paternal smoking</td>
<td>1.08 (1.04 to 1.13)</td>
<td>1.15 (1.11 to 1.19)</td>
</tr>
<tr>
<td></td>
<td>Current maternal smoking</td>
<td>1.12 (1.06 to 1.18)</td>
<td>1.20 (1.15 to 1.25)</td>
</tr>
<tr>
<td></td>
<td>Mother smoked first year of life</td>
<td>1.17 (1.10 to 1.24)</td>
<td>*</td>
</tr>
<tr>
<td>Eczema without wheeze</td>
<td>Current paternal smoking</td>
<td>1.06 (1.01 to 1.11)</td>
<td>1.15 (1.09 to 1.20)</td>
</tr>
<tr>
<td></td>
<td>Current maternal smoking</td>
<td>1.10 (1.03 to 1.17)</td>
<td>1.12 (1.05 to 1.18)</td>
</tr>
<tr>
<td></td>
<td>Mother smoked first year of life</td>
<td>1.11 (1.04 to 1.19)</td>
<td>*</td>
</tr>
<tr>
<td>Rhinoconjunctivitis without wheeze</td>
<td>Current paternal smoking</td>
<td>1.04 (0.99 to 1.09)</td>
<td>1.11 (1.07 to 1.16)</td>
</tr>
<tr>
<td></td>
<td>Current maternal smoking</td>
<td>1.05 (0.99 to 1.12)</td>
<td>1.11 (1.06 to 1.16)</td>
</tr>
<tr>
<td></td>
<td>Mother smoked first year of life</td>
<td>1.05 (0.98 to 1.13)</td>
<td>*</td>
</tr>
</tbody>
</table>

Data are OR (95% CI) and are adjusted for gender, language, region and Gross National Income (GNI) category. *Data not collected.
To assess the presence of a dose–response relationship, the number of cigarettes smoked were categorised as 0, 1–9, 10–19 and 20+ per day. In addition, combinations of maternal and paternal smoking were examined for both age groups, and combinations of maternal smoking in the child’s first year of life and current maternal smoking were examined in the 6–7-year age group.

Study sample
For the 6–7-year age group, data were submitted to the ISAAC International Data Centre (IIDC) from 165 centres in 65 countries with 421,543 participants. There were 21 centres in 16 countries (32,732 children) excluded as the response rate was <60% or there were <1,000 participants. There were a further 69 centres in 38 countries (168,404 children) that did not include the risk factor questionnaire. The final risk factor dataset for the younger age group comprised 75 centres from 32 countries with 220,407 children.

For the adolescent group, data from 242 centres in 98 countries with 814,837 participants were submitted to the IIDC. There were nine centres from six countries (16,152 adolescents) excluded as the response rate was <70% or there were <1,000 participants.

Country (% maternal smoking)
- Spain (40%)
- Chile (36%)
- Hungary (33%)
- Poland (32%)
- New Zealand (27%)
- Malta (25%)
- Venezuela (24%)
- Estonia (24%)
- Portugal (24%)
- Canada (24%)
- Uruguay (24%)
- Isle of Man (22%)
- Syrian Arab Republic (21%)
- Japan (19%)
- Belgium (19%)
- Lithuania (13%)
- Mexico (11%)
- Colombia (8%)
- Brazil (8%)
- South Korea (6%)
- Taiwan (5%)
- Kyrgyzstan (4%)
- Panama (4%)
- Indonesia (4%)
- Barbados (3%)
- Iran (3%)
- Singapore (3%)
- Malaysia (2%)
- Thailand (2%)
- Nigeria (1%)
- India (1%)
- Sultanate of Oman (1%)

Figure 1 ORs for the association between current maternal smoking and current wheeze by country in children aged 6–7 years. For every country, the percentage of children exposed to maternal smoking is stated in brackets.
There were a further 111 centres from 59 countries (437,086 adolescents) that did not include the risk factor questionnaire. Four centres did not collect data on current maternal and paternal smoking. The final risk factor dataset for the older age group comprised 118 centres from 53 countries with 350,654 adolescents. In addition, some centres did not collect data on specific risk factor questions or the response rate to the specific question was

**Figure 2** ORs for the association between current maternal smoking and current wheeze by country in adolescents aged 13–14 years (Tokelau excluded). For every country, the percentage of children exposed to maternal smoking is stated in brackets.
<70%. For the 6–7-year age group the following number of centres were excluded: current maternal smoking=1, current paternal smoking=2, mother smoked in child’s first year of life=3, amount smoked by mother=5, and amount smoked by father=6; and for the 13–14-year age group: current maternal smoking=1 and current paternal smoking=10.

**Statistical analysis**

To be included in the analysis, centres had to assess at least 1000 subjects (except Tokelau which was a complete country study) and to have a response rate of >60% for the children and >70% for the adolescents. Centres with minor deviations from protocol were included in the analyses and identified by the use of a footnote in the ISAAC publications. Odds ratios (ORs) were calculated with generalised linear mixed models, with a binomial distribution and logit link, and with centres being modelled as a random effect. Analyses were adjusted for gender, region of the world, language and gross national income. The regression models allowed for the sampling by schools by scaling the size of the sample by the design effect.

No imputation was done for missing data, as previous analyses had shown that little or no bias was introduced by limiting the size of the sample by the design effect.

To assess whether being in two smoke exposure groups, relative to being in one or the other, resulted in disproportionately high risk, the relative excess risk due to interaction and its 95% CIs were calculated. The population-attributable risk of current asthma symptoms for maternal smoking was calculated with the Mantel-Haenszel approach, with the adjusted relative risk (RR) and the proportion of subjects who were exposed.

**RESULTS**

Exposure values (6–7-year age group: current maternal smoking, current paternal smoking and mother smoking in first year of the child’s life; 13–14-year age group: current maternal smoking and current paternal smoking) and prevalence values (current symptoms of asthma, rhinoconjunctivitis and eczema) by centre are shown in online tables 1 (6–7 years old) and 2 (13–14 years old).

Table 1 shows the association between smoking by the parents and symptoms of asthma, rhinoconjunctivitis and eczema at 6–7 and 13–14 years of age. Smoking is associated with an increased risk of symptoms of asthma, eczema and rhinoconjunctivitis. Figures 1 and 2 show the adjusted ORs for the association between current maternal smoking and asthma symptoms for each age group by country and table 2 shows these associations by region. The ORs are higher for asthma outcomes than for symptoms of rhinoconjunctivitis and eczema. The magnitude of the ORs associated with symptoms of eczema and rhinoconjunctivitis are similar, but weaker than that seen with asthma outcomes. The ORs for the 6–7–year age group are very similar to those for the 13–14-year age group. In all analyses but one current maternal smoking is associated with higher ORs than current paternal smoking; the exception is symptoms of eczema without wheeze in the 13–14-year age group. In all the 6–7–year age group analyses maternal smoking in the child’s first year of life is associated with higher ORs than current maternal smoking. Furthermore, in both age groups severe symptoms of asthma are associated with higher ORs than current wheeze or asthma.

The association between all the smoking variables and symptoms of eczema without current asthma symptoms in both age groups are significant. The association between the smoking variables and symptoms of rhinoconjunctivitis without current asthma symptoms is significant for the 13–14-year age group, but not for the younger age group, although the point estimates are elevated.

For asthma in the 6–7–year age group (current wheeze, severe symptoms and asthma) there is clear evidence of a dose–response effect for current smoking by the father and mother (table 3). For example, the more cigarettes the mother currently smokes, the higher the risk of severe asthma symptoms (1–9 cigarettes/day, OR 1.27; 10–19 cigarettes/day, OR 1.55; 20+ cigarettes/day, OR 1.56). A dose–response effect is not seen with symptoms of eczema and rhinoconjunctivitis.

The risk of all outcome variables is greater when the mother only smokes than when the father only smokes (except for symptoms of rhinoconjunctivitis without wheeze in the 13–14-year age group) (table 4). When both the mother and father smoke the risk is increased by the expected amount; that is, the increased risk is almost exactly the same as that obtained by multiplying the OR when the mother only smokes with that when the father only smokes (except for symptoms of eczema without wheeze in the 13–14-year age group). In all the 6–7–year age group analyses maternal smoking in the first year of the child’s life is associated with higher ORs than current maternal smoking only, although there is also an effect of only current paternal smoking. There is no evidence of an interaction between these variables.

Table 5 examines the association between the combinations of smoking in the child’s first year of life and current maternal smoking and symptoms of asthma, rhinoconjunctivitis and eczema. Maternal smoking in the first year of the child’s life only is associated with greater risk of all symptoms than current maternal smoking only, although there is also an effect of only maternal smoking currently. Unexpectedly maternal smoking in the first year of life and currently is not associated with a further increase in risk compared with that seen with smoking in the first year of the child’s life. There is a significant negative relative
excess risk due to interaction, which indicates that the risk of symptoms from the combined exposure is less than one would have expected.

Population attributable risk (PAR) for the various exposures is given in online table 3. The PAR for current maternal smoking and the various asthma outcomes in both age groups range from 5.6% to 7.1%. Figure 3 shows the PAR for maternal and paternal smoking. For example, if the prevalence of maternal smoking is 20% and paternal smoking is 25%, the PAR is 8%.

**DISCUSSION**

**Main findings**

This study has shown that parental smoking is associated with an increased risk of diagnosis and symptoms of asthma, eczema and rhinoconjunctivitis in childhood. The magnitude of the association is greater for current smoking by mothers than fathers, which may be due to children and adolescents spending more time in the presence of their mothers than their fathers. However, smoking by the mother and the father are both important.

When we examined ‘severe asthma symptoms’, which is clinically more meaningful, the magnitude of the ORs was even greater than that seen with current wheeze. This result reinforces the role of ETS in the severity of asthma symptoms. Furthermore, this suggests that ETS is related to asthma rather than just trivial wheeze or other respiratory symptoms. The magnitudes of the relative risks observed in this study are modest, although consistent with other meta-analyses examining these relationships. However, given how prevalent smoking is in some countries, ETS exposure is an important contributor to the burden of asthma and allergies in those countries. If maternal smoking is causally related to asthma then it might account for 5–7% of asthma cases in the world.

Maternal smoking in the child’s first year of life was associated with a greater risk of symptoms for all outcome categories than current maternal smoking. Furthermore the magnitude of the risk associated with exposure to both smoking in the first year of life and current smoking was not greater than those who were only exposed to smoking in the first year of life. This indicates that early exposure to ETS is especially important. However, it needs to be recognised that we did not collect exposure data on smoking in pregnancy. As smoking in pregnancy and the first year of the child’s life are highly correlated, it is possible that maternal smoking in the child’s first year of life is a marker for smoking in pregnancy and that this is the more important factor. This study is not able to elucidate this.

We showed clear evidence of a dose–response effect for current maternal and paternal smoking on all the asthma outcomes; that is, the more the mother or father smoked, the greater the likelihood that the child or adolescent would develop

### Table 3

<table>
<thead>
<tr>
<th>Symptom</th>
<th>1–9 cigarettes</th>
<th>10–19 cigarettes</th>
<th>20+ cigarettes</th>
<th>Current maternal smoking</th>
<th>1–9 cigarettes</th>
<th>10–19 cigarettes</th>
<th>20+ cigarettes</th>
<th>RERI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Current wheeze</td>
<td>1.06 (1.01 to 1.12)</td>
<td>1.15 (1.08 to 1.22)</td>
<td>1.27 (1.19 to 1.35)</td>
<td>1.25 (1.17 to 1.34)</td>
<td>1.29 (1.20 to 1.38)</td>
<td>1.44 (1.31 to 1.59)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Severe asthma symptoms</td>
<td>1.05 (0.96 to 1.15)</td>
<td>1.12 (1.12 to 1.34)</td>
<td>1.42 (1.30 to 1.56)</td>
<td>1.27 (1.15 to 1.40)</td>
<td>1.35 (1.22 to 1.50)</td>
<td>1.56 (1.36 to 1.80)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Asthma ever</td>
<td>1.03 (0.97 to 1.10)</td>
<td>1.11 (1.04 to 1.18)</td>
<td>1.18 (1.11 to 1.26)</td>
<td>1.18 (1.09 to 1.27)</td>
<td>1.29 (1.20 to 1.40)</td>
<td>1.37 (1.24 to 1.52)</td>
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<tr>
<td>Current eczema symptoms</td>
<td>1.08 (1.02 to 1.15)</td>
<td>1.04 (0.97 to 1.11)</td>
<td>1.11 (1.04 to 1.20)</td>
<td>1.17 (1.08 to 1.26)</td>
<td>1.13 (1.04 to 1.23)</td>
<td>1.15 (1.02 to 1.30)</td>
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<td></td>
</tr>
<tr>
<td>Current rhinoconjunctivitis symptoms</td>
<td>1.04 (0.98 to 1.11)</td>
<td>1.06 (0.99 to 1.13)</td>
<td>1.10 (1.03 to 1.18)</td>
<td>1.18 (1.09 to 1.27)</td>
<td>1.02 (0.94 to 1.11)</td>
<td>1.14 (1.02 to 1.28)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Eczema symptoms without wheeze</td>
<td>1.08 (1.01 to 1.16)</td>
<td>1.01 (0.94 to 1.09)</td>
<td>1.02 (0.94 to 1.11)</td>
<td>1.07 (0.98 to 1.17)</td>
<td>1.11 (1.00 to 1.22)</td>
<td>1.08 (0.94 to 1.25)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Rhinoconjunctivitis symptoms without wheeze</td>
<td>1.04 (0.97 to 1.12)</td>
<td>0.98 (0.91 to 1.07)</td>
<td>1.00 (0.92 to 1.09)</td>
<td>1.07 (0.98 to 1.17)</td>
<td>0.92 (0.82 to 1.03)</td>
<td>1.05 (0.90 to 1.21)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Data are OR (95% CI) and are adjusted for gender, language, region and GNI category. Reference group is neither parent smokes.

### Table 4

<table>
<thead>
<tr>
<th>Symptom</th>
<th>6–7 years</th>
<th>13–14 years</th>
<th>RERI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Current wheeze</td>
<td>1.31 (1.22 to 1.41)</td>
<td>1.37 (1.29 to 1.45)</td>
<td>-0.071 (-0.182 to 0.040)</td>
</tr>
<tr>
<td>Severe asthma symptoms</td>
<td>1.31 (1.18 to 1.46)</td>
<td>1.46 (1.34 to 1.59)</td>
<td>-0.042 (-0.205 to 0.121)</td>
</tr>
<tr>
<td>Asthma ever</td>
<td>1.28 (1.19 to 1.39)</td>
<td>1.29 (1.22 to 1.38)</td>
<td>-0.057 (-0.168 to 0.054)</td>
</tr>
<tr>
<td>Current eczema symptoms</td>
<td>1.15 (1.06 to 1.24)</td>
<td>1.19 (1.11 to 1.27)</td>
<td>-0.029 (-0.136 to 0.078)</td>
</tr>
<tr>
<td>Current rhinoconjunctivitis symptoms</td>
<td>1.13 (1.04 to 1.23)</td>
<td>1.16 (1.09 to 1.24)</td>
<td>-0.035 (-0.141 to 0.070)</td>
</tr>
<tr>
<td>Eczema symptoms without wheeze</td>
<td>1.06 (0.97 to 1.16)</td>
<td>1.13 (1.05 to 1.22)</td>
<td>0.040 (-0.075 to 0.155)</td>
</tr>
<tr>
<td>Rhinoconjunctivitis symptoms without wheeze</td>
<td>1.08 (0.98 to 1.19)</td>
<td>1.06 (0.98 to 1.15)</td>
<td>-0.052 (-0.173 to 0.069)</td>
</tr>
</tbody>
</table>

Data are OR (95% CI) and are adjusted for gender, language, region and GNI category. Reference group is neither parent smokes.

RERI, relative excess risk due to interaction between the two smoking exposures.
asthma symptoms. The presence of a dose–response effect is strong evidence for a causal relationship.

We found a weak but consistent association between ETS and symptoms of eczema. A dose–response effect relationship was not seen between amount smoked and symptoms of eczema. In contrast to asthma, there have been fewer studies of the association between smoking and eczema. Smoking has been shown to increase the risk in adults in Sweden, but not in Norway. In children aged 3 years, in utero and postnatal exposure were not associated with eczema nor was an effect seen in children aged 0–12 years. However, in a study of children aged 6 years an increased risk of eczema was associated with urinary cotinine levels.

Finally we found that the association between ETS and symptoms of rhinoconjunctivitis was even weaker and for many of the analyses did not reach statistical significance. Furthermore, a dose–response relationship was not seen between the amount smoked and symptoms of rhinoconjunctivitis. Studies examining the association between smoking and rhinoconjunctivitis are also conflicting. One study reported a reduced risk of rhinoconjunctivitis for adult smokers and their children. Another showed a decreased risk of hayfever in adolescents who were exposed to tobacco in utero. However, in the ISAAC Phase Two Study there was an increased risk of rhinoconjunctivitis in children aged 9–11 years who had been exposed to maternal smoking during pregnancy.

**Strengths and limitations**

The strengths of this multicentre study were its power, size and multinational nature. The younger age group comprised 75 centres from 32 countries with 220,407 children and the adolescent group comprised 118 centres from 53 countries with 350,654 adolescents. As a result, we established the consistency of the association of ETS with the presence of asthma and other allergic disorders in various populations with different prevalences of asthma and allergies, different exposure levels and different cultures and economic development. Many previous studies have been conducted in developed countries and the meta-analyses reflect this. This study extends the meta-analyses by including many countries that are less developed and markedly increased the size and number of the studies examining these relationships. We mainly used self-reported symptoms rather than doctors’ diagnoses to avoid major diagnostic differences related to access to medical care, language and medical practice. Publication bias has been avoided in this study as the data from each centre in the study were included.

Limitations of this study also need to be considered. The outcomes and exposures were assessed by questionnaire, which were completed by the parents for the younger age group and by the participants in the adolescent age group. Although it is likely that the parent who completed the questionnaire would be more accurate about their own tobacco use than that smoked by the other parent, we did not record which parent completed the questionnaire. This might lead to misclassification, which would reduce the chance of identifying associations, but is unlikely to result in any systematic bias.

Selection bias is also unlikely as the participants in the adolescent age group. Although it is likely that the parent who completed the questionnaire would be more accurate about their own tobacco use than that smoked by the other parent, we did not record which parent completed the questionnaire. This might lead to misclassification, which would reduce the chance of identifying associations, but is unlikely to result in any systematic bias.

**Implications**

The association of ETS with asthma fulfills the criteria for a causal relationship. The association is moderately strong; the results were generally consistent across each of the centres; and a dose–response relationship was seen for asthma, but not eczema and rhinoconjunctivitis.

**Conclusion**

In summary, this study has confirmed the importance of maternal smoking, especially early in the child’s life, and the separate and additional effect of paternal smoking. The presence of a dose–response effect relationship with symptoms of asthma suggests that the relationship may be causal. However, for eczema and rhinoconjunctivitis causality is less certain.

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APPENDIX 1
ISAAC Phase Three Study Group

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The association between tobacco and the risk of asthma, rhinoconjunctivitis and eczema in children and adolescents: analyses from Phase Three of the ISAAC programme

Edwin A Mitchell, Richard Beasley, Ulrich Keil, Stephen Montefort, Joseph Odhiambo and and the ISAAC Phase Three Study Group

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