Mite, cat, and cockroach exposure, allergen sensitisation, and asthma in children: a case-control study of three schools

Richard Sporik, Susan P Squillace, Jim Mark Ingram, Gary Rakes, Richard W Honsinger, Thomas A E Platts-Mills

Abstract

Background—The amount of allergen necessary to sensitise genetically “at risk” children is unclear. The relation between allergen exposure and asthma is also uncertain.

Methods—To ensure a wide range of allergen exposures the data from case-control studies of asthma in children aged 12–14 years attending three schools in Los Alamos, New Mexico and Central Virginia were combined. Skin prick tests to indoor and outdoor allergens and bronchial hyperreactivity to histamine were assessed in children with and without symptoms of asthma. The concentration of mite, cat, and cockroach allergens in dust from the children’s homes was used as a marker of exposure.

Results—Three hundred and thirty two children (157 with asthmatic symptoms and 175 controls) were investigated. One hundred and eighty three were classified as atopic on the basis of allergen skin prick tests and 68 as asthmatic (symptoms plus bronchial responsiveness). The prevalence and degree of sensitisation to mite and cockroach, but not cat, was strongly associated in atopic children with increasing domestic concentrations of these allergens. Asthma was strongly associated with sensitisation to indoor allergens (p<0.01) and weakly to outdoor allergens (p = 0.026). There was an association between current asthma and the concentration of mite allergen amongst atopic children (p = 0.008) but not amongst those who were specifically mite sensitised (p = 0.16).

Conclusions—The domestic reservoir concentration of mite and cockroach, but not cat, allergen was closely related to the prevalence of sensitisation in atopic children. However, the prevalence of current asthma had a limited relationship to these allergen measurements, suggesting that other factors play a major part in determining which allergic individuals develop asthma.

(Thorax 1999;54:675–680)

Keywords: asthma; children; allergen exposure; atopy

A consistent feature of asthma in children and young adults is increased levels of IgE specific to indoor allergens. Evidence comes from cross sectional studies of children and young adults, studies from children and young adults presenting to the emergency department with asthma exacerbations, and from an “at risk” population studied prospectively.

Methods

Patients

Children attending three schools were studied: Buford Middle School, Charlottesville, Virginia; Henley Middle School, Albermarle County, Virginia; and Los Alamos Middle School, Los Alamos, New Mexico. The schools were selected because of their diverse geographical and socioeconomic settings, being located in city, rural, and high altitude communities, respectively. All children between 12 and 14 years of age were eligible for...
enrolment. Ethical permission for these studies was obtained from the Human Investigation Committee of the University of Virginia, the Los Alamos Medical Center, and the local School Boards.

**DESIGN OF STUDY**

The study consisted of two parts, a screening phase followed by the detailed investigation of a selected sample. The same protocols were used at each school. In phase I of the study a respiratory questionnaire was administered as part of a general science class. A total of 1621 questionnaires were completed (Buford 608, Henley 446, and Los Alamos 567, representing 95%, 94%, and 95% of the school populations, respectively). On the basis of responses to the questionnaire the children were divided into those with symptoms of asthma and those without (controls). All those with symptoms of asthma and an equal number of randomly chosen control children were invited to participate in the second phase of the study. Three hundred and thirty two (including 175 controls) underwent skin prick testing and their bronchial responsiveness to histamine was assessed. Skin reactivity to extracts of the house dust mites (*Dermatophagoides farinae* and *D. pteronyssinus*), cockroach, cat dander, and pollens (grass mix, ragweed mix, and a tree mix) (Miles, Spokane, WA) were assessed using a lancet technique. A positive reaction was recorded if the mean diameter of the skin weal was ≥4 mm larger than the negative control, and children were considered strongly sensitised if the skin weal was ≥8 mm. A DeVilbiss hand held nebuliser was used to assess bronchial responsiveness to inhaled histamine in the children according to the technique of Yan et al. Children were considered hyperresponsive if at or before the maximal cumulative dose of 3.9 µmol histamine the forced expiratory volume in one second (FEV₁) decreased by 20% of the post saline value. The dose response slope (percentage change in the final FEV₁, from the post saline baseline divided by the total cumulative dose of histamine given) was also calculated. Children were considered to have asthma if they reported wheezing in the previous year and showed bronchial responsiveness to 3.9 µmol or less of histamine.

Dust was collected by a standard procedure from the mattress, bedroom floor, living room floor and kitchen floor of 127 Buford houses, 68 Henley houses, and 108 Los Alamos houses between September 1992 and September 1993. The dust was sieved and weighed and, after aqueous extraction, the content of *Der p 1*, *Der f 1*, *Fel d 1*, and *Bl a 2* allergens (derived from the house dust mites *D. pteronyssinus* and *D. farinae*, cat, and the German cockroach *Blatella germanica*, respectively) were measured using two site monoclonal antibody enzyme linked immunoassays. In keeping with the recommendations of an international workshop, the results were expressed as micrograms of allergen per gram of sieved dust (µg/g). The limit of detection for group 1 mite allergens (*Der p 1* + *Der f 1*) was 0.2 µg/g, for *Fel d 1* was 0.5 µg/g, and for *Bl a 2* was 0.08 µg/g (2 units/g).

**STATISTICAL ANALYSIS**

Parametric and non-parametric analyses were performed. Results were expressed as geometric means and 95% confidence intervals. All p values were two tailed. As allergen concentrations were not normally distributed, results were log₁₀ transformed for analysis. To examine the degree of sensitisation with increasing allergen exposure without making assumptions about threshold levels for sensitisation, the highest domestic exposure to mite and cat allergen in atopic children was ranked and used to divide these atopic children into six groups; the first group contained the majority of children with undetectable allergen and consisted of 29 children, followed by five equal groups of 27 children. The association of allergen concentration and the prevalence of sensitisation and asthma was tested using the χ² test for trend based on the log₁₀ median allergen concentration in each group. Regression correlations between the log₁₀ (dose response slope + 3) and log₁₀ group 1 mite allergen concentration were also calculated (Stata, College Station, Texas, USA).

**Results**

**DOMESTIC ALLERGEN CONCENTRATION**

There was a wide range in the concentration of mite, cat, and cockroach allergens in the 303 homes studied (fig 1). Extremely low concentrations of mite allergen were seen in Los Alamos and high concentrations in the Henley School District. Cockroach allergen (*Bl a 2*) was much less common and was only consistently found in the homes of children attending Buford school. Cat allergen was detectable in most of the houses, although in lower concentrations in Buford.

**ALLERGEN EXPOSURE AND SENSITISATION**

One hundred and eighty three children were classified as atopic and 149 non-atopic on the basis of their skin test results. Dust samples were collected from the homes of 164 atopic

![Figure 1](http://thorax.bmj.com/)

*Figure 1* Cumulative prevalence of highest domestic cockroach, mite, and cat allergens (all schools combined). The bold lines represent the highest concentration found in the homes of atopic children while the remaining lines represent those found in the homes of non-atopic children. The lower panel shows the median (–), range, and 25–75th centiles for cockroach, mite, and cat allergen in Los Alamos (LA), Buford (BU) and Henley (HE).
and 139 non-atopic children. There was no difference in the concentration of allergens found in the homes of atopic and non-atopic children (mite, \( p = 0.3 \); cat, \( p = 0.5 \); cockroach, \( p = 0.1 \)). Figure 1 shows the cumulative frequency distributions of each allergen by atopic status.

The prevalence of allergen specific sensitisation in atopic children was associated with the child’s highest domestic concentration of mite (\( p < 0.0001 \)) and cockroach allergen (\( p = 0.0001 \)), but not with cat allergen (\( p = 0.3 \); table 1). No lower threshold for sensitisation was seen. The degree of sensitisation, as measured by the size of the skin prick response, was also associated with increasing concentrations of mite and cockroach allergen (table 1). Cockroach allergen appeared to be particularly effective at sensitising, with all atopic children exposed to more than 0.32 µg/g being sensitised.

### Table 1 Occurrence of sensitisation and asthma with increasing domestic allergen exposure

<table>
<thead>
<tr>
<th>Highest domestic allergen concentration (µg/g)</th>
<th>Percentage of atopic children sensitised specifically to each allergen</th>
<th>Percentage of children with asthma</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Strongly sensitised***</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Atopic</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mite</td>
<td></td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>0.2 (&lt;0.2-0.3)</td>
<td>38 (11/29)</td>
</tr>
<tr>
<td>2</td>
<td>0.42 (0.3-0.6)</td>
<td>41 (11/27)</td>
</tr>
<tr>
<td>3</td>
<td>0.92 (0.6-1.8)</td>
<td>33 (9/27)</td>
</tr>
<tr>
<td>4</td>
<td>5.0 (1.8-10.0)</td>
<td>63 (17/27)</td>
</tr>
<tr>
<td>5</td>
<td>17.1 (10.2-23.9)</td>
<td>70 (19/27)</td>
</tr>
<tr>
<td>6</td>
<td>38.2 (24.0-155)</td>
<td>78 (21/27)</td>
</tr>
</tbody>
</table>

*The highest domestic concentration of mite and cat allergen among atopic children was ranked and used to divide the population into six groups.
**Allergen weal diameter ≥ 4 mm.
***Allergen weal diameter ≥ 8 mm, this being a subgroup of those sensitised.

The three indoor allergens (including cat) in atopic children was also positively associated with the mean community concentration of all allergens (table 2). In those 136 children sensitised to indoor allergens, sensitisation to house dust mite was the most prevalent (71%), followed by cat (40%) and cockroach (35%). Multiple sensitisation was not uncommon; 65% of sensitised children were solely sensitised to one indoor allergen, 26% to two, and 10% to all three. The commonest pattern of sensitisation was isolated sensitisation to mite (35%) followed by isolated sensitisation to cat (15%).

### Table 2 Mean community allergen concentration (geometric mean, 95% confidence interval), and the occurrence of allergen sensitisation

<table>
<thead>
<tr>
<th>Allergen</th>
<th>Los Alamos (n = 108)</th>
<th>Buford (n = 127)</th>
<th>Henley (n = 68)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mite</td>
<td>0.5 (0.4 to 0.6)</td>
<td>5.9 (4.5 to 7.7)</td>
<td>16.6 (11.7 to 23.4)</td>
</tr>
<tr>
<td>&gt;2 µg/g</td>
<td>5 (3%)</td>
<td>93 (73%)</td>
<td>62 (91%)</td>
</tr>
<tr>
<td>(% atopic children)</td>
<td>24 (29%)</td>
<td>41 (72%)</td>
<td>29 (69%)</td>
</tr>
<tr>
<td>Cat</td>
<td>13.7 (9.2 to 20.4)</td>
<td>6.9 (4.6 to 10.3)</td>
<td>10.5 (5.4 to 20.3)</td>
</tr>
<tr>
<td>&gt;8 µg/g</td>
<td>56 (52%)</td>
<td>42 (33%)</td>
<td>33 (48%)</td>
</tr>
<tr>
<td>(% atopic children)</td>
<td>34 (40%)</td>
<td>8 (14%)</td>
<td>13 (31%)</td>
</tr>
<tr>
<td>Cockroach</td>
<td>0.04 (0.04 to 0.044)</td>
<td>0.13 (0.11 to 0.16)</td>
<td>0.08 (0.07 to 0.09)</td>
</tr>
<tr>
<td>&gt;0.08 µg/g (2 U/g)</td>
<td>4 (4%)</td>
<td>32 (25%)</td>
<td>2 (3%)</td>
</tr>
</tbody>
</table>

*Number of houses with allergen level greater than previously proposed “threshold” value.
**Number of specifically sensitised children among the children tested in each school, (expressed as a % of atopic children in each school).
cat, or cockroach exposure compared with controls (table 3). However, the proportion of children with bronchial hyperresponsiveness among the symptomatic children increased with increasing mite allergen concentration (p<0.006). By contrast, the proportion of children with bronchial hyperresponsiveness among the symptomatic children decreased, though not significantly, with increasing exposure to cat allergen (p = 0.15).

The proportion of atopic children with asthma symptoms and bronchial hyperresponsiveness increased with increasing concentrations of house dust mite allergen (p = 0.008; table 1). By contrast, no effect of mite allergen was seen in non-atopic children. There was no significant effect of cat exposure on asthma either for atopic (p = 0.85) or specifically allergic children (p = 0.98). The proportion of non-atopic children with asthma was low overall, but was slightly lower in children with higher exposure to cat allergen (p = 0.042; table 1). The effect of mite allergen exposure on atopic children was reflected by the increasing bronchial responsiveness to histamine (dose-response slope) with higher house dust mite allergen concentrations in atopic children (r = 0.26, p<0.001; fig 2). However, there was a wide scatter of results and a significant trend was also seen in non-atopic children (r = 0.19, p = 0.027). The regression lines crossed the dose-response slope value corresponding to a PD_{20} of 3.9 µmol at a mite allergen concentration of 4 µg/g for atopic children and 600 µg/g (extrapolated) for non-atopic children. In those atopic children sensitive to mite there was no significant association between the proportion with asthma and current exposure (p = 0.16), nor in the dose-response slope (mite sensitive, r = 0.19, p = 0.07; solely mite sensitive, n = 29, r = 0.17, p = 0.37; mite non-sensitive, r = 0.20, p = 0.09). Similarly, in those sensitised to cat no association with domestic exposure was seen (p = 0.98). There was also no relationship for those sensitised to cockroach and their domestic exposure (p = 1.0), although the number of sensitised children was small (table 1).

**Table 3 Numbers of children with symptoms and symptomatic bronchial hyperresponsiveness (BHR) by increasing exposure to domestic allergens**

<table>
<thead>
<tr>
<th>Allergen</th>
<th>µg/g median (range)</th>
<th>Symptoms and BHR</th>
<th>Symptoms</th>
<th>Control</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mite*</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>0.2 (&lt;0.2–0.3)</td>
<td>9</td>
<td>18</td>
<td>25</td>
</tr>
<tr>
<td>2</td>
<td>0.42 (0.3–0.6)</td>
<td>6</td>
<td>16</td>
<td>18</td>
</tr>
<tr>
<td>3</td>
<td>0.92 (0.62–1.8)</td>
<td>8</td>
<td>20</td>
<td>20</td>
</tr>
<tr>
<td>4</td>
<td>5.0 (1.8–10.0)</td>
<td>10</td>
<td>16</td>
<td>28</td>
</tr>
<tr>
<td>5</td>
<td>17.1 (10.2–23.9)</td>
<td>11</td>
<td>4</td>
<td>34</td>
</tr>
<tr>
<td>6</td>
<td>38.2 (24.0–155)</td>
<td>16</td>
<td>13</td>
<td>31</td>
</tr>
<tr>
<td>Cat*</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>0.6 (&lt;0.5–0.9)</td>
<td>17</td>
<td>12</td>
<td>33</td>
</tr>
<tr>
<td>2</td>
<td>1.5 (0.9–1.9)</td>
<td>8</td>
<td>14</td>
<td>23</td>
</tr>
<tr>
<td>3</td>
<td>3.2 (2.0–4.4)</td>
<td>12</td>
<td>14</td>
<td>27</td>
</tr>
<tr>
<td>4</td>
<td>9.3 (4.5–23.0)</td>
<td>12</td>
<td>11</td>
<td>27</td>
</tr>
<tr>
<td>5</td>
<td>64 (23.4–112)</td>
<td>9</td>
<td>11</td>
<td>27</td>
</tr>
<tr>
<td>6</td>
<td>270 (123–920)</td>
<td>8</td>
<td>20</td>
<td>24</td>
</tr>
<tr>
<td>Cockroach**</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>&lt;0.08 (&lt;0.08)</td>
<td>48</td>
<td>82</td>
<td>134</td>
</tr>
<tr>
<td>2</td>
<td>0.33 (&gt;0.08–15.6)</td>
<td>12</td>
<td>5</td>
<td>21</td>
</tr>
</tbody>
</table>

*p values analysed by χ² test for trend for mite and cat exposure groups (see table 1) and χ² for cockroach.

**Discussion**

In this multicentre case-control study of middle school children we found that the current domestic concentration of allergen was a major determinant of sensitisation to house dust mite and cockroach. However, this relationship between current exposure and sensitisation was not apparent for cat allergen. We have previously shown that asthma, whether defined as symptoms or as symptomatic bronchial hyperresponsiveness, is strongly associated with sensitisation to indoor allergens. What is important in the present analysis is that, although the relevant allergens were different in the three schools, the relationship between sensitisation and asthma was consistent. By contrast, when sensitised children were considered there was no strong relationship between asthma and the highest domestic allergen concentration.

An explanation for these differences in the exposure-sensitisation relationships could be that mite and cockroach allergens are only found in sufficient concentrations in the home, while cat allergen is also found in appreciable concentrations outside the home. Cat allergen should be considered not only a domestic but also a community allergen. We have previously shown in a prospective study that exposure to house dust mite during infancy is more strongly associated with sensitisation at the age of 11 years than exposure at age 11. It was therefore surprising that a single spot measurement of current exposure in the present study shows such an association, although the strength of association was lower (odds ratio ∼3 compared with 16). It is possible that current measurements of allergen in dust are both an index of exposure and also surrogate markers of more permanent features such as living at altitude or living in poverty.
By pooling data from three populations it was possible to explore the effects of a wide range of mite and cat allergen exposures. We have previously estimated the prevalence of current asthma in each of the schools studied. In two of the schools (Buford and Henley) asthma was strongly associated with mite sensitisation but was independent of current mite exposure. Los Alamos is an area with very low concentrations of house dust mite, a correspondingly low prevalence of mite sensitisation, where asthma is not associated with mite sensitisation. When data from Los Alamos were included in the analysis the results remained unchanged. In particular, asthma in mite sensitive children was not significantly associated with current mite exposure. However, for atopic children increasing exposure to mite allergen was strongly related to an increased prevalence of asthma. The lack of a major effect of current allergen exposure on the prevalence of asthma once sensitisation has occurred is an important observation. There are a number of possible explanations: (1) once sensitisation has occurred asthma is independent of allergen exposure; (2) there has been a misclassification of sensitisation; (3) there has been a misclassification of exposure; and (4) there are unmeasured confounders. Explanation (1) appears unlikely, given that symptoms can be exacerbated with exposure to inhaled allergen and asthma improves with total avoidance. (2) Allergen skin prick test with standardised extracts is a very sensitive way to detect allergen specific IgE. However, given the complex nature of many allergens an additional non-IgE mediated mechanism could explain the modest, but significant, effect of increasing allergen concentration amongst non-atopic children. (3) The highest reservoir concentration of allergen may not be a sensitive marker of the day to day changes that determine personal exposure. Indeed, the correlation between airborne or settling allergen, presumably a more direct measurement of allergen exposure, and reservoir measurements is not close. (4) There may be factors which protect sensitised individuals from asthma. Conversely, there are multiple other factors such as virus infection, endotoxin exposure, and air pollution that can contribute to the severity of symptoms in allergic individuals. These would all tend to interfere with the quantitative correlation between current exposure and symptoms.

There are a number of limitations to this study. While three large general populations of children were screened, skin testing and bronchial challenges were performed on a limited number of children, selected either for the presence or absence of respiratory symptoms. Thus the children studied include the population of interest and an equally sized, randomly chosen, control group, but not the entire population, which may have resulted in a selection bias. In addition, to obtain the wide range of exposures necessary to fully explore the exposure-response relationship it was necessary to study different populations. By pooling the data from three populations we have assumed that atopic children are a homogeneous group that behave similarly in diverse environments. Given that most of those not exposed to mite allergen lived in Los Alamos, and most of those exposed to cockroach lived in Buford, it is impossible to exclude “ecological” or other local factors confounding the results.

In summary, this study of children from three population based case-control studies confirms that the prevalence of sensitisation to mite and cockroach allergen is related to the degree of current allergen exposure. Asthma was also related to the degree of mite allergen exposure in atopic children, though not among those who were already sensitised to mite. The hypothesis that the prevalence of sensitisation and asthma could be reduced by allergen avoidance regimes in infancy and early childhood is being actively investigated by a number of groups. In keeping with recent results from Scandinavia, the concentration of cat allergen in the children's houses was not significantly related to either specific sensitisation or asthma. Reservoir measurements of mite allergen provide summary markers of cumulative exposure as indicated by the significant association between this marker and sensitisation. The results reported here may be due in part to the inadequacies of exposure assessment but are more likely to reflect the many confounders that influence the development of symptoms in allergic individuals.
22 Marks GB, Tovey ER, Toelle BG, et al. Mite allergen (Der p 1) concentration in houses and its relation to the presence and severity of asthma in a population of Sydney schoolchildren. J Allergy Clin Immunol 1995;96:49–56.
33 Munir AK, Einassorn R, Dreborg SK, Mite (Der p 1, Der f 1), cat (Fel d 1) and dog (Can f 1) allergens in dust from Swedish day-care centres. Clin Exp Allergy 1995;25:119–26.
Mite, cat, and cockroach exposure, allergen sensitisation, and asthma in children: a case-control study of three schools
Richard Sporik, Susan P Squillace, Jim Mark Ingram, Gary Rakes, Richard W Honsinger and Thomas A E Platts-Mills

Thorax 1999 54: 675-680
doi: 10.1136/thx.54.8.675

Updated information and services can be found at:
http://thorax.bmj.com/content/54/8/675

These include:
References
This article cites 38 articles, 2 of which you can access for free at:
http://thorax.bmj.com/content/54/8/675#BIBL

Email alerting service
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Topic Collections
Articles on similar topics can be found in the following collections
Child health (843)
Asthma (1782)
Epidemiologic studies (1829)

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/