

## ASTHMA

# Relationship between exposure to domestic allergens and bronchial hyperresponsiveness in non-sensitised, atopic asthmatic subjects

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**Background:** The effect of exposure to allergens not causing sensitisation in atopic asthmatic subjects has not previously been studied. A study was undertaken to assess the degree of asthma severity (measured by spirometry, airway reactivity and exhaled nitric oxide) in atopic asthmatic patients not sensitised to the domestic allergen to which they were exposed.

**Methods:** Dust samples were collected from the living room carpet and mattress in the homes of 248 subjects and dust mite, cat and dog allergen concentrations were measured. Spirometry, non-specific bronchial reactivity (BR), and exhaled nitric oxide (eNO) were ascertained. Patients' sensitisation status was assessed by skin prick testing.

**Results:** Adult atopic asthmatics not sensitised to mite but exposed to high levels of mite allergen had significantly more severe BR than subjects not exposed to high levels of mite (PD<sub>20</sub>, geometric mean (GM) 0.21 mg (95% CI 0.09 to 0.47) v 0.86 mg (95% CI 0.44 to 1.67), mean ratio difference 4.1 (95% CI 1.5 to 11.4), p=0.008). Subjects not sensitised but exposed to high levels of dog allergen also had significantly more severe BR than subjects not exposed (PD<sub>20</sub> GM 0.16 v 0.52 mg, mean ratio difference 3.3 (95% CI 1.2 to 9.2), p=0.01). The differences in BR between these groups were still significant after adjusting for confounding factors. This effect of greater airway reactivity was not seen in subjects exposed but not sensitised to cat allergens.

**Conclusion:** Atopic asthmatic subjects who are exposed to high levels of dust mite or dog allergens but not sensitised to these allergens have evidence of increased airway reactivity.

The inhalation of allergens in atopic asthmatic subjects in clinical studies is associated with bronchoconstriction. Subjects who have a late asthmatic response to allergen also show an exaggerated response to non-specific bronchoconstrictor agents (such as histamine or methacholine). This occurs after both high dose and repeated low dose allergen challenge.<sup>1–2</sup> An increase in non-specific airway reactivity also occurs after natural exposure to perennial<sup>3</sup> and seasonal<sup>4</sup> allergens, and the heightened airway responsiveness can occur without significant bronchoconstriction.<sup>2</sup>

Most observational and experimental research has focused on the effect of allergen exposure in subjects who are sensitised to that specific allergen. However, it is possible that increased airway responsiveness upon allergen exposure may in part be due to factors other than the generation of an allergic reaction—for example, some allergens are proteolytic enzymes<sup>5</sup> that cleave epithelial tight junctions<sup>6</sup> which may lead to increased permeability of the airways.

This study forms part of an ongoing project in which we have previously demonstrated the relationships between different physiological measures of asthma severity<sup>7</sup> and also assessed the effect of exposure to sensitising allergens in allergic subjects.<sup>8</sup> We have shown in these two studies that bronchial reactivity appeared to be more severe in atopic asthmatic subjects than in non-atopic asthmatic subjects and that exposure to allergens to which a subject is also sensitised is related to a more severe bronchial reactivity. We now report on the relationship between the natural exposure to domestic allergens and lung function, airway responsiveness, and exhaled nitric oxide in atopic asthmatic subjects who were *not* sensitised to the allergen to which they were exposed. Our null hypothesis was that exposure to allergen which is not

causing sensitisation does not affect the objective measures of asthma severity.

## METHODS

Subjects were recruited from the North West Lung Centre Medicines Evaluation Unit clinical trials database and outpatient clinics if the following criteria were met: physician diagnosed asthma; asthma symptoms in the previous 12 months; no exacerbation in the previous 4 weeks; sensitised to one or more common inhalant allergens (dust mite, dog and/or cat).

Subjects gave written informed consent following a full explanation of the study. The study protocol was approved by the local research ethics committee.

## Study procedures

Subjects underwent the following procedures having avoided their asthma medication as follows: short acting  $\beta$  agonists (6 hours); long acting  $\beta$  agonists, leukotriene antagonists and theophyllines (at least 12 hours).

## Measurement of exhaled nitric oxide (eNO)

eNO was measured before spirometry and bronchial challenge using a chemiluminescence analyser (LR 2000; Logan Research, Rochester, UK). Subjects inhaled fully to total lung capacity and then exhaled slowly (at a controlled rate of 6 l/min) to residual volume into a wide bore Teflon tube with minimal resistance. The response time (10–90%) was <0.6 seconds and the sampling rate was 250 ml/min. The

**Abbreviations:** BR, bronchial reactivity; eNO, exhaled nitric oxide; FEV<sub>1</sub>, forced expiratory volume in 1 second; FVC, forced vital capacity

nitric oxide reading was taken when the level had reached a plateau and the carbon dioxide concentration exceeded 5%. The mean of two readings was taken.

### Lung function testing

Forced expiratory volume in 1 second (FEV<sub>1</sub>) and forced vital capacity (FVC) were measured using a Microlab 3000 spirometer (MicroMedical, Rochester, UK). The best of three technically acceptable efforts was recorded.

### Assessment of airway reactivity

Bronchial challenge was performed if the subject's FEV<sub>1</sub> was  $\geq 60\%$  of the predicted value. A modified Yan technique bronchial challenge<sup>9</sup> was used to obtain a cumulative provocative dose of either histamine or methacholine causing a fall in FEV<sub>1</sub> of 20% or more (PD<sub>20</sub>FEV<sub>1</sub>). For technical and supply reasons, either histamine or methacholine was used for the provocation solution and the first and last concentrations differed slightly (histamine 3.13 mg/ml to 50 mg/ml, methacholine 1.56 mg/ml to 50 mg/ml). As previous studies<sup>10–11</sup> have shown that the two compounds produce similar results, dose equivalence was presumed for our study.

A Koko DigiDoser dosimeter (PDS Research, Louisville, CO, USA) and "646" nebuliser pots (Devilbiss, Somerset, PA, USA) were used for the challenge. Cumulative doubling doses of the provocation solution were given immediately after each measurement of lung function which was performed 1 minute after the dose of histamine/methacholine. The challenge was terminated when a fall in FEV<sub>1</sub> of 20% or greater was reached. The PD<sub>20</sub>FEV<sub>1</sub> was calculated (using cumulative doses) from the following formula: PD<sub>20</sub> = antilog[log D1 + (log D2 – log D1)(20 – R1)/(R2 – R1)]. The equation is derived from linear interpolation. In subjects who did not demonstrate a 20% fall in FEV<sub>1</sub> following the challenge, a PD<sub>20</sub> of 12 mg was entered as the result (one "doubling dose" above the final cumulative dose).

### Skin prick testing

Skin testing was performed by the prick method (*Dermatophagoides pteronyssinus*, cat and dog, negative and positive controls; Bayer, Elkhart, IN, USA). Sensitisation was defined as a wheal diameter at least 3 mm greater than the negative control.

### Home visits and assessment of allergen exposure

Dust samples were collected by vacuuming a 1 m<sup>2</sup> area of living room carpet and mattress for 2 minutes. Dust extracts were assayed for content of *Der p 1*, *Fel d 1*, and *Can f 1* using monoclonal antibody based enzyme linked immunosorbent assays as previously described.<sup>12–14</sup> The results were expressed as  $\mu\text{g}$  allergen/g fine dust.

### Statistical analysis

The statistical analysis was performed using SPSS 10.1 for Windows (SPSS, Chicago, IL, USA). The conventional 5% level of statistical significance was used. Results of PD<sub>20</sub>, eNO, and allergen concentrations were log transformed and are reported as geometric means (GM) with 95% confidence intervals (CI). Independent sample *t* tests were used for the comparative analysis of lung function data between any two groups.

A univariate analysis of variance was used to assess for any imbalance between the exposed/non-exposed groups due to confounding factors. The factors analysed were sex, age, smoking status, number of years with the diagnosis of asthma, use of inhaled steroids, dose of inhaled steroids if used, and the geometric mean levels of the other two confounding allergens (in carpet and mattress) in each group

**Table 1** Demographic details of study subjects (n = 248)

Characteristic	
Mean (range) age (years)	41 (11–68)
Sex (M/F)	105 (42%)/143 (58%)
Smoking status, n (%)	
Smokers	33 (13%)
Non-smokers	215 (87%)
Asthma treatment	
Receiving inhaled steroids, n (%)	149 (60%)
Mean, median (range)	658 $\mu\text{g}$ , 400 $\mu\text{g}$ (100–2000)
Not receiving inhaled steroids, n (%)	99 (40%)
Mean (range) FEV <sub>1</sub> % predicted	87.4% (24–132)
Allergen sensitisation (n (%) positive):	
Dust mite	185 (75%)
Cat	149 (60%)
Dog	124 (50%)
Allergen exposure (to high levels):	
Dust mite	109 (46%)
Cat	63 (25%)
Dog	52 (22%)

(for example, the levels of dog and cat allergens in the group of subjects not sensitised to house dust mite). A potential confounding factor was included in the multivariate analysis if it had a *p* value of  $\leq 0.1$ . Analyses of PD<sub>20</sub>, FEV<sub>1</sub> and eNO were adjusted for, using this same model, if confounding factors were found between the different groups.

To assess for bias between subjects receiving methacholine and histamine in the bronchial challenges, interaction between the method and any association was tested in a multivariate analysis.

Some subjects in our study did not undergo a bronchial challenge test, mostly because their lung function precluded the test while some refused the investigation. Technical problems prevented measurements of eNO in some subjects. However, post hoc power calculations showed that the study has approximately 80% power to detect differences of two doubling dilutions between groups for PD<sub>20</sub> (assuming a standard deviation of 0.7 for the log<sub>10</sub> values), 6.5 ppb between groups for eNO (assuming a standard deviation of 0.36 for the log<sub>10</sub> values), and 13% between groups for FEV<sub>1</sub> % predicted (assuming a standard deviation of 18).

## RESULTS

Two hundred and forty eight subjects were recruited into the study (table 1). Dust mite was the most common allergen causing sensitisation (n = 185 subjects). Subjects were classified as being exposed to high levels of domestic allergens based on previously proposed criteria if the allergen levels were  $>2 \mu\text{g/g}$  *Der p 1*,  $>10 \mu\text{g/g}$  *Fel d 1*, and  $>10 \mu\text{g/g}$  *Can f 1* in either site (carpet or mattress).<sup>15–16</sup> Dust mite was also the most common allergen to which subjects were exposed (46%).

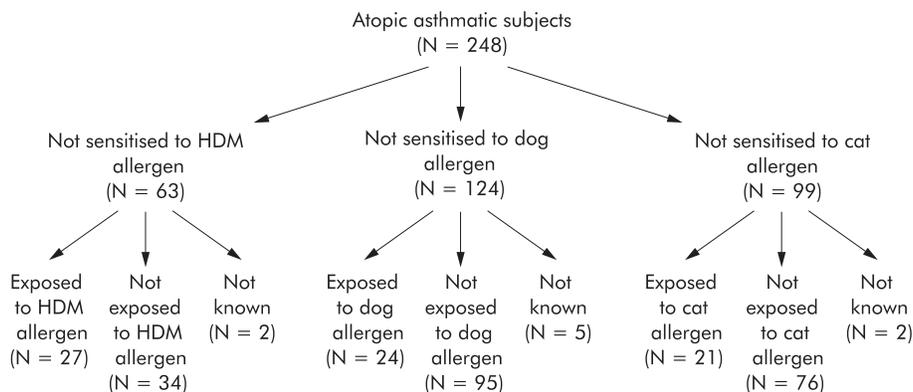
The distribution of patients in each subgroup is shown in fig 1. Before adjustment for confounding factors, the following results were obtained.

### *Der p 1*

Subjects not sensitised to dust mite had significantly greater airway reactivity (that is, lower log PD<sub>20</sub>) values when exposed to high levels of *Der p 1* (table 2). There was a trend towards higher eNO levels in the exposed group but the difference was not statistically significant (*p* = 0.09). There was no difference in FEV<sub>1</sub> % predicted between patients exposed and those not exposed to high levels of *Der p 1*.

### *Can f 1*

Subjects not sensitised to dogs had significantly greater airway reactivity when exposed to high levels of *Can f 1*



**Figure 1** Breakdown of subjects into subgroups (groups are not mutually exclusive).

(table 3), and this group also had significantly higher eNO levels. There was no difference in FEV<sub>1</sub> % predicted between the exposed and non-exposed groups.

**Fel d 1**

In subjects not sensitised to cat there was no difference in PD<sub>20</sub>, eNO, or FEV<sub>1</sub> % predicted between those exposed to high levels of that allergen and those not exposed.

Univariate analysis showed that, of the confounding factors considered (sex, age, smoking status, number of years with the diagnosis of asthma, use of inhaled steroids, dose of inhaled steroids if used, and the geometric mean levels of the other two allergens), only age was a significant associate for PD<sub>20</sub>, and only in the non-dog sensitised group. Age, inhaled steroid dose, and smoking status were significant associates for eNO. After adjusting for these, the difference in eNO levels between the *Der p 1* exposed and non-exposed groups in the non-house dust mite sensitised subjects and also the *Can f 1* exposed and non-exposed subjects in the non-dog sensitised subjects failed to reach significance ( $p = 0.07$  and  $p = 0.08$ , respectively).

If the diagnosis of sensitisation was re-defined as a wheal diameter of 2 mm, bronchial reactivity was still greater in subjects not sensitised but exposed to mite ( $p = 0.03$ ) and also subjects not sensitised but exposed to dog allergens ( $p = 0.01$ ). As the number of subjects was fairly small, a formal dose-response effect of allergen was not performed, but a modest but significant ( $R = -0.4$ ,  $p = 0.009$ ) correlation was seen between bronchial reactivity and log *Der p 1* level in the mattresses of subjects not sensitised to mite.  $\chi^2$  tests showed no significant difference between the actual and expected frequency of exposure to high levels of confounding

allergens for subjects not sensitised to dust mite—that is, in non-house dust mite sensitised subjects there was no difference in the frequency of high *Can f 1* or *Fel d 1* exposure in the two groups (exposed or not exposed to *Der p 1*). The same applied with the non-dog sensitised subjects—that is, there was no difference in the incidence of high *Der p 1* or *Fel d 1* exposure in the two groups (exposed or not exposed to *Can f 1* allergen). In the analysis of potential confounding allergens between the groups, the only significant finding was that subjects not sensitised to dust mite who were exposed to high levels of that allergen had a significantly higher level of *Can f 1* allergen in their mattress than subjects who were not exposed to *Der p 1* allergen (GM 1.45  $\mu\text{g/g}$  (95% CI 0.52 to 4.07) *v* 0.41  $\mu\text{g/g}$  (95% CI 0.18 to 0.31),  $p = 0.05$ ). However, after adjusting for the level of *Can f 1* allergen, there was still a significant difference between the *Der p 1* exposed and non-exposed groups in terms of their PD<sub>20</sub> (GM 0.24 mg (95% CI 0.11 to 0.52) *v* 0.71 mg (95% CI 0.37 to 1.34),  $p = 0.04$ ).

There was no difference in the expected frequencies of subjects who were sensitised to other domestic allergens which might affect the results—that is, in the non-house dust mite sensitised subjects there was no difference in the expected sensitisation frequency to dog or cat in the *Der p 1* exposed/non-exposed groups. This same analysis was performed for the non-dog and non-cat sensitised subjects.

There was no significant difference in bronchial challenge results between subjects receiving histamine (48% of total) and methacholine PD<sub>20</sub> (GM 0.29 mg (95% CI 0.20 to 0.42) *v* 0.48 mg (95% CI 0.32 to 0.73),  $p = 0.08$ ). Also, there was no significant difference in the numbers of subjects classified as having normal, mild, moderate, or severe bronchial reactivity between the two groups. The combined analyses adjusted for the method of challenge showed no effect of the method on

**Table 2** Pulmonary function data in subjects not sensitised to *Der p 1* (unadjusted data)

		Exposed to high levels of <i>Der p 1</i>	Not exposed to high levels of <i>Der p 1</i>
PD <sub>20</sub> (mg)	N	17	26
	GM (95% CI)	0.21 (0.09 to 0.47)	0.86 (0.44 to 1.67)
	Mean ratio difference (95% CI)	4.1 (1.5 to 11.4)	
	$p = 0.008$		
eNO (ppb)	N	25	29
	GM (95% CI)	13.9 (9.6 to 20.1)	9.3 (6.8 to 12.7)
	Mean ratio difference (95% CI)	0.7 (0.4 to 1.1)	
	$p = 0.09$		
FEV <sub>1</sub> (% predicted)	N	27	33
	Mean (95% CI)	87.2% (78.8 to 95.6)	94.5% (88.1 to 100.9)
	Mean difference (95% CI)	7.3 (-2.8 to 17.4)	
	$p = 0.15$		

**Table 3** Pulmonary function data in subjects not sensitised to *Can f 1* (unadjusted data)

		Exposed to high levels of <i>Can f 1</i>	Not exposed to high levels of <i>Can f 1</i>
PD <sub>20</sub> (mg)	N	15	67
	GM (95% CI)	0.16 (0.07 to 0.34)	0.52 (0.33 to 0.82)
	Mean ratio difference (95% CI)	3.3 (1.2 to 9.2)	
eNO (ppb)	N	20	83
	GM (95% CI)	14.6 (10.5 to 20.3)	9.8 (8.0 to 12.0)
	Mean ratio difference (95% CI)	1.2 (0.43 to 1.0)	
FEV <sub>1</sub> (% predicted)	N	24	94
	Mean (95% CI)	92.0 (84.3 to 99.7)	89.3 (84.8 to 93.8)
	Mean difference (95% CI)	-2.7 (-12.4 to 6.9)	
		p=0.53	

the outcomes, and no interaction was seen between the method and any of the exposed/not exposed groups. These data suggest that dose equivalence between the two challenge methods could be assumed.

## DISCUSSION

Our study has shown that airway reactivity was significantly greater in atopic asthmatic subjects who were not sensitised to dust mite or dog when they were exposed to high levels of these allergens. After correcting for potential confounding factors, FEV<sub>1</sub> % predicted and eNO were not significantly affected by exposure to high levels of non-sensitising allergens. This finding which has (to our knowledge) not previously been reported, may have significant implications in the interpretation of the effects of allergens in atopic asthma. The results suggest that exposure to high levels of domestic allergens may be associated with a deleterious effect on atopic asthma, even in subjects who are not sensitised to the specific allergen—that is, the effect is not consequent on the allergen's "allergenicity". A number of factors may have a role in causing this effect.

Several experimental studies have shown that some of the dust mite allergens (for example, *Der p 1*, 3, 6 and 9) have properties other than their allergenic effects. It has been shown that *Der p 1* is a cysteine protease digestive enzyme<sup>17</sup> and *Der p 9* is a serine protease that is capable of degrading collagen.<sup>18</sup> This enzymatic activity might cause airway damage independently from the allergenic effects by breaking down the tight junctions between epithelial cells. This action might cause direct inflammatory damage that could lead to some of the clinical and physiological features of asthma. Herbert *et al*<sup>19</sup> showed that *Der p 1* increases the permeability of bronchial mucosa to albumin, a feature that has been seen in acute asthma exacerbations.<sup>20</sup> It has also been suggested that *Der p 1* could activate mast cells without IgE involvement,<sup>21</sup> causing the release of inflammatory mediators. In vitro, *Der p 1* (by virtue of its proteolytic activity) also acts on epithelial cells, triggering a dose related release of the cytokine granulocyte macrophage colony-stimulating factor (GM-CSF).<sup>22</sup> This cytokine prolongs eosinophil survival, further enhancing the inflammatory process. Whether this occurs in vivo is yet to be ascertained. Finally, *Der p 1* may affect mucus secretion in the airway.<sup>23</sup> In the current study, the fact that increased airway reactivity was not seen in the subjects not sensitised to cat but exposed to high levels of *Fel d 1* might be due to the difference in properties of *Fel d 1*. This allergen does not exhibit the proteolytic activity of some of the *D pteronyssinus* allergens. These proteolytic enzymes have been found to cleave the tight junctions between cells in the epithelial barrier, potentially allowing the entry of allergens. *Fel d 3* has been identified as a member of the cysteine protease inhibitor family,<sup>24</sup> and

cysteine proteases have been implicated in enhancing IgE responses. Whether this is the reason for the difference in response between the two allergens is, however, debatable, as *Can f 1* and 2 may also contain a cysteine protease inhibitor.

As part of the current project involving this same cohort of subjects, we have previously shown that, despite having similar degrees of airway obstruction, atopic and non-atopic asthmatics have differing levels of airway reactivity and eNO.<sup>7</sup> In addition to the physiological differences, there are also contrasting pathological features. A previous study has shown that there are significant differences in the structure of the airways between atopic and non-atopic individuals.<sup>25</sup> These include differences in T lymphocytes and interleukin staining cells but, perhaps more importantly in relation to our study, the degree of epithelial damage was greater in the atopic group, with a negative correlation between epithelial integrity and both the numbers of eosinophils and CD25 positive cells. The authors did not speculate whether any of these features may have been caused by allergens, but it is possible that the differences between the atopic and non-atopic airways might mean that atopic asthmatics are more prone to damage from allergens. It has been proposed that the disruption to the epithelial barrier present in asthmatic airways may allow easier entry to the subepithelial layer (including smooth muscle) for bronchoconstrictor agents and possibly allergens.

In our study we did not measure the level of co-existing endotoxin in the dust samples. It is possible that the presence of endotoxin in the domestic environment may be a confounding factor that could affect asthma activity. For this to be the case, it would be expected that a higher concentration of endotoxin might be found in association with greater levels of domestic allergen exposure. Wickens *et al*<sup>26</sup> found a positive association between domestic endotoxin level and a number of factors including the presence of dogs in the house, while Michel *et al*<sup>27</sup> found no correlation between endotoxin and mite allergen concentrations. The effect of endotoxin has been examined in both experimental studies and in the domestic environment. The results of some studies are conflicting. The INGA study<sup>28</sup> found a positive association between exposure to mite allergens and the symptoms of wheeze and breathlessness, but no association between symptoms and endotoxin exposure. Other research has shown pro-inflammatory effects<sup>29</sup> and increased bronchial reactivity in non-asthmatic subjects,<sup>30</sup> as well as heightened reactivity<sup>31</sup> and increased airway obstruction in asthmatic subjects<sup>32</sup> after inhalation of endotoxin. Endotoxin may also augment the action of allergens in addition to having its own intrinsic inflammatory actions.

It is possible that a skin prick test threshold of 3 mm excludes some "sensitised" subjects and that some of the subjects apparently not sensitised to an allergen still have

enhanced reactivity when exposed to that allergen. However, the difference in bronchial reactivity was still present when sensitisation was re-defined as a wheal of 2 mm diameter or more.

Our results suggest that, within the atopic asthmatic population, exposure to high levels of dust mite and dog allergen is associated with increased airway reactivity even in subjects who are not sensitised to these allergens. Further experimental research may help to elucidate the cause of this phenomenon.

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