Gender differences in prevalence, diagnosis and incidence of allergic and non-allergic asthma: a population-based cohort

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ABSTRACT
Background Although women with severe non-allergic asthma may represent a substantial proportion of adults with asthma in clinical practice, gender differences in the incidence of allergic and non-allergic asthma have been little investigated in the general population.

Methods Gender differences in asthma prevalence, reported diagnosis and incidence were investigated in 9091 men and women randomly selected from the general population and followed up after 8–10 years as part of the European Community Respiratory Health Survey. The protocol included assessment of bronchial responsiveness, IgE specific to four common allergens and skin tests to nine allergens.

Results Asthma was 20% more frequent in women than in men over the age of 35 years. Possible under-diagnosis of asthma appeared to be particularly frequent among non-atopic individuals, but was as frequent in women as in men. The follow-up of subjects without asthma at baseline showed a higher incidence of asthma in women than in men (HR 1.94; 95% CI 1.40 to 2.68), which was not explained by differences in smoking, obesity or lung function. More than 60% of women and 30% of men with new-onset asthma were non-atopic. The incidence of non-allergic asthma was higher in women than in men throughout all the reproductive years (HR 3.51; 95% CI 2.21 to 5.58), whereas no gender difference was observed for the incidence of allergic asthma.

Conclusions This study shows that female sex is an independent risk factor for non-allergic asthma, and stresses the need for more careful assessment of possible non-allergic asthma in clinical practice, in men and women.

INTRODUCTION
The prevalence of asthma shows a sex reversal around puberty from a higher risk in boys early in life to a higher risk in girls after adolescence. This pattern has raised several hypotheses about the susceptibility to asthma of men and women, such as an effect of sex hormones, airway calibre, obesity, differences in exposure or diagnosis. Further studies are needed to appraise how much these determinants could explain the higher risk of asthma in women. Data are lacking on whether women remain at increased risk of asthma throughout all the reproductive years, and few studies have investigated the possibility of a differential diagnosis. Furthermore, most studies are not of a prospective nature that allows separating new-onset asthma from persistent asthma or relapse. In addition, new-onset of non-allergic asthma appears to be relatively frequent in adulthood. Clinical studies suggest that non-allergic (or ‘intrinsic’) asthma may be more severe and difficult to control than allergic asthma, and that women might be at increased risk of non-allergic asthma. However, most of our knowledge on non-allergic asthma comes from clinical studies which often include patients with more severe asthma, and little is known on non-allergic asthma in the general population.
We used data collected as part of the European Community Respiratory Health Survey (ECRHS) to estimate gender differences in the prevalence of asthma according to age; appraise possible differences in asthma diagnosis; prospectively investigate gender differences in the remission/persistence of asthma; and assess gender differences in the risk of new-onset asthma, accounting for potential confounders and considering allergic and non-allergic asthma separately, within the age range 20–55 years.

METHODS
Study design
The analysis is based on data collected in 29 centres from 14 countries as part of the ECRHS. The protocol and participation rates have been described elsewhere. Briefly, between 1991 and 1993, each participating centre randomly selected about 1500 men and 1500 women, representative of the age group 20–44 years, to answer a postal questionnaire (European Community Respiratory Health Survey I [ECRHS-I] stage 1; ‘screening’). A 20% random sample of respondents was then invited to a clinical investigation (ECRHS-I stage 2). Participants in stage 2 were eligible for the follow-up survey in 1998–2002 (ECRHS-II). At ECRHS-I stage 2 (‘baseline’) and follow-up, the protocol included assessment of respiratory symptoms via questionnaire and measurements of bronchial responsiveness and IgE specific to four common allergens. In each centre, the protocol was approved from the appropriate ethics committee, and written consent was obtained from each participant.

Analysis
Doctor-diagnosed asthma was defined as a positive answer to the questions ‘Have you ever had asthma?’ and ‘Was this confirmed by a doctor?’ Because only specific IgEs were assessed at follow-up, the main analysis was conducted with atopy defined as specific IgE≥0.55 kU/litre to any of the four common allergens tested. In a sensitivity analysis, atopic status was defined according to specific IgE and skin test reactivity measured at the baseline survey for nine common allergens. Atopic subjects with asthma were considered to have ‘allergic asthma’. Further details including cross tables are provided in the online supplement. All the analyses were conducted in Stata V8 (StataCorp 2001).

RESULTS
Gender differences in asthma prevalence and diagnosis
Participation of men and women at each step of the survey is described in the online supplement.

Prevalence
At ECRHS-I stage 1 (table 1, first column), large representative samples were screened to estimate asthma prevalence. There was no gender difference in the prevalence of asthma from age 20 to 35 years. However, women had a 20% higher risk of asthma than men after age 35 years. This pattern of results was consistent across the participating centres, despite large geographical variations in the prevalence of asthma.

To assess whether the magnitude of the gender difference had changed over time, we considered prevalence at ECRHS-II. Although the prevalence of asthma was higher at ECRHS-II, within each age-group the ORs for the gender difference were remarkably similar at each survey, showing no difference before age 35 and a 20% higher risk of asthma in women in the age group 36–44 years (table 1, third column). The higher risk of asthma in women appears to be further marked after age 45 years (OR 1.68; 95% CI 1.24 to 2.29).

Investigating possible differential diagnosis
We considered data collected at ECRHS-I stage 2 (including bronchial hyperresponsiveness [BHR] assessment) to assess whether the higher prevalence of asthma in women could be explained by differences in diagnosis.

Besides diagnosed asthma, women were also more likely than men to have asthma-like symptoms or asthma-like symptoms and BHR (table 1 and online supplement).

Furthermore, among subjects with respiratory symptoms and BHR (table 2), the likelihood of having received a diagnosis of asthma decreased with increasing smoking and was higher in atopic than in non-atopic subjects. However, in each group, it was similar or even lower in women than in men. As shown in table 2, in non-atopic subjects, only 16% of those with asthma-like symptoms and BHR reported a diagnosis of asthma. However, this low rate of diagnosis was similar in women and in men.

No major gender difference was observed for asthma treatment, apart from a more frequent use of inhaled steroids in women (online supplement, table E2).

Gender differences in the natural history of asthma
Forty-three per cent of men and 65% of women with a current diagnosis of asthma from the doctor at follow-up reported asthma onset in adulthood (online supplement, table E2).

Asthma remission, persistence and relapse in subjects with asthma at baseline
Women with current asthma at baseline were as likely as men to still have asthma at follow-up (asthma persistence: 68.1% in 215 women vs 74.1% in 135 men; p>0.20).

In subjects who had ever had asthma before the baseline survey, but without ‘current’ asthma at baseline, women were as likely as men to have current asthma at follow-up (asthma relapse after remission: 21.2% in 189 women vs 18.0% in 178 men; p>0.30).

Asthma incidence
The risk of developing asthma over the follow-up was investigated in the 4281 women and 3956 men without asthma at baseline (figure 1). The asthma incidence rate was higher in women than in men (table 3). There was no interaction with age (p for interaction =0.13). At baseline, women more frequently reported rhinitis, respiratory infections in childhood and maternal asthma compared with men. Women less frequently had positive specific IgE to dust mites and grass, had lower total IgE and smoked less than men. Women had lower body mass index (BMI) than men at baseline and at follow-up, but the mean increase in BMI between the two surveys was slightly higher in women than in men. Adjustment for the subject’s characteristics at baseline or follow-up had little effect on the gender difference in incidence (online supplement, table E3). The risk of developing asthma remained significantly higher in women than in men after adjustment for centre, maternal asthma, smoking, total IgE, atopic sensitisation, rhinitis, forced expiratory volume in 1 s (FEV1) and BMI at baseline (HR 1.94; 95% CI 1.40 to 2.68) as well as after additional adjustment for change in smoking status and change in BMI (HR 2.25; 95% CI 1.57 to 3.23). Further adjustment for occupational exposure over the follow-up to agents known to be related to occupational
Using skin test results for the nine common allergens to define atopy (online supplement, tables E4–E6).

### Discussion

In this large population-based cohort, the gender difference in the prevalence of asthma increased with increasing age and showed an increased risk of more than 20% in women than in men after age 35 years. The findings were consistent across the centres and during the two survey periods. Possible under-diagnosis of asthma appeared to be particularly frequent in non-atopic subjects, but was not more frequent in men than in women. In subjects without asthma at baseline, the risk of new-onset asthma over the follow-up was twofold higher in women than in men, and this was not explained by differences in smoking, lung function or obesity. More than 60% of the subjects reporting asthma-like symptoms plus bronchial hyperresponsiveness had no atopic sensitisation at follow-up. In men, 37% of incident non-atopic asthma were non-atopic. No gender difference was observed for the incidence of allergic asthma.

### Gender differences in non-allergic asthma

To our knowledge, this is the first study to demonstrate that the incidence of non-allergic asthma is higher in women than in men.
was a risk factor for adolescent wheeze in boys but not in girls. Mandhane et al reported that most of the incident cases were female, and atopy at age 10 was not associated with subsequent asthma onset. More recently, Mandhane et al reported that in the Dunedin birth cohort, atopy was a risk factor for adolescent wheeze in boys but not in girls. However, none of these studies have assessed the magnitude and significance of the gender difference in the incidence of allergic and non-allergic asthma. To our knowledge, our study is also the first to report a lack of gender difference in the incidence rates of allergic asthma. The incidence of allergic asthma appears to decrease with age. This decrease has to be interpreted with caution because subjects from younger cohorts contributed to a greater extent to the incidence estimate at a younger age, and there might be a cohort effect. However, the magnitude of the gender difference in the incidence of allergic asthma is unlikely to be biased by a cohort effect because the same increase in atopic sensitisation with younger age is observed in men and women (online supplement, table E5).

Possible explanations for gender differences in asthma incidence

Although most prospective studies in adults show a higher incidence of asthma in women than in men, few studies have simultaneously tested the possible effects of other relevant factors. Gender differences in asthma diagnosis have been demonstrated in children, but there are few data in adults. We did not find any suggestion of a more frequent diagnosis in women. Gender differences in environmental exposure have been put forward. In our study, women were at increased risk of non-allergic asthma, suggesting that exposure to allergens is unlikely to account for the higher incidence of asthma in women. Higher exposure of women to other bronchial irritants at home or at work cannot be excluded. Another possibility might be a greater susceptibility of women at the same level of exposure, which might, in turn, be related to their lower airway calibre. However, the incidence of asthma remained significantly higher in women than in men after adjustment for smoking, occupational exposure and lung function. In addition, the higher risk of asthma in women is supported by their increased risk of BHR, which cannot be explained by gender differences in airway calibre. Other factors are likely to be involved in the gender difference in asthma risk. The fact that the gender differences in asthma prevalence were consistent during the two survey periods and across countries with different levels of exposure to environmental factors suggests an effect of genetic and biological factors rather than socio-cultural and environmental factors.

Obesity appears to be a potential major risk factor for asthma development in women. In some studies, but not all, obesity was found to be a risk factor for incident asthma in women but not in men. Furthermore, obesity has been found to be more strongly associated with non-allergic asthma than with allergic asthma. A detailed analysis of the ECRHS suggested a potential differential diagnosis linked to the presence of obesity in
However, these studies have not considered allergic asthma. The number of person-years at risk, within each age group, were respectively 5043, 10 696 and 11 451 in men, and 5152, 10 935 and 11 317 in women. However, in the present analysis, the higher incidence of asthma in women could not be explained by differences in BMI or change in BMI. Although obesity, as such, seems unlikely to explain the higher risk of asthma in women, regulatory molecules secreted by the adipose tissues (such as leptin) might be involved in the gender differences in asthma.

A role for sex hormones is frequently suggested, but no study has been able to relate the switch in asthma prevalence occurring around puberty to changes in hormonal levels. In particular, a recent cohort of adolescents does not show any relationship between the pubertal stages and the incidence of asthma. However, in a cohort of children with asthma, a marked decrease in bronchial responsiveness was observed after age 11 years in boys, but not in girls. A possible relaxing effect of testosterone on airway smooth muscle, and an anti-inflammatory and immune-modulatory activity of the major adrenal androgen dehydroepiandrosterone have been put forward. Data from a murine model suggest that oestrogen may have a pro-inflammatory effect, while oestrogen in women already sensitised to allergens might speculate on whether such an anti-inflammatory effect of oestrogen in women already sensitised to allergens might decrease their likelihood of developing symptoms.

Study strengths and limitations
One of the strengths of this study lies in the quality of the ECRHS data. Standardised data for lung function, BHR and specific IgE were available at baseline and follow-up, and for a relatively large population-based sample of subjects with different levels of exposure to environmental factors. As regards the study limitations, loss to follow-up is a problem in any longitudinal study. However, there was no suggestion of any gender differential selection bias among subjects. The definition of asthma was based on the subject’s report of doctor-diagnosed asthma. This definition has been validated and found to be highly specific, but of rather low sensitivity. We cannot totally rule out a possible less frequent diagnosis of new-onset asthma in men than in women. Overlap of chronic obstructive pulmonary disease (COPD) and asthma is a concern in this type of analysis, particularly in non-atopic subjects. It has been suggested that COPD is more likely to be diagnosed as asthma in women than in men. However, only 12% of women with incident non-allergic asthma were heavy smokers, only 8% had a FEV₁/FVC ratio lower than 0.70, 75% were younger than 50 years old, and a higher incidence of non-allergic asthma was found in women after excluding smokers at baseline (online supplement, table E4). Furthermore, our data suggest in the process of antigen sensitisation per se, and an anti-inflammatory effect during the effector phase of the response to inhaled antigens. However, there are insufficient data to speculate on whether such an anti-inflammatory effect of oestrogen in women already sensitised to allergens might decrease their likelihood of developing symptoms.

Figure 2 Sex and age-specific incidence rates for allergic and non-allergic asthma. The number of person-years at risk, within each age group, were respectively 5043, 10 696 and 11 451 in men, and 5152, 10 935 and 11 317 in women.

Table 3 Incidence of asthma in women and men who reported they never had asthma at baseline

<table>
<thead>
<tr>
<th>Outcome at follow-up</th>
<th>Number of incident cases</th>
<th>Person-years</th>
<th>Incidence per 1000 person-years (95% CI)</th>
<th>HR* (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Women</td>
<td>Men</td>
<td>Women</td>
<td>Men</td>
</tr>
<tr>
<td>Asthma by age at baseline:</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>20–27 years</td>
<td>34</td>
<td>20</td>
<td>9118</td>
<td>8538</td>
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<tr>
<td>28–35 years</td>
<td>52</td>
<td>17</td>
<td>11 761</td>
<td>10 709</td>
</tr>
<tr>
<td>≥36 years</td>
<td>66</td>
<td>45</td>
<td>15 854</td>
<td>15 104</td>
</tr>
<tr>
<td>Allergic asthma</td>
<td>44</td>
<td>39</td>
<td>27 403</td>
<td>27 189</td>
</tr>
<tr>
<td>Non-allergic asthma</td>
<td>81</td>
<td>23</td>
<td>27 403</td>
<td>27 189</td>
</tr>
</tbody>
</table>

*Cox regression analysis (based on age at asthma onset) for incidence of doctor-diagnosed asthma in subjects who reported they never had asthma at baseline. Excludes incident cases with age at asthma onset missing or age at onset ≥5 years before baseline (see methods).
that asthma underdiagnosis is as frequent in women as in men. Because subjects were aged 20 years or more at inclusion, we cannot exclude the fact that a few subjects may have suffered from mild respiratory symptoms as early as childhood, and that these symptoms became severe enough in adulthood to be diagnosed as asthma. However, the results from a longitudinal birth cohort show that female sex is an independent predictor of adult-onset asthma after adjustment for wheeze in childhood. Furthermore, the higher incidence of asthma in women was still significant after excluding subjects with respiratory symptoms at baseline. As regards sensitisation, we cannot exclude that a few subjects with new-onset asthma classified as ‘non-atopic’ had ever been atopic and developed tolerance. Another possibility is that some subjects sensitised to other less common allergens may have been classified as ‘non-atopic’. However, similar findings were observed when baseline skin test sensitisation to any of nine common allergens was additionally used to classify incident cases as allergic and non-allergic asthma.

**Implications and conclusion**

Although women with severe non-allergic asthma may represent a substantial proportion of the adults with asthma in clinical practice, gender differences in the incidence of non-allergic asthma have been little investigated in the general population. In this large population-based cohort of adults, more allergic asthma have been little investigated in the general clinical practice, gender differences in the incidence of non-allergic asthma in adults. However, our results also suggest that this form of asthma is still poorly recognised in men and women, and stress the need for more careful assessment of potential non-allergic asthma in clinical practice. Women were found to be at higher risk of developing non-allergic asthma throughout all the reproductive years, whereas no gender difference was observed for allergic asthma. Overall, our findings suggest that biological factors that significantly differ between men and women are likely to be involved in the development of non-allergic asthma. Further research in this direction is needed.

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**Contributors**

Contributors that were involved in the study design and data collection in each participating centre are listed in the online supplement.

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**Competing interests**

None.

**Patient consent**

The article does not contain personal medical information about identifiable subjects.

**Ethics approval**

Local ethics committees and institutional review boards.

**Provenance and peer review**

Not commissioned; externally peer reviewed.

**Data sharing statement**

Researchers who wish to run analysis using the ECRHS data are invited to contact the coordinating centre in London. See the ECRHS website http://www.ecrhs.org.

**REFERENCES**

Tyrosine kinase inhibitor use in idiopathic pulmonary fibrosis

This randomised, multicentre, double-blind, placebo-controlled trial evaluated the efficacy and safety of four different doses of BIBF 1120, a tyrosine kinase inhibitor, in order to investigate its effect on progression of idiopathic pulmonary fibrosis (IPF).

The annual rate of decline in forced vital capacity (FVC), the primary end-point of this study, in the highest dose subgroup compared with the placebo subgroup failed to show statistical significance. However, the authors were able to demonstrate that the changes in FVC from baseline, total lung capacity, oxygen saturations, acute exacerbations and quality of life were statistically significantly better in the group receiving the highest dose of BIBF 1120. Serious adverse events were similar between groups, but lower in the highest dosing regimen compared with placebo. The most common reason for discontinuation of the medication was gastrointestinal upset.

Of note, only 53% of the patients recruited fulfilled the criteria for definite IPF with the remaining being diagnosed as probable, possible or no IPF (one patient). Discontinuation of BIBF 1120 was more evident among the group receiving the highest treatment dose relative to other groups (37.6%).

Although BIBF 1120 failed to demonstrate a decline in annual FVC rates, it was associated with numerous clinically significant benefits including reduced numbers of exacerbations and improved quality of life at the highest dose while proving to be safe. With limited options for treatment and the encouraging beneficial effects of BIBF 1120, further research is warranted on a carefully selected cohort of patients with a formal diagnosis of IPF.


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