

RESPIRATORY EPIDEMIOLOGY

Parental smoking in childhood and adult obstructive lung disease: results from the European Community Respiratory Health Survey

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Background: Early exposure to parental smoking appears to influence the development of the airways and predispose to respiratory symptoms. A study was undertaken to determine whether the consequences of parental smoking could be traced in adulthood.

Methods: Information from interviewer-led questionnaires was available for 18 922 subjects aged 20–44 years from random population samples in 37 areas participating in the European Community Respiratory Health Survey. Lung function data were available for 15 901 subjects.

Results: In men, father's smoking in childhood was associated with more respiratory symptoms (OR_{wheeze} 1.13 (95% CI 1.00 to 1.28); never smokers: OR_{wheeze} 1.21 (95% CI 0.96 to 1.50)) and there was a dose-dependent association between number of parents smoking and wheeze (one: OR 1.08 (95% CI 0.94 to 1.24); both: OR 1.24 (95% CI 1.05 to 1.47); $p_{\text{trend}} = 0.010$). A reduced ratio of forced expiratory volume in 1 second (FEV₁) to forced vital capacity (FVC) was related to father's smoking (−0.3% (95% CI −0.6 to 0)) and number of parents smoking ($p_{\text{trend}} < 0.001$) among men. In women, mother's smoking was associated with more respiratory symptoms and poorer lung function (OR_{wheeze} 1.15 (95% CI 1.01 to 1.31), never smokers: OR_{wheeze} 1.21 (95% CI 0.98–1.51); FEV₁ −24 ml (95% CI −45 to −3); FEV₁/FVC ratio −0.6% (95% CI −0.9 to −0.3)). These effects were possibly accounted for by maternal smoking in pregnancy (OR_{wheeze} 1.39 (95% CI 1.17 to 1.65); FEV₁ −23 ml (95% CI −52 to 7); FEV₁/FVC ratio −0.9% (95% CI −1.3 to −0.4)) as there was no association with paternal smoking among women (interaction by sex, $p < 0.05$). These results were homogeneous across centres.

Conclusion: Both intrauterine and environmental exposure to parental tobacco smoking was related to more respiratory symptoms and poorer lung function in adulthood in this multicultural study. The age window of particular vulnerability appeared to differ by sex, postnatal exposure being important only in men and a role for prenatal exposure being more evident in women.

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An increased risk for wheeze and asthma in children whose parents smoke is fairly well documented.^{1–4} Several studies have shown that intrauterine exposure to products from tobacco smoking in pregnancy reduces infant lung function.^{5–9} An independent effect of postnatal exposure to environmental tobacco smoke is suggested by an increased risk of asthma in children with fathers who smoke.¹ Even if some studies of children indicate that the harmful effects of parental smoking decrease when the children grow older,⁴ there is some evidence of permanent damage to the airways with reduced lung function^{10–11} and an increased risk for asthma^{12–13} or wheezing illness¹⁴ in adults exposed to parental smoking in childhood. Information on the long term consequences of parental smoking for adult respiratory health is, however, relatively scarce and we address this issue in the European Community Respiratory Health Survey (ECRHS).

There is some evidence that the airways of men and women respond differently to exposure to tobacco smoke products.^{3–6, 8, 11, 15, 16} This is plausible as there are differences between the male and female airways from early in fetal lung development and throughout life.^{17–18} Female lungs mature earlier with regard to surfactant production. Throughout life women have smaller lungs than men, but their lung architecture is more advantageous with a greater airway diameter in relation to the volume of the lung parenchyma. After puberty the relative advantage of the female airways is lost, possibly as a result of hormonal changes in women and

increased muscular strength among men which enhances lung function. Thus, in childhood airway hyperresponsiveness and asthma is more common among boys than girls, while this is reversed after puberty.^{17–19} It seems likely that the age windows for vulnerability to tobacco smoke products and to manifestation of symptoms could differ between the sexes. However, most studies do not give separate data for men and women. In this study we address possible sex differences in the effects of parental smoking on adult respiratory health. The ECRHS is large enough for analyses of interactions, and the international setting of the study to some extent allows for separation of heterogeneous effects of sociocultural sex differences and homogeneous effects related to biological sex differences.

METHODS

Data collection

The methodology for the ECRHS has been fully described elsewhere.^{20–21} Briefly, participating centres selected an area defined by pre-existing administrative boundaries with a population of at least 150 000. At least 1500 men and 1500 women aged 20–44 years were randomly selected from each centre. In stage I subjects were sent the ECRHS screening

Abbreviations: BHR, bronchial responsiveness; ECRHS, European Community Respiratory Health Survey; ETS, environmental tobacco smoke; FEV₁, forced expiratory volume in 1 second; FVC, forced vital capacity

Table 1 Parental smoking as reported by men and women aged 20–45 years

	Yes	No	Don't know	Not answered	Total
Did your father ever smoke regularly during your childhood?	11980 (61.5%)	5960 (31.5%)	641 (3.4%)	341 (1.8%)	18922
Did your mother ever smoke regularly during your childhood or before you were born?	5196 (27.5%)	13143 (69.5%)	247 (1.3%)	336 (1.8%)	18922
If "yes" (to mother smoking): When your mother was pregnant, in particular with you, did she smoke?	1994 (38.4%)	741 (14.2%)	2090 (40.1%)	371 (7.2%)	5196

questionnaire, a self-completed questionnaire about adult respiratory health. In stage II a smaller random sample of subjects who had completed the screening questionnaire was invited to attend for a more detailed interviewer-led questionnaire, lung function testing, and blood tests. The informed consent of all participants was obtained and the study was approved by all the involved ethics committees. This analysis includes data for 18 922 subjects from 37 centres representing 17 countries.

Specific IgE to house dust mite, cat, timothy grass and *Cladosporium* was measured in serum samples provided by 13 972 (74%) of the subjects. The test for specific IgE was considered to be positive if >0.35 kU/l. Details of the IgE measurements are described elsewhere.²² "Atopy" was defined as having specific IgE to cat, grass, house dust mite, and/or mould.

Forced expiratory volume in 1 second (FEV₁) and forced vital capacity (FVC) were recorded by a standard spirometric method.²³ Spirometric data were available for 15 901 (84%) subjects. The ratio between FEV₁ and FVC was calculated and used as a continuous variable unless otherwise explicitly stated. Metacholine challenge was performed with a dosimeter (Mefar, Brescia, Italy), providing information for 13 206 (70%) subjects. The degree of bronchial responsiveness (BHR) was expressed as the ECRHS slope.²⁴ Height was measured before measurement of lung function.

Information on parental smoking habits was collected using an interviewer-led questionnaire (table 1). Nearly all subjects gave information about maternal and paternal smoking, while a substantial number answered "don't know" to the question about maternal smoking in pregnancy. A variable describing whether one or both parents had smoked was created in order to provide a crude graded measure of exposure.

"Asthma" was defined as using current asthma medication or reporting asthma attacks during the previous 12 months.

"Three or more asthma symptoms" was defined as answering "yes" to three or more questions about asthma symptoms during the previous 12 months (wheeze, wheeze with shortness of breath, wheeze without having a cold, waking with tightness in chest, waking with shortness of breath, night cough, asthma attacks, asthma medication). "Chronic bronchitis" was defined as having both regular cough and regular phlegm.

Statistical analysis

Logistic regression models were used to assess the independent effects of parental smoking in childhood on adult respiratory symptoms, and adjustments were made for number of siblings, sex, age, body mass index, current passive smoking, current smoking (never, ex, and current smoking), current occupation (European Economic Community Status Groups-14),²⁵ and study centre. Linear regression models were used to analyse the effects of parental smoking on lung function, adjusting for the same confounders and for the interaction terms of sex with age, height, weight, and weight squared in analyses of all subjects, for age, height, weight and weight squared in sex-specific analyses. Potential heterogeneity between centres in the effect of each exposure variable on wheeze was studied by meta-analyses according to DerSimonian and Laird.²⁶ All analyses were carried out using the statistical software program Stata 7.0 (Stata Corporation, Texas, USA).

Differences according to sex, adult smoking status, and atopy were investigated. As all these aspects were important but sub-stratification by three variables is not feasible, the following strategy was chosen: Data for all subjects are presented to allow comparison with studies that do not stratify by sex; data for men and women are presented separately due to differences between the sexes in airways development. Non-atopic and atopic subjects differed with regard to associations with maternal smoking and these data

Table 2 Prevalence of parental smoking by country in study subjects born 1945–72

Country	No	Paternal smoking (%)*	Maternal smoking (%)*	Maternal smoking in pregnancy (%)*
Iceland	564	58	41	17
Norway	835	65	33	15
Sweden	1856	59	36	16
Estonia	431	63	10	2
Ireland	454	76	50	27
Great Britain	1579	68	43	20
Netherlands	1247	79	34	13
Belgium	1122	70	27	11
Germany	1983	67	27	5
Switzerland	853	57	21	4
France	2125	66	16	4
Spain	1942	71	3	1
Italy	894	71	17	5
New Zealand	1254	66	43	14
Australia	668	62	34	18
USA	723	63	44	24
Denmark	391	76	53	23
Total	18922	67	28	11

*Subjects answering "don't know" to question on paternal or maternal smoking are excluded.

are presented for men and women together as the associations with maternal smoking did not differ significantly by sex. Data for never smokers are presented in case of residual confounding by smoking.

RESULTS

In all, 67% of subjects reported that their father had smoked during their childhood. There were relatively small differences in this proportion between countries (table 2). Maternal smoking varied widely, from 3% to 17% in the centres in Spain, Estonia, France and Italy to over 40% in Denmark, Iceland and the English speaking centres. Similarly, maternal smoking in pregnancy varied greatly, from 1% in Spain to 27% in Ireland (table 2).

Considering men and women together (table 3), paternal smoking was not significantly associated with adult respiratory symptoms or lung function. Maternal smoking was associated with an increased risk for respiratory symptoms and with reduced FEV₁ and FEV₁/FVC ratio. The associations of maternal smoking with wheeze (fig 1) and with FEV₁/FVC ratio <75% were consistent between centres (p_{heterogeneity} = 0.4 and p_{heterogeneity} = 0.3, respectively). Among subjects who gave information about maternal smoking in pregnancy, this exposure was strongly associated with respiratory symptoms and with reduced FEV₁/FVC ratio (table 3) which was consistent between centres (wheeze: p_{heterogeneity} = 0.2). The association of parental smoking and measures of adult respiratory health increased in strength with number of smoking parents, a dose-response relationship that was significant for several symptoms, FEV₁ and FEV₁/FVC ratio (table 3). Adult asthma or BHR was not associated with any measure of parental smoking.

An association of adult respiratory symptoms and father's smoking was identified in men (table 4, fig 2) but not in women (table 5). This difference between the sexes was significant (wheeze: p_{interaction} = 0.033; wheeze without cold: p_{interaction} = 0.011) and consistent between centres (p_{heterogeneity} = 0.2 for the interaction effect of sex with paternal smoking on wheeze). Father's smoking was further associated with decreased lung function in men but not in women; interaction by sex was not statistically significant (FEV₁: p_{interaction} = 0.1; FEV₁/FVC ratio: p_{interaction} = 0.4). The risk for respiratory symptoms and poor lung function increased with an increasing number of parents smoking only in men (table 4). These differences in the sexes were also significant (wheeze: p_{interaction} = 0.024 (excluding maternal smoking in pregnancy); FEV₁: p_{interaction} = 0.035). Mother's smoking was associated with increased risk for respiratory symptoms and reduced lung function in women (table 5); the effects were particularly strong for maternal smoking during pregnancy. Most associations with maternal smoking in men did not reach statistical significance, although the estimates were only slightly smaller than in women.

When only never smokers were considered, the key findings were similar to those of the total population although many estimates did not reach statistical significance (table 6). In men, paternal smoking was associated with an increased risk for symptoms and an indicated decrease in the FEV₁/FVC ratio. In women, maternal smoking was associated with more symptoms and poorer lung function. These findings did not show a significant interaction with adult smoking status. The differences between the sexes in the associations of paternal smoking with symptoms and maternal smoking with symptoms and lung function were significant or of borderline significance (table 6). The full table of sex-specific effects of parental smoking stratified by current adult smoking status is available online at www.thoraxjnl.com/supplemental.

Table 3 Association of adult respiratory symptoms and lung function with parental smoking in childhood

	Paternal smoking		Maternal smoking		Maternal smoking in pregnancy		One parent smoking		Both parents smoking		Dose-response	
	Prevalence of symptom*	OR (95%CI)†	p value	OR (95%CI)†	p value	OR (95%CI)†	p value	OR (95%CI)†	p value	OR (95%CI)†	p value	p value
Wheeze	23% (4308/18688)	1.03 (0.94 to 1.13)	0.5	1.12 (1.02 to 1.23)	0.016	1.24 (1.09 to 1.42)	0.001	1.02 (0.92 to 1.12)	0.001	1.15 (1.02 to 1.29)	0.022	0.022
>3 asthma symptoms	16% (2861/18446)	1.08 (0.98 to 1.20)	0.1	1.14 (1.02 to 1.26)	0.016	1.28 (1.11 to 1.48)	0.001	1.03 (0.92 to 1.15)	0.004	1.22 (1.07 to 1.39)	0.004	0.004
Current asthma	5% (874/18626)	0.90 (0.76 to 1.06)	0.2	1.06 (0.89 to 1.25)	0.5	1.14 (0.89 to 1.44)	0.3	0.86 (0.72 to 1.03)	0.6	0.96 (0.77 to 1.18)	0.6	0.6
Chronic bronchitis	11% (1970/18638)	1.12 (0.99 to 1.27)	0.082	1.19 (1.05 to 1.35)	0.008	1.32 (1.11 to 1.57)	0.002	1.08 (0.94 to 1.25)	<0.001	1.34 (1.14 to 1.57)	<0.001	<0.001
		β coeff (95%CI)†	p value	β coeff (95%CI)†	p value	β coeff (95%CI)†	p value	β coeff (95%CI)†	p value	β coeff (95%CI)†	p value	p value
FEV ₁ (ml)		-6 (-23 to 11)	0.5	-24 (-42 to -5)	0.012	-24 (-50 to 3)	0.082	-7 (-26 to 11)	0.010	-31 (-53 to -8)	0.010	0.010
FVC (ml)		4 (-15 to 24)	0.7	-1 (-22 to 20)	0.9	19 (-11 to 50)	0.2	3 (-18 to 24)	0.7	4 (-22 to 30)	0.7	0.7
FEV ₁ /FVC ratio (%)		-0.2 (-0.4 to 0)	0.094	-0.5 (-0.8 to -0.3)	<0.001	-0.9 (-1.3 to -0.5)	<0.001	-0.2 (-0.4 to 0)	<0.001	-0.8 (-1.1 to -0.5)	<0.001	<0.001
BHR (ECRHS slope)		0.00 (-0.07 to 0.08)	0.9	0.04 (-0.5 to 0.12)	0.6	0.01 (-0.12 to 0.14)	0.9	-0.06 (-0.03 to 0.14)	0.5	0.03 (-0.08 to 0.13)	0.5	0.5

*The number of subjects answering each question varied, thus the denominators are different.

†Odds ratios with 95% confidence intervals as estimated by logistic regression adjusting for sex, age, body mass index, number of siblings, current smoking, current environmental tobacco smoking, current occupation, and study centre.

‡Analysed with adjustment for interaction of sex with height, age, weight and weight squared, number of siblings, current smoking, current environmental tobacco smoking, current occupation, and study centre.

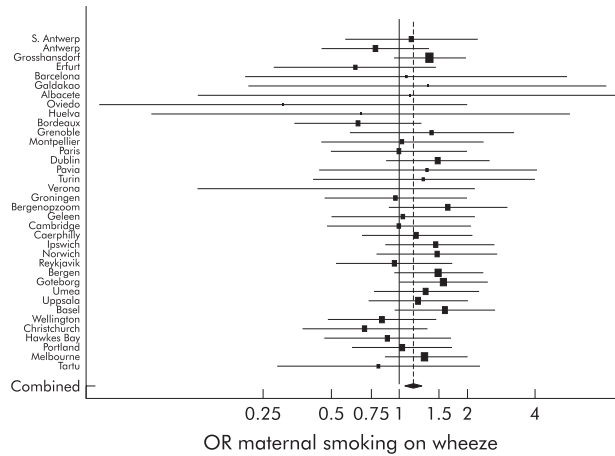


Figure 1 Odds ratios for the association of maternal smoking and adult wheeze by centre. Adjustment within centre for paternal smoking, adult passive smoking, adult smoking, number of siblings, sex, age, and occupation. For each centre horizontal lines indicate 95% CI. For combined odds ratio the diamond indicates 95% CI from model with centre as random effect. The size of each square is proportional to the sample size.

The effects of maternal smoking on respiratory symptoms differed with regard to atopic status, with a stronger effect of maternal smoking on some symptoms among non-atopic subjects (table 7; wheeze: $p_{\text{interaction}} = 0.024$). A corresponding difference by atopy was not found for lung function. Interactions by atopy in the associations of paternal smoking with respiratory symptoms or lung function were smaller and not significant. Atopy was a negative confounder for the associations between parental smoking and respiratory symptoms. Adjustment increased the strength of the associations but was not included in the final results because of the interaction effects noted above. Centre prevalence of wheeze was strongly correlated with centre prevalence of maternal smoking ($r = 0.57$, fig 3). The English speaking centres had a particularly high prevalence of wheeze and of maternal smoking. In the Latin countries, maternal smoking in pregnancy was uncommon and the prevalence of wheeze was relatively low in most centres.

DISCUSSION

Subjects whose parents had smoked experienced more respiratory symptoms and poorer lung function in adulthood consistently across 37 centres in the western world. Thus, parental tobacco smoking appears to influence the development of the child’s airways with permanent consequences in terms of poorer adult respiratory health. Our analysis indicated that both intrauterine exposure to maternal smoking during pregnancy and postnatal exposure to ETS was of importance. In men, exposure to father’s smoking in childhood was related to more respiratory symptoms and more airways obstruction in adulthood, and there was a dose-response relationship between the number of parents smoking and respiratory symptoms and lung function. There was therefore a convincing effect of postnatal exposure to ETS on adult lung health among men. The possible influence of intrauterine exposure could not be separated from postnatal effects of maternal smoking. In women, no convincing effect of postnatal exposure to paternal smoking could be identified, so the consistent associations of mother’s smoking with respiratory symptoms and poorer lung function in adult women may be related to the prenatal exposure to products from maternal smoking during pregnancy. This study therefore indicates that the age window for particular vulnerability to parental smoking differs between men and women.

Table 4 Association of adult respiratory symptoms and lung function with parental smoking in childhood in 7678 men

	Paternal smoking		Maternal smoking		Maternal smoking in pregnancy		One parent smoking		Both parents smoking		Dose-response	
	OR (95% CI)*	p value	OR (95% CI)*	p value	OR (95% CI)*	p value	OR (95% CI)*	p value	OR (95% CI)*	p value	β coeff (95% CI)†	p value
Wheeze	1.13 (1.00 to 1.28)	0.058	1.11 (0.97 to 1.26)	0.1	1.11 (0.95 to 1.28)	0.3	1.08 (0.94 to 1.24)	0.010	1.24 (1.05 to 1.47)	0.010	-40 (-77 to -3)	0.037
> 3 asthma symptoms	1.20 (1.03 to 1.39)	0.022	1.11 (0.95 to 1.29)	0.2	1.15 (0.91 to 1.48)	0.3	1.08 (0.91 to 1.27)	0.007	1.31 (1.08 to 1.59)	0.007	6 (-37 to 49)	0.8
Current asthma	0.95 (0.74 to 1.22)	0.7	1.02 (0.79 to 1.33)	0.9	0.98 (0.65 to 1.48)	0.9	0.86 (0.66 to 1.12)	0.8	0.99 (0.72 to 1.36)	0.8	-0.8 (-1.3 to -0.4)	<0.001
Chronic bronchitis	1.22 (1.02 to 1.45)	0.031	1.16 (0.97 to 1.39)	0.1	1.45 (1.12 to 1.88)	0.005	1.21 (0.99 to 1.47)	0.004	1.41 (1.12 to 1.77)	0.004	0.03 (-0.11 to 0.17)	0.6
	β coeff (95% CI)†	p value	β coeff (95% CI)†	p value	β coeff (95% CI)†	p value	β coeff (95% CI)†	p value	β coeff (95% CI)†	p value		
FEV ₁ (ml)	-15 (-42 to 13)	0.3	-22 (-52 to 7)	0.1	-31 (-78 to 16)	0.2	-14 (-43 to 16)	0.2	-40 (-77 to -3)	0.037		
FVC (ml)	4 (-27 to 36)	0.8	0 (-34 to 35)	1.0	27 (-27 to 81)	0.3	1 (-33 to 35)	0.3	6 (-37 to 49)	0.8		
FEV ₁ /FVC ratio (%)	-0.3 (-0.6 to 0.0)	0.082	-0.5 (-0.8 to -0.1)	0.012	-1.0 (-1.6 to -0.4)	<0.001	-0.2 (-0.6 to 0.1)	<0.001	-0.8 (-1.3 to -0.4)	<0.001		
BHR (ECRHS slope)	-0.1 (-0.1 to 0.1)	0.9	0.05 (-0.06 to 0.17)	0.4	0.05 (-0.14 to 0.23)	0.6	0.08 (-0.04 to 0.19)	0.6	0.03 (-0.11 to 0.17)	0.6		

*Odds ratios with 95% confidence intervals as estimated by logistic regression adjusting for age, body mass index, current smoking, current environmental tobacco smoking, current occupation, and study centre.
†Analysed with adjustment for height, age, weight, weight squared, number of siblings, current smoking, current environmental tobacco smoking, current occupation, and study centre.

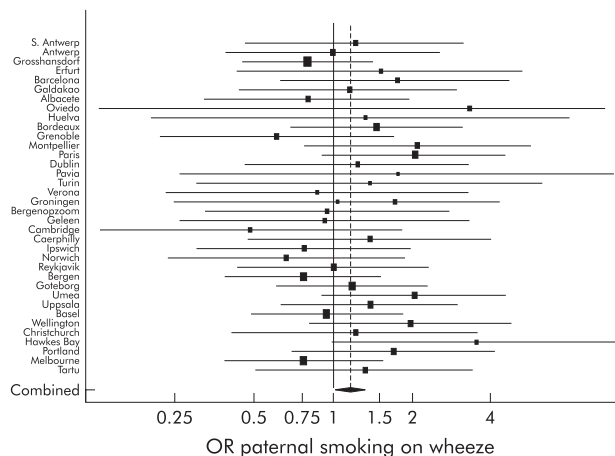


Figure 2 Odds ratios for the association of paternal smoking and adult wheeze among men by centre. Adjustment within centre for maternal smoking, adult passive smoking, adult smoking, number of siblings, age, and occupation. For each centre, horizontal lines indicate 95% CI. For combined odds ratio, diamond indicates 95% CI from model with centre as random effect. The size of each square is proportional to the sample size.

Recall bias is a potential problem in this study. However, parental smoking was significantly associated with reduced lung function also among non-wheezers. Nearly all participants (99% and 97%) reported whether their mother or father had smoked when they were children, even though they had been given an option of responding “don’t know” (an option used by 40% when asked about mother’s smoking in pregnancy). Symptomatic and non-symptomatic subjects might have recalled parental smoking differently but, for differential recall to explain our findings, non-atopic subjects rather than atopic subjects, and wheezers rather than diagnosed asthmatics, would have over-reported parental smoking. This seems unlikely. Thus, while we do not have information to validate the information on parental smoking, the role of differential recall bias seems to be limited. Information on whether the parents smoked or not, however, gives a very crude measure of exposure and, together with non-differential measurement error, this may be the reason why some main effects in subgroups did not reach statistical significance despite large sample sizes and near identical point estimates between subgroups. The small but consistent effects may therefore possibly reflect stronger underlying associations. The strengths of the study are the overall sample size that permit subgroup analyses and the broad range of populations studied in an international setting.

The main results were similar in never smokers to those in the total population sample, arguing against a role of residual confounding by current smoking habits and related lifestyle factors. The findings were also consistent between centres that, particularly during the post-war years, differed with regard to social, economic and cultural factors as illustrated by the wide variation in women’s smoking habits (table 2). This argues against a major role for confounding by smoking related lifestyle factors of the parents.

The quality of the information on maternal smoking in pregnancy is considered less reliable and is not an important basis for the conclusions of this paper. The 40% who reported that their mother had smoked when she was pregnant probably constitute a group with intrauterine exposure and, possibly, with heavier postnatal exposure. Those who answered “don’t know” may include a considerable number with intrauterine exposure as well. The lack of recall in this group could be suspected to be associated with better adult

Table 5 Association of adult respiratory symptoms and lung function with parental smoking in childhood in 7799 women

	Paternal smoking		Maternal smoking		Maternal smoking in pregnancy		One parent smoking		Both parents smoking		Dose-response	
	OR (95% CI)*	p value	OR (95% CI)*	p value	OR (95% CI)*	p value	OR (95% CI)*	p value	OR (95% CI)*	p value	β coeff (95% CI)†	p value
Wheeze	0.95 (0.83 to 1.08)	0.4	1.15 (1.01 to 1.31)	0.032	1.39 (1.17 to 1.65)	<0.001	0.96 (0.84 to 1.10)	0.3	1.09 (0.92 to 1.28)	0.001	0.04 (-0.09 to 0.17)	0.7
>3 asthma symptoms	1.00 (0.87 to 1.14)	0.9	1.16 (1.01 to 1.33)	0.040	1.42 (1.18 to 1.71)	<0.001	0.99 (0.85 to 1.16)	0.1	1.16 (0.97 to 1.39)	0.001	0 (-0.21 to 0.22)	0.2
Current asthma	0.84 (0.68 to 1.05)	0.1	1.11 (0.88 to 1.40)	0.4	1.28 (0.95 to 1.73)	0.1	0.85 (0.66 to 1.08)	0.6	0.94 (0.70 to 1.24)	0.083	5 (-1 to 30)	0.8
Chronic bronchitis	1.02 (0.85 to 1.23)	0.8	1.22 (1.01 to 1.46)	0.035	1.24 (0.97 to 1.58)	0.083	0.97 (0.79 to 1.19)	0.040	1.28 (1.01 to 1.61)	0.001	-0.7 (-1.1 to -0.3)	0.001
FEV ₁ (ml)	4 (-16 to 24)	0.7	-24 (-46 to -3)	0.025	-23 (-52 to 7)	0.1	0 (-21 to 22)	0.2	-19 (-46 to 7)	0.001	0.03 (-0.13 to 0.19)	0.7
FVC (ml)	5 (-18 to 28)	0.7	-3 (-27 to 21)	0.8	13 (-21 to 47)	0.5	5 (-19 to 30)	0.8	4 (-26 to 34)	0.001	-0.7 (-1.1 to -0.3)	0.001
FEV ₁ /FVC ratio (%)	-0.1 (-0.4 to 0.2)	0.6	-0.6 (-0.9 to -0.3)	<0.001	-0.9 (-1.3 to -0.4)	<0.001	-0.1 (-1.1 to 0.9)	0.001	-0.7 (-1.1 to -0.3)	0.001	0.03 (-0.13 to 0.19)	0.7
BHR (ECRHS slope)	0.02 (-0.10 to 0.14)	0.7	0.01 (-0.12 to 0.14)	0.9	-0.03 (-0.21 to 0.15)	0.7	0.04 (-0.09 to 0.17)	0.001	0.03 (-0.13 to 0.19)	0.001		

*Odds ratios with 95% confidence intervals as estimated by logistic regression adjusting for age, body mass index, current smoking, current environmental tobacco smoking, current occupation, and study centre.
 †Analysed with adjustment for height, age, weight, weight squared, number of siblings, current smoking, current environmental tobacco smoking, current occupation, and study centre.

Table 6 Association of adult respiratory symptoms and lung function with maternal smoking in childhood in never smoking men (n = 3043) and women (n = 3582)

	Men		Women	
	Paternal smoking	Maternal smoking	Paternal smoking	Maternal smoking
	OR (95% CI)*	OR (95% CI)*	OR (95% CI)*	OR (95% CI)*
Wheeze	1.21 (0.96 to 1.50)§	0.87 (0.68 to 1.10)§	0.93 (0.76 to 1.13)	1.21 (0.98 to 1.51)
>3 asthma symptoms	1.13 (0.87 to 1.47)	0.72 (0.53 to 0.97)‡	0.89 (0.72 to 1.10)	1.20 (0.95 to 1.52)
Asthma	1.08 (0.76 to 1.54)§	0.81 (0.54 to 1.23)	0.67 (0.49 to 0.91)	1.10 (0.78 to 1.55)
Chronic bronchitis	1.00 (0.70 to 1.42)	1.07 (0.73 to 1.56)	0.94 (0.70 to 1.25)	1.13 (0.82 to 1.56)
	β coeff (95% CI)†	β coeff (95% CI)†	β coeff (95% CI)†	β coeff (95% CI)†
FEV ₁ (ml)	3 (-38 to 43)	-2 (-49 to 44)	16 (-12 to 44)	-25 (-58 to 8)
FVC (ml)	15 (-32 to 62)	-10 (-64 to 44)	5 (-27 to 38)	4 (-34 to 41)
FEV ₁ /FVC ratio (%)	-0.2 (-0.7 to 0.3)	0.1 (-0.4 to 0.7)§	0.2 (-0.2 to 0.6)	-0.8 (-1.3 to -0.3)
BHR (ECRHS slope)	-0.07 (-0.23 to 0.09)	0.19 (0.01 to 0.38)	0.18 (0.01 to 0.35)	0.11 (-0.08 to 0.31)

*Odds ratios with 95% confidence intervals as estimated by logistic regression adjusting for age, body mass index, number of siblings, current environmental tobacco smoking, current occupation, and study centre.

†Analysed with adjustment for height, age, weight, weight squared, number of siblings, current environmental tobacco smoking, current occupation, and study centre.

‡Interaction effect by sex, $p < 0.05$.

§Interaction effect by sex, $p \leq 0.1$.

respiratory health, but more symptoms and poorer lung function were also observed in this group.

Our findings are in agreement with studies in children.¹⁻⁴ Based on meta-analyses of a large number of relevant studies, Cook and Strachan concluded that father's smoking—as well as mother's smoking—was related to respiratory symptoms in children, with strong evidence for a causal relationship.¹ The effect of paternal smoking, often not evident in smaller studies but convincing in their meta-analysis, argues for an independent effect of postnatal passive smoking. An in utero effect of maternal smoking during pregnancy has been indicated by several studies, most convincingly shown as changes in infant lung function.⁵⁻⁹ A few studies follow the effects of parental smoking into adult life. Upton *et al*¹⁰ observed a significant effect of maternal (but not paternal) smoking on lung function in 2295 adults, including a dose-response relationship with the daily number of cigarettes smoked. Masi *et al*¹¹ showed an effect of childhood ETS on lung function in young men, while

Strachan *et al*¹⁴ reported an effect of maternal smoking in pregnancy on wheezing illness before age 7 and at ages 17–33 years in the 1958 British cohort. Hu *et al*¹³ found an increased asthma risk in young adults related to paternal as well as maternal smoking, and a dose-response relationship with the number of parents smoking. Larsson *et al*¹² showed significant effects of childhood passive smoking on asthma among 3556 never smoking adults in Sweden, while Jenks *et al*²⁷ found no significant associations of self-reported asthma or wheeze with paternal or maternal smoking in a study of 1494 young adults.

In our study the vulnerability towards prenatal and postnatal exposure to tobacco smoke appeared to differ between men and women. In agreement with our findings in adults, several studies indicate a stronger effect of intrauterine exposure on female than on male newborns.^{4 5 8} It has been suggested that nicotine induces “masculinisation” of the airways with narrower airways, reduced airway compliance, and reduced elastic recoil.⁸ This view is supported by

Table 7 Association of adult asthma and lung function and maternal smoking in childhood according to atopic status (atopy defined as specific IgE to cat, grass, house dust mite and/or mould) (including 13791 subjects with information from blood tests as well as other variables)

	Non-atopics (N = 9170)		Atopics (N = 4383)	
	OR (95% CI)*	p value	OR (95% CI)*	p value
Wheeze	1.23 (1.08 to 1.40)	0.002	1.04 (0.89 to 1.21)	0.6
>3 asthma symptoms	1.24 (1.06 to 1.44)	0.007	1.11 (0.93 to 1.31)	0.2
Current asthma	1.11 (0.78 to 1.57)	0.6	1.15 (0.92 to 1.44)	0.2
Chronic bronchitis	1.21 (1.01 to 1.44)	0.032	1.17 (0.94 to 1.47)	0.2
	β coeff (95% CI)†	p value	β coeff (95% CI)†	p value
FEV ₁ (ml)	-21 (-45 to 2)	0.070	-19 (-55 to 17)	0.3
FVC (ml)	-3 (-30 to 24)	0.8	10 (-30 to 50)	0.6
FEV ₁ /FVC ratio (%)	-0.5 (-0.8 to -0.2)	0.003	-0.6 (-1.1 to -0.2)	0.008
BHR (ECRHS slope)	0.04 (-0.06 to 0.14)	0.5	-0.03 (-0.20 to 0.14)	0.7

*Odds ratios with 95% confidence intervals as estimated by logistic regression adjusting for sex, age, body mass index, number of siblings, current smoking, current environmental tobacco smoking, current occupation, and study centre.

†Analysed with adjustment for interaction of sex with height, age, weight and weight squared, number of siblings, current smoking, current environmental tobacco smoking, current occupation, and study centre.

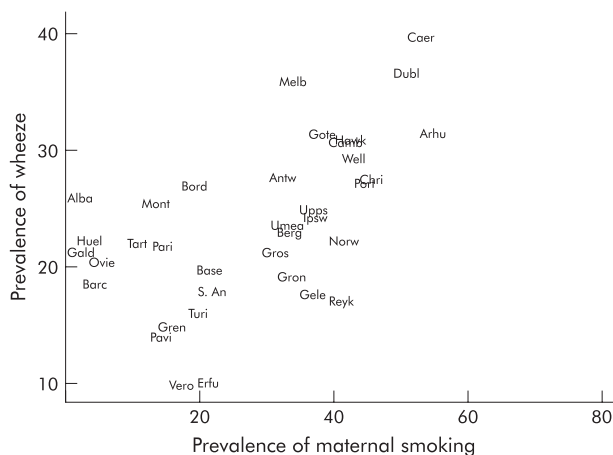


Figure 3 Prevalence of wheeze in subjects aged 20–44 years and prevalence of maternal smoking in 37 study centres in 17 countries.

animal studies showing structural changes in the fetal lungs including loss of elastic recoil, lung hypoplasia, and atelectasis related to in utero exposure to tobacco smoke products.^{28–30}

The stronger association between childhood ETS and obstructive lung disease in men, as shown in this study and by Hu *et al.*,¹³ could be related to boys having less mature lungs and relatively narrower airways during childhood. These factors are believed to contribute to the generally higher rates of pulmonary morbidity in boys than in girls, and could possibly also explain a higher susceptibility for permanent damage by exposure to ETS during this age window. It has been suggested that the association of paternal smoking and asthma among males could be explained by boys spending more time with their fathers than girls. However, in the multicultural setting of this study this association was homogeneous across centres (fig 2), so a biological explanation seems more plausible than a behavioural explanation that would tend to differ between cultures.

Parental smoking was associated with respiratory symptoms, reduced FEV₁, and increased airways obstruction, but not with diagnosed asthma and increased bronchial hyperresponsiveness. Exposure to tobacco smoke early in life may influence the development of airways structure (reflected in FEV₁ and FEV₁/FVC ratio) more than susceptibility to mucosal inflammation decades later (contributing relatively more to BHR), and this may contribute to a clinical picture slightly different from classical asthma.

Parental smoking was strongly associated with respiratory symptoms among non-atopic subjects. This suggests that parental smoking is an important causal factor for non-atopic asthma, a condition that deserves far more focus among researchers. Born in 1945–72, our study subjects are the offspring of the post-war generation of parents. Smoking among men was common all over the western world during the post-war years. The young women in the English speaking and the Northern European countries often took up the habit of smoking during this period, while smoking was still uncommon among women in the Latin countries. Differences in post-war smoking habits of women could partly explain the high prevalence of wheezy illness observed in the English speaking countries.

In conclusion, both mother's and father's smoking appears to cause permanent damage to the developing airways of the child, leading to more respiratory symptoms and impaired lung function in adult life. Exposure to ETS in childhood was related to more respiratory symptoms in men but not in

women, while maternal smoking in pregnancy increased the risk for obstructive lung disease in women and, possibly, also in men. We speculate that tobacco smoke products lead to structural changes in the developing lungs in females, particularly during fetal life, and in males to a larger extent postnatally. Other factors such as sex specific growth of the airways and hormonal factors may contribute to determine at what age and to what extent this susceptibility is manifested as respiratory symptoms. In young adults we found an increased risk for obstructive lung disease in both men and women whose parents had smoked, although this appeared to be related to exposure at different age windows during lung development.

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The full table of sex-specific effects of parental smoking stratified by current adult smoking status is available on the *Thorax* website at www.thoraxjnl.com/supplemental

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