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Impact of close-proximity air pollution on lung function in schoolchildren in the French West Indies

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Abstract

Background: High levels of asthma prevalence and severity of respiratory symptoms have been found in the Caribbean but little is known about the impact of air pollution in these regions. This study aimed to describe air pollution and measure the associations with child lung function in Guadeloupe (French West Indies).

Methods: Data from 30 randomly chosen elementary schools (8–13 years old) were obtained using a standardized protocol adapted from the ISAAC2 study. We considered two health outcomes: peak expiratory flow (PEF) before running and the variation in peak expiratory flow (Δ PEF) after running. The associations between pollutants and outcomes were investigated using several air pollution exposure models: i) medium-term exposure to close-proximity pollution both indoor and outdoor for ozone (O_3) and nitrogen dioxide (NO_2) and ii) short- and medium-term exposure to background pollution for O_3 , NO_2 , sulphur dioxide (SO_2) and small particulate matter (PM_{10}).

Results: Of 1,463 children, 277 (16%) were found to have asthma. A $1\text{-}\mu\text{g}/\text{m}^3$ increase in medium-term exposure to outdoor close-proximity pollution by O_3 was associated with a PEF decrease ($\beta = -0.32$; 95% CI: -0.61 ;- 0.03). No association was found with NO_2 regarding close-proximity pollution. The association between medium-term exposure to background pollution and PEF decrease was stronger in asthmatic children than in non-asthmatic children for O_3 . No reduction in PEF or Δ PEF was shown with NO_2 , SO_2 and PM_{10} pollutants but a significant association was found between PM_{10} and PEF increase.

Conclusions: Our results suggest that O_3 could have an acute effect on child lung function in the Caribbean even at a low concentration (below the WHO guidelines). Further research in the Caribbean is needed to confirm these findings.

Keywords: Asthma, ISAAC, Peak expiratory flow, Pollutants, Schoolchildren

Background

As with obesity and diabetes, the prevalence of asthma and allergic diseases has dramatically increased during the last three decades, particularly in children [1]. The possible explanations for this worldwide increase are either genetic variability or changes in environmental factor exposure. The genetic predisposition of each individual has

been shown to play a role in the development of allergic diseases, especially in the pathophysiological processes involved. With current knowledge of population genetics, a recent modification in genetic variability seems unlikely. However, the assumption of an interaction between individual susceptibility and environmental stimuli after birth is possible. This interaction could produce epigenetics that subsequently influence susceptibility to chronic inflammatory disease [2]. Shifts in environmental factors including exposure to air pollutants, allergens and infections might be associated with this increase.

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Several epidemiological studies have investigated the associations between exposure to urban air pollution and respiratory and allergic diseases among children, and have identified positive associations between background concentrations of nitrogen dioxide (NO₂) [3,4], sulphur dioxide (SO₂) [5,6], particulate matter (PM) [7,8] and ozone (O₃) [9,10].

Particles and NO₂ have shown more significant results. Negative associations between air pollution and pulmonary health were more pronounced in asthmatic children in this review [11]. Peak expiratory flow (PEF) was the most usual measurement for children's lung function, followed by the measure of forced expiratory volume in one second (FEV₁). In a meta-analysis, Weinmayr et al. reported for 51 studies (36 from Europe and 15 elsewhere mainly in USA) short-term effects of PM₁₀ and PM_{2.5} on respiratory symptoms and lung function mainly PEF [7]. For an increase of 10 µg/m³ of PM₁₀, they found a significant increase of 2.8% in asthma symptoms, an increase for cough (1.2%), and a decrease of PEF (-0.082 l/min). In addition, a higher decrease of PEF was found in asthmatic (-0.549 l/min) compared with others children (0.010 l/min). However, the impact of ozone on respiratory health was not studied in this meta-analysis. In healthy children living in UK, an increase of 1.0 µm² in the mean area of black carbon in airway macrophage, was associated with a decrease of 17% in FEV₁ [12]. In USA, a longitudinal study of children in 12 communities in California, reported a significant correlation between reduced lung function growth and the background concentration in PM₁₀ [13]. In Brazilia, Jacobson et al. in a panel study estimated the effect of current levels of outdoor pollution on peak expiratory flow [14]. The global effect was important for PM₁₀ with PEF reduction of 0.31 l/min (CI 95%: -0.56;-0.05), however, ozone was not studied in this analysis. More recently, a nationwide in Taiwan, was conducted on schoolchildren age 6–15 years (n = 1494) and showed that subchronic exposure to ambient PM_{2.5} and ozone leads to reduce lung capacity, and acute exposure to ozone decreases mid-expiratory flow (-0.123 l) [15].

The impact of background air pollution on respiratory diseases seems to vary in different parts of the world probably due to differences in the spatial and temporal variability of the composition and sources of air pollutants between different urban areas [16]. To our knowledge, little is known about the impact of air pollution in the Caribbean with its tropical wet and dry environments (defined by relatively constant temperatures throughout the year and seasonal variations dominated by rainfall), while levels of asthma prevalence and severe respiratory symptoms in children are mostly higher than in Northern Temperate Zones [17].

Population-based surveys of Caribbean children have reported that over 13% of participants admitted to having a

past or present diagnosis of asthma [18–20]. In Guadeloupe, which is a French West Indies department located in the Caribbean, previous studies also reported high asthma prevalence in children under 13 years old and a hospital admission risk related to asthma 1.9-fold higher than in Metropolitan France [19,21].

The present study aimed to provide data on air pollution exposure in Guadeloupe and to measure the associations between air pollution and lung function in elementary schoolchildren. The associations between the pollutants and lung function were assessed using several air pollution exposure models: i) medium-term exposure to close-proximity pollution both indoor and outdoor and ii) short- and medium-term exposure to background pollution.

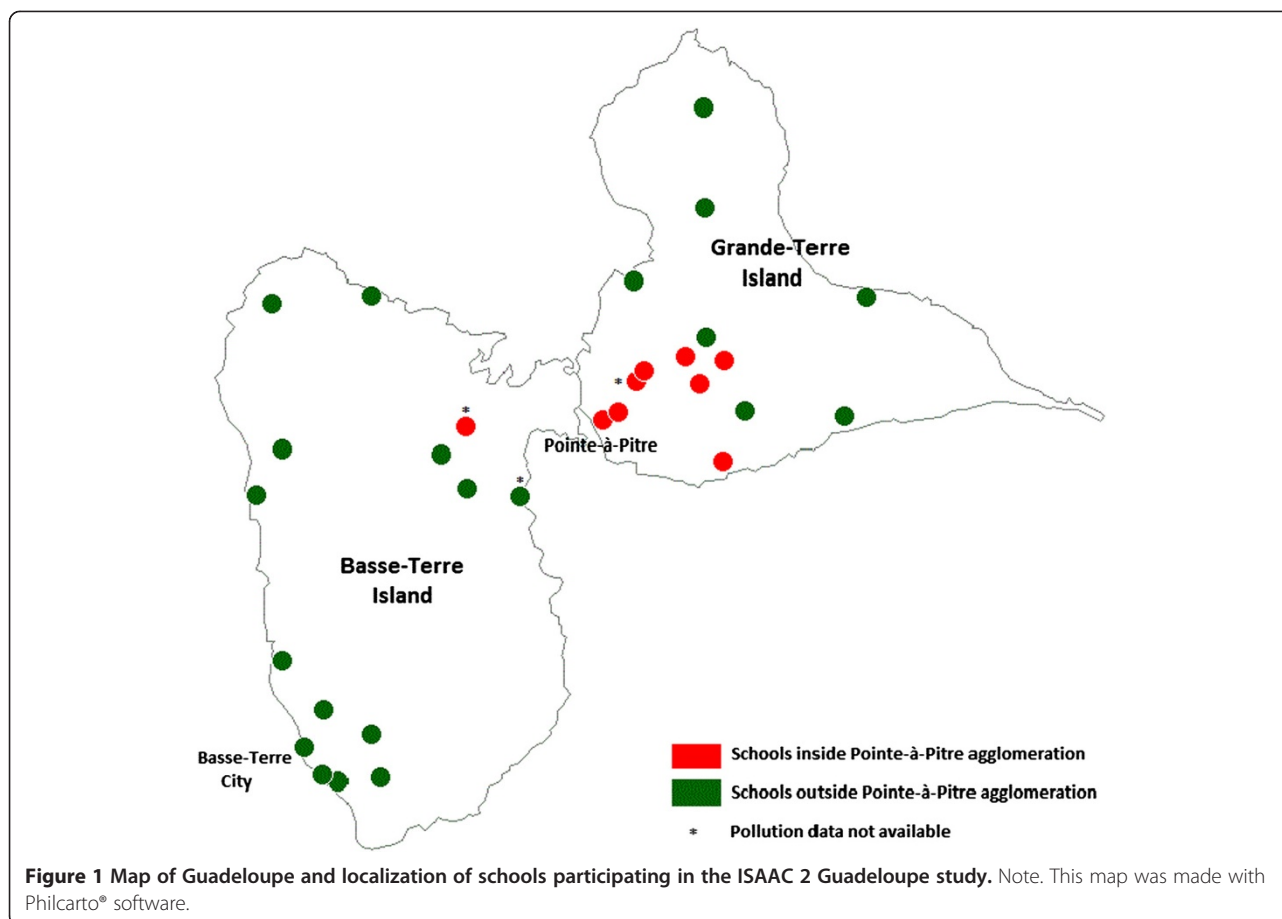
Methods

Participants and study design

This cross-sectional study was carried out between December 2008 and December 2009 using the second phase of the International Study of Asthma and Allergy in Childhood (ISAAC II) standardized protocol, which has been widely described in the literature [22]. A random representative sample of 30 elementary schools in Guadeloupe was chosen from a complete list of all elementary schools (Figure 1). As specified in the ISAAC II protocol [22], the minimum required sample size had to be above 1,000 children. Among the 30 elementary schools, completed valid data on clinical examination, questionnaire and pollution were available in 27 schools representing 1,436 children distributed in 91 classrooms. In three schools, pollution data could not be collected owing to a long strike in Guadeloupe (Figure 1).

The questionnaire contained the core ISAAC questions on respiratory and allergic diseases, how these were managed and potential risk factors. The clinical examination included the standardized protocol of the running test to assess exercise-induced asthma that was previously used in the French Six-City Study [8]. The detailed method has been described elsewhere [23]. The research protocol was approved by the National Ethics Board and all children and parents gave written informed consent. The timetable of the school visit for air pollution exposure assessment and the clinical examination of the children were randomly chosen.

During the study period, lung function using peak expiratory flow (PEF) was measured for each subject. Both background exposure and proximity exposure were used to assess personal exposure. Proximity pollution is an accurate measure of local pollution exposure. However, in our study, we only had the possibility to measure a part of the proximity pollution to which children were exposed (only at school) owing to economic and organizational constraints. For this reason, measuring background exposure was useful in our study.



Regarding medium-term exposure, concentrations of O_3 and NO_2 for close-proximity air pollution and concentrations of O_3 , NO_2 , and SO_2 and particulate matter with a diameter below to $10 \mu m$ (PM_{10}) for background air pollution were estimated for each classroom. All children in a classroom were assigned the same estimated pollution exposure. Regarding short-term exposure, the different lags of pollutant levels, corresponding to the measurement of pollutant levels the current day up to five days (D0, D1, D2, D3, D4, D5) before the measurement of PEF were estimated for each child.

Guadeloupe

Guadeloupe, a French overseas department located in the Caribbean (Figure 1), comprises two main islands: Basse-Terre Island and Grande-Terre, which are separated by a narrow sea channel. The land area represents 1,628 square kilometers with a density population of 249 inhabitants per square kilometers. The most important two cities are Basse-Terre, which is the capital of Guadeloupe, and Pointe-à-Pitre. Basse-Terre Island has a rugged terrain due to volcanic mountains, which explains why the main road network is only developed in Grande-Terre Island and around these two cities.

Health outcomes

Two health outcomes were investigated using the run test: baseline peak expiratory flow (PEF) before running and variation in peak expiratory flow (ΔPEF) after running. Baseline PEF was measured three times immediately before the run test with a Mini-Wright peak flow meter in accordance with the American Thoracic Society guidelines [24]. The maximum of these measures was retained and compared to the predicted value. The predicted value was assessed using the height value and the abacus built through the curve of Godfrey [25]. All the children with PEF superior or equal to 70% of the predicted value before running were invited to undergo a 6-min period of running. PEF was measured 5 and 10 min following the run test. The maximum of this second series of measurements was used to calculate the ΔPEF corresponding to the percentage decrease in PEF after exercise compared to the baseline PEF. Measurements of PEF were performed by a spirometry technician after training and a senior pulmonologist supervised the procedure. The interest of measuring the peak expiratory flow before running and the peak expiratory variation after it was to determine whether pollution had any impact on breathing at rest or on breathing after exercise,

respectively. In addition, Δ PEF represents exercise-induced asthma, which is considered to be a distinct asthma phenotype. Therefore, we exclude children with a PEF lower than 70% of the predicted value, i.e. probably the most vulnerable children.

The following allergic manifestations were considered: asthma was defined by an affirmative response to the question 'Has your child ever had asthma?'; atopic dermatitis by an affirmative response to the question 'Has your child ever had itchy rash affecting any of the following places: the folds of the elbows, behind the knees, in front of the ankles, under the buttocks, or around the neck, ears or eyes?'; and atopy by an affirmative response to at least one of the three questions 'Has your child ever had eczema?' or 'Has your child ever had hay fever?' or 'Has your child ever had another form of allergic rhinitis other than hay fever, i.e. caused by allergens other than pollens?'. In this study, we chose to analyze lifetime asthma in order to have a more specific diagnosis of asthma for children. The question on wheezing during the 12 past months is more sensitive than specific.

Assessment of exposure to air pollution

Data regarding medium-term exposure to close-proximity and background air pollution were provided by the Air Quality Monitoring Networks of Guadeloupe (GWAD'AIR Association) for a period of two consecutive weeks before the day the children received their clinical examination.

Medium-term exposure to close-proximity pollution by O_3 and NO_2 was measured in 27 participating schools. Levels of each pollutant were assessed using passive diffusion samplers in representative points of each classroom for indoor air pollution and near the playground for outdoor air pollution. The method has already been described elsewhere [26].

With regard to indoor pollution, the data showed there was little variation between classrooms in the same school. For this reason, the concentration average per school was used (i.e. 27 data points per pollutant). Background air pollution was measured using fixed stations located in three cities (Pointe-à-Pitre, Abymes and Baie-Mahault). According to the GWAD'AIR Association, background air pollution can be considered as uniform in all schools located in the agglomeration of Pointe-à-Pitre i.e. 7 schools recording pollution data (Figure 1).

The close-proximity exposure in the 27 schools reflected both spatial and temporal variations since schools were sampled at different points in time and space. For background pollution, the pollutant concentration was considered as the same in the area where the 7 schools were located. For this reason, the background exposure in the 7 schools represented only temporal variations.

The daily concentration of each air pollutant was calculated from the maximum daily 1-hour average for O_3 ,

NO_2 and SO_2 and the daily average for PM_{10} according to European guidelines (<http://www.airqualitynow.eu>). Medium-term exposure to background air pollution was defined by the average daily concentration for the two weeks before the clinical examination. Short-term background air pollution was defined by the daily average concentration of each pollutant on the clinical examination day and up to the five subsequent days corresponding to five different lags (D0, D1, D2, D3, D4, D5). These lags were classified into two categories: short delay exposure and cumulative short delay exposure, which indicates the mean exposure to pollutants in the five days before.

In summary, two analyses were performed: a first analysis regarding close-proximity pollution in 27 schools and a second analysis regarding background pollution in a restricted sample of 7 schools (Pointe-à-Pitre agglomeration). The variation in estimated pollution data could be both spatial and temporal in the first analysis whereas in the second analysis, it could only be temporal.

Particulate matter with a diameter inferior to $2.5 \mu m$ ($PM_{2.5}$) was not considered in this study because it was not monitored by the GWAD'AIR Association when the study started.

Statistical analysis

Data were analyzed using the statistical package SAS (Version 9.1; SAS Institute Inc., Cary, NC, USA). We built several linear mixed regressions to measure the associations between lung function (PEF and Δ PEF) and medium-term exposure to i) close-proximity pollution (indoor and outdoor) by O_3 and NO_2 and ii) background pollution by O_3 , NO_2 , SO_2 and PM_{10} . With respect to short-term exposure to background air pollution, we tested the associations between the lagged pollutant levels, defined previously, and lung function using a distributed-lag model. This approach reduces collinearity, decreases the number of parameters to be estimated via the polynomial function and allows estimation of specific lagged effects [27].

We fitted separate models for each pollutant. Variables such as sex, age, BMI, full-term birth, rainy season, temperature and relative humidity were considered as confounding factors and were forced in all models. The other explanatory variables were retained in multivariable models if the associations with the outcomes were statically significant or if they were a confounding factor. The interactions with "asthma" or "drugs to treat asthma before clinical examination" variables were tested in each model in order to check for a potential modification effect with the pollutants. Children were not independent observational units because they were nested within schools. For this reason, a random intercept was introduced in each model in order to take into account the

dependence in the outcome for children in the same school.

To summarize, the linear mixed model is defined as follows:

$$Y_{ci} = \beta_0 + \gamma_{0c} + \beta_1 \times \text{Pollutant}_{ic} + \beta \times X_{ic} + \varepsilon_{ic}$$

where Y_{ci} is the dependent variable (PEF or Δ PEF) with c which is the elementary school ($c = 1, \dots, k$) and i a child in an elementary school. β_0 is the fixed intercept and γ_{0c} is the random intercept at school level. β_1 is the fixed pollutant effect and Pollutant_{ic} is the average medium-term pollutant concentration for child i in school c . β is the coefficient vector of fixed effects for adjustment factors and X_{ic} is the variable matrix for the adjustment factors. The quantities ε_{ic} are random variables representing errors in the relationship.

The distributed-lag model to study short-term exposure to pollution is as follows:

$$Y_{cit} = \beta_0 + \gamma_{0c} + \zeta^T \times X_{ct} + \xi^T \times X_{ic} + \sum_{l=0}^5 \delta_l \times (Z_{ic,t-l}) + \varepsilon_{ic}$$

where t is the clinical examination day, ζ^T is the transpose vector of fixed effects for time-dependent variables (e.g. rainy season) with X_{ct} the matrix of these variables, and ξ^T is the transpose vector of fixed effects for individual variables with X_{ic} the matrix of these variables. Pollutant

effects were taken into account using $\sum_{l=0}^5 \delta_l \times (Z_{ic,t-l})$,

where pollutant effects that were delayed by up to five days and δ_l was constrained to follow a polynomial function of degree 2. $Z_{ic,t-l}$ is the pollutant concentration for day t at lag l for a child (i) in an elementary school (c).

Results

Children's characteristics

Children characteristics are presented in Table 1. Among the 1,436 children, the average age was 10.3 years (SD \pm 0.8; range: 8.2-12.9), 46.5% were males and 21.3% had a high BMI. Regarding their family situation, 52.4% of children lived with both parents. The majority of parents had a primary or secondary educational level (63.5%). With respect to environmental factors, the relative humidity and temperature were 68.6% (SD \pm 15.3; range: 25-95) and 27.8°C (SD \pm 1.9; range: 22-33) respectively. Data were collected during the rainy season in 44.5% of children. Exposure to tobacco smoke concerned 14.3% of children. The prevalence rates of asthma, atopy, atopic dermatitis, drugs used to treat asthma before clinical examination, family history of allergy and premature birth were 15.5%, 24.5%, 2.4%, 5.2%, 24.3% and

27.2% respectively. The values of PEF and Δ PEF were on average 272 L/min (SD \pm 47.8; range: 130-460) and minus 1% (SD \pm 9.9; range: -56%-97%) respectively. Mean height in the sample of children was 146.0 cm (SD \pm 9.4).

Educational level of parents and exposure to tobacco smoke had a high number of missing data (23.1% and 42.1%, respectively). For this reason, the latter were not taken into account in the models.

Exposure to air pollutants

Table 2 summarizes the distribution of air pollutant concentrations estimated for medium-term exposure to close-proximity air pollution (indoor and outdoor) and background air pollution.

The mean medium-term exposure to indoor close-proximity air pollution was 49.5 $\mu\text{g}/\text{m}^3$ (SD \pm 14.8 $\mu\text{g}/\text{m}^3$) for O_3 and 5.3 $\mu\text{g}/\text{m}^3$ (SD \pm 4.3 $\mu\text{g}/\text{m}^3$) for NO_2 , while the mean medium-term exposure to outdoor close-proximity air pollution was 55.3 $\mu\text{g}/\text{m}^3$ (SD \pm 16.4 $\mu\text{g}/\text{m}^3$) for O_3 and 5.3 $\mu\text{g}/\text{m}^3$ (SD \pm 3.5 $\mu\text{g}/\text{m}^3$) for NO_2 . Wide minimum-maximum intervals were found between the 27 participating schools (e.g. for indoor, O_3 range: 23.6-80.1 $\mu\text{g}/\text{m}^3$ and NO_2 range: 0.9-21.5 $\mu\text{g}/\text{m}^3$). A high correlation between indoor and outdoor close-proximity air pollution was identified with a Spearman's correlation coefficient of 0.68 for O_3 and 0.88 for NO_2 .

Data regarding medium-term background air pollution restricted to schools in the Pointe-à-Pitre agglomeration were 54.1 $\mu\text{g}/\text{m}^3$ (SD \pm 6.8 $\mu\text{g}/\text{m}^3$) for O_3 , 14.8 $\mu\text{g}/\text{m}^3$ (SD \pm 3.8 $\mu\text{g}/\text{m}^3$) for NO_2 , 4.7 $\mu\text{g}/\text{m}^3$ (SD \pm 3.7 $\mu\text{g}/\text{m}^3$) for SO_2 and 23.9 $\mu\text{g}/\text{m}^3$ (SD \pm 6.8 $\mu\text{g}/\text{m}^3$) for PM_{10} . There was no meaningful correlation between close-proximity and medium-term background air pollution.

Pollutant concentrations were lower than WHO guidelines for O_3 , NO_2 and SO_2 . However, five of the seven schools (i.e. 352 children) were exposed to more than 20 $\mu\text{g}/\text{m}^3$ of PM_{10} i.e. above WHO guidelines.

Associations between medium-term exposure to close-proximity pollution and health outcomes

Tables 3 and 4 show coefficients and 95% CI of PEF and Δ PEF outcomes associated with the 1-unit change of mean of each pollutant.

With respect to PEF health outcome, there was a statically significant association between the average concentration of outdoor close-proximity pollution by O_3 and PEF decrease, i.e. a 1- $\mu\text{g}/\text{m}^3$ increase in O_3 was associated with a reduction in PEF of 0.32 L/min ($\hat{\beta} = -0.32$; 95% CI: -0.61;-0.03). Conversely, no reduction in PEF was identified with indoor close-proximity pollution by O_3 . NO_2 was not significantly associated with PEF.

There was no association between Δ PEF and pollutants.

Table 1 Children from the 27 participating schools and the 7 schools in the Pointe-à-Pitre agglomeration

Variables	Participating schools including 1,436 children		Participating schools in Pointe-à-Pitre agglomeration including 506 children	
	% data provided	Mean ± SD or Frequency	% data provided	Mean ± SD or Frequency
Socio-demographic factors				
Age	100.0	10.3 ± 0.8	100.0	10.5 ± 0.7
Sex (male)	100.0	46.5	100.0	47.4
BMI	99.2		98.6	
<i>Low</i>		6.7		7.4
<i>Normal</i>		72.0		70.8
<i>High</i>		21.3		21.8
Family situation of children				
<i>with both parents</i>	88.6		88.3	
<i>with mother</i>		52.4		48.5
<i>with father</i>		43.4		47.9
<i>Other</i>		2.0		2.7
<i>Other</i>		2.2		0.9
Educational level of parents				
<i>Primary or secondary</i>	76.9		76.1	
<i>High school and university</i>		63.5		64.2
<i>High school and university</i>		36.5		35.8
Environnemental factors				
Relative humidity (unit: %)	100.0	68.6 ± 15.3	100.0	59.6 ± 15.2
Outside temperature (unit: °C)	100.0	27.8 ± 1.9	100.0	28.1 ± 1.4
Rainy saison (%)	100.0	44.5	100.0	35.4
Exposure to tobacco smoke (%)	57.9	14.3	61.7	12.2
Medical history				
Asthma (%)	100.0	15.5	100.0	16.6
Atopy (%)	100.0	24.5	100.0	23.7
Atopic dermatitis (%)	100.0	2.4	100.0	3.6
Drugs against asthma before clinical examination (%)	100.0	5.2	100.0	7.1
Family history of allergy (%)	100.0	24.3	100.0	25.5
Premature birth (%)	81.8	27.2	91.3	31.8
Lung function				
Peak expiratory flow before run (unit: L/min)	97.1	271.5 ± 47.8	96.8	274.2 ± 49.9
Variation of peak expiratory flow after run (unit: %)	80.6	-1.1 ± 9.9	78.7	-0.5 ± 12.0

Abbreviations: SD: standard deviation; BMI: Body Mass Index.

Association between short- and medium-term exposure to background pollution and health outcomes

Tables 5 and 6 present three models measuring medium-term (model 1 and 2) and short-term (model 3) effects of background pollution on PEF and Δ PEF outcomes in 7 schools.

In model 1, no significant association was found with O_3 , SO_2 and NO_2 . However, an interaction within the limits of significance was found between O_3 and asthma (model 2). This interaction meant that the effect of medium-term exposure to background pollution by O_3 and NO_2 on a PEF reduction was stronger in asthmatic children than in non-asthmatic ones. With regard to

PM_{10} , a significant association with PEF increase and Δ PEF increase (negative variation) was found (Tables 5 and 6).

In model 3, short delay (D0, D1, D2, D3, D4, D5) exposure and cumulative short delay exposure were not associated with PEF and Δ PEF.

Discussion

The asthma prevalence identified in this study was slightly higher than that reported in the few studies conducted in the Caribbean using comparable standardized ISAAC methodology [18,20]. The previous Guadeloupian study reported a prevalence of 14.1% among 5,094

Table 2 Concentration of medium-term air pollutants in schools participating in the study

Pollutants	WHO air quality guidelines 2005	Type of pollution	Number of schools	in $\mu\text{g}/\text{m}^3$			
				Mean	Standard deviation	Minimum	Maximum
O_3	100 $\mu\text{g}/\text{m}^3$ 8-hour mean	Indoor close-proximity	27	49.5	14.8	23.6	80.1
		Outdoor close-proximity	27	55.3	16.4	21.1	90.9
		Background	7	54.1	6.8	40.5	59.2
NO_2	40 $\mu\text{g}/\text{m}^3$ annual mean	Indoor close-proximity	27	5.3	4.3	0.9	21.5
		Outdoor close-proximity	27	5.3	3.5	0.9	15.3
		Background	7	14.8	3.8	11.5	22.1
SO_2	20 $\mu\text{g}/\text{m}^3$ 24-hour mean	Background	7	4.7	3.7	1.8	12.7
PM_{10}	20 $\mu\text{g}/\text{m}^3$ annual mean	Background	7	23.9	6.8	16.5	33.4

Abbreviations: O_3 , ozone; NO_2 , nitrogen dioxide; SO_2 , sulfur dioxide; PM_{10} , particles with aerodynamic diameter lower than 10 μm .

adolescents with a mean age of 12.9 years and another study in the Caribbean islands of Trinidad and Tobago reported 12.8% and 13.5% respectively, among 4,988 adolescents. Although the present study concerned younger children (a mean age of 10.3 among 1,436 children), our findings confirm the high asthma prevalence in the Caribbean compared with the prevalence worldwide [28]. Furthermore, the studies conducted in the Caribbean suggest small variations in asthma prevalence between the Caribbean islands and contrast with a recent study conducted in the Pacific where considerable variations were

found, the rates ranging from 5.8% in Samoa to 19.7% in the Tokelau Islands [29].

Many factors influence asthma such as genetic susceptibility and cultural differences [30]. A previous ISAAC phase two study conducted in Hong Kong and two cities in mainland China showed that the higher prevalence of asthma symptoms in Hong Kong could be explained by differences in environmental factors and diet [31]. The impact of environmental factors on respiratory diseases, particularly air pollution, seems to vary in different parts of the globe, probably owing to individual susceptibility.

Table 3 Results of linear mixed models to investigate associations between medium-term exposure to indoor and outdoor close-proximity air pollution and peak expiratory flow (n = 1133)

Pollutant	Indoor				Outdoor			
	O_3 (in $\mu\text{g}/\text{m}^3$)		NO_2 (in $\mu\text{g}/\text{m}^3$)		O_3 (in $\mu\text{g}/\text{m}^3$)		NO_2 (in $\mu\text{g}/\text{m}^3$)	
	β	(95% CI)	β	(95% CI)	β	(95% CI)	β	(95% CI)
Pollutant	-0.21	(-0.55; 0.13)	0.15	(-0.85; 1.16)	-0.32	(-0.61; -0.03)	-0.04	(-1.25; 1.17)
Sex (male vs girl)	5.73	(0.50; 10.96)	5.59	(0.36; 10.81)	5.75	(0.53; 10.97)	5.57	(0.35; 10.80)
Age	22.17	(18.58; 25.76)	22.03	(18.44; 25.61)	22.01	(18.43; 25.58)	22.04	(18.45; 25.62)
BMI								
High vs Normal	19.13	(12.74; 25.52)	19.01	(12.62; 25.41)	19.03	(12.66; 25.4)	18.97	(12.58; 25.36)
Low vs Normal	-18.60	(-28.58; -8.63)	-18.33	(-28.30; -8.36)	-18.78	(-28.74; -8.82)	-18.29	(-28.26; -8.32)
Premature birth (Yes vs No)	-5.32	(-11.19; 0.55)	-5.22	(-11.09; 0.65)	-5.35	(-11.21; 0.52)	-5.23	(-11.10; 0.64)
Rainy season (Yes vs No)	0.14	(-8.41; 8.70)	-0.85	(-10.23; 8.53)	1.64	(-6.24; 9.53)	-0.25	(-9.42; 8.92)
Outside temperature (unit: $^{\circ}\text{C}$)	1.82	(-0.05; 3.68)	2.10	(0.28; 3.92)	1.85	(0.09; 3.61)	2.10	(0.29; 3.92)
Relative humidity (unit: %)	0.27	(-0.08; 0.61)	0.26	(-0.12; 0.65)	0.19	(-0.15; 0.54)	0.29	(-0.08; 0.67)
Atopy (Yes vs No)	7.53	(1.44; 13.62)	7.50	(1.41; 13.60)	7.42	(1.33; 13.50)	7.52	(1.42; 13.61)
Asthma (Yes vs No)	-9.62	(-16.85; -2.39)	-9.60	(-16.83; -2.37)	-9.55	(-16.77; -2.33)	-9.63	(-16.86; -2.40)
Pointoise agglomeration (Yes vs No)	7.82	(-3.00; 18.64)	4.89	(-7.99; 17.76)	7.59	(-1.87; 17.06)	6.17	(-6.29; 18.62)
Variability of school random intercept: $\hat{\sigma}_{\text{VO}}^2$ (P value)	43.39 (0.02)		48.33 (0.01)		28.43 (0.09)		47.91 (0.01)	

Note. The peak expiratory flow (PEF) corresponds to baseline peak expiratory flow before running in children. The coefficients (β) and their 95% confidence interval are obtained using linear mixed models. Variables such as sex, age, body mass index, premature birth, rainy season, temperature and relative humidity are forced in all models. Other confounding factors are identified and included: asthma, atopy and Pointoise. The asthma pollution interaction was tested in each model but was not statistically significant.

Abbreviations: PEF, peak expiratory flow; O_3 , ozone; NO_2 , nitrogen dioxide; CI, confidence interval.

Table 4 Results of linear mixed models to investigate associations between medium-term exposure to indoor and outdoor close-proximity air pollution and peak expiratory flow variation (n = 938)

	Indoor				Outdoor			
	O ₃ (in µg/m ³)		NO ₂ (in µg/m ³)		O ₃ (in µg/m ³)		NO ₂ (in µg/m ³)	
	β	(95% CI)	β	(95% CI)	β	(95% CI)	β	(95% CI)
Pollutant	0.02	(-0.05; 0.09)	0.00	(-0.16; 0.17)	0.04	(-0.02; 0.10)	0.03	(-0.18; 0.24)
Sex (male vs girl)	-0.06	(-1.35; 1.24)	-0.04	(-1.33; 1.26)	-0.09	(-1.38; 1.21)	-0.03	(-1.33; 1.26)
Age	-0.21	(-1.11; 0.69)	-0.19	(-1.08; 0.71)	-0.22	(-1.11; 0.67)	-0.19	(-1.08; 0.71)
BMI								
<i>High vs Normal</i>	0.11	(-1.46; 1.68)	0.14	(-1.43; 1.70)	0.11	(-1.45; 1.67)	0.14	(-1.43; 1.70)
<i>Low vs Normal</i>	0.43	(-2.20; 3.07)	0.40	(-2.23; 3.04)	0.48	(-2.15; 3.11)	0.39	(-2.24; 3.02)
Premature birth (Yes vs No)	0.49	(-0.99; 1.97)	0.49	(-0.99; 1.97)	0.48	(-1.00; 1.96)	0.49	(-0.99; 1.97)
Rainy season (Yes vs No)	-0.24	(-1.99; 1.51)	-0.23	(-2.09; 1.62)	-0.42	(-2.08; 1.24)	-0.29	(-2.12; 1.53)
Outside temperature (unit: °C)	0.19	(-0.23; 0.62)	0.16	(-0.25; 0.58)	0.21	(-0.20; 0.61)	0.16	(-0.25; 0.58)
Relative humidity (unit: %)	-0.06	(-0.12; 0.01)	-0.06	(-0.12; 0.00)	-0.04	(-0.11; 0.02)	-0.06	(-0.12; 0.00)
Atopy (Yes vs No)	1.09	(-0.38; 2.57)	1.09	(-0.38; 2.57)	1.09	(-0.38; 2.57)	7.52	(-0.39; 2.56)
Variability of school random intercept: $\hat{\sigma}_{\gamma_0}^2$ (P value)	0.99 (0.10)		1.16 (0.07)		0.6 (0.22)		1.13 (0.08)	

Note. The peak expiratory flow (PEF) corresponds to baseline peak expiratory flow before running in children. The peak expiratory flow variation (Δ PEF) represents the percentage decrease in PEF after running compared to the retained baseline PEF. The coefficients (β) and their 95% confidence interval are obtained using linear mixed models. The variables such as sex, age, body mass index, premature birth, rainy season, temperature and relative humidity are forced in all models. Atopy are identified as confounding factor and included. The asthma pollution interaction was tested in each model but was not statistically significant. Abbreviations: PEF, peak expiratory flow; O₃, ozone; NO₂, nitrogen dioxide; CI, confidence interval.

To our knowledge, this study is the first to provide close-proximity and background pollution data in the Caribbean and to measure its impact on the respiratory lung function of elementary schoolchildren. The observed mean concentrations of PM₁₀ in 5 out of the 7 participating schools exceed the known threshold limit of 20 µg/m³ proposed by WHO air quality guidelines [32]. A recent French ministry report explained that the increase in PM₁₀ concentration was partly due to Saharan-Sahel dust carried by atmospheric circulation over the Caribbean area for five to seven months a year [33]. In contrast with PM₁₀, the observed mean concentrations of O₃, NO₂ and SO₂ were lower than the WHO known threshold limit values of 100 µg/m³ for O₃, 40 µg/m³ for NO₂, and 20 µg/m³ for SO₂. This low air pollution level is not striking with respect to the regional emission density of SO₂ and NO₂ in the Caribbean, which is still far less than that of many temperate-zone industrial countries. However for O₃, which is an atmospheric pollutant not directly emitted by car engines or by industrial operations but formed by the reaction of sunlight on air containing hydrocarbons and nitrogen oxides that react to form O₃, the relatively low concentration was not expected in Guadeloupe where temperature and sunlight intensity are high throughout the year. The impact of wind could explain these variations. In addition, another source of NO₂ and SO₂ emissions could be wood-burning in rural areas and fires in sugar cane plantations before harvesting begins.

The main strength of this study is that it assessed medium-term exposure to both indoor and outdoor close-proximity air pollution. Indeed, elementary schoolchildren spend a lot of time at school in many countries and almost 8 hours per day in an academic year in France. For this reason, this study provides an interesting insight into the effect of air pollution in schools. Medium-term exposure to both indoor and outdoor close-proximity air pollution was assessed by measuring the two-week cumulative effect of two major pollutants such as O₃ and NO₂ in a large random sample of Guadeloupean classrooms. Even with levels lower than WHO guidelines, the linear mixed model analyses showed significantly reduced lung function before exercise with outdoor exposure to O₃ after controlling for personal attributes and meteorological factors.

Most studies in schoolchildren have already provided evidence that exposure to O₃ reduces lung function after controlling for confounders but not at such a low level [34]. To our knowledge, only a recent study suggested that exposure to similar low levels of O₃ could have an impact on asthma. The authors reported an association between emergency pediatric department visits for asthma and relatively low O₃ concentrations [35]. Moreover, we showed a stronger association of medium-term exposure to background pollution by O₃ with PEF reduction in asthmatic children than in non-asthmatic children. This seems plausible because air pollution may act not only as a short-term trigger but also as a priming

Table 5 Results of linear mixed models with random intercept on schools to investigate associations between short and medium term background air pollution and peak expiratory flow (n = 425)

	O ₃ (in µg/m ³)		NO ₂ (in µg/m ³)		SO ₂ (in µg/m ³)		PM ₁₀ (in µg/m ³)	
	β	(95% CI)	β	(95% CI)	β	(95% CI)	β	(95% CI)
Medium-term exposure								
Model 1								
Pollutant	-0.44	(-2.61; 1.74)	-0.215	(-3.89; 3.46)	1.18	(-2.65; 5.00)	3.05	(0.04; 6.05)
Variability of school random intercept: $\hat{\sigma}_{\nu_0}^2$ (P value)	90.20	(0.05)	105.34	(0.11)	86.80	(0.04)	11.05	(0.50)
Model 2 (Interaction of pollutant with asthma)								
Non-asthmatic children	-0.13	(-2.32; 2.07)	-0.56	(-4.27; 3.15)	0.67	(-3.20; 4.54)	2.95	(0.12; 5.78)
Asthmatic children	-1.88	(-4.50; 0.74)	1.40	(-2.97; 5.78)	3.54	(-1.06; 8.14)	3.48	(0.31; 6.64)
Variability of school random intercept: $\hat{\sigma}_{\nu_0}^2$ (P value)	89.30	(0.04)	107.99	(0.19)	91.89	(0.04)	19.82	(0.50)
Short-term exposure								
Model 3								
D0	-0.24	(-5.09; 4.61)	0.54	(-2.44; 3.51)	0.41	(-35.98; 36.80)	0.14	(-1.82; 2.09)
D1	-0.52	(-1.62; 0.58)	0.11	(-2.19; 2.42)	-0.53	(-14.32; 13.27)	0.55	(-1.94; 3.04)
D2	-0.56	(-4.06; 2.94)	-0.15	(-2.54; 2.24)	-1.03	(-4.21; 2.14)	0.79	(-2.17; 3.75)
D3	-0.37	(-3.68; 2.94)	-0.27	(-2.68; 2.14)	-1.11	(-14.25; 12.04)	0.87	(-2.19; 3.94)
D4	0.05	(-0.63; 0.73)	-0.23	(-2.60; 2.15)	-0.74	(-18.06; 16.56)	0.79	(-2.02; 3.59)
D5	0.70	(-4.99; 6.39)	-0.03	(-3.06; 2.99)	0.04	(-15.52; 15.60)	0.53	(-1.86; 2.92)
Cumulative effect	-0.94	(-4.82; 2.95)	-0.03	(-14.01; 13.94)	-2.96	(-12.69; 6.77)	3.67	(-11.47; 18.81)
Variability of school random intercept: $\hat{\sigma}_{\nu_0}^2$ (P value)	0.00	(0.50)	0.00	(0.50)	0.00	(0.50)	0.00	(0.50)

Note. The peak expiratory flow (PEF) corresponds to baseline peak expiratory flow before running in children. The coefficients (β) and their 95% confidence interval were obtained using linear mixed models regarding models 1–2 and a distributed-lag model regarding model 3. Variables such as sex, age, body mass index, full-term birth, rainy season, temperature and relative humidity are forced in each model. Other confounding factors are included: atopy for model 1–2 and atopy, asthma and day of week for model 3.

Short-term exposure was defined by the average of each pollutant concentration on the current day and up to five days corresponding to five different lags (D0, D1, D2, D3, D4, D5). These lags were classified into two categories: short-delay exposure and cumulative short delay exposure, which indicates the mean exposure to pollutants on the five preceding days.

Abbreviations: PEF, peak expiratory flow; O₃, ozone; NO₂, nitrogen dioxide; SO₂, sulphur dioxide; PM10, small particulate matter; CI, confidence interval.

event inducing mechanisms of enhanced airway inflammation, which in turn result in subsequent bronchial hyperactivity [36].

Although our results concerning medium-term exposure to indoor pollutants suggest no meaningful association with a PEF, several studies have identified associations between air pollution in classrooms and the respiratory health of schoolchildren [37,38]. For instance, a recent French study assessing several pollutants (PM_{2.5}, NO₂ and three aldehydes) showed that poor air quality in classrooms was related to an increased prevalence of clinical manifestations of asthma and rhinitis [37].

The relatively low concentrations of SO₂ and NO₂ compared to those of quality guidelines could explain why no association was identified with a decrease in lung function. Indeed, we observed a factor close to 2 between air quality guidelines and the concentration mean of O₃ and a factor close to 3 and 4 with NO₂ and SO₂ respectively. With respect to PM₁₀, we found significant associations with lung function that increased for PEF and ΔPEF. This result is really puzzling because it is difficult to imagine that exposure to PM₁₀ has a

protective effect. A possible explanation is that we performed a total of 24 analyses and observed significant associations with outcomes just by chance. However, our results on O₃ point to a significant association not due to chance since O₃ associations were found with decreasing PEF both for medium-term exposure to outdoor close-proximity pollution and for background pollution (interaction with asthma). In addition, the use of passive diffusion samplers to measure pollutant concentrations in the schools is more reliable than the method using fixed stations, which underlines the pertinence of our findings about O₃ regarding medium-term exposure to outdoor close-proximity pollution.

Our study has some limitations. First, lung function was measured with a peak flow meter instead of spirometry, which provides more lung function parameters. However, peak flow meters are inexpensive, portable, easy to use and do not need to be plugged in. Moreover, PEF is still used in practice for asthma monitoring according to the guidelines in the Global Initiative for Asthma report [39] in epidemiological studies with trained technicians [40] and in clinical trials [41].

Table 6 Results of linear mixed models with random intercept on schools to investigate associations between short- and medium-term background air pollution and peak expiratory flow variation (n = 355)

	O ₃ (in µg/m ³)		NO ₂ (in µg/m ³)		SO ₂ (in µg/m ³)		PM ₁₀ (in µg/m ³)	
	β	(95% CI)	β	(95% CI)	β	(95% CI)	β	(95% CI)
Medium-term exposure								
Model 1								
Pollutant	-0.12	(-0.73; 0.49)	0.67	(-0.01; 1.35)	0.14	(-0.94; 1.22)	-1.16	(-1.95; -0.37)
Variability of school random intercept: $\hat{\sigma}_{\gamma_0}^2$ (P value)	6.42	(0.04)	0.93	(0.38)	6.69 (0.04)		0.00	(NS)
Model 2 (Interaction of pollutant with asthma)								
Non-asthmatic children	-0.14	(-0.76; -0.48)	0.66	(-1.04; 2.37)	0.17	(-0.91; 1.25)	-1.13	(0.16; -2.42)
Asthmatic children	0.00	(-0.74; 0.74)	0.65	(-0.34; 1.64)	-0.06	(-1.37; 1.25)	-1.28	(-0.36; -2.20)
Variability of school random intercept: $\hat{\sigma}_{\gamma_0}^2$ (P value)	6.34 (0.05)		0.97	(0.50)	6.63 (0.04)		0.00	(0.50)
Short-term exposure								
Model 3								
D0	-0.24	(-5.09; 4.61)	0.54	(-2.44; 3.51)	0.41	(-35.98; 36.80)	0.14	(-1.82; 2.09)
D1	-0.52	(-1.62; 0.58)	0.11	(-2.19; 2.42)	-0.53	(-14.32; 13.27)	0.55	(-1.94; 3.04)
D2	-0.56	(-4.06; 2.94)	-0.15	(-2.54; 2.24)	-1.03	(-4.21; 2.14)	0.79	(-2.17; 3.75)
D3	-0.37	(-3.68; 2.94)	-0.27	(-2.68; 2.14)	-1.11	(-14.25; 12.04)	0.87	(-2.19; 3.94)
D4	0.05	(-0.63; 0.73)	-0.23	(-2.60; 2.15)	-0.74	(-18.06; 16.56)	0.79	(-2.02; 3.59)
D5	0.70	(-4.99; 6.39)	-0.03	(-3.06; 2.99)	0.04	(-15.52; 15.60)	0.53	(-1.86; 2.92)
Cumulative effect	-0.94	(-4.82; 2.95)	-0.03	(-14.01; 13.94)	-2.96	(-12.69; 6.77)	3.67	(-11.47; 18.81)
Variability of school random intercept: $\hat{\sigma}_{\gamma_0}^2$ (P value)	0.00	(NS)	0.00	(NS)	0.00	(NS)	0.00	(NS)

Note. The peak expiratory flow (PEF) corresponds to baseline peak expiratory flow before running in children. The peak expiratory flow variation (Δ PEF) represents the percentage decrease in PEF after running compared to the baseline PEF. The coefficients (β) and their 95% confidence interval were obtained using linear mixed models regarding models 1–2 and a distributed-lag model regarding model 3. Variables such as sex, age, body mass index, full-term birth, rainy season, temperature and relative humidity are forced in each model. Other confounding factors are included: atopy for model 1–2 and atopy, asthma and day of week for model 3.

Short-term exposure was defined by the average of each pollutant concentration on the current day and up to five days before corresponding to five different lags (D0, D1, D2, D3, D4, D5). These lags were classified into two categories: short-delay exposure and cumulative short-delay exposure, which indicates the mean exposure to pollutants on the five preceding days.

Abbreviations: PEF, peak expiratory flow; O₃, ozone; NO₂, nitrogen dioxide; SO₂, sulphur dioxide; PM₁₀, small particulate matter; CI, confidence interval.

A second limitation is that the analyses of short- and medium-term exposure to background pollution were based on a small sample of 7 schools that did not reflect spatial variations. In addition, the lack of power could lead to a wide confidence interval and no significant association in the analyses. However, we were unable to collect data in the other participating schools because the fixed stations were located only in the agglomeration of Pointe-à-Pitre. With respect to close-proximity air pollution, a large sample size was used as recommended in the ISAAC 2 protocol, so robust analyses were possible.

The confounding role of medications for asthma may be considered as another potential limitation. Medication use in children may influence lung function and mask the effects of pollutants. For instance, asthmatic children taking a corticoid treatment on the days preceding the run test will be less susceptible to air pollutants than other children. In addition, the school environment may not have been the only factor responsible for decreasing lung function because some children may have been exposed to air pollution at home.

As in most studies of this type, unmeasured confounding is a real concern. Socio-economic factors may correlate with spatial variation in air pollution. Indeed, lower socio-economic groups are likely more exposed and more susceptible to air pollution. For this reason, they might have an effect on the parameters we measured. An analysis including socio-economic factors could highlight air pollution effects in lower socio-economic groups which could not be detected in this study.

A final limitation was the lack of PM_{2.5} measurements, as they were not being monitored by the GWA-D'AIR Association when the study started.

Conclusions

This observational exploratory study provided the opportunity to measure the impact of exposure to low concentration air pollution on lung function among a population of children with a high prevalence of asthma. It bridges the knowledge gap regarding pollution in the Caribbean and demonstrates a consistent association between medium-term exposure to low O₃ concentrations

under the WHO threshold levels and reduced lung function in children. However, additional research in the Caribbean is needed to confirm our results and to study the impact of other pollutants. For instance, our study was limited to measuring two indoor pollutants owing to logistical difficulties and economic constraints. In addition, further research is needed to measure the toxicological and biological effects of particles originating from Saharan-Sahel dust, as suggested in another study [42]. In the future, new statistical approaches combining pollutant mixtures will be useful to better understand the role of air pollutant concentrations on the development of asthma and allergies in the Caribbean [43].

Abbreviations

BMI: Body mass index; CI: Confidence interval; NO₂: Nitrogen dioxide; O₃: Ozone; PEF: Peak expiratory flow; PM₁₀: Particles with aerodynamic diameter lower than 10 µm; SD: Standard deviation; SO₂: Sulfur dioxide.

Competing interests

The authors declare they have no competing interests. The study was supported by the "Karu Asthme association".

Authors' contributions

BA wrote the paper; ChR designed and supervised the study and critically reviewed the manuscript. CeR performed data analysis and interpretation and helped to draft the manuscript. VR coordinated data analysis and critically reviewed the manuscript. MM, LC, XB, EC, JG, MG, GM and DL participated in the study coordination and critically reviewed the manuscript. All authors read and approved the final manuscript.

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References

- Anandan C, Nurmatov U, van Schayck OC, Sheikh A. Is the prevalence of asthma declining? Systematic review of epidemiological studies. *Allergy*. 2010;65:152–67.
- Renz H, Conrad M, Brand S, Teich R, Garn H, Pfeifferle PI. Allergic diseases, gene-environment interactions. *Allergy*. 2011;66 Suppl 95:10–2.
- Gauderman WJ, Avol E, Lumann F, Kuenzli N, Gilliland F, Peters J, et al. Childhood asthma and exposure to traffic and nitrogen dioxide. *Epidemiology*. 2005;16:737–43.
- Jerrett M, Shankardass K, Berhane K, Gauderman WJ, Kunzli N, Avol E, et al. Traffic-related air pollution and asthma onset in children: a prospective cohort study with individual exposure measurement. *Environ Health Perspect*. 2008;116:1433–8.
- Garcia-Marcos L, Guillen JJ, Dinwiddie R, Guillen A, Barbero P. The relative importance of socio-economic status, parental smoking and air pollution (SO₂) on asthma symptoms, spirometry and bronchodilator response in 11-year-old children. *Pediatr Allergy Immunol Off Publ Eur Soc Pediatr Allergy Immunol*. 1999;10:96–100.
- Pikhart H, Bobak M, Gorynski P, Wojtyniak B, Danova J, Celko MA, et al. Outdoor sulphur dioxide and respiratory symptoms in Czech and polish school children: a small-area study (SAVIAH). *Small-area variation in air pollution and health*. *Int Arch Occup Environ Health*. 2001;74:574–8.
- Weinmayr G, Romeo E, De Sario M, Weiland SK, Forastiere F. Short-term effects of PM₁₀ and NO₂ on respiratory health among children with asthma or asthma-like symptoms: a systematic review and meta-analysis. *Environ Health Perspect*. 2010;118:449–57.
- Penard-Morand C, Raheerison C, Charpin D, Kopferschmitt C, Lavaud F, Caillaud D, et al. Long-term exposure to close-proximity air pollution and asthma and allergies in urban children. *Eur Respir J Off J Eur Soc Clin Respir Physiol*. 2010;36:33–40.
- McConnell R, Berhane K, Gilliland F, London SJ, Islam T, Gauderman WJ, et al. Asthma in exercising children exposed to ozone: a cohort study. *Lancet*. 2002;359:386–91.
- Penard-Morand C, Charpin D, Raheerison C, Kopferschmitt C, Caillaud D, Lavaud F, et al. Long-term exposure to background air pollution related to respiratory and allergic health in schoolchildren. *Clin Exp Allergy J Br Soc Allergy Clin Immunol*. 2005;35:1279–87.
- Li S, Williams G, Jalaludin B, Baker P. Panel studies of air pollution on children's lung function and respiratory symptoms: a literature review. *J Asthma Off J Assoc Care Asthma*. 2012;49:895–910.
- Grigg J. Air pollution and children's respiratory health - gaps in the global evidence. *Clin Exp Allergy J Br Soc Allergy Clin Immunol*. 2011;41:1072–5.
- Gauderman WJ, Avol E, Gilliland F, Vora H, Thomas D, Berhane K, et al. The effect of air pollution on lung development from 10 to 18 years of age. *N Engl J Med*. 2004;351:1057–67.
- Jacobson Lda SV, Hacon Sde S, de Castro HA, Ignotti E, Artaxo P, Saldiva PHN, et al. Acute effects of particulate matter and black carbon from seasonal fires on peak expiratory flow of schoolchildren in the Brazilian Amazon. *PLoS One*. 2014;9:e104177.
- Chen C-H, Chan C-C, Chen B-Y, Cheng T-J, Leon Guo Y. Effects of particulate air pollution and ozone on lung function in non-asthmatic children. *Environ Res*. 2014;137C:40–8.
- Kim SB, Temiyasathit C, Chen VC, Park SK, Sattler M, Russell AG. Characterization of spatially homogeneous regions based on temporal patterns of fine particulate matter in the continental United States. *J Air Waste Manag Assoc*. 2008;58:965–75.
- Weiland SK, Husing A, Strachan DP, Rzehak P, Pearce N. Climate and the prevalence of symptoms of asthma, allergic rhinitis, and atopic eczema in children. *Occup Environ Med*. 2004;61:609–15.
- Monteil MA, Joseph G, Changkit C, Wheeler G, Antoine RM. Comparison of prevalence and severity of asthma among adolescents in the Caribbean islands of Trinidad and Tobago: results of a nationwide cross-sectional survey. *BMC Public Health*. 2005;5:96.
- Mounouchy MA, Cordeau L, Raheerison C. [Prevalence of asthma and related symptoms among adolescents in Guadeloupe: phase I of the ISAAC survey 2003]. *Rev Mal Respir*. 2009;26:944–51.
- Kahwa EK, Younger NO, Wint YB, Waldron NK, Hewitt HH, Knight-Madden JM, et al. The Jamaica asthma and allergies national prevalence survey: rationale and methods. *BMC Med Res Methodol*. 2010;10:29.
- Furhrman C, Nicolau J, Rey G, Solet J, Quénel P, Jouglu E, et al. Asthme et BPCO : taux d'hospitalisation et de mortalité dans les départements d'outre-mer et en France métropolitaine, 2005–2007. *Bull Dépidémiologie Hebd*. 2011;13–14:168–72.
- Weiland SK, Bjorksten B, Brunekreef B, Cookson WO, von Mutius E, Strachan DP. Phase II of the International Study of Asthma and Allergies in Childhood (ISAAC II): rationale and methods. *Eur Respir J Off J Eur Soc Clin Respir Physiol*. 2004;24:406–12.
- Addo-Yobo EO, Woodcock A, Allotey A, Baffoe-Bonnie B, Strachan D, Custovic A. Exercise-induced bronchospasm and atopy in Ghana: two surveys ten years apart. *PLoS Med*. 2007;4:e70.
- AJRCCM. Standardization of Spirometry, 1994 update. American thoracic society. *Am J Respir Crit Care Med*. 1995;152:1107–36.
- Godfrey S, Kamburoff PL, Nairn JR. Spirometry, lung volumes and airway resistance in normal children aged 5 to 18 years. *Br J Dis Chest*. 1970;64:15–24.
- Annesi-Maesano I, Moreau D, Caillaud D, Lavaud F, Le Moulec Y, Taytard A, et al. Residential proximity fine particles related to allergic sensitisation and asthma in primary school children. *Respir Med*. 2007;101:1721–9.
- Rondeau V, Berhane K, Thomas DC. A three-level model for binary time-series data: the effects of air pollution on school absences in the Southern California children's health study. *Stat Med*. 2005;24:1103–15.

28. Lai CK, Beasley R, Crane J, Foliaki S, Shah J, Weiland S. Global variation in the prevalence and severity of asthma symptoms: phase three of the International Study of Asthma and Allergies in Childhood (ISAAC). *Thorax*. 2009;64:476–83.
29. Foliaki S, Annesi-Maesano I, Daniel R, Fakakovikaetau T, Magatongia M, Tuuau-Potoi N, et al. Prevalence of symptoms of childhood asthma, allergic rhinoconjunctivitis and eczema in the Pacific: the International Study of Asthma and Allergies in Childhood (ISAAC). *Allergy*. 2007;62:259–64.
30. Murphy KR, Berger WE. The variability of asthma. *Curr Med Res Opin*. 2005;21:1519–26.
31. Wong GW, Ko FW, Hui DS, Fok TF, Carr D, von Mutius E, et al. Factors associated with difference in prevalence of asthma in children from three cities in China: multicentre epidemiological survey. *BMJ*. 2004;329:486.
32. WHO: Air quality guidelines for particulate matter, ozone, nitrogen dioxide and sulfur dioxide - Global update 2005 - Summary of risk assessment. WHO World Health Organization; 2006. http://www.euro.who.int/__data/assets/pdf_file/0005/78638/E90038.pdf.
33. MEDDTL: Air pollution issues: Context in Guadeloupe and in metropolitan France. Ministère de l'Ecologie et du Développement Durable des Transports et du Logement. available at: (last accessed). 2011, Volume]. http://www.developpement-durable.gouv.fr/IMG/pdf/ECO_CLIMAT_GUADELOUPE_HD.pdf
34. Chang YK, Wu CC, Lee LT, Lin RS, Yu YH, Chen YC. The short-term effects of air pollution on adolescent lung function in Taiwan. *Chemosphere*. 2012;87:26–30.
35. Strickland MJ, Darrow LA, Klein M, Flanders WD, Sarnat JA, Waller LA, et al. Short-term associations between ambient air pollutants and pediatric asthma emergency department visits. *Am J Respir Crit Care Med*. 2010;182:307–16.
36. Mortimer KM, Neas LM, Dockery DW, Redline S, Tager IB. The effect of air pollution on inner-city children with asthma. *Eur Respir J Off J Eur Soc Clin Respir Physiol*. 2002;19:699–705.
37. Annesi-Maesano I, Hulin M, Lavaud F, Raheison C, Kopferschmitt C, de Blay F, et al. Poor air quality in classrooms related to asthma and rhinitis in primary schoolchildren of the French 6 cities study. *Thorax*. 2012;67:682–8.
38. Mi YH, Norback D, Tao J, Mi YL, Ferm M. Current asthma and respiratory symptoms among pupils in Shanghai, China: influence of building ventilation, nitrogen dioxide, ozone, and formaldehyde in classrooms. *Indoor Air*. 2006;16:454–64.
39. GINA. Global Strategy for Asthma Management and Prevention Update. 2011.
40. Jacobson Lda S, Hacon Sde S, Castro HA, Ignotti E, Artaxo P, Ponce de Leon AC. Association between fine particulate matter and the peak expiratory flow of schoolchildren in the Brazilian subequatorial Amazon: a panel study. *Environ Res*. 2012;117:27–35.
41. Chew KS, Kamarudin H, Hashim CW. A randomized open-label trial on the use of budesonide/formoterol (Symbicort(R)) as an alternative reliever medication for mild to moderate asthmatic attacks. *Int J Emerg Med*. 2012;5:16.
42. Mallone S, Stafoggia M, Faustini A, Gobbi GP, Marconi A, Forastiere F. Saharan dust and associations between particulate matter and daily mortality in Rome, Italy. *Environ Health Perspect*. 2011;119:1409–14.
43. Billonnet C, Sherrill D, Annesi-Maesano I. Estimating the health effects of exposure to multi-pollutant mixture. *Ann Epidemiol*. 2012;22:126–41.

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