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RESEARCH

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Does the oxidative stress play a role in the associations between outdoor air pollution and persistent asthma in adults? Findings from the EGEA study

Anaïs Havet^{1,2*} , Zhen Li³, Farid Zerimech⁴, Margaux Sanchez⁵, Valérie Siroux⁶, Nicole Le Moual^{1,2}, Bert Brunekreef^{7,8}, Nino Künzli^{9,10}, Bénédicte Jacquemin^{1,2}, Raphaëlle Varraso^{1,2}, Régis Matran^{11†} and Rachel Nadif^{1,2†} 

Abstract

Background: Evidences that oxidative stress plays a role in the associations between outdoor air pollution and asthma are growing. We aimed to study the role of plasma fluorescent oxidation products levels (FIOPs; an oxidative stress-related biomarker), as potential mediators, in the associations between outdoor air pollution and persistent asthma.

Methods: Analyses were conducted in 204 adult asthmatics followed up in the French case-control and family study on asthma (EGEA; the Epidemiological study of the Genetic and Environmental factors of Asthma). Persistent asthma was defined as having current asthma at EGEA2 (baseline, 2003–2007) and EGEA3 (follow-up, 2011–2013). Exposures to nitrogen dioxide, nitrogen oxides, road traffic, particulate matter with a diameter $\leq 10 \mu\text{m}$ (PM_{10}) and $\leq 2.5 \mu\text{m}$ were estimated by ESCAPE models (2009–2010), and ozone (O_3) by IFEN models (2004). We used a mediation analysis to assess the mediated effect by FIOPs levels and the interaction between FIOPs levels and air pollution.

Results: FIOPs levels increased with PM_{10} and O_3 (adjusted $\beta = 0.04$ (95%CI 0.001–0.08), $a\beta = 0.04$ (95%CI 0.009–0.07) per $10 \mu\text{g}/\text{m}^3$, respectively), and the risk of persistent asthma increased with FIOPs levels (aOR = 1.81 (95%CI 1.08–3.02)). The risk of persistent asthma decreased with exposures to NO_2 , NO_x and $\text{PM}_{2.5}$ (aOR ranging from 0.62 to 0.94), and increased with exposures to PM_{10} , O_3 , $\text{O}_{3\text{-summer}}$ and road traffic, the greater effect being observed for O_3 (aOR = 1.78, 95% CI 0.73–4.37, per $10 \mu\text{g}/\text{m}^3$). Using mediation analysis, we observed a positive total effect (aOR = 2.16, 95%CI 0.70–11.9), a positive direct effect of O_3 on persistent asthma (OR = 1.68, 95%CI 0.57–7.25), and a positive indirect effect mediated by FIOPs levels (aOR = 1.28 (95%CI 1.01–2.29)) accounting for 41% of the total effect.

Conclusions: Our results add insights on the role of oxidative stress in the association between air pollution and persistent asthma.

Keywords: Epidemiology, Outdoor air pollution, Persistent asthma, Oxidative stress, Fluorescent oxidation products, Mediation analysis

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Background

Evidences that outdoor air pollution is associated with asthma incidence, and various asthma phenotypes as asthma control or asthma severity among asthmatics keep going up [1, 2], but the associations between outdoor air pollution and persistent asthma are very scarcely studied in adults. To our knowledge, only one study has investigated the associations between nitrogen dioxide (NO₂) exposure, traffic road and persistent asthma, and showed that living close to a major road was associated with persistent asthma in adults followed up over eight years [3].

The underlying biological mechanisms by which outdoor air pollution may affect respiratory health include inflammatory processes, immune response modulations, genetic modifications and oxidative stress damages, which are increasingly suggested. Asthma is an oxidative stress-related disease [4], and exposures to ozone (O₃), NO₂ and particulate matter (PM) have been found associated with oxidative stress [5]. Studying biomarkers is a useful approach to provide new insights into the biological mechanisms that drive the disease process, to predict the development and progression of a disease and to personalize intervention strategies [6, 7]. Among the various biological markers related to oxidative stress, plasma fluorescent oxidation products (FLOPs) levels are biomarkers of damages that reflect systemic oxidative stress [8] and are of growing interest in epidemiology. In prospective studies, high plasma FLOPs levels were positively associated with the incidence of coronary heart diseases (CHD) among men without previous cardiovascular events [9], and with the risk of future CHD in women [10]. Regarding asthma, among adults from the Epidemiological study of the Genetic and Environmental factors of Asthma (EGEA), we recently reported higher plasma FLOPs levels significantly associated with asthma attacks, poor asthma control and poor lung function [11]. Overall, better understanding the underlying biological mechanisms related to asthma, and discovering novel biomarkers is the first step towards improving asthma management. To our knowledge, only two studies have investigated the associations between environmental factors and FLOPs levels. Among a population of U.S. trucking industry employees, short-term exposure to occupational PM_{2.5} was unrelated to plasma FLOPs levels [12]. In the EGEA study, occupational exposure to irritant cleaning products and to low molecular weight agents, especially highly reactive chemicals, were significantly and positively associated with higher plasma FLOPs levels in men, and an association was suggested for irritant cleaning products in women [13]. To date, the role of plasma FLOPs levels in the association between outdoor air pollution and asthma has never been studied.

We hypothesized that oxidative stress is one of the underlying biological mechanisms involved in the association between outdoor air pollution and persistent

asthma. Among adults followed up in the EGEA study, we first studied the associations between outdoor air pollution (NO₂, nitrogen oxides (NO_x), particulate matter (PM), traffic load, traffic intensity, O₃ and O_{3-summer}), plasma FLOPs levels and persistent asthma. Then, according to the results, we investigated the role of plasma FLOPs levels as potential mediators in the association between outdoor air pollution and persistent asthma. We performed a mediation analysis which quantified both the mediated effect by FLOPs levels and the interaction between FLOPs levels and outdoor air pollution.

Methods

Study design

The EGEA is a cohort study based on an initial group of asthma cases recruited in chest clinics from five French cities (1991–1995) along with their first-degree relatives, and a group of controls (<https://egeanet.vjf.inserm.fr/>). The protocol and descriptive characteristics have been described previously [14, 15], and inclusion criteria used to define asthmatic cases and controls were described in Additional file 1. A 12-year follow-up of the initial cohort was conducted between 2003 and 2007 (EGEA2) [16], and 1571 adults aged ≥16 years had a complete examination. As a follow-up study of EGEA2, the third survey (EGEA3, 2011–2013, *n* = 1558) was conducted using self-completed questionnaire only. The EGEA collection was certified ISO 9001 since 2006 to 2018 [17]. Ethical approval was obtained from the relevant institutional review board committees (Cochin Port-Royal Hospital and Necker-Enfants Malades Hospital, Paris). All participants signed a written informed consent.

The analyses included 204 adults with current asthma at EGEA2, followed up at EGEA3, and with data on outdoor air pollution and plasma FLOPs levels (Additional file 2: Figure S1). In comparison to participants not included in the analyses, participants included had lower body mass index, lived longer at the same residential address and had lower exposure to NO₂, PM_{2.5}, O₃ and O_{3-summer} (all *P*-value ≤ 0.05, Additional file 1: Table S1). No differences were found regarding other variables.

Definition of persistent asthma

At EGEA2, the participants with ever asthma answered positively to at least one of the two following questions: “Have you ever had attacks of breathlessness at rest with wheezing?” or “Have you ever had asthma attacks?”, or were recruited as asthmatic cases at EGEA1. Among participants with ever asthma, those with current asthma reported asthma attacks or the use of asthma medication in the past twelve months. Participants with persistent asthma had current asthma at both EGEA2 and EGEA3, and those with remittent asthma had current asthma only at EGEA2. Participants with remittent asthma were

used as reference (see Additional file 1 and for further respiratory outcomes definitions).

Exposure assessment

Available air pollution data from ESCAPE (European Study of Cohorts for Air Pollution Effects) were NO₂, NO_x and particulate matter with a diameter ≤ 10 and ≤ 2.5 μm (PM₁₀ and PM_{2.5}), and those from IFEN (French Institute for the Environment) were O₃ and O_{3-summer}. Outdoor air pollution exposures were assigned to each participant's residential address.

Annual air pollution levels of NO₂ and particulate matter were derived from ESCAPE standardised models (www.escapeproject.eu). Briefly, the ESCAPE monitoring campaigns took place between 2009 and 2010, including 40 measurement sites for NO₂ and NO_x in Paris, Lyon, Grenoble and Marseille, and 20 particulate matter measurement sites in Paris and Grenoble. Land-use regression (LUR) models were developed and two indicators of road traffic exposure were also calculated. Traffic intensity on nearest road was defined as the number of motor vehicles circulating per day on the nearest road to the participant's home and was expressed in vehicles per day. Total traffic load was defined as the traffic load on all major roads based around a buffer of 100 m from the participant's home and was expressed by traffic intensity multiplied by road length. Back-extrapolation was used to transfer the current LUR models (2009–2010) to earlier years (2003–2007, EGEA2) (see Additional file 1 for more details). In our study, the estimation of outdoor air pollution by ESCAPE took place after EGEA2, and accordingly we also analysed the back-extrapolated pollution estimates in order to obtain a better temporality between outdoor air pollution and plasma collection. Back-extrapolated pollution data were available for NO₂ and NO_x in all cities, and for PM₁₀ in Paris. The spatial resolution was 50 m \times 50 m. In order to supplement the ESCAPE data set, we used O₃ and O_{3-summer} exposures from the IFEN (see Additional file 1 for more details). The O₃ estimate was the yearly mean ozone level in 2004 for each participant at the residential address and derived from a geo-statistical model as described previously [18]. The O_{3-summer} exposure was assessed from the monthly means from April to September. The spatial resolution was 4 km \times 4 km.

Measurement of plasma FLOPs levels

Plasma samples were collected in EGEA2 between 2003 and 2006 and stored immediately at -80 °C during 5.0 to 8.0 years until FLOPs measurements. Plasma FLOPs levels were measured as previously described [8, 13] (Additional file 1). Briefly, plasma was extracted into a mixture of ethanol/ether (3/1 v/v) and measured using a spectrofluorimeter (360 nm excitation wavelength, 430 nm emission

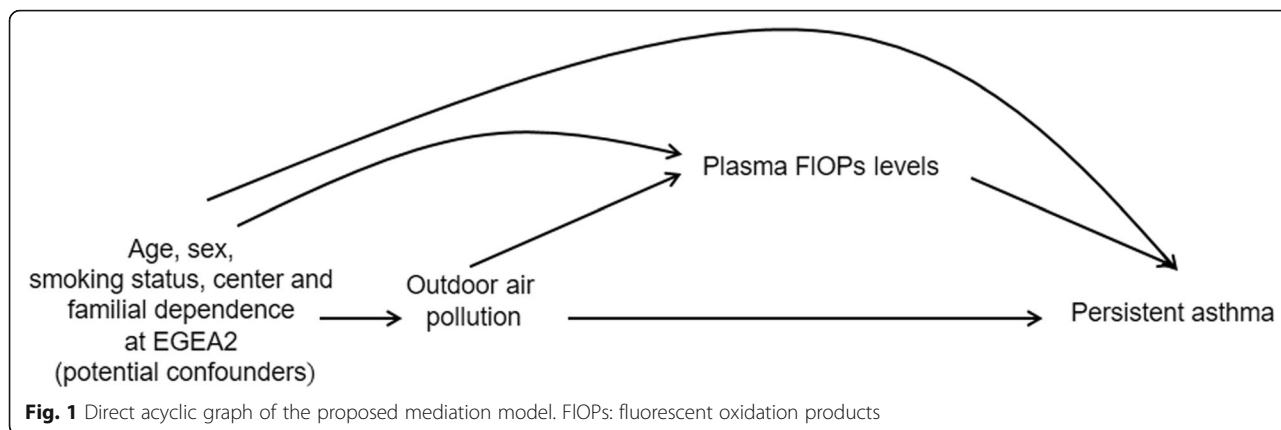
wavelength). Fluorescence was expressed as a unit of relative fluorescence intensity (RFU/mL) of plasma.

Statistical methods

Due to their skewed distribution, plasma FLOPs levels were log₁₀-transformed. Due to the familial dependence of the data, multivariate analyses (except mediation analyses) took into account dependence between observations. Linear regression models and logistic regression models with random effects on center and familial dependence were used to study the associations between outdoor air pollution with plasma FLOPs levels, and between outdoor air pollution and persistent asthma, respectively. To control a potential effect of short-term exposure to O₃ in the associations between O₃ with plasma FLOPs levels and persistent asthma, further adjustment for the season of plasma collection (EGEA2) was conducted. Logistic regression models using generalized estimated equations (GEEs) on familial dependence were performed to study associations between plasma FLOPs levels and persistent asthma. To study only the road traffic effect, estimates of associations between road traffic with persistent asthma or plasma FLOPs levels were also adjusted for background NO₂. To obtain a better temporality between outdoor air pollution assessed by ESCAPE and plasma collection, analyses with back-extrapolated pollution estimates were also performed.

We used a direct acyclic graph to represent our mediation model (Fig. 1), and conducted mediation analysis based on a counterfactual approach by using the CAUS ALMED procedure [19, 20] (see Additional file 1). Mediation analysis was performed for air pollutants associated with both persistent asthma and plasma FLOPs levels, regardless of the significance of the association between air pollutant and persistent asthma. Models did not include random effects on center and familial dependence. The four-way decomposition was used to investigate the proportions of total effect that were attributable to the controlled direct effect, to mediation (the pure indirect effect), to interaction (the reference interaction between pollutant and plasma FLOPs levels) and to both mediation and interaction (the mediated interaction) [21]. Percentages mediated and due to interaction were given. All these components were defined in Additional file 1.

All estimates were adjusted for age (continuous), sex and smoking status (never-, ex- or current smokers). We defined never-smoker as a participant who have never smoked in their life, ex-smoker as a participant who quit smoking for at least 4 weeks at EGEA2 and current smoker as a participant who was smoking at least one cigarette a day for more than 1 year at EGEA2. The results are expressed for an increase of 20 $\mu\text{g}/\text{m}^3$ of NO_x, 10 $\mu\text{g}/\text{m}^3$ of NO₂, PM₁₀, O₃ and O_{3-summer} and 5 $\mu\text{g}/\text{m}^3$ of PM_{2.5}. The results of traffic load are expressed for 4 million vehicles multiplied by meters per day on major



roads within a 100 m buffer, and those of traffic intensity for an increase of 5000 vehicles per day on major roads. We performed sensitivity analyses by excluding participants who lived at the same address for < 1 year. Statistical analyses were performed using SAS version 9.4 (SAS Institute, Cary, NC, USA).

Results

At EGEA2, the mean age of the 204 adults was 39 years, 48% were men, 24% were current smoker, 79% had persistent asthma, and the geometric mean (interquartile range) of plasma FIOPs levels was 92.0 (79.5–104.7) RFU/mL (Table 1). In comparison to participants with remittent asthma, participants with persistent asthma had more often poor controlled asthma and an asthma symptom score > 2, reported more often the use of inhaled corticosteroids, and had higher plasma FIOPs levels (all *P* values ≤ 0.02, Table 1). Plasma FIOPs levels increased with age (*p*-trend < 0.0001, Additional file 1: Table S2), were lower in never smokers than smokers (*p* = 0.02, Additional file 1: Table S2). No significant association was found between plasma FIOPs levels and other variables (Additional file 1: Table S2).

Associations between outdoor air pollution, plasma FIOPs levels and persistent asthma

Among all pollutants studied, plasma FIOPs levels increased by 1 RFU/mL with PM₁₀ and O₃ exposures (adjusted (a)β = 0.04, 95% CI 0.001–0.08, *p* = 0.03 and aβ = 0.04, 95% CI 0.009–0.07, *p* = 0.02 for an increase of 10 μg/m³ of O₃ and PM₁₀, respectively, Table 2). The results were similar after excluding participants who lived at the same address for less than 1 year. The association between PM₁₀ and plasma FIOPs levels did not remain significant with back-extrapolated data (aβ = 0.03, 95% CI -0.01–0.07, *p* = 0.22 for an increase of 10 μg/m³ of PM₁₀, Table 2). The risk of persistent asthma decreased not significantly with exposure to NO₂, NO_x and PM_{2.5} (aOR ranging from 0.62 to 0.94), and increased not

significantly with exposure to PM₁₀, O₃, O₃-summer and road traffic, the greater effect being observed for O₃ (aOR = 1.78, 95% CI 0.73–4.37, for an increase of 10 μg/m³ of O₃, model 2, Additional file 1: Table S3). Results were similar after excluding participants who lived at the same address for less than 1 year (Table 2 and Additional file 1: Table S3). Further adjustment for season of plasma collection did not change the results with O₃ (data not shown). The risk of persistent asthma increased with plasma FIOPs levels (unadjusted OR = 1.68, 95% CI 1.04–2.70, *p* = 0.03 for an increase of 1 interquartile range of FIOPs levels), and this association remained consistent after adjustment for age, sex and smoking status (aOR = 1.81, 95% CI 1.08–3.02, *p* = 0.02) for an increase of 1 interquartile range of FIOPs levels).

Ozone, FIOPs levels and persistent asthma: mediation analysis

Results were summarized in Table 3. We observed a positive total effect (OR_{TE} = 2.16, 95% CI 0.70–11.9, *P* > 0.05), a positive natural direct effect of O₃ on persistent asthma (OR_{CDE} = 1.68, 95% CI 0.57–7.25, *P* > 0.05), and a positive natural indirect effect mediated by plasma FIOPs levels (OR_{NIE} = 1.28, 95% CI 1.01–2.29, *P* = 0.04). Further decomposition of total effect showed that 50.3% of total effect was due to control direct effect, 8.6% due to the reference interaction, 26.5% due to mediated interaction and 14.6% due to the pure indirect effect. The percentage of total effect mediated by plasma FIOPs levels was 41.1% (26.5% + 14.6%). The percentage of total effect due to interaction was 35.1% (8.6% + 26.5%). These results were quite similar after excluding participants who lived at the same address for < 1 year (Additional file 1: Table S4). We also found that the controlled direct effect increased with plasma FIOPs levels (Additional file 1: Table S5).

Discussion

We studied and quantified the role of plasma FIOPs levels in the association between outdoor air pollution and persistent asthma. We found that plasma FIOPs

Table 1 Description of participants with current asthma at baseline (EGEA2) according to change in current asthma between EGEA2 to EGEA3

	All participants	Participants with remittent asthma	Participants with persistent asthma	P-value
N	204	43	161	
Age (years), mean \pm SD	39.3 \pm 16.9	36.7 \pm 15.3	40.0 \pm 17.3	0.26
Male, n (%)	99 (48.5)	16 (37.2)	83 (51.6)	0.09
Smoking status, n (%)				
Never-smoker	109 (53.4)	22 (51.2)	87 (54.0)	0.52
Ex-smoker	46 (22.6)	8 (18.6)	38 (23.6)	
Current smoker	49 (24.0)	13 (30.2)	36 (22.4)	
BMI (kg/m ²), mean \pm SD	n = 203 23.8 \pm 3.84	23.4 \pm 3.99	n = 160 23.9 \pm 3.81	0.40
Socioprofessional category, n (%)				
Unemployed	32 (15.7)	5 (11.6)	27 (16.8)	0.37
Manager	64 (31.4)	12 (27.9)	52 (32.3)	
Technician	88 (43.1)	19 (44.2)	69 (42.8)	
Manual worker	20 (9.8)	7 (16.3)	13 (8.1)	
Asthma onset years, mean \pm SD	N = 195 14.8 \pm 14.7	N = 39 16.6 \pm 14.6	N = 156 14.4 \pm 14.7	0.40
Poor controlled asthma ^a , n (%)	N = 190 102 (53.7)	N = 38 13 (34.2)	N = 152 89 (58.6)	0.007
Use of inhaled corticosteroids, n (%)	N = 200 132 (66.0)	N = 41 20 (48.8)	N = 159 112 (70.4)	0.009
Symptom score of asthma ^b , n (%)	N = 199	N = 40	N = 159	
0	20 (10.0)	9 (22.5)	11 (6.9)	0.005
1–2	30 (15.1)	8 (20.0)	22 (13.8)	
>2	149 (74.9)	23 (57.5)	126 (79.3)	
Plasma FLOPs levels (RFU/mL), GM (q1–q3)	92.0 (79.5–104.7)	85.6 (76.0–96.5)	93.8 (79.9–104.8)	0.02
Residence time (years), mean \pm SD	11.7 \pm 10.2 (0–42)	10.3 \pm 8.90 (0–31)	12.1 \pm 10.5 (0–42)	0.31
Air pollutant levels and traffic metrics ^c , mean \pm SD				
NO ₂	27.9 \pm 13.9	29.7 \pm 15.3	27.4 \pm 13.5	0.34
NO _x	49.7 \pm 32.7	53.2 \pm 38.4	48.8 \pm 31.0	0.44
PM ₁₀	25.0 \pm 4.07	25.1 \pm 3.71	24.9 \pm 4.18	0.88
PM _{2.5}	15.0 \pm 2.04	15.5 \pm 1.96	14.9 \pm 2.05	0.20
Traffic load	1,755,353 \pm 4,718,630	1,672,062 \pm 4,007,561	1,777,599 \pm 4,902,148	0.90
Traffic intensity	7307 \pm 13,109	7027 \pm 11,360	7382 \pm 13,569	0.87
O ₃	43.9 \pm 4.62	43.1 \pm 4.08	44.2 \pm 4.74	0.15
O ₃ -summer	65.4 \pm 6.20	64.7 \pm 6.41	65.6 \pm 6.14	0.38

SD standard deviation, BMI body mass index, FLOPs fluorescent oxidation products, GM geometric mean; q1–q3: 25th and 75th percentiles of the GM

^aDefined according to GINA 2015 guidelines and from participants with partly controlled asthma or uncontrolled asthma (see Additional file 1)

^bMore details in Additional file 1

^cPollutant levels in $\mu\text{g}/\text{m}^3$, traffic load in vehicles/day x meters, and traffic intensity in vehicles/day

levels increased with PM₁₀ and O₃ exposures, and the risk of persistent asthma increased with plasma FLOPs levels. We also found that the risk of persistent asthma increased with O₃ exposure, but not significantly. We therefore performed mediation analysis to investigate the role of plasma FLOPs levels in the association

between O₃ exposure and persistent asthma. We found positive indirect effect mediated by plasma FLOPs levels accounting for 41% of the total effect, and O₃ effect on persistent asthma increased with plasma FLOPs levels. Due to the limited sample size and borderline significant findings, and to the other biomarkers related to

Table 2 Associations between outdoor air pollution and plasma FIOPs levels

		NO ₂	NO _x	PM ₁₀ ^a	PM _{2.5} ^a	Total traffic load on major roads in a 100-m buffer of the home ^b	Traffic intensity at the road nearest to a participant's home ^b	O ₃	O ₃ -summer
Model 1	n	204	204	146	146	204	204	204	204
	β crude	0.007	0.006	0.04	0.03	0.008	0.0004	0.03	0.02
	(95% CI)	(− 0.004, 0.02)	(− 0.003, 0.02)	(0.001,0.08)	(− 0.009, 0.07)	(− 0.005,0.02)	(− 0.005,0.006)	(− 0.004, 0.06)	(− 0.03, 0.07)
	P	0.19	0.15	0.04	0.15	0.24	0.88	0.06	0.36
Model 2	n	204	204	146	146	204	204	204	204
	β adjusted	0.007	0.007	0.04	0.03	0.006	0.0002	0.04	0.02
	(95% CI)	(− 0.003, 0.02)	(− 0.002, 0.02)	(0.001,0.08)	(− 0.008, 0.07)	(− 0.006,0.02)	(− 0.005,0.006)	(0.009,0.07)	(− 0.02, 0.06)
	P	0.18	0.16	0.03	0.18	0.30	0.94	0.02	0.45
Model 3	n	204	204	72					
	β adjusted	0.007	0.006	0.03					
	(95% CI)	(− 0.002, 0.02)	(− 0.002, 0.01)	(− 0.01, 0.07)					
	P	0.14	0.12	0.22 ^c					
Model 4	n	186	186	133	133	186	186	186	186
	β adjusted	0.009	0.008	0.05	0.02	0.003	0.0005	0.05	0.02
	(95% CI)	(− 0.003, 0.02)	(− 0.002, 0.02)	(0.006,0.09)	(− 0.02,0.06)	(− 0.01,0.02)	(− 0.005,0.006)	(0.01,0.08)	(− 0.03, 0.07)
	P	0.14	0.14	0.04	0.24	0.63	0.86	0.01	0.37

The linear regression models were conducted with random effects on familial dependence and center. FIOPs levels were log₁₀-transformed. NO₂, nitrogen dioxide; NO_x, nitrogen oxides; PM₁₀, particulate matter with a diameter ≤ 10 μm; PM_{2.5}, particulate matter with a diameter ≤ 5 μm. Results are expressed per 20 μg/m³ increase of NO_x exposure, per 10 μg/m³ increase of NO₂, PM₁₀ O₃ and O₃-summer exposures, per 5 μg/m³ increase of PM_{2.5} exposure, per 4 million vehicles x meters per day increase of total traffic load, per 5000 vehicles per day increase of traffic intensity. Model 1: unadjusted. Model 2: adjusted for age, sex and smoking status. Model 3: with back-extrapolated pollution and adjusted for age, sex and smoking status. Model 4: after excluding the participants living at the same residential address <1 year

^aNot back-extrapolated PM were only estimated in Paris and in Grenoble, and back extrapolated PM₁₀ only in Paris

^bEstimates were also adjusted for background NO₂

^cThe linear regression model was only conducted with random effects on familial dependence because back-extrapolated PM₁₀ were estimated only in Paris

oxidative stress potentially involved in this association, the results need to be interpreted with caution.

To our knowledge, this study is the first one investigating the associations between outdoor air pollution and plasma FIOPs levels. FIOPs levels are biomarkers of damages related to oxidative stress reflecting a mixture of oxidation products from lipids, proteins and DNA [6]. These

Table 3 Results of mediation analysis between O₃, persistent asthma and plasma FIOPs levels using the CAUSALMED procedure (n = 204)

OR	Estimate	95% CI
Odds Ratio Total Effect	2.16	0.70–11.9
Odds Ratio Natural Direct Effect	1.68	0.57–7.25
Odds Ratio Natural Indirect Effect	1.28	1.01–2.29
Decomposition of the total effect		
Four-way	Percent	
Controlled direct	50.3	
Reference interaction	8.6	
Mediated interaction	26.5	
Pure indirect	14.6	

Models were adjusted for age, sex and smoking habits. The exposure was O₃. The mediators were plasma FIOPs levels. The outcome was persistent asthma

biomarkers are of growing interest in epidemiological studies because they are stable, easily measurable and applicable in large-scale human studies [9]. We found that plasma FIOPs levels increased with long-term exposure to O₃, an irritant gas with a strong oxidative potential [5]. To date, no study has investigated the associations between O₃ exposure and plasma FIOPs levels, and a recent review reported positive and significant associations between short-term exposure to O₃ and 8-isoprostane, another biomarker related to oxidative stress [22]. We did not have data to properly assess by which extent acute exposure to O₃ (over the past hours/days) could have biased our results. Nevertheless, the adjustment for the season of plasma collection did no change the results. As plasma FIOPs levels are biomarkers of damages that reflect cumulative oxidative stress, we hypothesized to find associations with long-term exposure to air pollution rather than with short-term air pollution. Further studies are needed to differentiate the effects of short-term and long-term exposure to air pollution, and in particular O₃, on plasma FIOPs levels. We also found that plasma FIOPs levels increased with non-back extrapolated PM₁₀. The way and the strength of the association with back-extrapolated PM₁₀ were the same, but the association was not significant due to the small effective. Previously, we found that 8-

isoprostane in exhaled breath condensate, a matrix close to the lungs, increased significantly with $PM_{2.5}$ exposure in the EGEA study [23]. 8-isoprostane is a biomarker of damages related to oxidative stress, and a specific product of lipid peroxidation. In the present analysis, plasma FLOPs levels increased with $PM_{2.5}$ exposure but the association was not significant. Fine and ultrafine particulates are known to be more harmful by penetrating deeper into the lungs and inducing damages due to oxidative stress both at the airways and systemic compartment [24]. The discrepancies in the results could be partly explained by difference in the sample sizes, in the composition and concentration of the particulates, and by difference in underlying mechanism related to the studied biomarker. In a previous work conducted among participants without asthma in the EGEA study, occupational exposure to irritant cleaning/disinfecting agents increased plasma FLOPs levels [13]. Overall, all these results underlined that occupational exposure to irritants, and outdoor air pollution exposure, especially exposure to O_3 and PM_{10} , were associated with higher plasma FLOPs levels, and outdoor air pollution increased oxidative stress at both lung and systemic level.

We studied persistent asthma in association with O_3 , PM and plasma FLOPs levels. Asthma is a chronic and heterogeneous disease defined by various overlapping phenotypes, including the phenotype “persistent asthma” [25]. Although persistent asthma is still under-studied in epidemiology, it reflects the activity and the evolution of asthma over time. Most associations between pollution and persistent asthma were close to 1; the risk of persistent asthma decreased with $PM_{2.5}$ and increased with O_3 and $O_{3\text{-summer}}$ exposures. The unexpected result observed for $PM_{2.5}$ may be partly due to the lack of back-extrapolated data leading to an inverse temporality between $PM_{2.5}$ and persistent asthma, or to random effect or residual bias. To date, only one study investigated the associations between exposure to outdoor air pollution and with persistent asthma in adults, and showed that living within 200 m of a major road was associated with persistent asthma in middle-aged Tasmanian participants [3]. Like us, the authors defined “current asthma” as “any episode of asthma or use of asthma medication during the last 12 months”. We also found that the risk of persistent asthma increased with plasma FLOPs levels. Our definition of “current asthma” included the report of asthma attack and the use of respiratory treatment in the last twelve months, and interestingly in adults of the EGEA study, asthma attacks, any asthma treatment and use of inhaled corticosteroids in the past 12 months were positively associated with plasma FLOPs levels [11]. Interestingly, leukocyte telomere length, which reflects oxidative-stress damages to DNA [26], was shorter in participants who had

persistent asthma from childhood into adult as compared to those who had adolescent or adult-onset asthma [27]. From a larger sample size, it would be now interesting to study the associations between outdoor air pollution, plasma FLOPs levels and asthma incidence.

We acknowledge that performing a mediation analysis despite the non-significant association between O_3 and persistent asthma may open a debate. We based our decision on the biological hypothesis that oxidative stress is one mechanism by which outdoor air pollution affects respiratory health. Although the association between O_3 and persistent asthma was non-significant, the strength of the association between O_3 and persistent asthma, and previous results obtained in the EGEA study guided our decision. Indeed, O_3 was associated with severe asthma [18], uncontrolled asthma [28] and current asthma [23] in adults.

The CAUSALMED procedure is a recent tool available in SAS software to estimate causal mediation effects from observational data [19]. This procedure is advisable without prior knowledge about the lack of the interaction [29]. In mediation analysis, the random effects on center and familial dependence were not taken into in models, explaining the differences of results between the association between O_3 and persistent asthma studied outside the mediation analysis and the direct effect. Beyond direct and indirect effects, the four-way decomposition shed insights into the role of both mediation and interaction in the associations between O_3 and plasma FLOPs levels with persistent asthma. Overall, our mediation analysis may suggest a not negligible effect of O_3 on persistent asthma through plasma FLOPs levels, and more precisely that effect of O_3 on persistent asthma increased with plasma FLOPs levels. Due to the small sample size, the estimates from mediation analysis may be imprecise and the results should therefore be interpreted with caution. Further studies, with a larger study sample, using a model including other biomarkers related to oxidative stress or biomarkers related to other pathways, as well as the interrelations between these biomarkers, would be helpful to better understand the underlying biological mechanisms between outdoor air pollution and asthma.

Conclusion

For the first time in adults, we found that plasma FLOPs levels increased with O_3 and PM_{10} exposures, and the risk of persistent asthma increased with plasma FLOPs levels. Overall, our results add insights into the potential role of plasma FLOPs levels in the association between O_3 and persistent asthma, and add new evidence on the role of oxidative stress in the association between outdoor air pollution and asthma.

Supplementary information

Supplementary information accompanies this paper at <https://doi.org/10.1186/s12940-019-0532-0>.

Additional file 1: Table S1. Description of participants included and not included in analyses. **Table S2.** Associations between plasma FLOPs levels and characteristics of participants. **Table S3.** Associations between outdoor air pollution and persistent asthma. **Table S4.** Results of mediation analysis using the CAUSALMED procedure among participants who lived at the same address for > 1 year ($n = 186$). **Table S5.** Controlled direct effect according to quantiles of plasma FLOPs levels ($n = 204$).

Additional file 2: Figure S1. Flow chart of the studied population.

Abbreviations

CI: confidence interval; EGEA: Epidemiological study of the Genetic and Environmental factors of Asthma; ESCAPE: European Study of Cohorts for Air Pollution Effects; FLOPs: fluorescent oxidation products; IFEN: French Institute of Environment; NO₂: nitrogen dioxide; NO_x: nitrogen oxides; O₃: ozone; O_{3-s}: summer ozone; OR: odds ratio; PM₁₀: particulate matter with a diameter $\leq 10 \mu\text{m}$; PM_{2.5}: particulate matter with a diameter $\leq 2.5 \mu\text{m}$

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Authors' contributions

AH, RM and RN designed and conducted the research; MS, VS, NLM, BB, NK and BJ: provided essential reagents or provided essential materials; FZ performed biological analyses; AH, ZL, RV and RN: analyzed data or performed statistical analysis; AH, RM and RN: wrote the manuscript and had primary responsibility for final content; all authors read, edited and approved the final manuscript.

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Availability of data and materials

Due to third party restrictions, EGEA data are not publicly available. Please see the following URL for more information: <https://egeanet.vjf.inserm.fr/index.php/en/contacts-en>. Interested researchers should contact egea.cohorte@inserm.fr with further questions regarding data access.

Ethics approval and consent to participate

Ethical approval was obtained from the relevant institutional review board committees (Cochin Port-Royal Hospital and Necker-Enfants Malades Hospital, Paris).

Consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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References

- Guarnieri M, Balmes JR. Outdoor air pollution and asthma. *Lancet*. 2015; 383(9928):1581–92.
- Kar Kurt O, Zhang J, Pinkerton KE. Pulmonary health effects of air pollution. *Curr Opin Pulm Med*. 2016;22(2):138–43.
- Bowatte G, Lodge CJ, Knibbs LD, Erbas B, Perret JL, Jalaludin B, et al. Traffic related air pollution and development and persistence of asthma and low lung function. *Environ Int*. 2018;113:170–6.
- Kleniewska P, Pawliczak R. The participation of oxidative stress in the pathogenesis of bronchial asthma. *Biomed Pharmacother*. 2017;94:100–8.
- Lodovici M, Bigagli E. Oxidative stress and air pollution exposure. *J Toxicol*. 2011;2011:487074.
- Margaritelis NV, Copley JN, Paschalis V, Veskoukis AS, Theodorou AA, Kyparos A, Nikolaidis MG. Going retro: oxidative stress biomarkers in modern redox biology. *Free Radic Biol Med*. 2016;98:2–12.
- Guerra S. New asthma biomarkers: shorter telomeres, longer disease? *Am J Respir Crit Care Med*. 2014;190(4):356–8.
- Wu T, Willett WC, Rifai N, Rimm EB. Plasma fluorescent oxidation products as potential markers of oxidative stress for epidemiologic studies. *Am J Epidemiol*. 2007;166(5):552–60.
- Wu T, Rifai N, Willett WC, Rimm EB. Plasma fluorescent oxidation products: independent predictors of coronary heart disease in men. *Am J Epidemiol*. 2007;166(5):544–51.

10. Jensen MK, Wang Y, Rimm EB, Townsend MK, Willett W, Wu T. Fluorescent oxidation products and risk of coronary heart disease: a prospective study in women. *J. Am Heart Assoc.* 2013;2(5):e000195.
11. Andrianjafimasy M, Zerimech F, Akiki Z, Huyvaert H, Le Moual N, Siroux V, et al. Oxidative stress biomarkers and asthma characteristics in adults of the EGEE study. *Eur Respir J.* 2017;50(6):1701193.
12. Hart JE, Wu T, Laden F, Garshick E. Plasma fluorescent oxidation products and short-term occupational particulate exposures. *Am J Ind Med.* 2012; 55(10):953–60.
13. Dumas O, Matran R, Zerimech F, Decoster B, Huyvaert H, Ahmed I, et al. Occupational exposures and fluorescent oxidation products in 723 adults of the EGEE study. *Eur Respir J.* 2015;46(1):258–61.
14. Kauffmann F, Dizier MH, Annesi-Maesano I, Bousquet J, Charpin D, Demenais F, et al. EGEE (epidemiological study on the genetics and environment of asthma, bronchial hyperresponsiveness and atopy) – descriptive characteristics. *Clin Exp Allergy.* 1999;29(Suppl 4):17–21.
15. Kauffmann F, Dizier MH. EGEE (Epidemiological study on the Genetics and Environment of Asthma, bronchial hyperresponsiveness and atopy)—design issues. EGEE Co-operative Group. *Clin Exp Allergy J Br Soc Allergy Clin Immunol.* 1995;25 Suppl 2:19–22.
16. Bouzigon E, Nadif R, Le Moual N, Dizier MH, Aschard H, Boudier A, et al. Facteurs génétiques et environnementaux de l’asthme et de l’allergie : synthèse des résultats de l’étude EGEE. *Rev Mal Respir.* 2015;32(8):822–40.
17. Nadif R, Bouzigon E, Le Moual N, Siroux V. EGEE collection: a biobank devoted to asthma and asthma-related phenotypes. *Open J Bioresources.* 2017;4(1):5.
18. Rage E, Siroux V, Künzli N, Pin I, Kauffmann F. Air pollution and asthma severity in adults. *Occup Environ Med.* 2009;66(3):182–8.
19. SAS/STAT® 14.3 User’s Guide The CAUSALMED Procedure. <https://support.sas.com/documentation/onlinedoc/stat/143/causalmed.pdf>.
20. Valeri L, Vanderweele TJ. Mediation Analysis Allowing for Exposure – Mediator Interactions and Causal Interpretation: Theoretical Assumptions and Implementation With SAS and SPSS Macros. *Psychol Methods* 2013; 18(2):137–150. 2.
21. VanderWeele TJ. A unification of mediation and interaction: a four-way decomposition. *Epidemiology.* 2014;25(5):749–61.
22. Bromberg PA. Mechanisms of the acute effects of inhaled ozone in humans. *Biochim Biophys Acta.* 2016;1860(12):2771–81.
23. Havet A, Zerimech F, Sanchez M, Siroux V, Le Moual N, Brunekreef B, et al. Outdoor air pollution, exhaled 8-isoprostane and current asthma in adults: the EGEE study. *Eur Respir J.* 2018;51(4):1702036.
24. Valavanidis A, Fiotakis K, Vlachogianni T. Airborne particulate matter and human health: toxicological assessment and importance of size and composition of particles for oxidative damage and carcinogenic mechanisms. *Journal of Environmental Science and Health Part C.* 2008; 26(4):339–62.
25. Beasley R, Semprini A, Mitchell EA. Risk factors for asthma: is prevention possible? *Lancet.* 2015;386(9998):1075–85.
26. Clemente DBP, Vrijheid M, Martens DS, Bustamante M, Chatzi L, Danileviciute A, et al. Prenatal and childhood traffic-related air pollution exposure and telomere length in European children: the HELIX project. *Environ Health Perspect.* 2019;127(8):87001.
27. Belsky DW, Shalev I, Sears MR, Hancox RJ, Lee Harrington H, Houts R, et al. Is chronic asthma associated with shorter leukocyte telomere length at midlife? *Am J Respir Crit Care Med.* 2014;190(4):384–39.
28. Jacquemin B, Kauffmann F, Pin I, Le Moual N, Bousquet J, Gormand F, et al. Air pollution and asthma control in the epidemiological study on the genetics and environment of asthma. *J Epidemiol Community Health.* 2012; 66(9):796–802.
29. Yung YF, Lamm M and Zhang W. Causal mediation analysis with the CAUSALMED procedure <https://www.sas.com/content/dam/SAS/support/en/sas-global-forum-proceedings/2018/1991-2018.pdf>.

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