

Effects of air pollution on acute exacerbation of chronic obstructive pulmonary disease: a descriptive retrospective study (pol-AECOPD)

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Purpose: Acute exacerbation of COPD (AECOPD) is among the most frequent causes for hospital admission, causing morbidity and mortality. Infection is the most frequent cause, and studies on pollution have shown higher hospital admission and mortality with inconsistent results. The objective was to identify if there is a change in levels of particulate matter (PM) during the days leading up to the symptom onset.

Patients and methods: A retrospective study was carried out on medical records of patients with AECOPD from a University Hospital. PM values of the consultation day, onset symptoms, and up to three previous days were recorded. Moreover, clinical presentation, laboratory findings, treatments, and hospital outcomes were recorded.

Results: A total of 250 medical records were included, mean age of 77 years, hospital stay mean of 6.7 days, 26.8% with no previous exposure was identified, coexistence with asthma was 5%, Obstructive Sleep Apnea Syndrome 15%, Pulmonary Hypertension 34%, antibiotic use 62%, ICU admission of 14% with non-invasive mechanical ventilation of 68%, and in-patient mortality of 2.4%. PM 2.5 of 48 hrs before onset symptoms median was 20.1 $\mu\text{g}/\text{m}^3$ versus 15 and 16.5 for the day of symptoms and 3 days prior to onset symptoms ($p < 0.001$). PM 10 of 46.65 $\mu\text{g}/\text{m}^3$, versus 39 and 35.6, respectively ($p < 0.001$). Expectoration OR 4.74; Purulence OR 6.58; Pleuritic pain OR 3.62; Antibiotic use OR 2.87, and corticoids use OR 2.62, all with $p < 0.05$.

Conclusions: Patients with AECOPD have a higher median of particulate matter 48 hrs prior to symptomatic onset, as well as greater use of antibiotics and corticosteroids.

Keywords: pulmonary disease, chronic obstructive, particulate matter, air pollutants

Introduction

Chronic obstructive pulmonary disease (COPD) represents one of the main causes of morbidity and mortality worldwide. According to the latest report of the World Health Organization,¹ it is estimated that in 2008 in the United States 15 million people suffered from it, as well as 210 million people worldwide, with an associated mortality of 3 million people.² COPD generates not only respiratory complications but also cardiovascular ones,³ and even osteoporosis.⁴ The deterioration in symptoms beyond the daily variation that leads to the use of additional therapy is known as Acute Exacerbation of COPD (AECOPD), being the most frequent cause of medical hospital admission in the United Kingdom with almost 15.9% of the medical hospitalizations.⁵⁻⁷ In 22% of the patients with very severe COPD, exacerbation recurs, on average, 50 days after the first episode⁵ and generating a 25% deterioration of lung function,⁸ thus increasing mortality, making it five times

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higher in the tenth episode when compared to the first,⁶ and up to 50% when in the Intensive Care Unit.⁹ The cause of AECOPD can be infectious (40–60%)⁶ and non-infectious (smoking, pulmonary thromboembolism, and air pollution or exposure to particulate matter, PM).¹⁰

Regarding exposure to PM, people exposed to environmental pollution have a greater proportion of lung dysfunction with an excess risk of 64%, which when combined with smoking increases to 129%.¹¹ Likewise, it is related to higher hospital admission,¹² deterioration of FEV1,¹³ and mortality.¹⁴ However, assuming that environmental exposure can exacerbate COPD, the objective of this study was to examine whether the levels of PM below 2.5 (PM2.5) and 10 microns (PM10) in the days before the symptomatic onset in adult patients with AECOPD show any change, and if that is associated with clinical presentation or outcome of an Anthonisen classification, use of antibiotic, hospitalization days, admission to ICU, and mortality.

Materials and methods

The protocol was approved by the institutional review board of Universidad Nacional de Colombia Number 004-033-17. Patient consent to review their medical review records was not required by the board, and thus no identifying information such as name, Identification number, and Medical number were used, and it was conducted in accordance with the principles of the Declaration of Helsinki.

Study design and participants

A retrospective study was conducted with medical records of adult patients hospitalized at the Hospital Universitario Nacional de Colombia (Bogotá, Colombia), with an admissions diagnosis of AECOPD (ICD 10: J441 to J449) between the months of April 2016 and April 2017 and residents in places with air quality measurement. Exposure higher to the daily average levels recommended by the WHO is of 25 $\mu\text{g}/\text{m}^3$ and 50 $\mu\text{g}/\text{m}^3$ for PM2.5 and PM10, respectively.¹⁵ Computer records of the medical records with the following inclusion criteria were taken: patient over 18 years of age with a spirometric diagnosis of COPD who came in due to respiratory symptoms that suggested acute exacerbation of COPD, from a borough that has air quality measurement. Patients with other diagnoses that explained the clinical profile, missing data referring to diagnosis or treatment, and residents in areas where there is not a meteorological station into a kilometer radius or does not have PM measurements were excluded.

Sampling

If they met the inclusion criteria, medical histories were selected by non-probabilistic convenience sampling in order of arrival. A sample size of 200 was calculated for mean differences of independent samples in order to detect mean differences between levels of PM2.5 and PM10 of 0.2 $\mu\text{g}/\text{m}^3/24\text{hrs}$, with a standard deviation of 0.5 $\mu\text{g}/\text{m}^3/24\text{hrs}$, a statistical power of 80% and 95% confidence, calculated using the Epidat software (Version 4.1, Oct 2014, PAHO-WHO). Twenty percent of the estimated losses were added due to missing data, totaling 240. The record was made in EXCEL and the analysis in EPIDAT.

Bias control

PM2.5 and PM10 values from the day of consultation, day of symptom onset, and three days prior to it were recorded. To achieve this, during hospital admission, each patient had to register their place of residence, the nearest meteorological station was located, and the average values for 24 hrs of PM2.5 and PM10 were recorded. In addition, variables of clinical presentation were recorded such as increased dyspnea, expectoration and sputum change, pleuritic pain, fever and altered state of consciousness, clinical examinations and, as outcome variables, the use of antibiotics, corticosteroids, admission to the Intensive Care Unit (ICU), days at the ICU, ventilatory support requirement, new indication for oxygen therapy, and in-patient mortality. The bias control was made by selecting those exposed in an automated way using the software. Temperature, precipitation, humidity, and alternative pollutants were registered, but these had rarely been collected, about 10%; therefore, these were excluded from the analysis.

A univariate analysis in proportions with 95% confidence intervals and central tendency and dispersion measurements considered normal distributions by Shapiro–Wilk test with a 0.05 significance that characterizes the exacerbation of COPD according to person, time, and place variables was performed. A median difference was made using the Kruskal–Wallis statistic with a significance less than 0.05. Finally, to determine the secondary objectives and control of confounding variables, a multivariate logistic regression analysis was carried out in search of the association between qualitative variables (Anthonisen Group, use of antibiotic and admission to ICU), and PM10 and 2.5 levels registered at the time of symptom onset, three days prior to the consultation and a cumulative total of four days with control of confounding variables, including variables that show statistical significance defined

by the Wald statistic with a significance below 0.05. All graphs and analyses were made using the GraphPad Prism software (GraphPad Prism version 7.00 for Windows, GraphPad Software, La Jolla, California, USA) through license GPS-1,030,895-EFOT-DAF4F. The final document was written according to the STROBE initiative¹⁶

Results

We reviewed 1549 medical records, finally including 250 (see Figure 1). The demographic characteristics can be seen in Table 1. Twenty-eight percent of the patients did not have previous spirometric diagnosis; coexistence with asthma

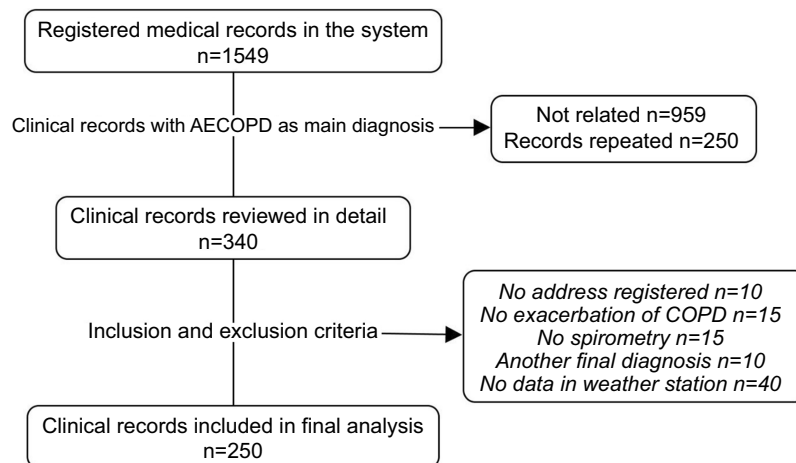


Figure 1 Selection algorithm of medical records. AECOPD: Acute exacerbations of chronic obstructive cardiopulmonary disease. Flow-chart of clinical records included.

Table 1 Clinical-demographic characteristics of the patients included in the study

Characteristic	Frequency (n)	Proportion (%)	Mean	IC95%
Age (years)			77.4	[76.1–78.8]
Symptom duration (days)			7.8	[7.09–8.62]
Hospitalization duration (days)			6.7	[6.23–7.25]
Female	147	58.8		[52.8–64.9]
GOLD A	17	6.8		
B	21	8.4		
C	56	22.4		
D	90	36.0		
Asthma	13	5.2		[3.0–8.7]
Interstitial disease	13	5.2		[3.0–8.7]
PHT	87	34.8		
OSA	37	14.8		[10.9–19.8]
Atrial fibrillation	37	14.8		[10.9–19.8]
Arterial hypertension	155	62.0		[56.0–68.0]
Diabetes mellitus	46	18.4		[14.1–23.7]
Smoking	120	48.0		[42.0–54.3]
No documented exposure	67	26.80		[21.7–32.7]
Anthonisen I	103	41.2		[35.4–47.5]
II	78	31.2		[25.8–37.3]
III	69	27.6		[22.5–33.5]

Notes: Clinical characteristics table at hospital admission with proportions for categorical, and dispersion measures and central tendency for the continuous ones. Anthonisen criteria which comprise three patient-reported items: increased dyspnea, increased sputum volume, and increased sputum purulence. Type I exacerbation – all three symptoms; Type II exacerbation – two of the three clinical symptoms; Type III exacerbation – one of the three clinical symptoms.

Abbreviations: GOLD, Global Initiative for Chronic Obstructive Lung Disease; PHT, pulmonary hypertension; OSA, Obstructive Sleep Apnea Syndrome.

occurred in 5%; one-third of the patients presented pulmonary hypertension (PHT), with an average systolic pulmonary artery pressure (SPAP) of 57 mmHg measured by transthoracic echocardiography and generating pulmonary heart disease in 15% of them. PHT was more frequent among GOLD D patients (34%) when compared with GOLD A and B which had 4% and 3%, respectively ($p=0.043$) (Figure 2), as well as higher SPAP-values in GOLD D patients ($p=0.023$). In 26%, no previous exposure to smoking or combustion of biomass was identified. Regarding the clinical presentation, frequent symptoms were increased dyspnea 95% ($n=238$), increased expectoration 89% ($n=174$), and increased cough 215 (85%); fever only occurred in 42% of the cases ($n=105$). Regarding clinical tests, the mean of pCO_2 was 37.5 mmHg, HCO_3^- of 27.3 mEq/Lt. Likewise, 20.8% ($n=52$) presented anemia according to WHO criteria,¹⁷ and 7.2% ($n=18$) presented polyglobulia. On the other hand, antibiotic was used in 62% of the patients, predominantly beta-lactam antibiotics, requiring change in 13.7% of the cases due to non-responsiveness in 40% of them ($n=9$), but indicating therapeutic failure in 5.6% ($n=9$) of all patients undergoing antibiotic use. Patients admitted without previous use of home oxygen, 40% ($n=46$), were prescribed this therapy after hospital discharge and in-patient mortality was 1.2 per 100 people per year (See Table 2).

Environmental pollution

As high as 11.6% of the data were not recorded for the different stations. Regarding PM values, the median of PM_{2.5} recorded for 24 hrs was 16.4 $\mu\text{g}/\text{m}^3$ with a range of 0.9 to 49.3 $\mu\text{g}/\text{m}^3$, with 30% of the values above those established. For PM₁₀, the median was 37.7 $\mu\text{g}/\text{m}^3$ with a

range of 6.7 to 103 $\mu\text{g}/\text{m}^3$ and records higher than those of WHO for 32.3% of the cases. Likewise, the median of the daily averages of PM_{2.5} in the 48 hrs prior to the symptom onset was 20.1 $\mu\text{g}/\text{m}^3$, compared with 15 and 16.5 $\mu\text{g}/\text{m}^3$ for the day of the symptoms and the three previous initial days, differences that were statistically significant ($p<0.001$). In the case of PM₁₀, the median of these levels on the second day before the symptom onset was 46.65 $\mu\text{g}/\text{m}^3$, compared with 39 and 35.6 on the third day after symptom onset ($p<0.001$) (see Figure 3).

Additionally, during the day of symptom onset, patients with Anthonisen I or II presented 1.4 times more PM_{2.5} values greater than 25 $\mu\text{g}/\text{m}^3$ (OR 2.45, 95% CI [1.13–5.31], $p=0.019$). Likewise, high levels of PM_{2.5} were associated with increased expectoration, purulence, pleuritic pain, and antibiotic and corticosteroid use (Table 3). Regarding values for the day of consultation, no statistically significant associations were found between exposure to high levels of PM_{2.5} and symptoms, treatment, or outcomes. On the other hand, it was found that patients who were started on corticosteroids or those who were discharged with a new order for an oxygen therapy were associated with Anthonisen I and II (OR 3.33, 95% CI [1.81–6.30] and OR 2.24; % [1.20–4.17], respectively).

Discussion

The role of environmental pollution in the exacerbation of COPD is increasing. In recent years, a growing number of studies have been published with the intention of evaluating the impact of exposure to PM_{2.5} and PM₁₀ in AECOPD, evidencing deterioration of FEV₁ and higher mortality. However, for the increase in hospital admissions, the results have been inconsistent.^{12,18} The main objective of this study was to identify a variation in PM levels prior to symptoms since this aspect has been insufficiently studied. A significant finding was the evidence of change in the 48 hrs prior to the symptom onset, suggesting a possible role in the pathogenesis of AECOPD, but prospective studies are required.

In addition to the effect on the symptom onset, there was also an association with the increase in expectoration, purulence, and even pain referred to as pleuritic, which leads to a higher probability of being classified as Anthonisen I-II and, therefore, an increased use of antibiotics is needed in 70% of the patients exposed to PM_{2.5} greater than 25 $\mu\text{g}/\text{m}^3$ compared to a 60% of the antibiotic use in unexposed individuals ($p=0.016$);

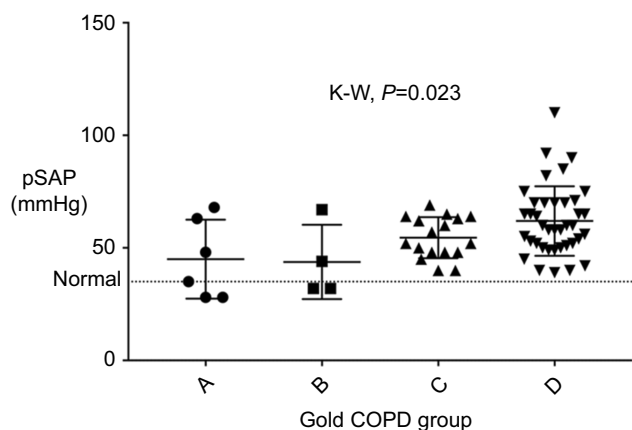


Figure 2 Frequency and distribution of pSAP-values according to COPD severity. The distribution of the value medians for systolic pressure of the pulmonary artery (pSAP) according to the severity of the COPD classified by GOLD is shown; p -values are shown for the Kruskal–Wallis statistic (KW).

Table 2 Course and clinical outcome of hospitalized patients

Characteristic	Frequency (n)	Proportion (%)
Intensive care unit	35	14.0
Ventilation	22	62.8
• NIMV	15	68.1
Vasoactive	7	31.8
Community-acquired pneumonia	52	20.8
ANTIBIOTIC USE	160	64.0
• Ampicillin/sulbactam	95	59.4
• Piperacillin/tazobactam	35	21.9
• Doxycycline	17	10.6
• Clarithromycin	4	2.5
Combination	59	36.8
• Clarithromycin	55	93.2
Modified	22	13.7
• Piperacillin/tazobactam	7	31.8
• Meropenem	2	9
Inhalation therapy		
• Beta-agonist	171	68.4
• Albuterol	150	87.7
• Salmeterol	12	7.0
Anticholinergic	223	89.2
• Ipratropium. B	203	91.0
• Tiotropium	20	8.9
Corticosteroids	121	48.4
• Intravenous	66	54.5
• Oral	102	84.2
Home oxygen	46	39.7
Mortality	6	2.4

Notes: Table with proportions of pharmacological and non-pharmacological treatments received by patients during their stay in the Hospital as well as their outcomes.
Abbreviation: NIMV, non-invasive mechanical ventilation.

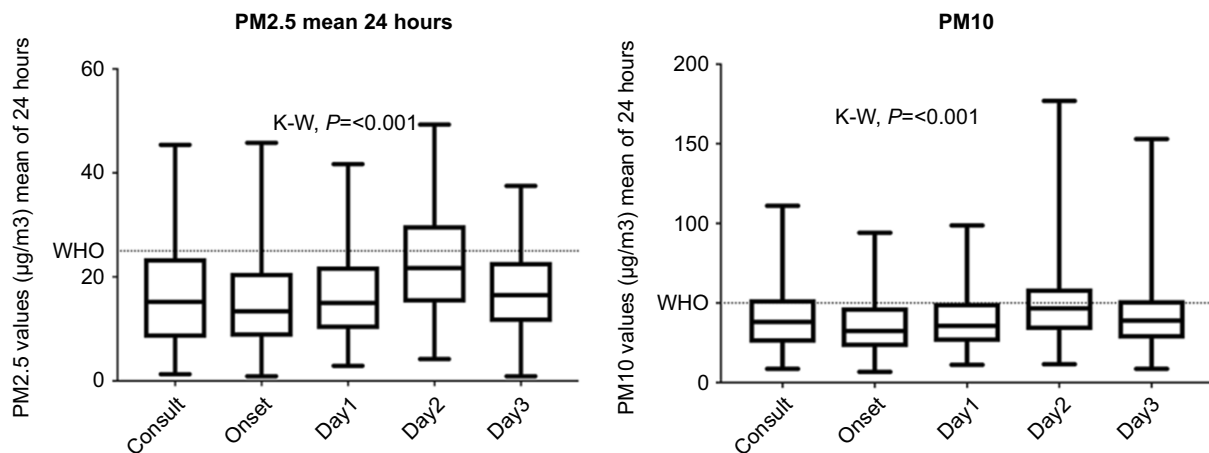


Figure 3 (A, B) PM2.5 and PM10 values, according to the day of consultation and days before the symptom onset. The medians are represented for mean values of 24 hrs for PM2.5 and PM10, according to the day of consultation, day of symptom onset (DaySx), the day before the symptom onset (Day1), two days before (Day2), and three days before the symptom onset (Day3). Dotted line: Upper limit recommended by the WHO. Kruskal–Wallis (K–W) statistic for median differences. Significant p -value<0.05.

Table 3 Association between values of PM2.5 on the day of consultation and onset of symptoms and clinical presentation and clinical course of acute exacerbation of COPD, using a multivariate model

Variable	PM 2.5						PM10						
	Consultation day			Symptom onset day			Consultation day			Symptom onset day			
	OR	95% CI OR	p	OR	95% CI OR	p	OR	95% CI OR	p	OR	95% CI OR	p	
	Lower	Upper		Lower	Upper		Lower	Upper		Lower	Upper		
Cough	2.13	0.66	0.206	1.33	0.37	4.72	0.661	1.43	0.60	3.41	0.46	3.21	0.687
Expectoration	1.59	0.71	0.256	4.74	1.02	21,90	0,046*	1.71	0.88	3.34	1.40	3.01	0.389
Purulence	0.89	0.44	0.742	6.58	2,51	17,20	0,022*	1.50	0.82	2.73	0.98	1.95	0.961
Pleuritic pain	0.86	0.32	0.767	3.62	1,27	10,37	0,016*	0.91	0.39	2.08	0.59	1.66	0.317
Pneumonia	1.15	0.50	0.747	1.21	0.47	3.10	0.692	1.08	0.52	2.21	0.98	2.23	0.955
Antibiotic	1.34	0.63	0.443	2.87	1,17	7,07	0,022*	0.63	0.34	1.16	1.73	3.64	0.151
Corticosteroids	0.49	0.23	0.057	2.62	1,07	6,44	0,035*	0.64	0.35	1.17	1.89	3.82	0.076
ICU	0.45	0.16	0.144	7.34	1,08	49,70	0,041*	1.45	0.79	2.67	1.84	3.91	0.113
Mortality	1.79	0.19	0.608	1.22	0.09	16,11	0.879	0.98	0.14	6.72	3.48	25.10	0.217

Notes: *Values that showed statistical significance using Wald statistic with a statistical significance of p<0.05. 95% CI OR: 95% Confidence Interval for the odds ratio. Table with values of OR (Exp (B)) for multivariate logistic analysis including exposure to values higher than those recommended by the WHO of PM2.5 and PM10 for the days of consultation and the day of symptom onset. The previous one, two, and three days were not recorded since there was no significant values in them. The odds in bold are statistically significant.

Abbreviation: ICU, intensive care unit.

the aforementioned is due to a collinearity effect. It should be noted that, in general, the use of antibiotics was 64%, while the one reported in previous studies was between 74.8% and 86%.¹⁹ Likewise, diagnosing pneumonia in patients with exacerbation of COPD of 20.8% is of interest, since their coexistence is associated with greater use of health resources, due to a long hospital stay, admission to the ICU and the performance of in-patient procedures. Such association has been described in 36.1% of the first episodes of exacerbation and in 25.6% of the subsequent hospitalizations, increasing even the risk of in-patient mortality (adjusted HR 1.20, 95% CI [1.17–1.24]).²⁰ The association between PM, symptomatology, and antibiotic onset could be due to the fact that PM inhibits phagocytosis and increases free radicals with a subsequent lesion of the bronchial epithelium, increased inflammatory response and production of Tumor Necrosis Factor-alpha and Interleukin-1, which perpetuate inflammation affecting mucociliary movement, favoring viral and bacterial infection.²¹ On the other hand, it was found that exposure to values higher than 25mcgr/m³ of PM2.5, by making it more likely to be classified as Anthonisen I or II, also increases the probability of using corticosteroids as a treatment, since there is an interaction between the three variables.

On the other hand, of the patients hospitalized due to AECOPD, 16% (n=35) required admission to the Intensive Care Unit, of which approximately 70% received invasive mechanical ventilation; similar to the findings of those reported in the literature.²² The median stay was 4 days, which is less than that reported in previous studies which were of 6 and 8 days.^{23,24} Regarding environmental pollution, patients admitted to the ICU were more likely to have exposure to high levels of PM2.5 compared to those who were not admitted into the ICU; however, in these patients, there was no increased requirement for ventilatory or vasoactive support. The previously reported in-patient mortality due to exacerbation of COPD was of 7.5% and 11%,^{22–25} in this study, it was 2.4%.

Another finding is the identification of patients with pulmonary tuberculosis in approximately 3.2% (n=8) of the cases. This association is noteworthy because tuberculosis can cause exacerbation, and it is also a possible etiology of COPD since the prevalence of COPD in patients with pulmonary tuberculosis is higher than that of non-infected patients, being 30.7% versus 13.9%, respectively.^{26,27} Following that train of thought, the

study of tuberculous infection in patients with AECOPD in endemic countries is recommended.

Finally, the weaknesses of the study are those typical of retrospective studies, where memory bias could affect the dates of symptom onset, especially in patients with prolonged symptoms. Likewise, the data record can omit important details. On the other hand, 11% of the PM values were not recorded in the different stations. However, estimating this loss, the sample size was increased by 20%. Influenza epidemics were not taken into account in the statistical analysis. No microbiological study was performed for any case in order to discern between infectious or non-infectious exacerbation. Medical criteria were taken into account, so the findings described here only serve to generate hypotheses and do not express a real cause-effect relationship. On the other hand, the strength of the study is the guidance toward a more clinical scope, the impact of environmental pollution, beyond hospital admissions or mortality, being able to roughly explain a possible clinical effect and, especially, an association with greater use of antibiotics. Furthermore, it shows a possible change in the levels of external contaminants prior to the symptomatic profile, a finding not previously described in the literature.

Conclusions

According to the findings of this study, patients hospitalized for AECOPD have increased levels of PM2.5 and PM10 in the 48 hrs prior to the symptomatic onset. In addition, elevated levels of PM during the day of symptom onset are associated with a higher finding of increased sputum volume, purulence, pleuritic pain, and use of antibiotics and corticosteroids. Taking into account the limitations of this study, prospective studies should be conducted to evaluate the associations presented here.

Responsibility for data

For efficacy data: Dr Jairo Morantes-Caballero has personally reviewed the efficacy data, understands the statistical methods employed for efficacy analysis, and confirms an understanding of this analysis, that the methods are clearly described, and that they are a fair way to report the results. For safety data: Dr Jairo Morantes-Caballero has personally reviewed the safety data. He understands the statistical methods employed for safety analysis and confirms that he understands this analysis, that the methods are clearly described, and that they are a fair way to report the results. For

study design: Dr Jairo Morantes-Caballero confirms that the study objectives and procedures are honestly disclosed. Moreover, he has reviewed study execution data and confirms that procedures were followed to an extent that convinces all authors, that the results are valid and generalizable to a population similar to that enrolled in this study.

Acknowledgments

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Author contributions

All authors contributed to data analysis, drafting or revising the article, gave final approval of the version to be published, and agree to be accountable for all aspects of the work.

Disclosure

The authors report no conflicts of interest in this work.

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