Relationship between blood eosinophils, clinical characteristics, and mortality in patients with COPD

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Abstract: In patients with COPD, there is controversy regarding the association of blood eosinophil (Eos) levels with 1) exacerbation frequency and 2) the effect of inhaled corticosteroids for prevention of exacerbations. To determine whether Eos define subgroups of patients exhibiting attributes of COPD clinical phenotypes, we compared clinical features and mortality rates in COPD patients from the Initiatives BPCO French cohort categorized using different thresholds of blood Eos levels. The following data were collected at inclusion: medical and smoking history, occupational exposures, dyspnea, cough and sputum production, exacerbations in the previous year, history of allergy and asthma, nasal symptoms, body mass index, St George Respiratory Questionnaire (SGRQ) total score, post-bronchodilator spirometry, comorbidities, and medications. Three-year survival between groups was compared using Kaplan–Meier analysis. Three sets of analyses were performed to compare patients with $\geq 2\%$ versus $< 2\%$, $\geq 3\%$ versus $< 3\%$, and $\geq 4\%$ versus $< 4\%$ Eos. Eos was available in 458 patients (mean age: 62 years, 72% male, mean forced expiratory volume in 1 second: 51% pred), including 235 patients with Eos $\geq 2\%$ (49%), 149 with Eos $\geq 3\%$ (33%), and 90 with Eos $\geq 4\%$ (20%). For all cutoffs, there was no difference between Eos+ and Eos– groups in univariate analyses except for diabetes and SGRQ score (more frequent and more impaired, respectively, in lower Eos categories). In particular, there was no difference in exacerbation rate, history of asthma, or three-year survival. In conclusion, regardless of the cutoff, Eos+ COPD patients exhibited no specific characteristic in terms of symptoms, lung function, exacerbation rate, and prognosis. These findings suggest that the association of higher Eos with exacerbations reported in previous studies could be population specific, which does not support generalizing the use of Eos as a biomarker for COPD phenotyping.

Keywords: COPD, eosinophils, survival, exacerbations, quality of life

Introduction

Several studies have been conducted to better understand the heterogeneity of patients with COPD and to identify different phenotypes and endotypes. More specifically, identifying clinically relevant phenotypes with specific responses to treatments is an important goal for current and future research, in order to allow proper treatment personalization based on the predicted benefit/risk ratio of each available drug class in each individual patient.¹ Biomarkers represent an important avenue of research in this area. Several studies suggest that, in patients with COPD, high sputum and blood eosinophil (Eos) counts are associated with specific clinical phenotypes defined by 1) more frequent exacerbations and 2) better response to inhaled corticosteroids (ICS) for exacerbation prevention.^{2–6} Additionally, ICS therapy has been suggested to reduce

the rate of decline in forced expiratory volume in 1 second (FEV_1) in patients with high blood Eos counts.⁷ However, in that study, higher Eos was associated with a decreased preventive effect of ICS on exacerbation risk. Contradictory data have also been found regarding the yield of Eos to predict outcomes and response to oral corticosteroids during acute exacerbations, a relation being shown in some,^{8,9} but not all, studies.¹⁰ Data on the relations between Eos count and long-term survival in COPD patients are scarce.¹¹

To determine whether Eos define subgroups of patients exhibiting attributes of COPD clinical phenotypes, possible associations between Eos and clinical characteristics and prognosis in COPD patients were further explored in a French cohort of subjects with spirometry-confirmed COPD.

Methods

Data from the Initiatives BPCO French cohort of smokers and ex-smokers (>10 pack-years), aged >40 years with spirometry-confirmed COPD ($n=1,128$ when data were extracted), were analyzed. As previously described,¹² patients from this cohort are recruited by respiratory physicians from tertiary care university hospital centers. A current main diagnosis of asthma is an exclusion criterion, but a past history of asthma in childhood or early adulthood is allowed. Patients were divided into Eos+ and Eos– groups based on blood Eos level, using several thresholds ($\geq$ and $<2\%$, 3% , and 4%). Differences in mortality rates and clinical characteristics were assessed using chi-square tests or Fisher's exact tests, as appropriate, for discrete variables, and Wilcoxon or Kruskal–Wallis tests for quantitative variables. The following data collected at inclusion were compared between groups: medical and smoking history, occupational exposures, dyspnea, cough and sputum production, exacerbations in the previous year, history of allergy and asthma, nasal symptoms, body mass index, St George Respiratory Questionnaire (SGRQ) total score, post-bronchodilator spirometry, physician-diagnosed comorbidities, and medications. Vital status was available for all patients. Overall survival was assessed using the Kaplan–Meier method.

Ethics approval

The study was approved by the Ethics Committee of Versailles, France (number: 04–479) for the protection of human beings involved in biomedical research. All patients provided written consent.

Results

Blood Eos count was available in 458 patients (mean age: 62 years, 72% male, mean FEV_1 : 51% pred), who did not

differ from those with no Eos count in terms of clinical and lung function characteristics or outcomes (not shown). The median (interquartile range [IQR]) Eos level was 1.90 (0.9–3.4)%. The population comprised 223 patients with Eos $\geq 2\%$ (49%), 149 with Eos $\geq 3\%$ (33%), and 90 with Eos $\geq 4\%$ (20%). Table 1 presents comparisons between Eos+ ($n=223$; 49%) and Eos– ($n=235$; 51%) patients using the 2% threshold. Median follow-up was 48 (33; 104) months and was not different between Eos+ and Eos– patients ($P=0.64$). All-cause mortality rate was 13% in Eos+ versus 17% in Eos– ($P=0.23$), and there was no significant difference in terms of 3-year survival (Figure 1). Diabetes was less prevalent in Eos+ patients (8% vs 17% in Eos–, $P=0.01$). Further, SGRQ total score was lower in Eos+ patients (median, 40 units vs 48 units in Eos–, $P=0.007$), reflecting better quality of life. There was no other difference between Eos+ and Eos– participants regarding clinical characteristics, lung function, comorbidities, and treatments. Eos+ patients were not more prone to exacerbations (median, 1 exacerbation/patient/year in both groups, $P=0.247$), which could not be explained by a more frequent use of ICS or history of asthma (14% vs 13.5% in Eos–, $P=0.86$) (Table 1). All analyses were repeated using thresholds of 3% and 4% to categorize patients, which provided comparable results (not shown). Even when the population was divided into four groups $<2\%$ (235 patients), $\geq 2\%$, and $<3\%$ (74 patients), $\geq 3\%$ and $<4\%$ (59 patients), and $\geq 4\%$ (90 patients), the only significant differences regarded SGRQ total score and diabetes (Table 2). There was also no significant difference in terms of 3-year survival, among these four groups (Figure 2).

Discussion

The main finding from this real-life cohort study is the lack of noticeable difference in prognosis and in most clinical and lung function features between COPD patients with higher versus lower blood Eos levels, for all tested thresholds of Eos (2%, 3%, and 4%).

In this population, half of the participants had Eos $\geq 2\%$, which is comparable to what was observed, for example, in the WISDOM study¹³ but less than in several other studies of patients with COPD identified from the general population¹⁴ or clinical trials. This finding underscores the heterogeneity of COPD patients recruited in different cohorts. Although patients with a current primary diagnosis of asthma were not recruited in the present cohort, patients with a past history of asthma represented 13% of the population but did not exhibit higher Eos levels (median: 1.9, IQR 0.9–3.4 in both patients with and without associated asthma). In addition, their clinical characteristics and outcomes were similar to

Table 1 Characteristics of 458 patients with COPD according to the 2% blood eosinophil cutoff

Variables	Eos $\geq 2\%$ (N=223)		Eos $< 2\%$ (N=235)		P-values
		Missing values		Missing values	
Sex, M/F	72.6% (162)/27.4% (61)	0	71.5% (168)/28.5% (67)	0	0.783
Age, years	62 (55–70)	0	62 (55–70)	0	0.715
BMI (kg/m ²)	25.3 (21.9–29.4)	0	24.2 (21.2–28.4)	0	0.093
Obesity (BMI > 30 kg/m ²)	22.0% (49)	0	18.3% (43)	0	0.326
Smoking habits		7		5	0.542
Former smoker	67.6% (146)		62.6% (144)		
Current smoker	29.6% (64)		34.3% (79)		
Never smoker	2.8% (6)		3.0% (7)		
Cumulative smoking (pack-years)	36.0 (24.0–54.0)	22	37.1 (22.5–52.5)	25	0.704
History of asthma	13.5% (30)	15	14.0% (33)	13	0.855
Hay fever	9.9% (22)	0	12.3% (29)	0	0.400
Eczema	7.6% (17)	0	8.1% (19)	0	0.854
Rhinitis/sinusitis	17.5% (39)	0	20.4% (48)	0	0.423
Occupational exposures	27.8% (62)	0	32.3% (76)	0	0.290
Chronic cough and sputum production	65.9% (147)	14	71.9% (169)	0	0.166
Exacerbation rate (per patient-year)	1.0 (0.0–2.0)	5	1.0 (0.0–3.0)	7	0.247
Severe (hospitalized) exacerbation rate (per patient-year)	0.0 (0.0–0.0)	5	0.0 (0.0–1.0)	7	0.174
mMRC dyspnea grade	2 (1–2)	18	2 (1–3)	21	0.211
Ischemic heart disease	11.2% (25)	0	11.5% (27)	0	0.925
Chronic heart failure	11.2% (25)	0	13.2% (31)	0	0.518
Diabetes mellitus	8.1% (18)	0	16.6% (39)	0	0.006
SGRQ total score	40 (30–56)	33	48 (32–63)	30	0.007
FEV ₁ % predicted	52 (37–68)	0	51 (34–70)	0	0.658
ICS outside fixed-dose combinations	21.5% (48)	5	23.0% (54)	10	0.709
ICS + long-acting beta-agonist	41.7% (93)	5	36.2% (85)	10	0.225
Long-acting antimuscarinic agents	30.5% (68)	5	34.0% (80)	10	0.299
Oral steroids	5.1% (12)	5	2.2% (5)	10	0.120
Follow-up duration (months)	45 (33–100)	1	51 (29–107)	1	0.641
Death rate	13.0% (29)	1	17.0% (40)	1	0.230

Notes: Data are expressed as the median (quartile 1–quartile 3) or % (n). Data were assessed using chi-square tests or Fisher's exact tests, as appropriate, for discrete variables, and Wilcoxon tests for quantitative variables.

Abbreviations: BMI, body mass index; Eos, eosinophils; FEV₁, forced expiratory volume in 1 second; ICS, inhaled corticosteroids; mMRC, modified Medical Research Council; SGRQ, St George Respiratory Questionnaire.

that of other patients.¹⁵ This may relate to a tendency of investigators to refrain from including patients with higher Eos counts (Eos was $< 4\%$ in $> 80\%$ of the population) in COPD cohorts, even when this is not an exclusion criterion.

Figure 1 Kaplan–Meier analysis for comparison of survival between COPD patients with high versus low eosinophils using the 2% blood eosinophil cutoff.

A similar hypothesis may be proposed to explain why Eos (threshold: 2%) did not predict response to ICS in the Flame trial,¹⁰ where patients with blood Eos $> 600/\text{mm}^3$ could not be recruited, while higher Eos was significantly associated with response to ICS in several post hoc analyses from previous trials.^{3,5,6} Discrepancies between study results may also relate to differences in COPD severity: for instance, our population had less severe airflow obstruction than patients from the Evaluation of COPD Longitudinally to Identify Predictive Surrogate Endpoints (ECLIPSE) cohort (mean FEV₁: 51% pred vs 44% pred). Conversely, patients from the Copenhagen City Heart Study clinical COPD cohort (n=203)⁴ had similar FEV₁ levels compared to our population. In this later population, there was a higher rate of severe exacerbations in patients with an Eos level $\geq 2\%$, while this relation was surprisingly in the opposite direction for moderate exacerbations.⁴ In our population, neither moderate nor severe (ie, hospitalized) exacerbations were different between Eos+ and Eos– groups. Another point to consider when interpreting study results is that, in many cases, Eos count was measured

Table 2 Characteristics of 458 patients with COPD according to four different blood eosinophil cutoffs: <2%, ≥2% and <3%, ≥3% and <4%, and ≥4%

Variables	<2% (N=235)		≥2% and <3% (N=74)		≥3% and <4% (N=59)		≥4% (N=90)		P-values
		Missing values		Missing values		Missing values		Missing values	
Sex, M/F	71.5% (168)/ 28.5% (67)	0	73.0% (54)/ 27.0% (20)	0	71.2% (42)/ 28.8% (17)	0	73.3% (66)/ 26.7% (24)	0	0.983
Age, years	62 (55–70)	0	61 (55–69)	0	61 (54–69)	0	63 (56–72)	0	0.712
BMI (kg/m ²)	24 (21–28)	0	25 (23–31)	0	Min 17.26 (22–29)	0	24 (20–29)	0	0.074
Obesity (BMI >30 kg/m ²)	18.3% (43)	0	27.0% (20)	0	22.0% (13)	0	17.8% (16)	0	0.372
Smoking habits		5		1		4		2	0.484
Former smoker	62.6% (144)		72.6% (53)		61.8% (34)		67.0% (59)		
Current smoker	34.3% (79)		27.4% (20)		32.7% (18)		29.5% (26)		
Never smoker	3.0% (7)		0.0% (0)		5.5% (3)		3.4% (3)		
Cumulative smoking (pack-years)	37.1 (22.5–52.5)	25	38.0 (24.0–50.0)	5	39.0 (28.5–55.0)	6	31.0 (20.0–55.5)	11	0.379
History of asthma	14.0% (33)	0	13.5% (10)	0	11.9% (7)	0	14.4% (13)	0	0.972
Hay fever	12.3% (29)	0	8.1% (6)	0	5.1% (3)	0	14.4% (13)	0	0.240
Eczema	8.1% (19)	0	4.1% (3)	0	8.5% (5)	0	10.0% (9)	0	0.531
Rhinitis/sinusitis	20.4% (48)	0	17.6% (13)	0	18.6% (11)	0	16.7% (15)	0	0.866
Occupational exposures	32.3% (76)	0	23.0% (17)	0	22.0% (13)	0	35.6% (32)	0	0.142
Chronic cough and sputum production	71.9% (169)	0	59.5% (44)	0	62.7% (37)	0	73.3% (66)	0	0.113
Exacerbation rate (per patient-year)	1	7	1	1	1	2	1	2	0.581
Severe (hospitalized) exacerbation rate (per patient-year)	0.5	7	0.3	1	0.5	2	0.4	2	0.195
mMRC dyspnea grade	2	21	2	7	2	5	2	6	0.665
Ischemic heart disease	11.5 (27)	0	6.8% (5)	0	13.6% (8)	0	13.3% (12)	0	0.533
Chronic heart failure	13.2% (31)	0	12.2% (9)	0	8.5% (5)	0	12.2% (11)	0	0.807
Diabetes mellitus	16.6% (39)	0	12.2% (9)	0	5.1% (3)	0	6.7% (6)	0	0.024
SGRQ total score	48	30	43	16	40	9	36	8	0.047
FEV ₁ % predicted	51	0	53	0	54	0	51	0	0.878
ICS outside fixed-dose combinations	23.0% (54)	0	23.0% (17)	0	15.3% (9)	0	24.4% (22)	0	0.570
ICS + long-acting beta-agonist	36.2% (85)	0	43.2% (32)	0	44.1% (26)	0	38.9% (35)	0	0.575
Long-acting antimuscarinic agents	34.0% (80)	0	28.4% (21)	0	35.6% (21)	0	28.9% (26)	0	0.655
Oral steroids	5.1% (12)	0	2.7% (2)	0	3.4% (2)	0	1.1% (1)	0	0.408
Follow-up duration (months)	51	0	46	0	53	0	45	0	0.885
Death rate	17.1 (40)	1	17.6% (13)	0	8.5% (5)	0	12.4% (11)	1	0.306

Notes: Data are expressed as the median (quartile 1–quartile 3) or % (n). Data were assessed using chi-square tests or Fisher's exact tests, as appropriate, for discrete variables, and Wilcoxon tests for quantitative variables.

Abbreviations: BMI, body mass index; Eos, eosinophils; FEV₁, forced expiratory volume in 1 second; ICS, inhaled corticosteroids; mMRC, modified Medical Research Council; SGRQ, St George Respiratory Questionnaire.

Figure 2 Kaplan–Meier analysis for comparison of survival between COPD patients with four different blood eosinophil cutoffs: <2%, ≥2% and <3%, ≥3% and <4%, and ≥4%.

only once, while it appears to vary above and below the 2% cutoff in up to half of the COPD subjects.¹⁶ Finally, it may be hypothesized that the Eos cutoff influences the results. However, the 2% threshold is the most extensively studied at present,^{2,6} and analyses with a 3% threshold (considering the Eos distribution in this cohort with a Q3 lower limit at 3.4%) did not change our conclusions. Even the analyses performed with a 4% threshold did not provide different results.

In our COPD population as in the ECLIPSE cohort,¹⁶ COPD patients with higher Eos counts (≥2%) had significantly lower SGRQ scores, suggesting less impact of COPD despite similar lung function abnormalities. The reason for this is not fully understood. The higher prevalence of diabetes in patients with lower Eos counts is also difficult to explain.

One limitation of the study is that blood Eos were available in only slightly less than half of the population included at the time of data extraction. However, patients with and without available blood Eos levels did not differ in terms of clinical and lung function characteristics or outcomes (not shown), suggesting that Eos measurement was missing at random. Another limitation is the unavailability of absolute Eos counts, which were not recorded in this cohort. However, it is unlikely that they would have provided different results given the tight concordance between analyses performed with the 2%, 3%, and 4% cutoffs.

Given the contradictory data from our study and others regarding the association between Eos and exacerbation risk, exploring the potential predictive value of other biomarkers appears necessary. However, results of a combined analysis of the SPIROMICS and COPD gene studies have been disappointing and lead the authors to conclude that, although some blood biomarkers were significantly associated with the occurrence of exacerbations, none was robust between cohorts; in addition, biomarkers added little altogether to the predictive value of clinical features for exacerbations.¹⁷

Conclusion

In this COPD cohort, Eos+ patients (regardless the cutoff chosen) exhibited no specific clinical characteristic, especially regarding symptoms, lung function, exacerbations, and, most importantly, prognosis. Health-related quality of life was better only in Eos+ patients. These findings differ from that of several other studies, which may relate to differences in patients' populations and underlines that Eos count may not be a generalizable biomarker to define clinical COPD phenotypes.

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