



**HAL**  
open science

## **Blood basophils and asthma among participants from CONSTANCES, the French population-based cohort**

Rachel Nadif, Joseph Henny, Tajidine Tsiavia, Céline Ribet, Marcel Goldberg,  
Marie Zins, Laurent Orsi, Nicolas Roche

► **To cite this version:**

Rachel Nadif, Joseph Henny, Tajidine Tsiavia, Céline Ribet, Marcel Goldberg, et al.. Blood basophils and asthma among participants from CONSTANCES, the French population-based cohort. *Allergy*, 2024, 10.1111/all.16093 . hal-04567467

**HAL Id: hal-04567467**

**<https://hal.science/hal-04567467>**

Submitted on 3 May 2024

**HAL** is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

## LETTER

# Blood basophils and asthma among participants from CONSTANCES, the French population-based cohort

To the Editor,

Along with eosinophils and neutrophils, basophils have initiator and effector functions in allergy and inflammation.<sup>1</sup> However, their role in asthma remains largely unexplored. In adults from CONSTANCES, a population-based cohort, blood inflammatory phenotypes based on eosinophils and neutrophils were associated with distinct clinical expressions of asthma.<sup>2</sup> In the present study, we hypothesized that basophils provide additional information to disentangle the heterogeneity of asthma in the same cohort ([https://www.constances.fr/index\\_EN.php](https://www.constances.fr/index_EN.php)).

At baseline, adults with data on ever-asthma and white blood cell counts were included ( $n=161,796$ , Figure S1). Four years later, 2896 participants were followed-up, including 1223 with CA (Figure S1, Tables 1 and S1). Current asthma (CA) was defined by asthma attacks, symptoms, or treatments in the last 12 months. Blood eosinophil and neutrophil cut-offs defined paucigranulocytic, neutrophilic, eosinophilic, and mixed inflammatory phenotypes.<sup>2</sup> The 75th percentile of the basophil distribution among the 161,796 participants was defined as a cut-off. Regression models adjusted (a) for age, sex, smoking status, education level, French deprivation index (Fdep), treatment, and body mass index were used (main model), and several supplementary analyses were performed (Supporting information).

Among the 161,796 participants, basophil count ranged from  $0.0027$  to  $0.43 \times 10^9/L$  and differed by asthma status (a) means:  $0.0431$ ,  $0.0438$ , and  $0.0456 \times 10^9/L$  in never-, non-current and CA respectively,  $p_{\text{trend}} < 10^{-4}$ ). Basophil count ranged from  $0.0027$  to  $0.43 \times 10^9/L$  among never-asthmatics (median [Q1–Q3] of  $0.038$  [ $0.0258$ – $0.0531$ ]  $\times 10^9/L$ ), and from  $0.0029$  to  $0.301 \times 10^9/L$  among current asthmatics ( $0.0402$  [ $0.028$ – $0.057$ ]  $\times 10^9/L$ , Figure S2, Kuiper's test for homogeneity of distributions,  $p < 10^{-4}$ ). Among participants with CA ( $n=15,206$ , Table 1), basophil count increased with age, and differed according to smoking, Fdep and treatment (Table S2). Both at baseline and at follow-up, significant correlations were found between basophil and eosinophil (a)  $r=0.30/0.39$  at baseline/follow-up) and neutrophil (a)  $r=0.19/0.20$ ) counts, and basophil count differed according to inflammatory phenotype (Table S3,  $p < 10^{-4}$ ). At follow-up, 83.7% of the participants with low ( $n=857$ ) and 63.1% of the participants with high ( $n=366$ ) basophil count at baseline were in the same basophil class ( $\kappa=0.466$  95%CI [ $0.412$ – $0.519$ ]). At baseline, high basophil count was associated with a higher risk of woken by an attack of cough, and specifically among

those with the eosinophilic phenotype (Figure 1). Results did not change whatever the cut-off used or after excluding participants with chronic bronchitis (Table S4), after additional adjustments or in sensitivity analyses (Table S5). No associations were found with the other asthma characteristics (not shown). At follow-up, participants with high basophil count at baseline were at higher risk of having eosinophilic and mixed phenotypes, CA, and woken with chest tightness (Figure 2), and associations persisted after adjustment on eosinophil count, additional adjustments or in sensitivity analyses (Table S6).

This study is the first to explore blood basophils levels in association with clinical and biological characteristics of asthma and their evolution among adults with non-severe asthma from a large population-based cohort. We acknowledge that assessing asthma by questionnaire could be considered as a weakness, and we do not exclude some bias selection due to the lost to follow-up participants. We also acknowledge that the technique used to measure basophils may be viewed as a limitation. The analytical performance of the analyzers used routinely in laboratory medicine for measuring basophils is lower than that of analyzers using flow cytometry, leading to under or over estimation of basophil counts with some analyzers due to the presence of pathological leukocytes.<sup>3</sup> The CONSTANCES population does not include, or only marginally, individuals with serious hematological diseases, and we applied a two-step procedure to be as specific as possible to capture participants with asthma. The lack of skin prick tests and immunoglobulin E measurements did not allow to investigate associations between basophils and allergic asthma. We found increased blood basophil count in asthmatics as compared to never-asthmatics and correlations between basophil and eosinophil counts that agree with two previous studies that investigated sputum basophils in asthmatic patients compared with healthy controls.<sup>4,5</sup> Increased risks of subsequent eosinophilic/mixed asthma and CA among participants with high basophil counts at baseline, suggest more active T2-oriented disease. Increased frequency of nocturnal cough and woken with chest tightness might reflect a specific clinical phenotype or poorer disease control. Accordingly, children with acute asthma and high blood basophil counts have been shown to be more likely to experience additional exacerbations during the follow-up.<sup>6</sup> In conclusion, even if these results warrant further investigations, they suggest an actual role of basophils as actors and/or potential biomarkers in asthma.

This is an open access article under the terms of the [Creative Commons Attribution-NonCommercial-NoDerivs](https://creativecommons.org/licenses/by-nc-nd/4.0/) License, which permits use and distribution in any medium, provided the original work is properly cited, the use is non-commercial and no modifications or adaptations are made.

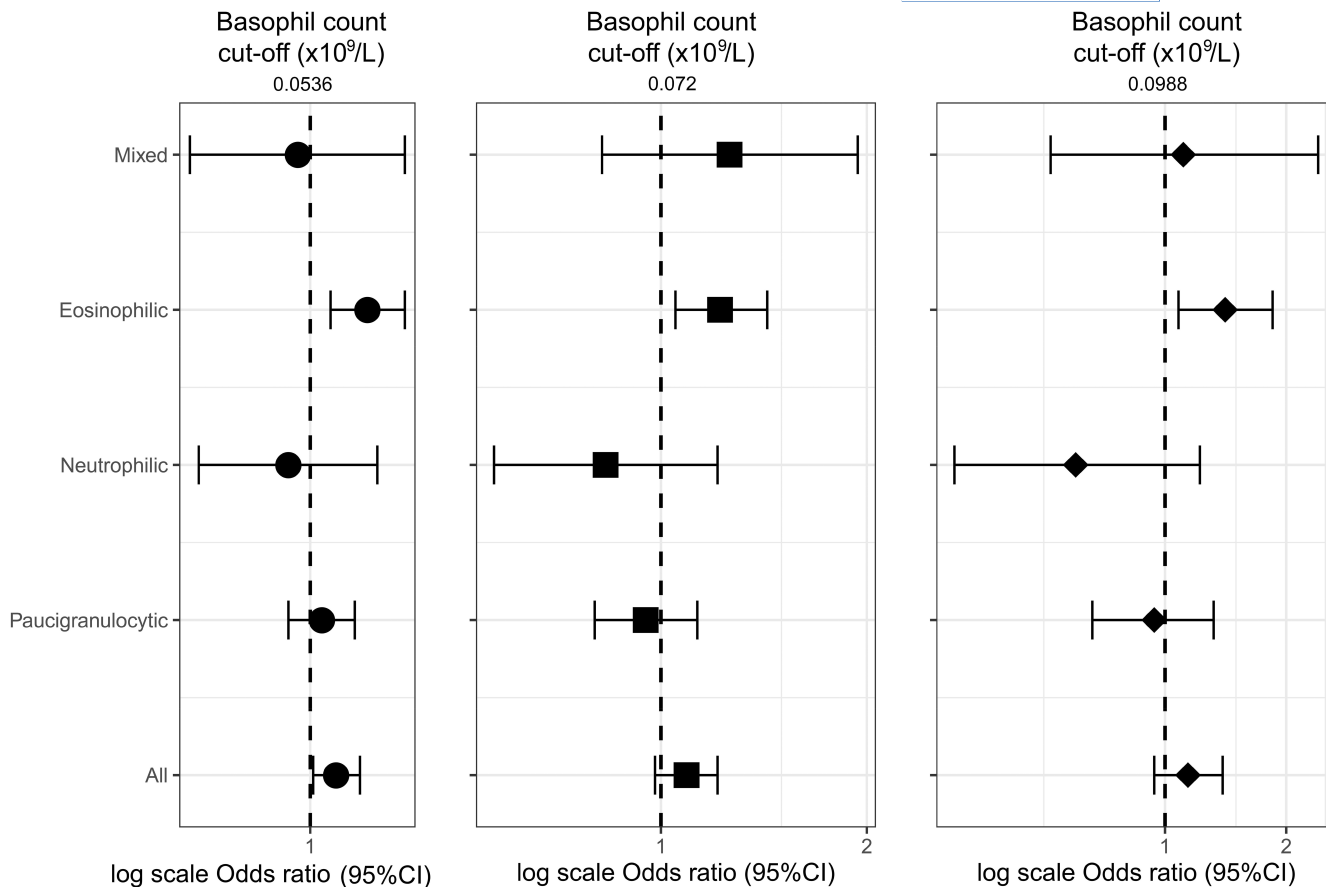
© 2024 The Authors. *Allergy* published by European Academy of Allergy and Clinical Immunology and John Wiley & Sons Ltd.

	At baseline n = 15,206	At follow-up n = 2896
Age, year, mean ± SD	44.4 ± 13.6	51.7 ± 13.1
Sex, women, n (%)	8604 (56.6)	1583 (54.7)
Smoking habits, n (%)	n = 14,702	n = 2582 <sup>a</sup>
Never smokers	6654 (45.3)	/
Ex-smokers	4843 (32.9)	/
Current smokers	3205 (21.8)	237 (9.2)
Body Mass Index (BMI), kg/m <sup>2</sup> , n (%)	n = 15,076	
<18.5	403 (2.70)	741 (25.6)
18.5–25	7417 (49.2)	1093 (37.7)
25–30	4770 (31.6)	681 (23.5)
≥30	2486 (16.5)	381 (13.2)
Education level, n (%)	n = 15,040	n = 2872
Lower than high school	3575 (23.8)	565 (19.7)
High school	2553 (17.0)	431 (15.0)
College	4168 (27.7)	834 (29.0)
University	4744 (31.5)	1042 (36.3)
FDep, n (%)	n = 15,025	n = 2,896
FDep quintile 1, less deprived	3015 (19.8)	582 (20.1)
FDep quintile 2	2984 (19.6)	577 (19.9)
FDep quintile 3	3104 (20.4)	612 (21.1)
FDep quintile 4	3048 (20.1)	545 (18.8)
FDep quintile 5	3054 (20.1)	580 (20.0)
Basophil cut ≥0.0536 × 10 <sup>9</sup> /L	4320 (28.4)	371 (30.3)
Asthma attacks (last 12 months), n (%)	n = 14,155	n = 2397
	5404 (38.2)	676 (28.2)
Asthma symptom score, n (%)	n = 14,264	n = 2761
0	2095 (14.7)	1138 (41.2)
1	4843 (34.0)	685 (24.8)
2	3269 (22.9)	431 (16.7)
3	1916 (13.4)	258 (9.3)
4	1223 (8.60)	126 (4.6)
5	918 (6.40)	93 (3.4)
Nocturnal symptoms (last 12 months), n (%)		
Woken with chest tightness	n = 14,989	n = 2836
	6546 (43.7)	831 (29.3)
Woken by an attack of shortness of breath	n = 14,929	n = 2839
	2240 (15.0)	269 (9.5)
Woken by an attack of coughing	n = 14,993	n = 2833
	6145 (41.0)	784 (27.7)
Any asthma treatment, n (%)	n = 14,698	n = 2427
	7052 (48.0)	1067 (44.0)

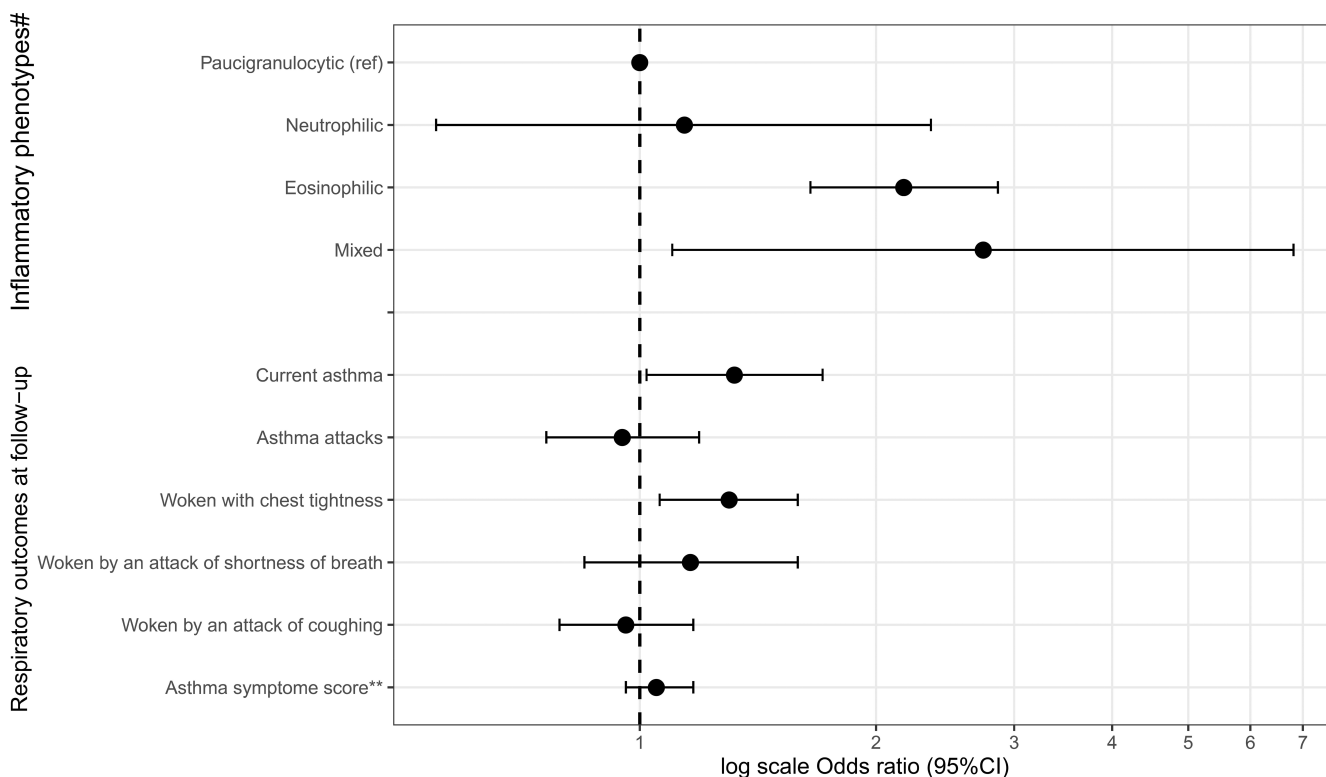
TABLE 1 Characteristics of the 15,206 participants with current asthma at baseline and of the 2286 participants at follow-up.

Abbreviations: Fdep, the French deprivation index; SD, standard deviation.

<sup>a</sup>The question has changed to "Do you currently smoke (excluding electronic cigarettes)? Yes (at least once a day) or No or occasionally".



**FIGURE 1** Associations between basophil count cut-offs and woken by an attack of coughing among participants with current asthma, and according to inflammatory profiles. Adjusted odds ratios were estimated using binary logistic models adjusted on age, sex, smoking, body mass index, education, FDep, and asthma treatment.



**FIGURE 2** Associations between basophil count cut-off of 0.0536 at baseline and asthma characteristics 4 years later among participants with current asthma. Adjusted odds ratios were estimated using logistic or negative binomial models adjusted on age, sex, smoking, body mass index, education, FDep, and asthma treatment at baseline. non ordinal polytomous regression. \*\*, Mean Score Ratio (MSR) were estimated using negative binomial regression.

## AUTHOR CONTRIBUTIONS

JH, NR, and RN designed and conducted the research; CR, MG, and MZ provided essential materials; RN, TT, and LO analyzed data and performed statistical analyses; RN wrote the manuscript and had primary responsibility for final content; all authors read, edited, and approved the final manuscript.

## ACKNOWLEDGMENTS

The authors thank the “Caisse nationale d'assurance maladie” (CNAM) and the “Centres d'examens de santé” of the French Social Security which are collecting a large part of the data, as well as the “Caisse nationale d'assurance vieillesse”, ClinSearch, Asqualab and Eurocell in charge of the data quality control. The authors thank all those who participated to the setting of the study and on the various aspects of the examinations involved: interviewers, technicians for lung function testing, coders, those involved in quality control, data and sample management and all the staffs from the inclusion centres (HPCs). They are indebted to all the participating individuals without whom the study would not have been possible. The authors also thank S Le Got, S Lemonnier, A Ozguler, C Ribet from Inserm UMS11. The authors are also grateful to the “Groupe Respiratoire CONSTANCES”: MC Delmas, O Dumas, V Giraud, Y Iwatsubo, B Leynaert, N Le Moual, R Nadif, T Perez, N. Roche, R Varraso.

## FUNDING INFORMATION


The Constances cohort study was supported and funded by the French national health insurance fund (« Caisse nationale d'assurance maladie », Cnam). Constances is an National infrastructure for biology and health (“Infrastructure nationale en biologie et santé”) and benefits from a grant from the French national agency for research (ANR-11-INBS-0002). Constances is also partly funded to a small extent by industrial companies, notably in the healthcare sector, within the framework of Public-Private Partnerships (PPP). None of these funding sources had any role in the design of the study, collection and analysis of data or decision to publish.


## CONFLICT OF INTEREST STATEMENT

The authors declare that they have no relevant conflicts of interest.

## DATA AVAILABILITY STATEMENT

The data that support the findings of this study are available on request from the corresponding author. The data are not publicly available due to privacy or ethical restrictions.

Rachel Nadif<sup>1</sup>   
Joseph Henny<sup>2</sup>  
Tajidine Tsiavia<sup>1</sup>  
Céline Ribet<sup>2</sup>

Marcel Goldberg<sup>2</sup>  
Marie Zins<sup>2,3</sup>  
Laurent Orsi<sup>1</sup>   
Nicolas Roche<sup>1,4</sup>

<sup>1</sup>Université Paris-Saclay, UVSQ, Univ. Paris-Sud, Inserm, Équipe d'Épidémiologie respiratoire intégrative, CESP, Villejuif, France

<sup>2</sup>Université de Paris Cité, Université Paris-Saclay, UVSQ, Inserm, UMS 11, Cohortes Épidémiologiques en population, Villejuif, France

<sup>3</sup>Faculty of Medicine, University of Paris, Paris, France

<sup>4</sup>APHP Centre-Université de Paris, Hôpital et Institut Cochin, Service de Pneumologie, Paris, France

## Correspondence

Rachel Nadif, Université Paris-Saclay, UVSQ, Univ. Paris-Sud, Inserm, Équipe d'Épidémiologie respiratoire intégrative, CESP, 94807 Villejuif, France.  
Email: [rachel.nadif@inserm.fr](mailto:rachel.nadif@inserm.fr)

## ORCID

Rachel Nadif  <https://orcid.org/0000-0003-4938-9339>

Laurent Orsi  <https://orcid.org/0000-0002-9332-3620>

## REFERENCES

1. Radtke D, Voehringer D. Granulocyte development, tissue recruitment, and function during allergic inflammation. *Eur J Immunol*. 2023;53(8):e2249977.
2. Tsiavia T, Henny J, Goldberg M, et al. Blood inflammatory phenotypes were associated with distinct clinical expressions of asthma in adults from a large population-based cohort. *EBioMedicine*. 2022;76:103875.
3. Amundsen EK, Henriksson CE, Holthe MR, Urdal P. Is the blood basophil count sufficiently precise, accurate, and specific?: three automated hematology instruments and flow cytometry compared. *Am J Clin Pathol*. 2012;137:86-92.
4. Suzuki Y, Wakahara K, Nishio T, Ito S, Hasegawa Y. Airway basophils are increased and activated in eosinophilic asthma. *Allergy*. 2017;72(10):1532-1539.
5. Brooks CR, van Dalen CJ, Hermans IF, Gibson PG, Simpson JL, Douwes J. Sputum basophils are increased in eosinophilic asthma compared with non-eosinophilic asthma phenotypes. *Allergy*. 2017;72(10):1583-1586.
6. Leffler J, Jones AC, Hollams EM, et al. Basophil counts in PBMC populations during childhood acute wheeze/asthma are associated with future exacerbations. *J Allergy Clin Immunol*. 2018;142(5):1639-1641.e5.

## SUPPORTING INFORMATION

Additional supporting information can be found online in the Supporting Information section at the end of this article.