

*Guillaume Lefèvre, MD, PhD<sup>a,b,c,d</sup>*

*Amélie Leurs, MD<sup>c</sup>*

*Jean-Baptiste Gibier, MD, PhD<sup>a,e</sup>*

*Matthieu Groh, MD<sup>a,f</sup>*

*Jean-Emmanuel Kahn, MD, PhD<sup>a,g</sup>*

<sup>a</sup>Centre de Référence National des Syndromes Hyperéosinophiliques (CEREO), Lille, France

<sup>b</sup>Institut d'Immunologie, Univ. Lille, CHU Lille, Lille, France

<sup>c</sup>Département de Médecine Interne et Immunologie Clinique, Centre de Référence des Maladies Auto-immunes Systémiques Rares du Nord et Nord-Ouest de France (Ce-RAINO), Univ. Lille, CHU Lille, Lille, France

<sup>d</sup>U1286—Infinite—Institute for Translational Research in Inflammation, Univ. Lille, Inserm, CHU Lille, Lille, France

<sup>e</sup>Institut de Pathologie, Centre de Biologie Pathologie, Univ. Lille, CHU Lille, Lille, France

<sup>f</sup>Département de Médecine Interne, Hôpital Foch, Suresnes, France

<sup>g</sup>Service de Médecine Interne, Assistance Publique-Hôpitaux de Paris, Hôpital Ambroise Paré, Boulogne Billancourt, France.

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Corresponding author: Guillaume Lefèvre, MD, PhD, Institut d'Immunologie, Centre de Biologie-Pathologie-Génétique, CEREO, Centre de Référence Maladies Rares des Syndromes Hyperéosinophiliques, CHRU Lille, F-59037 Lille cedex, France. E-mail: guillaume.lefeuvre@chru-lille.fr.

## REFERENCES

- Nasser M, Thivolet-Béjui F, Cottin V. Idiopathic non-necrotizing eosinophilic vasculitis limited to the lung: part of a complex spectrum. *J Allergy Clin Immunol Pract* 2020;8:2454-5.
- Nasser M, Thivolet-Béjui F, Sève P, Cottin V. Lung-limited or lung-dominant variant of eosinophilic granulomatosis with polyangiitis. *J Allergy Clin Immunol Pract* 2020;8:2092-5.
- Lefèvre G, Leurs A, Gibier J-B, Copin M-C, Staumont-Sallé D, Dezoteux F, et al. "Idiopathic eosinophilic vasculitis": another side of hypereosinophilic syndrome? A comprehensive analysis of 117 cases in asthma-free patients. *J Allergy Clin Immunol Pract* 2020;8:1329-40.e3.
- Comarmond C, Pagnoux C, Khellaf M, Cordier J-F, Hamidou M, Viallard J-F, et al. Eosinophilic granulomatosis with polyangiitis (Churg-Strauss): clinical characteristics and long-term followup of the 383 patients enrolled in the French Vasculitis Study Group cohort. *Arthritis Rheum* 2013;65:270-81.
- Leurs A, Chenivesse C, Lopez B, Gibier J-B, Clément G, Groh M, et al. C-reactive protein as a diagnostic tool in differential diagnosis of hypereosinophilic syndrome and ANCA-negative eosinophilic granulomatosis with polyangiitis. *J Allergy Clin Immunol Pract* 2019;7:1347-51.

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## Analyzing environmental control studies by the achieved decrease in exposure



### To the Editor:

In his recent editorial, Eggleston<sup>1</sup> discussed the limitations of analyzing environmental control trials by only comparing the active intervention group and the control group—where the control group, knowing that it is in a study concerning a potentially harmful environmental substance, also decreases its exposure—and he noted that information is to be gained by also sorting data according to the actual decrease in allergen achieved. Specifically, a study of mouse-allergen remediation,<sup>2</sup> in which

there were similar reductions in household allergen exposure and asthma activity in both the active and control groups, showed decreased disease activity when the data were sorted by the decrease in the final allergen level, with each 50% decrease in bedroom floor mouse allergen level associated with a further reduction in asthma symptoms, beta-agonist use, and emergency department visits for asthma. A recent follow-up of that study cohort<sup>3</sup> also showed that a reduction in mouse allergen exposure by 75% or more, whether in the active remediation group or the control group, was associated with a greater increase over 1 year in prebronchodilator forced expiratory volume in 1 second and in pre- and postbronchodilator forced expiratory flow at 25% to 75% of forced vital capacity.

It should be noted that this approach has also been applied to dust mite allergens. In his comprehensive book *Dust Mites*, Coll-off<sup>4</sup> revisited the Cochrane meta-analysis on house dust mite control measures for asthma.<sup>5</sup> That meta-analysis, comparing all combined treatment groups with all combined control groups, had concluded that there were no statistically significant differences in number of patients improved, asthma symptom scores, or medication usage. However, the studies in that meta-analysis used different methods, of differing effectiveness, in their effort to decrease mite allergen exposure. When Colloff separated those studies that showed clinical improvement from those that did not, he found that only in the former had there been a significant decrease in allergen levels in the active group compared with the controls.

Allergen avoidance studies can thus yield 2 distinct types of information: the effectiveness of measures to reduce allergen levels, which can be obtained by comparing active and control groups, and the clinical effects of such reduced allergen exposure, which can be obtained by comparing those with decreased exposure with those without such a decrease. Environmental control studies should be looked at with both questions in mind.

*Jeffrey D. Miller, MD<sup>a,b</sup>*

<sup>a</sup>Mission: Allergy, Inc., Hawleyville, Conn

<sup>b</sup>Department of Pediatrics, New York Medical College, Valhalla, NY.

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Corresponding author: Jeffrey D. Miller, MD, Mission: Allergy, Inc., 28 Hawleyville Rd., Hawleyville, CT 06877. E-mail: [JeffreyMillerMD@comcast.net](mailto:JeffreyMillerMD@comcast.net).

## REFERENCES

- Eggleston PE. A new approach to environmental control trials. *J Allergy Clin Immunol Pract* 2020;8:603-4.
- Matsui EC, Perzanowski M, Peng R, Wise RA, Balcer-Whaley S, Newman M, et al. Effect of an integrated pest management intervention on asthma symptoms among mouse-sensitized children and adolescents with asthma: a randomized clinical trial. *JAMA* 2017;317:1027-36.
- Grant T, Phipatanakul W, Perzanowski M, Balcer-Whaley S, Peng R, Curtin-Brosnan J, et al. Reduction in mouse allergen exposure is associated with greater lung function growth. *J Allergy Clin Immunol* 2020;145:646-53.
- Colloff M. *Dust Mites*. The Netherlands: Springer; 2009391-6.
- Götzsche PC, Johansen HK. House dust mite control measures for asthma. *Cochrane Database Syst Rev* 2008;(2):CD001187.

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