



Testing bronchodilator responsiveness

To the Editor:

The recent paper in the European Respiratory Journal by Janson et al. [1] on testing bronchodilator responsiveness suggests that it has no value in distinguishing asthma from COPD. The authors correctly state that "there are many different ways of defining bronchodilator reversibility." However, they do not then mention any of them other than using the change in forced expiratory volume in 1 s (FEV₁) standardised by the start value. This can lead to a sex and size bias in assessing bronchodilator response [2]. One method, that was recommended by the European Respiratory Society many years ago [3], standardises the change in FEV₁ by the subject's predicted value and not their start value. Using this method it has been found that a change in FEV₁ of 8% of predicted or more due to a bronchodilator was associated with a survival advantage [2]. This approach avoided all the pitfalls around the clinical diagnosis of COPD versus asthma. Previously it has been found that a change in FEV₁ of 4% of predicted in COPD patients was associated with subjects being able to appreciate that their breathlessness was improved [4].

Before the respiratory community dismisses testing bronchodilator responsiveness based on the evidence of Janson *et al.* [1], it needs to look at expressing any change due to a bronchodilator as a percent of the subject's predicted value, as this has been shown to be free from potential sex and size bias and is better at distinguishing important clinical end-points.

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A paper in the *ERJ* suggests that testing bronchodilator responsiveness is not helpful but the authors have not used the most appropriate method http://bit.ly/2MJR3KM

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From the authors:

We thank M.R. Miller for his comments on our paper regarding bronchodilator reversibility in asthma and COPD [1]. We agree that it is important to look at different ways of defining bronchodilator reversibility. In our analysis, we investigated both flow-related bronchodilator reversibility, defined by the change in forced expiratory volume in 1 s (FEV₁), and volume-related bronchodilator reversibility, defined

by the change in forced vital capacity. We also looked at both the change in lung function parameters expressed as percent of the baseline value and the change in ${\rm FEV_1}$ standardised by the subject's predicted value. The latter was evaluated to control for the sex, age and height dependency of lung function. The results when reversibility was expressed as percent of the predicted value (in supplementary tables E3 and E4) [1] were the same as when reversibility was expressed as percent of the baseline value. Our interpretation was therefore that, in the present study, neither flow-related nor volume-related bronchodilator reversibility were independently associated with the symptom burden, health status or dyspnoea in the COPD population.

It should be noted that our study was population-based and thus it may better reflect real life conditions, such as those encountered in general practice. However, we agree that cohort studies on patients, which include a higher number of subjects with a severe COPD, could yield different results that may be more applicable to decision-making in specialist practice. Furthermore, as our analysis was cross-sectional, we could not assess a possible association between bronchodilator responsiveness and prognosis over time. We therefore agree with M.R. Miller that further studies are needed before the respiratory community can dismiss testing of bronchodilator responsiveness in COPD.

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Neither flow-related nor volume-related bronchodilator reversibility were independently associated with the symptom burden, health status or dyspnoea in the COPD population http://bit.ly/2rigD1r

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Reference

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