



Dynamic compliance and reactance in older non-smokers with asthma and fixed airflow obstruction

Timothy Durack^{1,2,3}, David G. Chapman^{1,2,4}, Sandra Rutting^{1,2,5}, Cindy Thamrin^{2,3},
Gregory G. King^{1,2,3,5} and Katrina O. Tonga^{1,2,3,6,7}

¹The Department of Respiratory Medicine, Royal North Shore Hospital, St Leonards, Australia. ²Airway Physiology and Imaging Group and The Woolcock Emphysema Centre, The Woolcock Institute of Medical Research, Glebe, Australia. ³Faculty of Medicine and Health, The University of Sydney, Sydney, Australia. ⁴Discipline of Medical Sciences, University of Technology Sydney, Broadway, Australia. ⁵NHMRC Centre of Excellence in Severe Asthma, New Lambton Heights, Australia. ⁶The Department of Thoracic and Lung Transplant Medicine, St Vincent's Hospital, Darlinghurst, Australia. ⁷Faculty of Medicine, St Vincent's Clinical School, The University of New South Wales, Sydney, Australia.

Katrina Tonga (katrina.tonga@sydney.edu.au)



Shareable abstract (@ERSpublications)

In older non-smokers with asthma and fixed airflow obstruction, X_5 measured by oscillometry reflects dynamic rather than static compliance. Distinguishing between dynamic and static compliance is important as they are due to different factors. <https://bit.ly/3th0uEr>

Cite this article as: Durack T, Chapman DG, Rutting S, *et al.* Dynamic compliance and reactance in older non-smokers with asthma and fixed airflow obstruction. *Eur Respir J* 2021; 58: 2004400 [DOI: 10.1183/13993003.04400-2020].

This single-page version can be shared freely online.

Copyright ©The authors 2021. For reproduction rights and permissions contact permissions@ersnet.org

Received: 3 Dec 2020
Accepted: 5 April 2021

To the Editor:

In asthma, abnormal mechanical properties of the airways and lung tissue leads to airway narrowing and changes in ventilation distribution [1]. Ventilation distribution is determined by the variation of time constants [2], the product of resistance and static compliance of individual lung units. Ventilation distribution is heterogeneous in healthy lungs, but even more so in airway diseases, including asthma [3]. This is because time constants are often even more heterogeneous in disease due to changes in resistances and compliances [3, 4]. Lung compliance measured under dynamic conditions, *e.g.* during breathing (dynamic compliance, or C_{dyn}), is sensitive to these heterogeneities in time constants. C_{dyn} decreases relative to static compliance (C_{stat}) with increasing ventilation heterogeneity [2, 3, 5], due to diversion of ventilation from lung units with longer to those with shorter time constants [6]. Hence C_{dyn} is a measure of lung function in relation to ventilation heterogeneity under tidal breathing conditions, that complements spirometry. However, C_{dyn} is not used clinically because it requires invasive oesophageal pressure measurements.

Copyright ©The authors 2021. For reproduction rights and permissions contact permissions@ersnet.org