Chronic obstructive pulmonary disease

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Competing interests None.

Ethics approval Institutional Review Board of University of Pittsburgh, Partners Health Care (Boston), participating NETT centres, Boston VA, Norway, Costa Rica, and Lovelace Respiratory Institute.

Contributors Manuscript preparation: JMB, JCC; data analysis and study design: JMB, YT, SB, TJM, SB, NB, JPZ, MES-O, LA, MHC, BH, AAL, FJ, EF, SD, EKS, JCC; data collection: YT, TJM, MES, LA, AAL, PB, AG, WHA, DAL, EKS, JCC; statistical analysis: JMB, KH, MHC.

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The increasing importance of innate antimicrobials

The protein short palate, lung and nasal epithelium clone 1 (SPLUNC1) is found in airway epithelium. It is known to have reduced expression in chronic airways diseases and in smokers. This study aimed to demonstrate its antimicrobial properties against *Mycoplasma pneumoniae* infection. The investigators compared the inflammatory and antibacterial responses to *M pneumoniae* infection in transgenic mice deficient in expressing this protein with the responses in mice overexpressing the protein.

The overall results showed SPLUNC1 has antibacterial effects by inhibiting bacterial adherence proteins, thereby inhibiting M pneumoniae growth. Following M pneumoniae infection, a reduction in tissue inflammation and increase in neutrophil elastase production was seen in those mice with expression of SPLUNC1. Neutrophil elastase is important in relation to infection and was shown to reduce M pneumoniae growth when incubated with human sputum neutrophil elastase.

This study shows the potential antibacterial and immunomodulatory functions of SPLUNC1, which may help in the development of novel treatments for chronic airway diseases. In an ever-increasing climate of drug resistance, it emphasises the importance of focusing on host endogenous antimicrobial responses.

► **Fabienne G**, Di YP, Smith SK. SPLUNC1 promotes lung innate defense against mycoplasma pneumoniae in mice. *Am J Path* 2011:178:2159—67.

A Loughnan

Correspondence to A Loughnan, Homerton University Hospital, London, UK; alice.loughnan@homerton.nhs.uk

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