

LETTERS

Interstitial lung disease guideline

It is a pity that so many eminent societies have sponsored, and *Thorax* has published, a supplement entitled "Guidelines on Interstitial Lung Disease"¹ which is incomplete because no mention is made of children. Interstitial lung disease is a problem at all ages.²⁻⁴ Indeed, genetic disorders such as surfactant protein C deficiency are relevant in adults and children.⁵ The supplement should have been entitled "Guidelines on Interstitial Disease in Adults" or, far better, brought together specialists in interstitial lung disease across all ages to achieve the truly comprehensive guideline that the present title erroneously implies.

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Pulmonary rehabilitation and interstitial lung disease

The recent guideline on interstitial lung disease (ILD)¹ has a welcome emphasis on best supportive care, including pulmonary rehabilitation. However, we were disappointed that the guideline states that "... there are no randomised controlled trials of pulmonary rehabilitation" and, as a result, ascribes a low level of evidence (C) to this intervention.

As the authors indicate, the guideline was developed during a time of rapid change and growth in the body of scientific evidence pertaining to management of ILD. Pulmonary rehabilitation is no exception. Last year we published a randomised controlled trial of exercise training for ILD in this journal, which demonstrated short-term improvements in dyspnoea and exercise tolerance.² The gain in exercise tolerance

was smaller than previously reported in chronic obstructive pulmonary disease, but was accompanied by improvements in quality of life. Also in 2008, Nishiyama and colleagues³ reported similar findings in a randomised controlled trial of patients with idiopathic pulmonary fibrosis who were diagnosed according to the consensus statement. These findings have since been synthesised in a meta-analysis.⁴

The guideline will be an important aid to diagnosis and management for people with ILD across many settings and countries. However, as the authors point out, there are few data on which to base recommendations in many areas. We suggest that pulmonary rehabilitation is an area where recent evidence may be helpful. Although the benefits attributable to pulmonary rehabilitation may be small and short-lived, there are few treatments which have successfully impacted on symptoms and quality of life in this patient group. We would hope that the growing evidence pertaining to pulmonary rehabilitation for ILD might be included in future editions of this document.

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Authors' reply

Thank you for this concise and helpful statement. Plainly, there will be much more to say on this question when the guidelines are eventually revised.

As you may be aware, guideline statements must be approved (in this case by the BTS Standards of Care Committee) and a further time period is then needed for preparation of a guideline supplement. The two studies to which you refer appeared only a month or two before the final publication of the guideline document and their exclusion from consideration was unavoidable. The BTS guideline group had concluded their deliberations very much earlier. Post hoc changes in guideline statements

cannot be made by individuals at the proof reading stage.

However, this does highlight a problem with guideline statements: recommendations can be dated within a matter of months. Interstitial lung disease is currently changing rapidly as a speciality and, as the BTS guidelines may not be revised for a further 10 years, there is a strong case for brief updates every 2 years in which changes to the evidence base are summarised. This possibility will be explored.

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Statins and cancer in patients with asthma

Imamura and colleagues¹ found that pravastatin attenuated allergic airway inflammation through suppression of interleukin 17 in the lungs of ovalbumin-sensitised mice. However, in the accompanying editorial,² Rubin pointed out that, in clinical practice, it is unlikely that adding a statin to an appropriate dose of inhaled corticosteroids might provide any additional benefit for patients with asthma, highlighting that in this setting statin therapy can represent a "snake oil panacea". We concur with Rubin,² and further suggest that statin drugs might actually be harmful in patients with asthma.

In healthy individuals, immune responses to allergens include a dominant regulatory element. There is mounting evidence that the function of regulatory T cells (Tregs) may be defective in patients with allergy and asthma.³ Indeed, as Imamura and colleagues reported,¹ there is a reciprocal developmental pathway for the generation of pathogenic Th17 cells and protective Tregs in the immune system, depending on the state of the innate immune system.

On the other hand, some of the well-known immunomodulatory effects of statins are mediated through an increase in the peripheral numbers and functionality of Tregs⁴ by the induction of the transcription factor forkhead box P3. However, an increase in Treg numbers and functionality may impair the host antitumour immunity via the suppression of tumour-specific effector T cell responses and the development of immune tolerance to neoplastic cells.⁴

Interestingly, epidemiological evidence suggests that a history of allergy is associated with a decreased overall risk of cancer.⁵ It is plausible that the defective function of Tregs in subjects with allergic disease could reduce the cancer risk by