LETTERS TO THE EDITOR

Inhibition of T cell proliferation by human alveolar macrophages

The demonstration by Upham et al. that human alveolar macrophages selectively inhibit proliferation of T cells by secretion of unidentified effector molecules raises the question as to whether pathological processes in the lung characterised by extensive macrophage recruitment or activation can have a systemic effect on T cell development.

It has been shown in several studies that patients with pulmonary tuberculosis who are not infected by HIV often show a lymphopenia principally affecting the CD4+ T cells. This appears to be a transient phenomenon as the CD4+ count reverts to normal after successful therapy. A similar transient CD4+ lymphopenia has also been observed after antigenic bronchial provocation in asthmatic subjects.

It would be of great interest to determine whether this systemic phenomenon is due to the same mechanism as the one described by Upham et al., whether it accounts (at least in part) for the so-called "idiopathic CD4+ T lymphopenia syndrome," and whether it affects the balance between Th1 and Th2 cells which may be critical to the pathogenesis of both asthma and tuberculosis.

JOHN M GRANGE
Imperial College School of Medicine, Duchess Street, London SW3 6LY, UK

Adenosine and adenosine antagonist in asthma

I read with interest the excellent update on adenosine by Polosa and Holgate.1 An important use of this challenge agent is demonstrated and adenosine antagonism as a potential treatment for asthma is revisited. However, the role of adenosine as a mediator of asthma is somewhat inconsistent with several functional observations.

Besides the fact that adenosine has dual effects in many systems, data are available particularly involving the pharmacology of enprofylline (3-methyl xanthine)—which suggest that the therapeutic efficacy of theophylline (1,3-dimethyl xanthine) in asthma may not reflect adenosine antagonism. This latter aspect is significant because theophylline, at therapeutic concentrations, effectively antagonizes adenosine (at receptors and functionally in vivo).

Qualitatively different from theophylline, enprofylline does not antagonise the physiological/pathophysiological actions of adenosine yet enprofylline and theophylline share several pharmacological actions including cardiac stimulation, microvascular anti-exudative activity, and a range of smooth muscle relaxant effects although enprofylline is consistently about three times more potent than theophylline.2 Equally, enprofylline is about three times more potent than theophylline in asthma as a bronchodilator,3 as an inhibitor of histamine-induced bronchoconstriction,4 as an inhibiting effect on mast cell release,5 and in maintenance therapy.6 Indeed, it is only under artificial conditions when asthmatic subjects inhale adenosine that theophylline provides greater protection than enprofylline.

In contrast to its efficacy in the treatment of asthma, enprofylline lacks several well known clinical effects of theophylline such as diuretic activity, CNS arousal effects, free fatty acid releasing effects, and gastric secretory effects.7 This distinct human pharmacology is evidence for the clinically effective adenosine antagonism of theophylline and indicates that enprofylline tonically suppresses volume and acidity of gastric secretion, natriuresis, and inotropy of cardiac stimulation, microvascular anti-exudative activity, and a range of smooth muscle relaxant effects although enprofylline is consistently about three times more potent than theophylline.2 Equally, enprofylline is about three times more potent than theophylline in asthma as a bronchodilator,3 as an inhibitor of histamine-induced bronchoconstriction,4 as an inhibiting effect on mast cell release,5 and in maintenance therapy.6 Indeed, it is only under artificial conditions when asthmatic subjects inhale adenosine that theophylline provides greater protection than enprofylline.

If the clinical efficacy of the xanthines in asthma cannot be explained by adenosine antagonism, phosphodiesterase inhibition may offer an alternative explanation but, unfortunately, there are also doubts about this.8—hence the widely promoted non-xanthine phosphodiesterase IV inhibitors cannot rely on theophylline for any predictable clinical efficacy. Perhaps both adenosine antagonism and phosphodiesterase inhibition are examples of how theoretically attractive mechanisms may prevent unbiased exploration of truly important in vivo modes of action of anti-asthma drugs.

Incidentally, enprofylline was discovered by unexpected observations in complex biosystems.9 Such exploratory in vivo work, if allowed, will continue to be a source of novel drugs; when successful, one should not be surprised to learn that the discovered class of drug was not predicted by reductionist research paradigms. The new efficacious compounds may thus unveil novel mechanisms—for example, omeprazole and the acid pump—or, as with the experimental drug enprofylline, the new properties will seriously question the therapeutic relevance of a widely held mechanism.

CARL PERSSON
Department of Clinical Pharmacology, University Hospital of Lund, S-22185 Lund, Sweden

AUTHORS' REPLY

Studies in a variety of species5,8 indicate that a substantial proportion of the recirculating T cell population is sequestered for significant periods during transit through the lung vascular bed, and many of these cells extravasate and move into the lung interstitium. The initial trapping of T cells in transit is due, at least in part, to local endothelial expression of inflammation associated molecules such as ICAM-1. This process is selective for recently activated T cells, and T lymphoblasts generated at immunoinflammatory foci distal to the lung readily enter the lung and therefore contribute to the local immunological milieu.

Moreover, the extremely large size of this peripheral lung T cell population indicates that this is a physiological process which operates continuously in normal individuals, and it is conceivable (in our view highly likely) that it is further amplified


Reasonably digestible reviews of recent clinical trials in the management of lung cancer are rare and, in general terms, this 11 chapter book is welcome. The emphasis here is on the use of chemotherapy and radiotherapy in the management of lung cancer, with eight of the 11 chapters considering these aspects, and the other three are concerned with chemoprevention, palliative medicine, and molecular biology.

Nine of the 11 authors are from the USA. The chapters take the form of a traditional review and are reasonably well set out with an average of about 50 references for each topic. The strengths of the book are the comprehensive assessment of new drug therapies, with separate chapters for paclitaxel/carboplatin, gemcitabine, and docetaxel in non-small cell lung cancer, and a separate chapter on novel drugs for small cell lung cancer, including the topoisomerase-1 inhibitors, carboplatin, and the taxanes.

Sadly, the volume lacks an adequate introduction by the Editor, which would have been useful if it had been able to point out the “major messages” from each of the chapters—for example, by bringing out the importance of the recent meta-analysis of trials of prophylactic cranial irradiation in responding small cell lung cancer, or the superiority of standard chemotherapy regimens over low dose oral etoposide in this disease. Surgery gets no mention at all, and nor does endobronchial therapy. This is a pity since there have been major advances in our understanding of the role of endobronchial treatments, and the literature, particularly that relating to brachytherapy, is badly in need of review. Likewise, I found the chapter on palliative medicine disappointing with no consideration of psychosocial problems or some important major physical symptoms such as cough and pleural disease, and a misplaced discussion here of the meta-analysis of chemotherapy in advanced lung cancer.

The best chapter, in my view, was that by Wagner on radiation therapy in small cell lung cancer which was a well set out discussion of the attempts that have been made to optimise local control by altering the timing and fractionation of thoracic radiotherapy, together with an up to date discussion on prophylactic cranial irradiation. The book is just about up to date enough to include the results of the important MRC study on continuous hyperfractionated accelerated radiotherapy for non-small cell disease (CHART), which must now be considered as one of the few studies on radiotherapy recently to have shown an improvement in survival compared with local control.

This book will not appeal to the non-specialist, though it would be a useful starting point for doctors or groups who want an up to date background account as a preliminary to designing their own studies or choosing a pattern of management for their patients. Inevitably, in a fast moving field such as the assessment of new and existing therapies for lung cancer, a book like this will rapidly become out of date and, as with guidelines, I would estimate that “an update of this update” will probably be needed within a couple of years.—MFM


When asked to review this book last year I devised my own “sightindex” to assess its worth to me as a respiratory physician with the responsibility for a tuberculosis service. First a sit down to get acquainted with each other. It is big, attractive, well laid out and easy to grasp, but somewhat let down by the index. The first sections on history and epidemiology are as interesting as a British Medical Journal Christmas issue, but potentially more expensive to read in the bath—I enjoyed them. The 28 colour photos are cheerful and useful, except three brown-on-brown endoperoxidase stains which make the eyelids droop, presumably the reason why photograph 23 of the eye is presented upside down.

During the year it sat on my shelf four colleagues borrowed the book and said it was very useful. My personal “sightindex” was 10, nearly equalling my most popular text book. I scored the usefulness of each sightsection on a scale from O (no value) to 3 (excellent). I searched for M szulgai (useful about soft tissue infection and antibiotic sensitivity but little regarding lung infection, scores 2/3), how to give BCG in the thigh (nothing, 0/3), management of multi-drug resistant tuberculosis (excellent, 3/3), renal tuberculosis (excellent review and helpful discussion of the role of nephrectomy and oral antituberculous, 3/5) M boris (good review and references, 3/3); management of BCG complications (subcutaneous abscess not mentioned, vague advice on therapy, 0/3), medical and surgical management of tuberculosis empyema (most useful, 3/5); TB in prisons—some local institutional advice or “tax supported exposure chambers for tuberculosis” (useful but review limited to problem in USA, some unrealistic recommendations, 2/3); advice on standard drug therapy (useful summary of ATS recommendations but not of drug dosages, 2/3); and directly observed therapy (again no summary of intermittent dosages, 2/3).

So the borrow index was 4, my “sightindex” was 10, and usefulness score 66%. That’s pretty good (BCG apart), and I am very pleased to have it available. So should you, if you have an interest in or responsibility for a tuberculosis service.—JTM

New Drugs for Asthma

A two-day conference on “New Drugs for Asthma” will be held at the National Heart and Lung Institute, Imperial College School of Medicine, London on 16 and 17 June 1998. For further details please contact Caroline Elliott at IBC UK Conferences Ltd, Biomedical Division, Gilmoora House, 57–61 Mor-timer Street, London W1N 8JX, UK. Tel: +44 (0)171 453 2701; Fax: +44 (0)171 631 3214; email caroline.elliott@ibcuk.co.uk.

7 Xiaow Jin. PhD Dissertation, University of Vir-ginia School of Medicine, 1996.
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CARL PERSSON

Thorax 1998 53: 437
doi: 10.1136/thx.53.5.437a

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