Preventing hospital admissions for COPD: role of physical activity

M D L Morgan

There is increasing evidence that non-pharmacological interventions including physical activity may prevent hospital admissions for COPD.

A hospital admission or readmission for an exacerbation of chronic obstructive pulmonary disease (COPD) is bad news for everyone. For the patient it may signal the beginning of the terminal phase of the illness. For the health services it is a significant component of the cost of care for a condition that is increasingly burdensome. In recent years knowledge has been growing about the important role of exacerbation in the deteriorating progress of COPD. The last few years of life with COPD may be characterised by repeated episodes of illness culminating in hospital admission. Exacerbations only lead to hospital admission in about 16% of cases, but their increasing frequency is associated with declining state of health.

In UK hospitals emergency inpatient admission is the largest component of the total cost of respiratory disease to the NHS that amounts to over £2.5 billion.1 In recent years attention has focused on reducing the cost and impact of hospital admissions for COPD. The emphasis of the strategy to date has been on admission avoidance and early supported discharge schemes.2 3 These have had some success in curtailing admissions and reducing lengths of hospital stay, and are beginning to be introduced more widely into clinical practice. Contrary to popular perception, COPD is not a stable condition and there are inevitably day to day variations in both symptoms and ability to function. The periodic episodes of worsening are known as exacerbations, although a precise agreed definition is lacking. However, the meaning of the term is generally understood and a working appreciation of the impact of these events is becoming clear. Not all episodes of exacerbation lead to hospital admission, but if these can be contained or prevented by the patient or primary care services, then the burden of COPD will be reduced. A study of the factors that can prevent hospital admission in these circumstances would therefore make an important and valuable contribution.

This issue of Thorax contains the third major publication from a group in Barcelona (EFRAM) that has been examining the factors associated with hospital admission.4 This latest paper is the report of a prospective study that examines the risk factors for readmission to hospital for exacerbations of COPD. In previous publications the group have described the prevalence and relative risk of modifiable risk factors for hospital admission.5 6 The initial study population was a sample of 404 patients who were admitted to hospital in Barcelona over a period of 1 year.7 During the admission the patients completed a questionnaire and 353 later underwent spirometric tests and arterial blood gas measurements. Identification of risk factors that were potentially amenable to modification showed that lack of rehabilitation and poor inhaler technique were the most frequent associations. Other important factors were continued smoking, inadequate oxygen prescription, and lack of influenza immunisation. In the second paper the authors reported a case-control study of the factors associated with hospital admission in a subsample of the original cohort.8 The control subjects were patients with COPD who had had a previous admission but were stable at the time of the comparison. The only apparent risk factors found in 86 pairs of cases and controls were a previous history of three or more admissions, lower FEV1, and underprescription of oxygen. Perversely, continued smoking appeared to convey an advantage and inhaled corticosteroids offered no benefit. Some other potentially modifiable factors including rehabilitation could not be tested because of the very small numbers receiving it. In their conclusion the authors acknowledged that the case-control methodology had limitations in terms of selection bias and small numbers and that a prospective cohort study was required.

The latest paper describes a prospective study of the risk of readmission in a cohort of 340 patients from the original study population who were followed for over 1 year after their index admission.9 At the end of the study 63% of the patients had been readmitted and 29% had died, suggesting that previous admission is an important risk factor. Other expected risk factors were lower FEV1, and hypoxaemia. One unexpected finding was that a history of physical activity was associated with a 46% reduction in the risk of admission. The activity profile was obtained through self-reporting and no objective test of exercise capacity was made. Nevertheless, the association was very strong. Once again the authors could not test the influence of rehabilitation because the numbers were too small. Other factors weakly associated with increased risk of admission were supervision by a respiratory specialist, oral corticosteroids, and anticholinergic drugs. These latter associations are unlikely to be causal and probably reflect confounding by severity of disease.

Similar findings were reported by the British Thoracic Society and Royal College of Physicians’ audit of admission for acute exacerbation of COPD.10 In this audit of 1400 admissions two thirds of the patients had had a previous admission for COPD and one third had had a similar episode in the previous 4 months. In a follow up audit of readmission within 3 months, poor performance status was a predictor of mortality at the first admission but not readmission.11 Once again the best predictors of admission were low FEV1, and previous admission.

The pattern of hospital readmission may be affected by both patient and healthcare delivery factors. It could be that general practitioners find it easier to admit the patient to hospital than to deal with the exacerbation at home. However, high rates of readmission to hospital are common to both studies from different healthcare systems. The implication of the latest EFRAM study is that the risk of readmission to hospital for COPD can be reduced by improving spontaneous domestic activity and thereby breaking the cycle of hospital dependency. This may be true but, so far, trials of pulmonary rehabilitation that improve exercise capacity have been unable to show a reduction in hospital admission. However, they have shown a reduction in the length of stay once admitted.12 It is possible that rehabilitation may reduce hospital admissions but studies with an appropriately sensitive design have not been performed. In addition, the capacity to provide rehabilitation in most countries is so poor that it has never been testable. In the British Thoracic Society audit only 3% of the patients were recommended for rehabilitation, while in Spain only 14% of the study group had received it. An alternative explanation for the results of the study is that patients with less severe COPD simply feel better, do more, cope better, and are not admitted so frequently. It is known that physical performance, as reflected by a walking test and functional performance questionnaire, is a strong
Malignant pleural mesothelioma has become a mainstream cancer. This is partly due to the increasing incidence, but is also a result of the advances being made in its treatment. This summer the American Society of Clinical Oncology Annual Meeting plenary session included a clinical research paper on malignant pleural mesothelioma for the first time. In fact, this may well have been the first oral presentation on the disease at this important international meeting. The reason for the increased interest is that the study presented is the largest phase III randomised trial reported in malignant pleural mesothelioma. The trial, which recruited internationally and was led by researchers at the University of Chicago, showed a positive clinical benefit for an experimental arm based on a novel chemotherapy drug. But what do these data mean for respiratory physicians, oncologists and patients, and is this a definitive result? What effect, if any, does this trial result have on the ongoing UK mesothelioma trial?

The University of Chicago multicentre trial compared a combination of pemetrexed and cisplatin chemotherapy with a control arm of single agent cisplatin. Pemetrexed is a new cytotoxic drug that inhibits several folate dependent reactions that are essential for cell proliferation, hence its previous name “multitargeted antifolate”. Its primary target is thymidylate synthase, but it also inhibits folate dependent enzymes involved in purine synthesis. It is related to the existing cytotoxic drugs methotrexate, 5-fluorouracil, and raltitrexed. Phase I and II data had suggested a dose for a 21 day cycle of 500–600 mg/m² pemetrexed and 75–100 mg/m² cisplatin, with both drugs being administered on the same day. The investigators of the phase III trial chose doses of 75 mg/m² cisplatin and 500 mg/m² pemetrexed for the experimental arm. Patients randomised to the control received 75 mg/m² cisplatin. All patients were treated every 3 weeks.
A total of 472 patients with malignant pleural mesothelioma were recruited between 1998 and 2002. All had good performance status (Karnofsky score 70–100%). After an initial accrual period, four drug-related deaths from febrile neutropenia were noted (three in the experimental arm and one in the control arm). These deaths were linked to raised homocysteine levels and it was decided to give all patients folic acid supplements and vitamin B12 to counter this. This measure appeared to reduce the toxicity of the chemotherapy in both arms of the study. The researchers reported that, for the complete cohort of patients, the chemotherapy therapy significantly lengthened the time to disease progression (5.7 months vs 3.9 months; \( p=0.001 \)) and overall survival (12.1 months vs 9.3 months; \( p=0.020 \)). For patients receiving full vitamin supplementation, the overall survival for those treated with pemetrexed and cisplatin was 13.3 months compared with 10.0 months for the control patients (\( p=0.051 \)). The study organisers concluded that the combination of pemetrexed and cisplatin with folic acid and vitamin B12 should now be considered the “standard front line therapy for patients with malignant pleural mesothelioma”.

In the UK the British Thoracic Society has recently completed the pilot phase of a randomised trial of chemotherapy for patients with malignant pleural mesothelioma. The trial—known as “MESO-I”—started as a feasibility study because the investigators wanted to determine which of two quality of life instruments was more appropriate. MESO-I contained a multiple randomisation option such that patients and their oncologists could choose to be randomised between active symptom control (ASC) versus one of two chemotherapy regimens or between the chemotherapy regimens only. The chemotherapy regimens chosen were single agent vinorelbine for 12 weeks and MVP (mitomycin C, vinblastine and cisplatin) for four 21 day cycles. These regimens were chosen because they both give a response rate of approximately 20% and have proven quality of life benefit in a substantial proportion of patients.

Following completion of the pilot study, the trial has been granted full support from Cancer Research UK and the National Cancer Research Network and is now designated “MSO1”. This three arm phase III trial aims to randomise 840 patients with malignant pleural mesothelioma into one of three arms: ASC without chemotherapy; ASC with vinorelbine chemotherapy; and ASC with MVP chemotherapy. The main end points of MSO1 are overall survival, symptom palliation, quality of life, toxicity, response, and recurrence. One hundred and fifty eight patients from the pilot study who were randomised between all three arms will be included in the MSO1 analysis.

The important question is whether the MSO1 trial is still ethical in the light of the new data on pemetrexed with cisplatin. I think the answer is “yes”. The University of Chicago trial, although promising and an important step in the advancement of knowledge of mesothelioma, can be criticised. Firstly, what was the rationale for the control arm? Few physicians would recommend single agent cisplatin in a dose of 75 mg/m\(^2\)/m to a patient with mesothelioma: the response rate is likely to be low and toxicity—especially in patients with constitutional symptoms—can be appreciable. Choosing a control arm of limited efficacy may have made the pemetrexed and cisplatin combination appear more effective than it was. Indeed, from the quality of life data currently available, the symptom scores of patients treated with cisplatin 75 mg/m\(^2\)/m appeared to worsen on treatment, thus exaggerating the palliative benefit of the experimental treatment and emphasising why trials including a “no chemotherapy” arm may be appropriate.

Secondly, the investigators concluded that pemetrexed with cisplatin, folic acid and vitamin B12 should be the “standard front line therapy” for patients with malignant pleural mesothelioma. However, the data for patients given this exact combination showed that the improvement in overall survival compared with the question control arm only achieved borderline statistical significance (\( p=0.051 \)).

These criticisms weaken the argument that pemetrexed with cisplatin and vitamin supplementation should be standard treatment, although the combination is certainly an option for fitter patients. A randomised trial including a “no chemotherapy” arm remains reasonable, and the pilot phase of this UK trial has shown that patients are willing to be randomised into such a trial. The MSO1 trial, with its comprehensive set of end points, should define the role of palliative chemotherapy in malignant pleural mesothelioma. It demands our full support, as do other trials examining new treatments for this once neglected group of cancer patients.

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Author’s affiliation
J P C Steele, Department of Medical Oncology, St Bartholomew’s Hospital, London E1A 7BE, UK; jeremy.steele@bartsandthelondon.nhs.uk

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The new BTS/SIGN asthma
guidelines: where evidence leads the way

B G Higgins, J G Douglas

Announcing the publication of the new BTS/SIGN asthma
guidelines as a supplement to this issue of Thorax.
measures which many have long felt should be beneficial, such as reduction of exposure to house dust mite or avoidance of pet allergen. However, those reviewing this literature will feel that what is necessary is more conclusive evidence, be it positive or negative. This also applies to the data on alternative medical therapies where high quality studies are few. Even where there is good evidence of efficacy, recommendations are not necessarily clear cut. Immunotherapy can improve asthma but there are insufficient data to assess its value relative to conventional pharmacological treatment.

These are issues which require further research and, indeed, this is one of the great secondary benefits of producing formal evidence-based guidelines. The extensive review of the literature which is part of the process has revealed many areas where more evidence is needed before clear guidance can be given. The gaps in our knowledge are sometimes surprising. For example, despite the number of pharmacological studies carried out in asthma, we still do not know the threshold at which inhaled steroids should be introduced and we have no evidence to help decide which treatment strategy to try first at step 4. Some will regard it as a failure that there are not more grade A recommendations, but it would be better to regard this as a challenge. We have the opportunity to take this as a starting point from which to analyse the major gaps in our knowledge and develop appropriate research to address these. This process has already started with an initiative led by the Asthma Taskforce, administered by the National Asthma Campaign.

A further consideration for the future is the concept of developing a “living guideline”. This would involve a regular—probably annual—review of the literature and revision of the guideline where appropriate. It is extremely difficult to keep a guideline both up to date and yet also grounded on firmly established evidence, but the current system of major revision every few years may lean too far away from the former aim.

In the meantime we believe that this new joint BTS/SIGN guideline represents the best synthesis of available evidence and practical advice on the clinical management of asthma. Implementing the recommendations should lead to improved care for our patients but, in addition, we would be delighted if this guideline acted as a stimulus to improving the evidence base available in the future.

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Authors’ affiliations
B G Higgins, Freeman Hospital, Newcastle upon Tyne, UK

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B G Higgins and J G Douglas

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