1 Guidance

1.1 Current evidence on the safety and efficacy of lung volume reduction surgery for advanced emphysema appears adequate to support the use of this procedure provided that the normal arrangements are in place for consent, audit and clinical governance.

1.2 Clinicians wishing to use lung volume reduction surgery for advanced emphysema should ensure that patients are fully informed about the risks of the procedure and the likelihood of deterioration in the longer term. Use of NICE's information for the public is recommended.

1.3 Patient selection is important because mortality is increased in patients with the most seriously compromised lung function. NICE has issued a clinical guideline on chronic obstructive pulmonary disease.

1.4 The procedure should be undertaken by a multidisciplinary team that includes a respiratory physician, specialists in pulmonary rehabilitation
and a thoracic surgeon.

2 The procedure

2.1 Indications

2.1.1 Emphysema is a chronic lung disease. The walls of the air sacs (alveoli) in the lung weaken and disintegrate, leaving behind abnormally large air spaces that remain filled with air even when the patient breathes out. These air spaces may coalesce to form larger air-filled sacs called bullae. The surface area of the alveoli is decreased, so there is less space for the exchange of oxygen and carbon dioxide. This leads to reduced levels of oxygen in the blood. The most common symptoms of emphysema are shortness of breath (dyspnoea), coughing, fatigue and weight loss.

2.1.2 Emphysema often co-exists with chronic bronchitis. Both of these conditions may be described by the more general term of chronic obstructive pulmonary disease (COPD).

2.1.3 Treatment for COPD involves a multidisciplinary approach, which may include education, exercise, breathing retraining, smoking cessation, oral and inhaled medication, oxygen therapy, and lung transplantation. Lung volume reduction surgery may be an option for patients with severe symptoms for whom conservative treatments have proved inadequate.

2.2 Outline of the procedure

2.2.1 Lung volume reduction surgery is a palliative treatment that aims to remove the least functional part of the lungs. Computed tomography (CT) and perfusion scanning are used to identify the diseased lung tissue. The diseased part of the lung can be accessed by various techniques including median sternotomy, video-assisted thoracoscopic and thoracotomy. The first two are the most common techniques. Median sternotomy involves cutting through the sternum to open the chest. The video-assisted procedure involves making a number of small incisions in both sides of the chest to allow the insertion of instruments.
into the chest between the ribs. A thoracotomy involves making an incision between the ribs on one side of the chest and separating the ribs to access the lung.

2.2.2 The aim of the surgery is to reduce the volume of the lung. This is done by using a surgical stapling device to cut and seal the tissue, laser ablation to shrink lung volume, or a combination of both. Once the tissue has been removed, the lung is re-inflated and the chest closed.

2.3 Efficacy

2.3.1 Evidence on efficacy indicates that in certain patients lung function, exercise performance and quality of life are improved in the short term after lung volume reduction surgery. These results have been relatively consistent across study designs and were confirmed in the National Emphysema Treatment Trial, a recent large-scale randomised controlled trial comparing surgery with medical therapy.

2.3.2 The National Emphysema Treatment Trial randomised 1218 patients, of whom 580 underwent surgery. At 24 months, exercise capacity had improved in 15% (54/371) of patients in the surgery group compared with 3% (10/378) of patients in the medical group (p < 0.001). Quality of life had also improved in the surgical group (121/371) as compared with the medical group (34/378) at 24 months (33% versus 9%, p < 0.001). However, the trial found no difference in overall mortality between the two groups (0.11 deaths per person-year, risk ratio 1.01, p = 0.90). For more details, refer to the Sources of evidence section.

2.3.3 The Specialist Advisors considered that the procedure is beneficial for a select proportion of patients, but the benefit tends to decline with time.

2.4 Safety

2.4.1 The most common complication was persistent air leak from the lung. In one study of 250 patients, 45% (113/250) of patients experienced prolonged air leaks lasting more than 7 days, with 8 of these patients (3%) requiring a subsequent operation. Other complications in this series
included pneumonia 10% (24/250), in-hospital mortality 5% (12/250), myocardial infarction 2% (5/250), deep vein thrombosis 2% (4/250), small bowel obstruction 2% (6/250) and phrenic nerve injury < 1% (2/250). For more details, refer to the Sources of evidence section.

2.4.2 Complications include those that may arise from pre-existing co-morbidities as well as those that are directly due to the surgery.

2.4.3 The Specialist Advisors considered that the risks of surgery were well known. They listed the main complications as being air leaks, chest infections and respiratory failure.

2.5 Other comments

2.5.1 It was noted that endobronchial techniques are being used increasingly as an alternative to this procedure.

3 Further information

Sources of evidence

The evidence considered by the Interventional Procedures Advisory Committee is described in the overview to this guidance.

Information for patients

NICE has produced information on this procedure for patients and carers ('Understanding NICE guidance). It explains the nature of the procedure and the guidance issued by NICE, and has been written with patient consent in mind.

Update information

Minor changes since publication

January 2012: minor maintenance.
Endorsing organisation

This guidance has been endorsed by Healthcare Improvement Scotland.