Air leaks, pneumatic thorax, and chest drains

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Air leaks can be defined as any extrusion of air from normal gas-filled cavities including the upper airway, sinuses, tracheobronchial tree, and gastrointestinal (GI) tract. Clinical conditions of relevance in anaesthesia and critical care include pneumothorax, pneumomediastinum, pneumopericardium, pneumoperitoneum, and subcutaneous emphysema. This review will cover those related to the chest; of these, pneumothorax is the most common serious complication. In critically ill patients, the diagnosis of pneumothorax is often complicated by other disease processes and the limitations of bedside imaging.

Pneumothorax is the presence of air in the pleural cavity with associated lung collapse. It is classified into spontaneous (occurring without an obvious preceding event), traumatic (direct or indirect), and iatrogenic. Spontaneous is the commonest condition in general medicine and is sub-classified as: (i) primary spontaneous pneumothorax (PSP) occurring in the absence of obvious lung disease and (ii) secondary spontaneous pneumothorax (SSP) complicating a pre-existing lung disease.

Most pneumothoraces occurring during anaesthesia and in the critically ill are classified into two categories: (i) pneumothorax secondary to barotrauma and (ii) traumatic pneumothorax as a result of thoracic injury from trauma, surgery, or other interventions.

Pathophysiology

Although patients with PSP do not have clinically apparent lung disease, sub-pleural bullae are found in 76–100% of cases during video-assisted thorascopic surgery. The mechanism of bulla formation is unclear. Even among non-smokers with a history of pneumothorax, 81% have bullae. In smokers, a likely explanation is that degradation of elastic fibres in the lung occurs, induced by the smoking-related influx of neutrophils and macrophages. This degradation causes an imbalance in the protease–antiprotease and oxidant–antioxidant systems. After bullae have formed, inflammation-induced obstruction of the small airways increases alveolar pressure, resulting in an air leak into the lung interstitium. The air then moves to the hilum, causing pneumomediastinum and, when the mediastinal pressure increases, rupture of the mediastinal pleura occurs causing pneumothorax.

In patients requiring intensive care, a pneumothorax is often caused by barotrauma associated with mechanical ventilation in the presence of reduced lung compliance. The initial process in barotrauma is the production of perivascular interstitial emphysema. When the pressure gradient between the alveoli and the interstitium exceeds a critical level, alveoli rupture occurs and air enters into the interstitium. The pressure at which this occurs varies with severity of lung injury and is associated with the use of excessive tidal volumes during ventilation.

The over-distension of the non-dependent areas of the lung and rupture of the mediastinal pleura explain why anterior, medial, and sub-pulmonary pneumothoraces are more common in ARDS. Restrictive ventilatory strategies are helpful in preventing such air leaks.

A pre-existing pneumothorax is likely to enlarge and tension with the application of positive pressure ventilation, the use of nitrous oxide anaesthesia, and any reduction in barometric pressure (e.g. high altitude/air transport). Similarly, such changes in pressure may cause the development of a pneumothorax in patients with a pre-existing air leak.

Causes of pneumothorax

Primary spontaneous pneumothorax occurs more often in smokers and patients with Marfanoid habitus. The common causes of secondary pneumothorax are summarized in Table 1.

Table 1.

<table>
<thead>
<tr>
<th>Causes of Pneumothorax</th>
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<tr>
<td>Marfanoid habitus</td>
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<td>Smoking</td>
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<td>High altitude/air transport</td>
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Key points

A plain PA or AP radiograph generally underestimates the size of a pneumothorax. Simple aspiration is recommended as a first-line treatment for all primary pneumothoraces requiring intervention in the spontaneously breathing patient without CPAP. Patients who develop a pneumothorax while on positive pressure ventilation or CPAP should be treated with a chest drain unless immediate weaning from positive pressure ventilation is possible. A thoracic surgical opinion should be sought in cases of persistent large volume air leaks or failure of the lung to re-expand significantly within 4 days. Except in rare cases of tension, air emphysema and pneumomediastinum are not life-threatening and need observation only. Tension pneumothorax is an unusual but a recognized cause of respiratory and cardiovascular compromise during anaesthesia and surgery.
Clinical features

Typically, most awake spontaneously breathing patients have pleuritic chest pain and dyspnoea. In patients with underlying lung disease, dyspnoea is severe and significant hypoxaemia can occur, even with a small pneumothorax. Arterial blood gas measurements typically show an increase in the alveolar–arterial oxygen gradient and acute respiratory alkalosis. The diagnosis of pneumothorax in critical illness or in the ventilated patient is suggested by deterioration in the patient’s respiratory and cardiovascular variables, and physical examination. It is confirmed by radiological investigation or ultrasound. Patients with a small pneumothorax (<15% of the hemithorax) often have a normal physical finding on examination. Tachycardia is the most common physical finding. In patients with a larger pneumothorax, examination shows decreased movement of the chest wall, a hyperresonant percussion note, tracheal shift, and decreased or absent breath sounds on the affected side. The physical findings are often subtle and may be masked by the underlying lung disease, particularly in patients with chronic obstructive pulmonary disease. Systemic hypotension and central cyanosis should raise the suspicion of a tension pneumothorax. Ventilator parameters will show decreased tidal volumes and compliance.

Chronic lung conditions give rise to adhesions between parietal and visceral pleura restricting lung collapse. In such situations, a pneumothorax may be loculated and localized rather than spreading throughout the pleural space. In established adult respiratory distress syndrome (ARDS), a pneumothorax is often present without the lung completely collapsing as a result of the stiff, non-compliant nature of the lungs that are filled with fluid and cellular debris and associated pleural inflammation. Similarly, such affected lungs may be slow to re-expand (Fig. 1). Therefore, a tension pneumothorax may exist without total lung collapse or mediastinal shift. Once a pneumothorax has occurred, the high pressures generated during mechanical ventilation cause the pneumothorax to tension producing respiratory and haemodynamic effects. If suspected, it should be confirmed with a chest X-ray or other investigations without delay.

Table 1 Common causes of secondary pneumothorax

<table>
<thead>
<tr>
<th>Less common</th>
<th>Disease processes</th>
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<tr>
<td>Necrotizing pneumonias</td>
<td>Iatrogenic procedures</td>
</tr>
<tr>
<td>Sarcoïdosis</td>
<td>Positive pressure ventilation</td>
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<tr>
<td>Connective tissue diseases</td>
<td>Central venous catheterization</td>
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<tr>
<td>Idiopathic pulmonary fibrosis</td>
<td>Surgical procedures in the thorax,</td>
</tr>
<tr>
<td>Marfan’s syndrome</td>
<td>or lower neck</td>
</tr>
<tr>
<td>Lung cancer</td>
<td>Tracheostomy</td>
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<td>Sarcoma</td>
<td>Thoracoscropy/laparoscopy</td>
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Subcutaneous emphysema, pneumomediastinum, and pneumopericardium

Subcutaneous emphysema is characterized by painless swelling of the tissues because of air tracking along tissue planes. It is commonly seen over the chest wall around drain sites, in the head and neck. Palpation elicits a characteristic tissue paper feeling beneath the fingers. Air may track deeper into the mediastinum, retroperitoneum, scrotum, and down into the limbs. It is rarely a problem clinically but may herald the presence of a pneumothorax or other visceral damage. If very tense and causing distress to the patient, it can be drained by skin incision with a needle or pointed blade. Radiologically, it is seen as translucent areas after the line of anatomical structures (Fig. 1).

Pneumomediastinum is caused by similar mechanisms to pneumothorax. Radiologically, it is seen as black lucent air collections along the line of tissue planes and visceral structures. In addition, it may indicate perforation of the GI tract (e.g. after oesophageal instrumentation). It is also seen with perforation of pharynx, duodenum, colon, and rectum with tracking of air into mediastinum. Pneumothorax is associated with, but not generally caused by, pneumomediastinum. Except in rare cases of tension, pneumomediastinum is not a life-threatening condition. It is a predictable effect of positive pressure ventilation through a misplaced tracheostomy tube in the paratracheal tissues. If ventilation is continued, the latter is typically followed by a unilateral or bilateral pneumothoraces.

The Macklin effect is a pathophysiological process summed up in three steps: alveolar rupture, air dissection along bronchovascular sheaths, and spreading of this pulmonary interstitial emphysema...
into the mediastinum. This leads to pneumomediastinum in conditions such as neonatal respiratory distress syndromes, acute asthma, positive-pressure mechanical ventilation, and Valsalva manoeuvres (e.g. parturition, Boerhaave’s syndrome, and epileptic seizures). Venous air embolism from such mechanisms is well described as a terminal event in neonates with neonatal respiratory distress syndrome (RDS).

Pneumopericardium is an uncommon but potentially life-threatening condition. It is often caused by penetrating chest trauma, invasive procedures (e.g. laparoscopy and tracheostomy), and infections (e.g. lung abscess), but it can also occur during coughing, the Heimlich manoeuvre, the Valsalva manoeuvre, and mechanical ventilation. In tension pneumopericardium, rapid resuscitation and emergency pericardiocentesis may be required.

**Barotrauma and mechanical ventilation**

An estimated 4–15% of critically ill patients on ventilators develop barotrauma, manifesting as abnormal air collections in the chest. Underlying lung disease, such as pneumonia and especially ARDS, increases the risk significantly. The major factors associated with development of barotrauma include a peak inspiratory pressure >40 cm H₂O, the use of positive end-expiratory pressure (PEEP), and an inappropriately large tidal volume. Restrictive ventilation strategies should help in prevention.

**Distinction of pneumothorax from emphysematous bullae**

The bullae of emphysema can be very large and, when situated in the periphery of the lung, can mimic a loculated pneumothorax. A chest drain inserted into a bulla in the mistaken belief that it is a pneumothorax is not uncommon. The lack of a lung edge, the round nature of the bulla, and the presence of multiple bullae elsewhere in the lung are all clues to the diagnosis. In difficult cases, computed tomography (CT) is helpful in distinguishing between the two. Another classical differential diagnosis not be missed is air-filled stomach or bowel in the chest secondary to diaphragmatic hernia.

**Investigations**

**Chest X-ray**

The classical appearance in the upright position is the presence of radiolucent air and the absence of lung markings between the shrunken lung and the parietal pleura (Fig. 2). In the supine ventilated patient, gravity and the effects lung disease often give rise to a different appearance of the so-called ‘supine pneumothorax’. The pneumothorax is usually anteromedial or sub-pulmonic causing lucent upper quadrants of the abdomen, sharp superior surfaces of the diaphragm, the deep sulcus sign (Fig. 3), and visualization of the inferior surface of consolidated lung. Less often, the pneumothorax is apical, lateral (displaces the minor fissure from the chest wall), or posteromedial. False-positive appearances may occur from skin folds, overlying tubing/dressing/lines, and prior chest tube tracks.

Small pneumothoraces are significant in the ventilated patient as they can acutely enlarge and tension. Signs of tension include mediastinal shift, loss of compliance, cardiovascular system (CVS) instability, high central venous pressure (CVP), displacement of the anterior junction line, azygoesophageal recess, and flattening of heart and vascular shadows.
The radiographic signs of pneumomediastinum depend on the depiction of normal anatomical structures that are outlined by the air as it leaves the mediastinum. If there is sufficient air, the thymus can become elevated to produce the thymic sail sign. Air anterior to the pericardium (pneumoprecardium) is a frequent manifestation; it may be seen surrounding the pulmonary artery and the major branches of the aorta. Air collections in the pericardium are seen within the outline of the pericardium.

The plain posterior–anterior (PA) radiograph is a poor method of quantifying the size of a pneumothorax as it usually underestimates it. The volume of a pneumothorax approximates to the ratio of the cube of the lung diameter to the hemithorax diameter; hence, a pneumothorax of 2 cm on a PA film would occupy 49% of the hemithorax volume. This is recommended as a guide to aid decisions on intervention by British Thoracic Society.10

Ultrasound

Ultrasoundography has been shown to have high sensitivity (95%), specificity (100%), and diagnostic effectiveness (98%) for pneumothorax when compared with CT as a standard. It is useful for detecting small collections not seen on plain films and the extent of the air collection can be estimated by tracking the presence of the ‘sliding lung sign’ over the chest wall. Experience in the use of ultrasound for this indication is required to be confident in its application. It is difficult or impossible to visualize pleural structures with ultrasound through surgical emphysema.

CT scan

Indications for CT scanning include: differentiation of a pneumothorax from complex bullous lung disease; the diagnosis of supine pneumothorax; when aberrant chest tube placement is suspected; and when the plain chest radiograph is obscured by surgical emphysema. It allows definitive diagnosis of other pleural and lung pathologies and should be considered early when doubt exists. Overall, it is the gold standard for such imaging (see Fig. 2) but is difficult and potentially dangerous in the critically ill patient, and impractical for frequent repeated imaging.

Management of pneumothorax

General

Observation should be the treatment of choice for primary spontaneous small closed pneumothoraces without significant breathlessness, in a spontaneously breathing patient. Inhalation of high concentrations of oxygen may speed the resolution of a pneumothorax by reducing the partial pressure of nitrogen in the pulmonary capillaries. This should increase the pressure gradient between the pleural cavity and pleural capillaries, so increasing the absorption of air from the pleural cavity. The rate of re-absorption of spontaneous pneumothoraces is 1.25–1.8% of the volume of hemithorax every 24 h.11

Symptomatic patients should not be left without intervention regardless of the size of the pneumothorax on a chest radiograph. Other considerations include the need for positive pressure ventilation, impending anaesthesia and surgery (nitrous oxide diffuses into air collections and increases pressure/volume), transport in or outside the hospital, and altitude changes (including air transport).

Aspiration

Simple aspiration is recommended as first-line treatment for all primary pneumothoraces requiring intervention but is less likely to succeed in secondary pneumothoraces. In the latter situation, it is only recommended as an initial treatment in small (<2 cm) pneumothoraces in minimally breathless patients.

Pleural aspiration is performed after strict aseptic precautions with the patient in the supine position. The aspiration is carried out in the fourth inter-costal space in the anterior axillary line. The site is infiltrated with lidocaine and an 18-G i.v. cannula is inserted into the pleural cavity. The needle is withdrawn and three-way stopcock connected to the i.v. cannula. A 50 ml syringe and i.v. tubing with its end under a water seal are connected to the cannula through the three-way tap. Air is aspirated and expelled by means of the tubing and its volume noted. The end point of the procedure is a feeling of resistance to aspiration or if the patient begins to cough excessively. The patient will usually get a sensation that the lung has expanded. The i.v. cannula is withdrawn and entry site sealed. Avoid using a needle alone, as the lung will become lacerated as it expands towards the needle.

Advanced Trauma Life Support guidelines recommend the use of a cannula of 3–6 cm long to perform needle thoracocentesis for life-threatening tension pneumothorax. However, in 57% of patients with tension pneumothorax, the thickness of the chest wall has been found to be >3 cm. Therefore, it is recommended that a cannula length of at least 4.5 cm should be used in needle thoracocentesis of tension pneumothoraces.12 The cannula should be left in place until bubbling is confirmed in a formal chest drain with underwater seal to indicate proper function of the intercostal tube. We encourage clinicians, wherever possible, to ascertain by imaging that a pneumothorax is present before inserting such cannulae. Otherwise, it is inevitable that the needle will enter the lung and produce an air leak.

Chest drains and closed underwater systems

If simple aspiration of any pneumothorax is unsuccessful in controlling symptoms, an intercostal tube should be inserted. Intercostal tube drainage is recommended in secondary pneumothorax except in patients who are not breathless and have a very small (<1 cm or apical) pneumothorax.

Small bore drains are as effective for air drainage as large bore drains and are more comfortable for patients. If there is associated blood, a large bore drain will be required. There are no large
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Mechanically abraded/stripped). Both techniques produce fibrosis

Fig 4 The ‘safe triangle’ for inserting a chest drain.

randomized, controlled trials directly comparing small and large

bore drains.

The most common position for chest tube insertion is in the

mid-axillary line, through the ‘safe triangle’ illustrated in Figure 4.

This position minimizes risk to underlying structures such as the

viscera and internal mammary artery and avoids damage to muscle

and breast tissue resulting in unsightly scarring. A more posterior

position may be chosen if suggested by the presence of a loculated

collection. While this is relatively safe, it is not the preferred site

as it is more uncomfortable for the patient to lie on after insertion

and there is more risk of the drain kinking.

For apical pneumothoraces, the second intercostal space in the

mid-clavicular line is sometimes chosen; it is not recommended

routinely, as it may be uncomfortable for the patient, may leave an

unsightly scar, and internal mammary vessels are at risk. If the

drain is to be inserted into a loculated pleural collection, the pos-

ition of insertion will be dictated by the site of the locule as deter-

mined by imaging. A common mistake is to insert drains too low

in the chest risking damage to the diaphragm, liver, spleen, and

heart.

The drain should not be removed until bubbling has ceased, and

chest radiography demonstrates lung re-inflation. There is no evidence

that clamping a chest drain at the time of its removal is beneficial.

The use of high-volume/low-pressure suction pumps has been

advocated in cases of non-resolving pneumothorax or after chemi-

cal pleurodesis; however, there is no evidence to support its

routine use in the initial treatment of spontaneous pneumothorax.

If suction is required, this should be performed through the under-

water seal at a level of 10–20 cm H2O. A high-volume pump is

required to cope with a large leak.

Prevention strategies

There are two methods used to prevent recurrence of pneu-

mothoraces: (i) medical pleurodesis (installing talc/bleomycin by a

drain) and (ii) surgical pleurodesis (parietal pleura is mechanically abraded/stripped). Both techniques produce fibrosis

and scarring so that the visceral pleura adheres to the chest wall

obliterating the pleural space and prevents further pneumothorax.

Bronchopleural fistula

A bronchopleural fistula is a communication between the bronchial
tree and pleural space. Clinically, it may be best described as a

persistent air leak or a failure to re-inflate the lung despite chest
tube drainage for 24 h. Causes include chest trauma, complications

diagnostic or therapeutic procedures (e.g. thoracic surgery with

a failure of suture/staple line), chest drains inserted into the lung

parenchyma, and complications of mechanical ventilation. The

main problems with a large fistula in a ventilated patient are the

loss of delivered tidal volume, inability to apply PEEP, persistent

lung collapse, and delayed weaning from assisted ventilation.

Management strategies include general conservative measures

such as large bore chest drains (multiple if necessary) and the use

of drainage system with adequate capabilities. In mechanically

ventilated patients, the goal is to maintain adequate ventilation and

oxigenation while reducing the fistula flow to allow the leak to

heal. This includes reducing inspiratory pressures, tidal volumes,

respiratory rate, PEEP, and inspiratory times, and accepting per-

missive hypercapnia and lower oxygen saturations. Most air leaks

will settle spontaneously over a few days if the patient can be

weaned onto spontaneous respiration without high levels of con-

tinuous positive airways pressure (CPAP). The size of the air leak

is critical; small tears or punctures will heal quickly while larger

structural damage to the lung or a major bronchus will not settle

with conservative management, particularly if high inflation pres-

sures are required for associated lung injury.

The use of other modes of ventilation including high-frequency

ventilation, oscillation, and differential lung ventilation through

double-lumen tubes has been reported. For proximal leaks, fibreop-

tic bronchoscopy and direct application of sealants (e.g. cyanoacry-

late, fibrin agents, gelform) have been tried with limited success.

Refractory cases need surgical repair of the air leak by thoraco-

plasty, lung resection/stapling, pleural abrasion/decortication, or

other techniques.

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Please see multiple choice questions 7–10