Bedside ultrasonography for diagnosis of pneumothorax

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Abstract: Ultrasonography (US) has found its way into the critical care and emergency settings for the evaluation of acute respiratory failure conditions in recent years. It is useful for the diagnosis of varieties of abnormalities involving pleura and lung such as pleural effusion, alveolar interstitial syndrome, and pneumothorax (PTX). In addition to its reproducibility and timeliness, US has high sensitivity and specificity for the diagnosis of these conditions. The most widely used method for bedside evaluation of PTX is chest X-ray (CXR). However, the diagnostic sensitivity of CXR in detecting PTX is limited especially in occult PTX and when the patient is assumed supine position. Computed tomography (CT) is the gold standard in the evaluation of PTX, but is limited by its high radiation exposure and safety concerns in transporting critically ill patients. In this paper we review current advances in PTX diagnosis using US.

Keywords: Critical care; ultrasonography (US); pneumothorax (PTX); lung point

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Introduction

Pneumothorax (PTX) is an emergency that requires urgent management to avoid catastrophic consequences. PTX is also an important cause of respiratory failure in the emergency department and an annual rate of PTX is estimated at 22.7 cases for 100,000 population (1). Timely and accurate confirmation or exclusion of PTX is of paramount importance in emergency and critical care settings. The plain chest X-ray (CXR) is a traditional method for the evaluation of PTX. However, CXR is shown to have low sensitivity in detecting intrapleural air in trauma patients especially in supine position (2,3). Although computed tomography (CT) is the gold standard diagnostic test for the PTX, it is well limited by its high exposure to radiation and sometimes it is unsafe to transport these unstable patients. Chest ultrasonography (US) has found its way into the emergency and critical care setting for more than three decades and it is now gaining momentum in the evaluation of acute respiratory emergencies including but not limited to PTX. Diagnosis of PTX with US was first reported in 1986 by Rantanen (4). Recently numerous clinical studies have shown that PTX can be diagnosed by US with high sensitivity and specificity (5-8).

Normal findings in chest US

The pleural line is the basic structure that should be identified in the image of chest US. The so-called lung sliding identified at pleural line of the normal lung shows a cyclic movement between the visceral and parietal pleura with spontaneous respiration (Figure 1). It is the basic sign we need to identify when performing chest US. When lung sliding is observed, PTX can be ruled out. Since the structure above the pleural line is static during respiration, it produces parallel lines that look like waves in M-mode (e.g., the horizontal line represents time and static structure produces parallel lines). Beneath the pleural line, the cyclic movement of lung with respiration produces sand-like appearance. Pioneers in chest US termed this a “seashore sign” that represents the normal findings in chest US (Figure 2).

Besides the lung sliding, the most common artefacts in lung ultrasound are A-line (Figure 3) and B-line (Figure 4). The A-line artefact is horizontal artifactual repetitions of
A-lines are often presented in normal lung, indicating that the lung is well aerated. Otherwise, A-line can also be found in PTX when lung sliding is absent.

B-line is the vertical line that is perpendicular to the pleural line and its appearance represents fluid accumulation in the alveoli. The B-line is also known as comet-tail artefact which is formed by repetitive reflections of the ultrasound wave arising from the pleural line. B-line is characteristic of long and laser-like appearance that never fades at the distal image. B-line can be typically found in interstitial syndrome and occasionally in dependent regions of normal lung. Because PTX can be ruled out by the presence of B-line, it is discussed in this section. There is a trick in identification of B-line in modern bedside ultrasound machines. Most modern machines have the “multi-beam” (MB) function, which presents the ability of a probe to get rid of all sorts of angled ‘beams’ of sound across the field to try to get around echoic interfaces and other obstacles that might interfere with the distal image. This MB function helps the probe getting rid of artefacts from echoic objects in the near field. However, it is not desirable to get rid of artefacts in near field because in chest US the B-line is just the artefact that we want in the near field. Therefore the MB function should be switched off when the purpose is to identify B-line...
Diagnosis of PTX

PTX is typically confirmed by the presence of the following ultrasonographic findings: a stratosphere sign, abolished lung sliding or lung pulsing, the absence of B lines, and the presence of lung point. Lung point can be found at the boundary where normal lung sliding and PTX coexist, and it is a specific sign of PTX (see more descriptions and videos below). Pathophysiologically, PTX is the detachment of visceral and parietal pleura by entrapped air in the pleural space. All ultrasonographic signs should be comprehended with the understanding of this pathological process of PTX.

A hallmark sign that can be found in PTX is the stratosphere sign (Figure 5). In the pleural line the lung sliding is abolished and the sand-like appearance beneath the pleural line is replaced by parallel lines which is termed stratosphere or barcode sign. However, this finding can also be identified in conditions in which there is no relative movement between visceral and parietal pleura, such as apnea and pleuritis. PTX abolishes B-line, which is very sensitive that the presence of B-line can help to rule out PTX in scanned regions. Lung sliding has the same value in determining PTX that PTX can be 100% ruled out by the presence of lung sliding (10). In conditions with less movement between visceral and parietal pleura, lung pulse can be employed to rule out PTX. Lung pulse is produced by transmission of cardiac pulse to pleural line, given that the visceral and parietal pleura contact with each other (Figure 6). Volpicelli et al. gives important value to the lung pulse sign by stating that “even in the absence of lung sliding and B lines, visualization of a lung pulse rules out pneumothorax” (12). This is important because in clinical emergency, the ruling out of PTX can immediately direct clinicians to other culprits of a drop in oxygenation or respiratory distress. One useful method to determine PTX is to examine the contralateral side of suspected lung region. Figure 7 shows healthy lung with evident lung sliding, while Figure 8 shows the absence of lung sliding and pulsing, replaced by multiple A-lines. These two video clips were obtained from both lungs of the same patient in the third intercostal space at mid-clavicular line.

Lung point is thought to be a 100% specific sign for
PTX, though the sensitivity is less satisfactory by the fact that lung-point cannot be found in every patient with PTX (15). Typically, lung-point is identified at the junction where visceral and parietal pleura contact with each other (Figure 9). In B-mode ultrasound, the stratosphere and seashore patterns alternate with each other. However, there is a tricky sign that may confuse beginners in determining the presence of lung point. In our previous work we identified this sign for the first time but it is thought to be universe in normal lung. We termed it the physiological lung point (17). This sign distinguished from the true lung point in that there is no barcode sign in the abnormal region. Instead, the cardiac pulse can be identified in this region. We proposed that physiological lung point could be identified at the junction of lung pleura and mediastinal soft tissue. “Pseudo-lung point” is another sign that should be distinguished from true lung point. The sign can be found in lung contusions, and again there is no stratosphere pattern (18).

After confirmation of the presence of PTX, the next important issue is to quantification of the volume of PTX. The latter is important in that it may determine whether conservative or surgical procedure needs to be instituted. Volpicelli et al. found that the location of lung point can reliably classify PTX size as compared to CXR (19). In animal models of PTX, Oveland et al. demonstrated that chest US can accurately assess the progression of PTX during mechanical ventilation (20). In a case study reported by our group, we identified a double lung point sign in an infant and the PTX was further confirmed by chest CT scan (21). The PTX can be managed with conservative method. To the best of our knowledge, there are three cases of double lung point having been reported and one common feature of them is that all of the patients were managed by conservative method due to small size of PTX (22-24). As a result, we propose that the identification of PTX with double lung point indicate limited volume of PTX. If the patient presents severe respiratory compromise, other potential causes should be thoroughly sought for.

Diagnostic accuracy of chest US

Due to its wide availability in emergency and critical care setting, US has been investigated in numerous studies for its diagnostic accuracy in the determination of PTX (25-31). Recently, two independent study groups conducted systematic reviews of these articles (32,33). Ebrahimie et al.’s study included 28 studies and showed that the pooled sensitivity and specificity of chest US were 0.87 (95% CI: 0.81-0.92) and 0.99 (95% CI: 0.98-0.99), respectively (33). Additionally, they performed meta-regression and the result showed that the diagnostic accuracy of US in diagnosing PTX performed by the emergency physician was higher than by non-emergency physician. Higher pooled sensitivity (0.90; 95% CI: 0.83-0.98) was found in non-trauma
setting, but lower specificity (0.97; 95% CI: 0.95–0.99) was reported in non-trauma setting. Alrajab et al.’s study included only 13 original research articles, probably due to more strict selection criteria (32). Their study showed that US had a pooled sensitivity of 78.6% (95% CI: 68.1–98.1) and a specificity of 98.4% (95% CI: 97.3–99.5). They also performed meta-regression and subgroup analyses to investigate heterogeneity, which indicated that consecutive sampling of patients provided higher sensitivity results than convenience sampling.

Conclusions

US represents a novel approach for the evaluation of PTX, with advantages of timeliness, high accuracy and high reliability. The US skills should be incorporated into the standard training programs of physicians working at emergency and critical care settings. Because lung ultrasound is relatively new, there are many areas of active research. For example, new signs continue to be reported and defined. The technique to quantification of PTX size is still under investigation. Furthermore, the diagnosis of PTX under special conditions such as mechanical ventilation and patients with large pulmonary bullae are being reported (34). Atypical signs such as “physiological lung point” and “pseudo-lung point” should be noted and carefully distinguished from true lung point.

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None.

Footnote

Conflicts of Interest: The authors have no conflicts of interest to declare.

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