

Sonographic Bedside Quantification of Pleural Effusion Compared to Computed Tomography Volumetry in ICU Patients

OPEN
ACCESS

Authors

Ulf Karl-Martin Teichgräber^{1,3}, Judith Hackbarth^{2,3}

Affiliations

- 1 Institut für Diagnostische und Interventionelle Radiologie, Universitätsklinikum Jena, Jena, Germany
- 2 Thoraxklinik, Abteilung für Anästhesiologie und Intensivmedizin, Universitätsklinikum Heidelberg, Heidelberg
- 3 Institut für Diagnostische und Interventionelle Radiologie, Charité Universitätsmedizin Berlin, Berlin, Germany

Key words

pleural effusion, sonography, volumetric assessment, chest CT

received 21.12.2017

revised 27.07.2018

accepted 30.08.2018

Bibliography

DOI <https://doi.org/10.1055/a-0747-6416>

Ultrasound Int Open 2018; 4: E131–E135

© Georg Thieme Verlag KG Stuttgart · New York

ISSN 2199-7152

Correspondence

Prof. Ulf Karl-Martin Teichgräber, MD, MBA
Institut für Diagnostische und Interventionelle Radiologie
Universitätsklinikum Jena
Am Klinikum 1
07747 Jena
Germany
Tel.: +49/03641/9324 831, Fax: +49/03641/9324 832
ulf.teichgraeber@med.uni-jena.de

ABSTRACT

Objectives To date, the reliability of ultrasound for the quantitative assessment of pleural effusion has been limited. In the following study, an easy and cost-effective bedside ultrasound method was developed and investigated for specific use in the intensive care unit (ICU).

Methods 22 patients (median age: 58.5 years, range: 37–88 years, 14 men and 8 women) with a total of 31 pleural effusions were examined in the ICU. The inclusion criterion was complete visualization of the effusion on chest computed tomography (CT). The ultrasound (US) examination was performed less than 6 h after the diagnostic CT scan. The pleural effusion volume was calculated volumetrically from the CT scan data. Within 4.58 +/- 2.87 h after the CT scan, all patients were re-examined with US in the ICU. The fluid crescent's thickness was measured between each intercostal space (ICS) with the patient in a supine position and a 30° inclination of the torso. The US measurements were compared to the calculated CT volumes using regression analysis, resulting in the following formula: $V = 13.330 \times \text{ICS}_6$ (V = volume of the effusion [ml]; ICS_6 = sonographic measurement of the thickness of the liquid crescent [mm] in the sixth ICS).

Results A significant correlation between the sonographically measured and the CT-calculated volumes was best observed for the sixth ICS ($R^2 = 0.589$; $\text{ICC} = 0.7469$ with $p < 0.0001$ and a 95% CI of 0.5364–0.8705).

Conclusion The sonographic assessment of pleural effusions in a supine position and a 30° inclination of the torso is feasible for the volumetric estimation of pleural effusion. This is especially true for ICU patients with severe primary diseases and orthopnea who are unable to sit upright or lie flat.

Introduction

Chest sonography is a viable bedside method to verify free fluid and to differentiate contained effusions, pleural peel, atelectasis, diaphragmatic elevation, and other lesions [1–5]. To date, there is no reliable method for the fast quantification of pleural effusions in intensive care unit (ICU) patients as a diagnostic basis for puncture or drainage.

In 1994, Eibenberger et al. developed a sonographic method for the volumetric quantification of pleural effusions in a strictly supine position [6]. In this study, the extent of the effusion was measured sonographically in each intercostal space. From these measurements and the effusion volumes, which were quantified by puncture, a formula was developed to estimate effusion volumes.

Such methods are feasible in ICU patients because they are rarely able to sit upright [6, 7]. This is especially true for consciously sedated and ventilated patients. Patients in respiratory distress who are not intubated and often ventilated suffer from orthopnea and are unable to lie flat on their back. Schmidt et al. developed a method for the estimation of pleural effusion volumes in patients with a 30° inclination of the torso [8]. The pleural effusion volume was defined as the sum of the basal expanse of the free fluid between the diaphragm, the inferior lobe of the lung, and the maximum craniocaudal expanse of the effusion multiplied by a factor of 70. This method is adequate and feasible for ICU patients with a slight inclination of the torso. However, this approach requires multiple measurements and is therefore relatively time-consuming and less feasible in an ICU setting.

The aim of this prospective non-randomized study was to develop a simple and fast method for the bedside sonographic quantification of pleural effusions in ICU patients.

Materials and Methods

Patient cohort

Overall, 22 patients (median age: 58.5 years, range: 37–88 years, 14 male and 8 female patients) with a total of 31 pleural effusions were sonographically examined in a university hospital. A chest CT was performed on all patients as part of the clinical and diagnostic routine prior to the ultrasound exam. All CT scans were then assessed and evaluated by an experienced radiologist. If pleural effusions were detected, the patient was re-examined with an additional sonogram. The inclusion criteria demanded that the chest CT and the ultrasound exam be performed less than six hours apart and that the patient had not yet been treated with effusion puncture or drainage prior to the ultrasound exam. Finally, only patients with uncontained effusions were included in this study.

Sonographic measurements of pleural effusion

The sonographic exam was performed while the patient's torso was positioned at a 30° incline. Intercostal spaces (ICSs) 4–9 were traced in the posterior axillary line (PAL) with a 3.5-MHz curved linear transducer head of an ultrasound unit type EUB 405 (Hitachi Medical Systems, Tokyo, Japan). The ICSs were identified on the basis of anatomical landmarks such as the clavulae and vertebral bodies and used as acoustic windows. The transducer head was tilted until the effusion became visible on the ultrasound screen. The expanse of the fluid moat between the visceral pleura and the parietal pleura was measured in millimeters. The most caudal portion of the effusion was chosen for each measurement (► Fig. 1).

Computed tomographic volumetry

The patients were selected on the basis of routine diagnostic CT scans, which depicted the entire pleural effusion. All scans were performed on a 4-slice spiral CT scanner with a slice thickness of 5 mm and a pitch of 2. The pleural effusion volume was calculated using semi-automated volumetric software with threshold analysis and contour limiting on an Easyvision workstation (Philips Healthcare, Best, the Netherlands). A threshold analysis was used

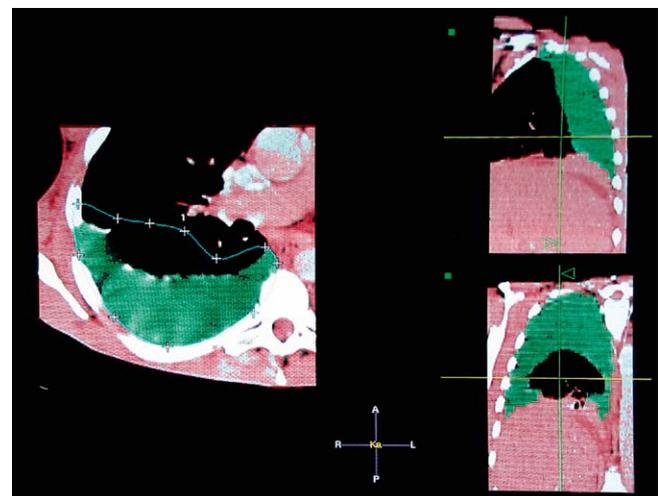
to subtract the air and skeletal structures from the calculated volume. Each image slice was examined and evaluated individually. The area of effusion was delineated with the cursor in each image slice by the radiologist (► Fig. 2). The computer program then calculated the total volume of the pleural effusion from the sum of each demarcated region and slice thickness.

Statistical analysis

The sonographically measured values in millimeters were compared to the CT-calculated volumes in a scatter plot. The CT volumes and the maximum extent of the effusion were compared using linear regression analysis. The regression line equations were then used to calculate the pleural effusion volumes, which were then compared to the corresponding CT-calculated volumes. An interclass correlation coefficient (ICC) was determined for each comparison. On the basis of the regression line function of the ICS



► Fig. 1 Measurement of the liquid crescent of the pleural effusion during sonography.



► Fig. 2 CT reconstruction in 3 planes: Region marked with cursor (green: volume considered in total volume calculation) in a single CT image slice after threshold analysis (red: volume not considered for total volume calculation).

with the highest ICC and the smallest two-way confidence interval (CI), a formula was developed to calculate the pleural effusion volumes with the sonographic image data.

Results

To assess and measure the maximum interpleural expanse of the effusions, a sonographic window at the sixth ICS in the PAL is ideal. Both the CI and ICC for the sixth ICS as sonographic windows confirm this hypothesis in ► **Fig. 3**.

The sixth ICS regression line equation is as follows:

$$\text{Volume}_{\text{CT}} (\text{ml}) = 13.330 \times \text{section}_{\text{ultrasound}} (\text{mm}) + 27.134$$

The regression coefficient for the regression line of the sixth ICS was $r = 0.767$. The coefficient of determination was $R^2 = 0.589$ with a 95% CI of 9.017-17.643 of the regression line. The ICC for the sixth ICS was 0.7487 ($p < 0.00001$; 95% CI 0.5393-0.8714).

Because the absolute coefficient loses its mathematical relevance in practice, the absolute coefficient is negligible in favor of a simplified formula:

$$V_{\text{effusion in CT}} (\text{ml}) = 13.330 \times \text{thickness}_{\text{liquid head in the sixth ICS}} (\text{mm})$$

We have listed the rounded values with which the estimation of the pleural effusion volume is more comfortable and considered the standard deviation in ► **Table 1**.

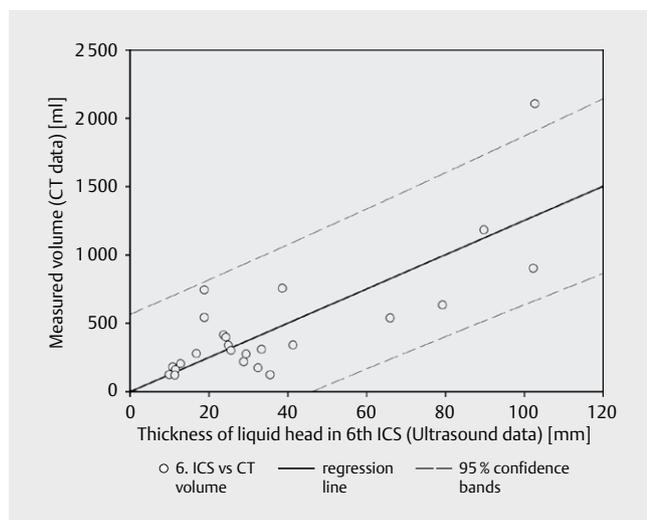
Discussion

In the verification of pleural effusions, sonography (sensitivity: 100%, specificity: 99.7%) is superior to chest X-ray in an erect patient (sensitivity: 71%, specificity: 98.5%) [1, 9]. Moreover, an upright chest X-ray is impossible for most ICU patients with severe primary diseases and impaired physical stamina. The sonographic identification of pleural effusions is simple; the effusions appear

echo-free, are sharply delineated, and show dorsal echo enhancement [2, 3]. Since atelectasis, diaphragmatic elevation, tumors, and pleural sheaths cause no difficulty with respect to differentiation on ultrasound images as in conventional fluoroscopic images, larger effusions can be diagnosed effortlessly [1, 2, 5]. An effusion volume of at least 150 ml is necessary for the diagnosis of an uncontained pleural effusion on a standard X-ray in an erect patient [10]. However, 5-ml effusion volumes in the basal and laterodorsal compartment between the ribcage and the diaphragm suffice for a reliable sonographic diagnosis [10]. The ultrasound examination can be performed bedside and can be repeated easily and quickly for control purposes. Compared to conventional chest X-ray, sonography is the more precise method and is also readily available in most ICUs [6].

In this study, a sonogram was performed on patients whose torso was inclined by 30°. The image data were collected during expiration. In 1994, Eibenberger et al. examined 51 patients with pleural effusions in a strictly supine position at maximum inspiration. In this study, the expanse of the pleural effusion was measured sonographically. Measurements were taken between the parietal and visceral pleura. In the second step, the effusion was punctured in 200-ml increments until the entire effusion was drained. Sonographic measurements were taken between each puncture from the same angle. All 331 sonographic measurements were then compared to the drained volumes using regression analysis. Statistical analysis rendered the following formula: $y = 47.6x - 837$ (the formula proposed in this paper is $V = 13.330 \times \text{ICR6}$) with y equaling the approximated effusion volume (ml) and x being the maximum expanse of the effusion (mm). The arithmetic median standard deviation was ± 224 ml [6].

In a further study, Balik et al. examined 81 patients with pleural effusions in a supine position with a torso elevation of 15°. The maximum separation between the parietal and visceral pleura (Sep) at maximum expiration was measured. Thoracocentesis was performed



► **Fig. 3** Regression diagram for the 6th ICS: measured volume (CT data) over thickness of liquid head in the 6th ICS (ultrasound data).

► **Table 1** Estimation of effusion volume with sonographically measured liquid crescent.

Thickness of liquid crescent in sonography (mm)	Estimated volume of the effusion (ml)	Standard deviation (ml)
5	70	20
10	130	40
15	200	60
20	270	90
30	400	130
40	530	170
50	670	220
60	800	260
70	930	300
80	1070	350
90	1200	390
100	1330	430

and the total volume of the drained effusion was determined. The pleural effusion was calculated as follows: $V \text{ (ml)} = 20 \cdot \text{Sep (mm)}$. The mean predictive error of V in this method was $158 \text{ ml} \pm 160.6 \text{ ml}$ [11]. For comparison, in our study, the arithmetic median standard deviation was slightly worse at $\pm 204.85 \text{ ml}$.

However, these values should be seen in a clinical context. An exact quantification, i. e., 260 ml exactly, is most likely less important than the ability to determine if an effusion volume is in the range of 200 ml or 300 ml.

Ventilated lateral lung compartments may mask small effusions and thus impede reliable measurement of the effusion, especially during deep inspiration [12]. Therefore, sonographic measurements were performed during expiration. The patients were asked to breathe calmly and normally, with the goal of achieving comparability between spontaneous breathing and ventilated patients.

In this study, CT volumetry was used as a reference volume (the gold standard). We based this decision on studies that showed good correlation between the actual organ volumes and CT-calculated volumes [13–17]. It needs to be further evaluated whether CT calculations render more precise results than drainage in the quantification of effusion volumes. After the puncture of a pleural effusion, a residual volume may remain within the interpleural space and could thus lead to an underestimation of the total effusion volume. With CT volumetry, the entire effusion volume is imaged. However, this method can lead to other sources of error, such as the breath-induced movement of the thoracic organs. Since all CT scans were performed in a spiral CT, the patients were asked to hold their breath for 20 to 40 s. The scan technique errors caused by breath-induced movements are negligible [18]. Moreover, outlining the effusion with the cursor in each image slice can induce user-related errors [18]. Through the predefinition of a density threshold in CT excluding air and bone from the soft tissue image, merely intercostal musculature and atelectasis had to be differentiated from the effusion. The intercostal musculature was easy to differentiate due to its anatomic location; atelectases were precluded from the volume due to differences in density on the CT images. Differentiation from the diaphragm and liver in the abdominal slices of the CT images was also achieved using the anatomic location and density.

Drainage of pleural effusions for the diagnostic determination of effusion volume is an invasive method that leads to risks such as bleeding, infection, and pneumothorax. The formula we describe herein for the estimation of pleural effusion volumes with sonography is a dimension-free function describing a regression line. This allows the estimation of an effusion volume from a sonographically measured distance, i. e., the maximum expanse of the effusion measured from the sixth ICS. Further validation of the formula with a larger patient cohort and differentiation between ventilated and spontaneously breathing patients are necessary. To ensure simplicity and clinical feasibility, we assembled rounded measurements of the maximum expanse of the effusion to simplify the clinical quantification (► **Table 1**).

This technique can be used for bedside examinations in most clinical settings due to the wide availability and mobility of modern ultrasound units [19, 20]. Sonography of pleural effusions is a feasible method for critically ill patients who are mechanically

ventilated [11, 21, 22]. Moreover, a sonographic examination does not expose the patient or physician to ionizing radiation [19]. The ultrasound volumetric estimation of pleural effusions permits both determination of an indication for drainage and determination of the ideal puncture point [6, 8, 23, 24]. Furthermore, sonography is a cost-efficient method [19].

Our results are comparable to those noted in the cited studies, which used drainage as the gold standard for quantification. In contrast to the existing literature, we took advantage of the accuracy of CT volumetry and avoided the disadvantages and inaccuracies of drainage.

In conclusion, the ultrasound volumetric estimation of pleural effusion volumes is easy, cost-efficient, and clinically feasible using the proposed method. This method is ideal in critically ill ICU patients for bedside determination of an indication for pleural effusion diagnostics and drainage.

Conflict of Interest

The authors declare no conflict of interest.

References

- [1] Mathis G. Thoraxsonography—Part I: Chest wall and pleura. *Ultrasound Med Biol* 1997; 23: 1131–1139
- [2] Pernice H, Braun B. Sonographische Differenzierung pulmonaler Verschattungen. *Prax Klin Pneumol* 1979; 33: 1132–1137
- [3] Matalon TA, Neiman HL, Mintzer RA. Noncardiac chest sonography. The state of the art. *Chest* 1983; 83: 675–678
- [4] Rosenberg ER. Ultrasound in the assessment of pleural densities. *Chest* 1983; 84: 283–285
- [5] Doust BD, Baum JK, Maklad NF et al. Ultrasonic evaluation of pleural opacities. *Radiology* 1975; 114: 135–140
- [6] Eibenberger KL, Dock WI, Ammann ME et al. Quantification of pleural effusions: sonography versus radiography. *Radiology* 1994; 191: 681–684
- [7] Lorenz J, Börner N, Nikolaus HP. [Sonographic volumetry of pleural effusions]. *Ultraschall Med* 1988; 9: 212–215
- [8] Schmidt O, Simon S, Schmitt R et al. Volumetrie von Pleuraergüssen bei multimorbiden, postoperativen Patienten einer operativen Intensivstation, Vergleich von Sonographie und Thoraxbettaufnahme. *Zentralbl Chir* 2000; 125: 375–379
- [9] Schwerk WB, Riestler KP, Hess F. Real-Time- Ultraschalltomographie von Pleuraergüssen und pleuranahen intrathorakalen Raumforderungen. *Respiration* 1980; 39: 219–228
- [10] Grymiski J, Krakowka P, Lypacewicz G. The diagnosis of pleural effusion by ultrasonic and radiologic techniques. *Chest* 1976; 70: 33–37
- [11] Balik M, Plasil P, Waldauf P et al. Ultrasound estimation of volume of pleural fluid in mechanically ventilated patients. *Intensive Care Med* 2006; 32: 318–321
- [12] Börner N, Kelbel C, Lorenz J, Weilemann LS, Meyer J. Sonographische Volumenbestimmung und Drainage von Pleuraergüssen. *Ultraschall in Klinik und Praxis* 1987; 2: 148–152
- [13] Moss AA, Friedman MA, Brito AC. Determination of liver, kidney, and spleen volumes by computed tomography: an experimental study in dogs. *J Comput Assist Tomogr* 1981; 5: 12–14

- [14] Heymsfield SB, Fulenwider T, Nordlinger B et al. Accurate measurement of liver, kidney, and spleen volume and mass by computerized axial tomography. *Ann Intern Med* 1979; 90: 185–187
- [15] Fritschy P, Robotti G, Schneekloth G et al. Measurement of liver volume by ultrasound and computed tomography. *J Clin Ultrasound* 1983; 11: 299–303
- [16] Breiman RS, Beck JW, Korobkin M et al. Volume determinations using computed tomography. *AJR Am J Roentgenol* 1982; 138: 329–333
- [17] Brenner DE, Whitley NO, Houk TL et al. Volume determinations in computed tomography. *Jama* 1982; 247: 1299–1302
- [18] Lemke AJ, Hosten N, Neumann K et al. CT-Volumetrie der Leber vor der Transplantation. *Röfo Fortschr Geb Röntgenstr Neuen Bildgeb Verfahr* 1997; 166: 18–23
- [19] Patel MC, Flower CD. Radiology in the management of pleural disease. *Eur Radiol* 1997; 7: 1454–1462
- [20] Hirsch JH, Rogers JV, Mack LA. Real-time sonography of pleural opacities. *AJR Am J Roentgenol* 1981; 136: 297–303
- [21] Vignon P, Chastagner C, Berkane V et al. Quantitative assessment of pleural effusion in critically ill patients by means of ultrasonography. *Crit Care Med* 2005; 33: 1757–1763
- [22] Roch A, Bojan M, Michelet P et al. Usefulness of ultrasonography in predicting pleural effusions >500 mL in patients receiving mechanical ventilation. *Chest* 2005; 127: 224–232
- [23] Opacic M, Bilic A, Ljubicic N et al. Thoracocentesis under ultrasonographic control. *Acta Med Iugosl* 1991; 45: 71–75
- [24] Kohan JM, Poe RH, Israel RH et al. Value of chest ultrasonography versus decubitus roentgenography for thoracentesis. *Am Rev Respir Dis* 1986; 133: 1124–1126