VIDEO-ASSISTED THORACOSCOPIC SURGERY IN A 35-DAY-OLD INFANT WITH MULTILOCULATED THORACIC EMPYEMA: CASE REPORT

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Summary

The authors report a clinical case of a 35-day-old infant with multiloculated thoracic empyema who was successfully operated on by the surgical team. The infant was transferred from a regional medical facility with fever, low-productive cough, and dyspnea. The disease had started 7 days earlier, but despite 5 days of antibiotic treatment in the hospital of his place of residence, the child’s condition continued to deteriorate, so he was transferred to the pediatric intensive care unit of our institution. After admission, a CT scan revealed massive closed fluid collections with subpleural localization in the right hemithorax and in the projection of the anterior mediastinum associated with subtotal atelectasis of the right lung. Closed pleurisy without signs of lung tissue destruction, suspicion of intrathoracic cystic lymphangioma. The patient underwent video-assisted thoracoscopic surgery, during which several purulent localized collections were identified, which were opened and drained with a volume of approximately 200 ml of viscous purulent fluid. The postoperative period was difficult but favorable. 6 months later, CT and scintigraphy showed some insignificant residual lung changes and diffuse perfusion changes, confirming the slow regression of the septic inflammatory process of the pleural cavity after resolution.

The authors conclude that the presented case highlights the difficulties of imaging differential diagnosis and suggests that VATS is an effective and safe treatment option for infants with pleural empyema, especially in multiloculated fibrinopurulent forms, allowing a favorable outcome with a short duration of thoracostomy with tube and comparatively short recovery and hospitalization periods.

Key words: Empyema; Multiloculated; Lung; VATS; Infant.

Thoracic empyema is the accumulation of pus in the pleural cavity, which is quite common in children, usually as a complication of bacterial pneumonia. Thoracic empyema is less common in the neonatal period or in infants due to the immaturity of the immune system, which limits the localization of infection to the pleural space and the ability of the pleura to produce sufficient exudate [1]. The incidence of pleural empyema in children has increased significantly in the last 20 years in developed countries, being a complication of bacterial pneumonia in 0.6 % [2], representing approximately 3.5-12.5 cases per 100,000 children, with a morbidity and mortality rate of up to 10 %. At the same time, pleural empyema is a rare entity in the neonatal period with an estimated incidence of 20-440 cases per 100,000 neonatal admissions. These data have led to a significant focus of research on this infection, the analysis of local epidemiologic data, as well as the evaluation of some factors that could determine the progression of pneumonia to complicated forms with empyema, which are currently uncertain [3, 4, 5, 6].

The main goal of treatment of thoracic empyema is to limit the manifestations of sepsis by evacuating and sterilizing the pleural space, thereby restoring circulation to limit the manifestations of sepsis by evacuating and sterilizing the pleural space, thereby restoring circulation...
Labs: RBC – 3.0 x 10¹²/L; Hb concentration – 105 g/L, marked leukocytosis (leukocytes – 49x10⁹/L) with deviation to the left of the leukocyte formula (neutrophils: unsegmented – 20 %, segmented – 47 %, metamyelocytes 6 %, myelocytes 8 %, lymphocytes – 10 %, eosinophils – 0 %, monocytes 9 %, toxic granulation +++), platelets was 568 x 10⁹/L, total protein was 49.0 g/l. Arterial blood gas test showed: pH – 7.35; pCO₂-36.6 mmHg; pO₂-61.2 mmHg. Blood biochemistry test found: Alanine aminotransferase – 9 U/L, aspartate aminotransferase – 13 U/L; creatinine – 64 mmol/l; fibrinogen – 6.60 g/l; Na⁺ – 134 mmol/l; K⁺ – 3.8 mmol/kg, Ca²⁺ – 2.26 mmol/l, Cl⁻ – 106 %, glucose – 4.2 mmol/l, ESR – 30 mm/hour. Blood culture – negative.

Computed tomography showed massive closed fluid collections with subpleural localization in the right hemithorax and at the level of the anterior mediastinum associated with subtotal atelectasis of the right lung. Closed pleurisy without evidence of lung tissue destruction; intrathoracic cystic lymphangioma? Moderate displacement of the mediastinum to the left.

Fig. 1. Child U.M., aged 36 days. Computed tomography performed preoperatively on the 9th day after the onset of the disease (explanations in the text)

Tc⁹⁹m MAA lung scintigraphy performed preoperatively showed that the right lung was reduced in size, with an unclear and irregular contour, with uneven distribution of the radio-pharmaceutical and with a sudden diffuse decrease in pulmonary blood flow, with multiple areas of no pulmonary perfusion in the upper segments. Left lung with no perfusion changes.

Fig. 2. Child U.M., Preoperative perfusion lung scintigraphy (explanations in the text)

After the administered treatment, including antibiotic therapy, although the fever and the intensity of the toxic syndrome decreased in intensity, the persistence of dyspnea and subcostal retraction were determined, which, along with some diagnostic imaging problems, served as indications for video-assisted thoracoscopy.

After treatment, including antibiotic therapy, although the fever and toxic syndrome decreased in intensity, persistent dyspnea and subcostal retraction were noted, which, along with some diagnostic imaging problems, served as an indication for video-assisted thoracoscopy.

Examination with a 3-mm thoracoscope (0°; Richard Wolf, Germany) revealed free purulent fluid, viscous, yellow-green in color, which was aspirated in a volume of approximately 200 ml, as well as several purulent collections localized by adhesions, which were later removed. The pleura was apparently thickened with considerable fibrin deposits, which were removed. The pleural cavity was
lavaged with aminocaproic acid, Betadine solution and drained with a 12 F tube.

The postoperative period was difficult, the child was on assisted ventilation for the first 4 hours postoperatively, after which spontaneous breathing was restored. The chest tube was removed on postoperative day 10. Bacteriological results from the purulent pleural fluid showed no growth, while S.aureus, Corynebacter spp. and Candida were detected in the throat swab.

The patient was discharged on postoperative day 25 (hospital day 32) in satisfactory condition. The follow-up radiologic examination performed at discharge showed normal lung volume with an enhanced and distorted lung pattern without free air and fluid in the pleural cavity. Reaction of the interlobular pleura in regression. The pneumonic infiltrate was practically completely resolved. The thickening of the paracostal pleura on the right side is preserved. The mediastinum is normal. The scintigraphy performed at discharge showed uneven distribution of the radiopharmaceutical in the right lung, unclear contour of the upper zone with restoration of the medio-basal contour and pulmonary blood flow. On the left side there were no changes in pulmonary perfusion.

6 months after discharge, a favorable clinical evolution was observed, the child grew according to his age without any symptoms. Although some insignificant residual lung changes were detected on CT (Fig. 5), diffuse perfusion changes of the left lung were observed on lung scintigraphy (Fig. 6). These data confirm the slow regression of residual changes after resolution of the septic inflammatory process of the pleural cavity in infants.
Discussions. Empyema is a purulent collection of the pleural space of various etiologies and is a serious, potentially fatal complication of pneumonia, chest trauma, infected hemothorax or surgery, after lung resection or iatrogenically during thoracentesis or intercostal tube insertion. It is characterized by effusions and fibrinopurulent collections, which can lead to pulmonary constriction and resistance to drainage [11, 12].

Although the incidence of pneumonia has decreased with the introduction of the pneumococcal conjugate vaccine, several reports have indicated an increase in the incidence of thoracic empyema in children in recent decades [13]. According to some authors, pleural effusions and parapneumonic empyema are found in 10% to 40% of all children hospitalized with bacterial pneumonia, with the main pathogens being serotype 1 and serotype 3 Streptococcus, both of which are included in the 13-valent pneumococcal conjugate vaccine (PCV13), which has been used for standard vaccination of infants from 2 months of age in Germany since 2009 to prevent invasive pneumococcal disease [14, 15, 16]. Of note, the overall detection rate of bacteria in blood and/or pleural fluid is approximately 29% – 34%. According to some studies, the majority (87%) of the bacteria detected belonged to the group of gram-positive aerobic cocci, with S. pneumoniae, S. pyogenes and S. aureus being the most common of the 41 species reported, while the gram-negative aerobic bacteria (P. aeruginosa, E. coli, Klebsiella pneumoniae, Francisella tularensis) were found quite rarely in 5.7% and anaerobic bacteria in 2.6% [17, 18]. Few cases of empyema in young children following SARS-CoV-2 infection have been reported, including unique cases of co-infection with SARS-CoV-2 and S. aureus in a premature neonate complicated by thoracic empyema [19].

The concept that bacteria invade the visceral pleura to develop an infected parapneumonic effusion has recently been challenged because of differences between the typical bacteriology of pneumonia and thoracic empyema, and because many patients with empyema have no evidence of underlying pneumonia. In this regard, it has been suggested that in some cases pleural empyema and pneumonia should be considered as separate entities, as the mechanisms and sources of bacterial invasion are unclear [18, 20, 21].

According to the American Thoracic Society (1962), the course of pleural infection is characterized by three distinct phases. The first (exudative) phase is characterized by a sterile, transparent exudate resulting from increased permeability of the visceral pleura. The second phase (fibrinopurulent) is associated with infection of the exudate by the initial invading pathogen and is characterized by the deposition of fibrin on the surface of the visceral and parietal pleura, leading to the formation of loculations and/or adhesions. The third phase (organization or consolidation) is characterized by the growth of fibroblasts and the formation of granulation tissue and a non-elastic lung lining [22, 23].

Chest radiography, ultrasonography, and chest computed tomography allow visualization of the presence of massive pleural effusion and the area of atelectasis or consolidation of the affected lung [6], which also requires differential diagnosis with some cystic congenital malformations [1]. In the presented case, although the CT scan allowed evaluation of the lung parenchyma and provided an accurate anatomical representation of the pathology, the differential diagnosis of multilocular effusion with a bulky intrathoracic cystic process was difficult.

Although thoracic empyema was known as early as Hippocrates, real progress in the treatment of this pathology occurred only in the mid-19th century, when Roe (1844) introduced repeated aspiration instead of simple knife incision. Later, Goodfellow and de Morgan (1859) proposed intercostal drainage [24].

The medical-surgical management of thoracic empyema is quite controversial and further studies are needed to investigate the efficacy of chest tube drainage as a first-line treatment [6, 25], with some studies reporting failure rates between 38% and 47%, and failure of first-line treatment being associated with increased mortality [26].

Some authors indicate the effectiveness of intrapleural use of fibrinolytics in the treatment of pleural empyema in adults and children [27], including neonates [19]. According to some studies, intrapleural fibrinolytic therapy in the early stages of the disease with both tissue plasminogen activator and streptokinase was safe and successful in patients with complicated parapneumonic pleurisy and empyema [28]. However, the role of fibrinolytic therapy in the treatment of pediatric pleural empyema is not clear, and the safety profile and conflicting results remain uncertain and concerning due to adverse effects and complications such as bleeding, allergic reactions, severe pain. The recent Cochrane systematic review showed that fibrinolytic therapy...
is associated with longer duration of chest tube drainage, longer duration of fever, higher rates of need for analgesia, and increased total cost of care [29, 30, 31, 32].

In cases of inefficiency of first-line treatment with worsening of pleural empyema and progression to the fibropurulent or organizing phase, surgical debridement is required, with open thoracotomy being the conventional approach, a method associated with pain and significant postoperative morbidity [33]. Recent studies have shown that VATS performed in the early stage of the disease is technically easier, allows better drainage and has a higher probability of achieving full lung expansion, as well as being a method that significantly improves the recovery time and shortens the inpatient treatment period, including in young children [31, 34]. Although minimally invasive surgical techniques may offer additional treatment options, their exact role in the treatment of pleural empyema is still unknown and requires new complex studies in the future [35].

Therefore, the presented case highlights the difficulties of imaging differential diagnosis and suggests that VATS is an effective and safe treatment option for infants with pleural empyema, especially in multiloculated fibrinopurulent forms, which allows a favorable evolution with a low duration of thoracostomy with tube and comparatively short recovery and hospitalization periods.

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References:

ВІДЕОАСИСТОВАНА ТОРАКОСКОПІЧНА ОПЕРАЦІЯ У 35-ДЕНЬНОГО НЕМОВЛЯТИ З МУЛЬТИЛОКАЛІЗОВАНОЮ ЕМПІЄМОЮ ПЛЕВРАЛЬНОЇ ПОРожнини: КЛІНИЧНИЙ ВИПАДОК

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Резюме.
Автори повідомляють про клінічний випадок відеоасистованої еміпіємою плевральної порожнини у 35-денньої дитини, з приводу чого вона була успішно пропіонарена. Немовля було переведено з регіонального медичного закладу з лихоманкою, малопродуктивним кашлем і задишкою. Заходження почалося 7 днів тому, але, незважаючи на 5 днів лікування антибіотиками у лікарні за місцем проживання, стан дитини продовжував погіршуватися, з приводу чого його перевели в дитяче відділення інтенсивної терапії нашого закладу. Після госпіtalізації на КТ виявлено масові зміни в підпільній локалізації плеврального гнійного середовища, зміни, що були реальні та держані з відмінними близько 200 мл в’язкої гнійної рідини. Після операційної передачі був здійснений хірургічний вміс, при цьому відновлювалась картина землянки та ділянка, яка відповідала кількісним показникам нормального агрегату порожнини.

Ключові слова: відеоасистована еміпієма; плевра; лікування; ВАТО; немовля.