Clinical analysis of extralobar pulmonary sequestration with torsion in children: report of 6 cases

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Abstract

Background: Extralobar pulmonary sequestration is an uncommon congenital pulmonary malformation. Clinically, pedicle torsion of extralobar pulmonary sequestration is extremely rare. Due to inadequate awareness of its atypical presentation and imaging characteristics, clinical diagnosis is very difficult, and it is extremely easy to misdiagnose.

Case presentation: There were 6 children (3 males and 3 females), aged 3–12 years old. The main clinical symptoms of the children were abdominal and chest pain (3 cases), abdominal pain (1 case), chest pain (1 case), and vomiting and abdominal distension (1 case). Two cases were accompanied by fever. Preoperative ultrasound revealed a well-bordered mass with soft-tissue density, accompanied by pleural effusion. On contrast-enhanced computed tomography scans, the mass showed no obvious enhancement. A blood supply was only present in 1 case, and there was no feeding artery shown in the other 5 cases. Extralobar pulmonary sequestration with haemorrhagic infarction was pathologically confirmed. On postoperative days 2–6, the children were discharged uneventfully. There were no complications during the median follow-up of 4 months.

Conclusions: Torsed extralobar pulmonary sequestration usually occurs in childhood or adolescence, with abdominal and/or chest pain as the primary symptoms. Imaging examination shows a well-defined soft-tissue mass without enhancement. The feeding vessel is not clearly displayed in the mass, and extralobar pulmonary sequestration is accompanied by varying amounts of pleural effusion. Video-assisted thoracoscopic surgical resection is associated with excellent prognosis.

Keywords: Extralobar, Pulmonary sequestration, Pedicle torsion, Children

Background

Pulmonary sequestration (PS) is an uncommon congenital pulmonary malformation, accounting for 1.5% of all congenital pulmonary malformations [1]. It is characterized by a nonfunctioning mass of lung tissue that shows no normal communication with the tracheobronchial tree and receives its vascular supply from the systemic circulation [2]. PS is anatomically divided into intralobar pulmonary sequestrations (ILS) and extralobar pulmonary sequestrations (ELS) according to whether there is a complete visceral pleura. ELS comprises 25% of all pulmonary sequestrations [3]. The majority of ELS is located between the lower lobe of the lung and the diaphragm [4–6], and its feeding artery mainly originates from the thoracic aorta, abdominal aorta, or other vessels in the systemic circulation. Venous drainage reaches the right atrium via the azygos vein, hemiazygos vein or vena cava [7]. The overall incidence of ELS is very low, and concurrent pedicle torsion is extremely rare. To date, only 13 paediatric cases have been reported in the English literature. To improve the diagnosis of ELS with
| Patient No | Age (y) | Sex  | Weight (kg) | Chief complaint                  | Location | Imaging examination for diagnosis | Preoperative diagnosis                                          | Feeding artery on image | Combined deformity | Pathological diagnosis                                                                 |
|------------|---------|------|-------------|----------------------------------|----------|-----------------------------------|---------------------------------------------------------------|------------------------|------------------|-------------------------------------------------------------------------------|
| 1          | 7       | Male | 22.4        | Vomiting and abdominal distension| Left     | Enhanced CT scan and ultrasound   | Neurogenic tumour/ELS, intestinal obstruction, pleural effusion| −                      | −                | ELS with haemorrhage, necrosis and myofibroblast proliferation                |
| 2          | 3       | Male | 15.4        | Abdominal pain and fever         | Left     | Enhanced CT scan and ultrasound   | Lung consolidation, pleural effusion                          | −                      | −                | ELS with haemorrhage and necrosis                                              |
| 3          | 5       | Male | 25          | Chest pain                       | Right    | Enhanced CT scan                  | Neurogenic tumour/ELS                                         | −                      | −                | ELS with haemorrhage and necrosis                                              |
| 4          | 6       | Female | 22.1     | Chest and abdominal pain         | Right    | Enhanced CT scan and ultrasound   | Torsion ELS, pleural effusion                                 | +                      | −                | ELS with haemorrhage, necrosis and myofibroblast proliferation                |
| 5          | 10      | Female | 34.6     | Chest and abdominal pain, fever  | Right    | Enhanced CT scan and ultrasound   | ELS, pleural effusion, severe pneumonia                      | −                      | −                | ELS with haemorrhage and necrosis                                              |
| 6          | 12      | Female | 38.4     | Chest and abdominal pain         | Right    | Enhanced CT scan                  | Torsion ELS, pleural effusion                                 | −                      | −                | ELS with haemorrhage and necrosis                                              |

* Year, kg kilogramme, ELS Extralobar pulmonary sequestration, CT Computed tomography
* “+” means presence and “−” means absence or unclear diagnosis
pedicle torsion by paediatricians, herein, the clinical data of 6 children with ELS with torsion of the pedicle in our hospital were analysed.

**Case presentation**

**Patients’ general information**

Six cases comprising 3 males and 3 females were studied, with an age range of 3–12 years and a weight range of 15.4–38.4 kg. The ELS was located on the left side in 2 cases and on the right side in 4 cases. All children were free of comorbidities. The general information of these 6 children is summarized in Table 1.

**Clinical manifestations**

The initial symptoms of the 6 children on admission included abdominal pain and chest pain in three cases, abdominal pain in one case, chest pain in one case, and abdominal distension and vomiting in one case, and two of them had fever. The course of illness ranged from 1 to 10 days (median, 5 days).

**Imaging examinations and diagnosis**

Preoperative colour Doppler ultrasound and chest computed tomography (CT) examination revealed a well-bordered mass with soft-tissue density between the diaphragm and the lower lobe of the lung accompanied by a small amount of pleural effusion in 2 cases, medium pleural effusion in 3 cases and large pleural effusion in 1 case. On contrast-enhanced CT scans, the mass showed no obvious enhancement, the blood supply was only presented in 1 case, and no feeding artery was shown in the other 5 cases. On serial imaging, the pleural effusion and the mass diameter increased progressively in one child (Fig. 1).

A total of 3 patients were diagnosed as pulmonary sequestration or pulmonary sequestration with torsion by CT-scan or Colour Doppler ultrasound before surgery, and the remaining 3 cases were not clearly diagnosed by CT-scan or ultrasonography. One patient with imaging findings of pleural malignancy was finally diagnosed by thoracoscopy. A thoracoscope was used for exploration in one undiagnosed children. Pathological examination of the resected tissue specimens confirmed extralobar pulmonary sequestration with haemorrhagic infarction. Of the six cases with pulmonary sequestration, two were accompanied by myofibroblast proliferation.

**Treatment and prognosis**

Eventually, all 6 children underwent thoroscopic pulmonary sequestration resection. After entering the chest cavity, a dark red or purplish-red mass with intact capsule, congestion and necrosis was seen, accompanied by varying amounts of bloody pleural effusion in the pleural cavity. The feeding artery that originated from the intercostal artery in 3 cases and the descending aorta in 3 cases was observed intraoperatively. The ELS vascular pedicle was twisted several times at its origin. The vascular pedicles of all lesions were clipped with metal clips and then cut off. The median procedure time was 60 min (range, 60–120 min), and intraoperative haemorrhage ranged from 1 to 10 ml (median, 4 ml). The children did not have any intraoperative or postoperative complications, and their symptoms were relieved immediately after surgery. On postoperative days 2–6, they were discharged home uneventfully. There were no complications during a median follow-up of 4 months (range, 0.4–8.7 months).

**Discussion**

Symptomatic ELS caused by pedicle torsion is extremely uncommon in clinical practice, and early diagnosis is challenging. It has been reported that only 1 case of vascular pedicle torsion occurred in 13 cases of ELS treated in a single centre from 2000 to 2009 [8]. In the past 14 years, a total of 32 cases of ELS were diagnosed in our hospital, of which only 6 cases were combined with torsion of the vascular pedicle, accounting for 18.75% of cases. To date, a total of 13 paediatric cases of ELS with torsion have been reported in the English literature [4–6, 8–17]. To the best of our knowledge, this is the largest number of cases reported in the literature. Among the 13 cases of ELS reported in the previous literature, including 9 males and 4 females, 12 cases were located on the left side and 1 case on the right side. The patients were between 4 and 13 years old. The left/right ELS ratio was 1:2 in our centre’s cases. The proportion of ELS on the right side was larger in our study, while previous literature found that ELS tended to occur on the left side [4–7], which may be one of the reasons for the low preoperative diagnosis rate. In our cases, one child was an adolescent, three were school age, and two were preschool age. The male to female ratio was 1:1. The age range in our study was similar to that reported in the literature, but the sex ratio differed. ELS is associated with other congenital anomalies, such as congenital diaphragmatic hernia, congenital pulmonary airway malformation or congenital heart disease. However, it is very interesting that none of the 6 children in our study had other congenital malformations.

Clinically, ELS with torsion shows no specific manifestations or symptoms, and it is difficult to diagnose. The literature showed that chest pain or discomfort was recorded as the main symptom among adult patients [18, 19], whereas abdominal pain was the primary clinical manifestation in children [5, 8–11, 13–16], followed by chest pain [6, 12–15]. Some children had fever [10,
Abdominal pain seems to be the primary symptom of ELS with torsion in children. 80% of ELS in children is located between the lower lobe of the lung and the diaphragm, typically in the left hemithorax [9]. The type of abdominal pain is likely similar to the pain experienced by patients with right lower lobe pneumonia [9], which makes it easy to misdiagnose [5]. The scope of the CT scan should include the lower thorax to the level of the pulmonary veins when evaluating acute abdominal pain to rule out the possibility of lung disease [14]. The cause of the abdominal pain may be due to local inflammation caused by ischaemic necrosis after torsion of the ELS vascular pedicle. The children's symptoms disappeared immediately after removing the lesion [4, 12, 13]; however, antibiotic treatment could not have achieved this effect [13].

Imaging examinations, such as Doppler ultrasound, CT and magnetic resonance imaging (MRI), can play an important role in confirming the diagnosis. The discovery that the blood supply artery originates from the branch of a systemic artery rather than a pulmonary artery is strong evidence for the diagnosis of ELS [6, 14]. Unfortunately, torsion of the vascular pedicle hinders the visibility of the blood supply, resulting in unrecognizable blood supply vessels and atypical imaging findings [14]. Therefore, it is difficult for radiologists to diagnose ELS correctly before surgery. Additionally, there were atypical clinical symptoms in the children. These issues lead to higher rates of delayed diagnosis or misdiagnosis [9]. The preoperative diagnosis rate of ELS presenting with torsion is extremely low in childhood and adolescence. Among the 13 reported paediatric cases of torsed ELS in the literature, only 4 cases were diagnosed before the operation [4, 6, 10, 11]. In the early years, owing to a lack of awareness about the imaging characteristics of ELS with torsion among radiologists in our hospital, the first 3 cases were not identified before the procedure. After studying, analysing and summarizing the imaging characteristics of enhanced CT among radiologists, the last 3 cases were accurately identified before surgery. On contrast-enhanced CT scan, the main imaging signs of ELS with torsion are as follows: the mass has no obvious enhancement or only edge enhancement, and its feeding artery is unclear or not displayed accompanied by pleural effusion [5, 9–11, 14–16, 19]. Additionally, a rapid increase in pleural effusion or the mass size in a short period of time is considered to be one of the characteristic manifestations of acute torsion. Compared with CT, MRI seems to have more advantages in distinguishing ELS with pedicle torsion: lack of enhancement.

Fig. 1 Axial Contrast-enhanced CT image of the chest A indicates that a well-bordered, soft-tissue density mass without enhancement in the right thoracic cavity, coronal CT image B shows a well-defined soft tissue mass between the diaphragm and the spine, video-assisted thoracoscopy C reveals a purplish-red mass with congestion and necrosis in the pleural cavity, and D shows a twisted vascular pedicle.
Consent for publication
All of the authors agree to the publication of the article.

Competing interests
The authors declare that they have no competing interests.

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References
1. Li J, Jiang Y, Xiao J, Liang G. Extralobar pulmonary sequestration with a cyst: a case report. Ann Trans Med. 2020;8(15):969. https://doi.org/10.21037/atm-20-3815.
2. Huang D, Habudinc A, Yuan M, Yang G, Cheng K, Luo D, et al. The clinical management of extralobar pulmonary sequestration in children. Pediatr Pulmonol. 2021;56(7):2322–7. https://doi.org/10.1002/ppul.25433.
3. Gabelloni M, Faggioni L, Accogli S, Aringhieri G, Neri E. Pulmonary sequestration: what the radiologist should know. Clin Imaging. 2021;73:61–72. https://doi.org/10.1016/j.clinimag.2020.11.040.
4. Walcott J, Abdessalam S, Timmons Z, Winningham P, Beavers A. A rare case of torsion and infarction of an extralobar pulmonary sequestration with MR, CT, and surgical correlation: Radiol Case Rep. 2021;16(12):3931–6. https://doi.org/10.1016/j.radcr.2021.09.045.
5. Zucker EJ, Tracy DA, Chwals WJ, Solky AC, Lee YE. Paediatric torsed extralobar sequestration containing calcification: imaging findings with pathological correlation. Clin Radiol. 2013;68(1):94–7. https://doi.org/10.1016/j.crad.2012.05.008.
6. Yang L, Yang G. Extralobar pulmonary sequestration with a complication of torsion: a case report and literature review. Medicine (Baltimore). 2020;99(29):e21104. https://doi.org/10.1097/MD.00000000000021104.
7. Zhang N, Zeng Q, Chen CH, Yu J, Zhang X. Distribution, diagnosis, and treatment of pulmonary sequestration: report of 208 cases. J Pediatr Surg. 2019;54(3):1286–92. https://doi.org/10.1016/j.jpedsurg.2018.08.054.
8. Lima MA, Randi B, Gargano T, Tani G, Pession A, Gregoni G. Extralobar pulmonary sequestration presenting with torsion and associated hydrothorax. Eur J Pediatr Surg. 2010;20(3):208–10. https://doi.org/10.1055/s-0029-1241837.
9. Chen W, Wagner L, Boyd T, Nagarajan R, Dasgupta R. Extralobar pulmonary sequestration presenting with torsion: a case report and review of literature. J Pediatr Surg. 2011;46(10):2025–8. https://doi.org/10.1016/j.jpedsurg.2011.07.017.
10. Choe I, Goo HW. Extralobar pulmonary sequestration with hemorrhagic infarction in a child: preoperative imaging diagnosis and pathological correlation. Korean J Radiol. 2015;16(3):662–7. https://doi.org/10.3348/kjr.2015.16.3.662.
11. Gawilzta M, Hirsch W, Weißier M, Ritter L, Metzger RP. Torsion of extralobar lung sequestration-lack of contrast medium enhancement could facilitate MRI-based diagnosis. Klin Padiatr. 2014;226(1):38–9. https://doi.org/10.1055/s-0033-1351319.
12. Huang EY, Monforte HL, Shaul DB. Extralobar pulmonary sequestration presenting with torsion. Pediatr Surg Int. 2004;20(3):218–20. https://doi.org/10.1007/s00383-004-1156-0.
13. Kirkendall ES, Guiot AB. Torsed pulmonary sequestration presenting with gastrointestinal manifestations. Clin Pediatr (Phila). 2013;52(10):961–4. https://doi.org/10.1177/0009922812453197.
14. Shah R, Carver TW, Rivard DC. Torsed pulmonary sequestration presenting as a painful chest mass. Pediatr Radiol. 2010;40(8):1434–5. https://doi.org/10.1007/s00247-010-1558-1.
15. Son SA, Do YY, Kim YE, Lee SM, Lee DH. Infarction of torsed extralobar pulmonary sequestration in adolescence. Gen Thorac Cardiovasc Surg. 2020;68(1):77–80. https://doi.org/10.1007/s11748-019-01105-7.
16. Uchida DA, Moore KR, Wood KE, Pysher TJ, Downey EC. Infarction of an extralobar pulmonary sequestration in a young child: diagnosis and excision by video-assisted thoracoscopic surgery. J Laparoendosc Adv Surg Tech A. 2010;20(4):399–401. https://doi.org/10.1089/lap.2009.0217.
17. Yokota R, Sakamoto K, Ukahawa H, Takeshita M, Yoshimitsu K. Torsion of right lung sequestration mimicking a posterior mediastinal mass.

Abbreviations
PS: Pulmonary sequestration; ELS: Extralobar pulmonary sequestration; ILPS: Intralobar pulmonary sequestration; CT: Computed tomography; MRI: Magnetic resonance imaging.

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Author contributions
The manuscript was written by YT. YW participated in the surgeries as the chief surgeon, collected the data of children and analysed the data. FZ revised the manuscript and participated in the surgeries. JH and QZ were responsible for proofreading and editing. All authors read and approved the final manuscript.

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Availability of data and materials
The data were presented in the main manuscript.

Declarations

Ethics approval and consent to participate
This is a retrospective study, so there is no need for ethical approval.

in the peripheral portion of the lesion with haemorrhaging within the mass, and vascular pedicle was well visualized [10].

The exact reason for torsion of ELS remains unclear. The literature suggests that activity or respiratory exertion may be the predisposing factor for vascular pedicle torsion; for example, some patients had performed vigorous activity (track competition, tennis) prior to the onset of illness [4]. Unlike the literature, reviewing the medical history of the children in our cases revealed that they did not have a history of similar vigorous activity before the onset. An effective treatment for ELS is surgery. Minimally invasive video-assisted thoracoscopic surgery is a safe and effective treatment for PS and has considerable long-term effects [20]. Therefore, thoracoscopic remains the preferred procedure for both the diagnosis and treatment of this disease, especially for bilateral lesions [7, 8, 13–16, 21].

Conclusions
The ELS with torsion usually occurs in children or adolescents and has abdominal and/or chest pain as the primary symptom. Imaging examination shows a well-defined mass comprised of soft tissue between the lower lobe of the lung and the diaphragm. Contrast-enhanced CT shows that each mass had no enhancement or only marginal enhancement. The feeding vessel was not clearly displayed, and it was accompanied by varying amounts of pleural effusion. Video-assisted thoracoscopic surgical resection for ELS is associated with excellent prognosis.

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presenting as acute abdomen: usefulness of MR imaging. Radiol Case Rep. 2019;14(5):551–4. https://doi.org/10.1016/j.radcr.2019.02.008.

18. Kanauchi N, Kato H, Endo M, Okazaki T. Torsion of extralobar pulmonary sequestration. Eur J Cardiothorac Surg. 2011;39(3):e31. https://doi.org/10.1016/j.ejcts.2010.11.002.

19. Takeuchi K, Ono A, Yamada A, Toyooka M, Takahashi T, Shirakawa Y, et al. Two adult cases of extralobar pulmonary sequestration: a non-complicated case and a necrotic case with torsion. Pol J Radiol. 2014;79:145–9. https://doi.org/10.12659/PRJ890662.

20. Wang S, Li Y, Wang J. Video-assisted thoracoscopic surgery for pulmonary sequestrations: series of 35 consecutive patients in a single center. Thorac Cardiovasc Surg. 2019;67(1):73–8. https://doi.org/10.1055/s-0038-1668596.

21. Bleve C, Conigli ML, Biondini D, Ceccarelli PL, Giarraputo L, Savastano S, et al. Thoracoscopic treatment of a rare bilateral extralobar lung sequestration in a 3-years old girl. Pediatr Med Chir. 2021. https://doi.org/10.4081/pmc.2020.237.

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