Congenital pyriform sinus fistula presenting as a neck abscess in a newborn

A case report

Yishu Teng, PhD, Shuling Huang, MD, Guowei Chen, PhD, Zhixiong Xian, MD, Saihong Han, MD, Lan Li, PhD

1. Introduction

Patients with congenital pyriform sinus fistula (CPSF) generally present with recurrent neck swelling, pain, and/or a fistulous orifice with purulent discharge in the anterior or lateral neck region. CPSFs tend to be misdiagnosed as acute suppurative thyroiditis, neck cellulitis, neck abscess, or another disorder, such as a thyroglossal cyst or second branchial cleft.[1] A relatively rare clinical manifestation in neonates is a non-infectious, compressible neck mass.[2] The possibility of this disease should be kept in mind in a patient with an acute suppurative thyroid inflammatory response, especially on the left side and with repeated episodes, and timely examination is needed for a definitive diagnosis.[3,4]

Several imaging investigations, including barium swallow X-ray, computed tomography (CT), magnetic resonance imaging (MRI), suspension laryngoscopy, and esophagoscopy, are generally used to confirm the diagnosis.[5]

2. Case report

A male neonate was admitted to our Department of Pediatrics directly from the delivery room 25 minutes after birth following discovery of a cystic mass with extensive skin swelling in the left side of the neck (Fig. 1). He had a birth weight of 3660g and 1-minute and 5-minute Apgar scores of 10. The mother was gravida 2, para 1, and gave birth at 40 weeks' and 3 days' gestation. Her amniotic fluid was clear. The diagnosis was suspected to be a space-occupying lesion in the larynx. On physical examination, the neonate had a clear mind, normal reflexes, pink cheeks, and a swelling in the left neck with a soft palpable 5 × 5-cm mass with an unclear boundary. His skin temperature and respiration rate were normal. Auscultation of both lungs revealed coarse breath sounds without rales and normal heart sounds.

A chest X-ray suggested an increase in lung markings on both sides. No abnormality was found on color Doppler ultrasonography of the head. Echocardiography revealed a left-to-right (horizontal) shunt, an atrial septal defect, and a patent ductus arteriosus. A B-mode ultrasonographic examination of the left neck showed an anterior cervical cystic mass of indeterminate nature.
side of the neck revealed an anterior cervical cystic mass of indeterminate nature. Routine blood, urine, stool, electrolyte, and liver and kidney function tests were unremarkable and myocardial enzyme levels were normal. Blood culture showed no bacterial growth. A B-mode ultrasonogram obtained at 39 weeks’ gestation had revealed a 2.8 × 2.6-cm anechoic mass in the left side of the neck with the esophagus and trachea shifted to the right. A further B-mode ultrasonogram obtained one day after birth revealed a 5.1 × 3.8 × 3.1-cm cystic mass extending from the posterior pharynx and left side of the trachea to the front of the ascending aorta beneath, with no obvious displacement of the trachea but forward and outward movement of the jugular vein. The mass was observed to be regular in shape with a visible capsule and a clear boundary. No obvious blood flow signals were found in the mass on color Doppler flow imaging. On the 6th day after birth, B-mode ultrasonography showed a 6.6 × 5.0 × 3.9-cm cystic mass in the deep cervical fascial space on the left side; the mass had a clear boundary, an irregular shape, and a thick cystic wall, with dense spot-like echoes and gas-like echoes in the cystic cavity but no substantial space-occupying echoes. The upper lateral margin of the mass was adjacent to the left submandibular and parotid glands, and the lower margin extended to the thymus gland; it was adjacent to the left carotid sheath on the deep side with no wrapping and adjacent to the esophagus and left thyroid lobe on the medial side. A portion of the mass extended through the posterior part of the trachea to the inside of the right carotid sheath. Color Doppler flow imaging revealed spot-like blood flow signals in the wall of the cyst but no obvious blood flow signals in the mass. Axial contrast-enhanced CT scan image showing an abscess cavity with an irregularly enhanced cystic wall (Fig. 2).

Under the surgical microscope, the fistula and surrounding mucosa of the pyriform sinus were burned using a CO2 laser at a frequency of 2 W in continuous output mode. An area of basal mucosa in the pyriform sinus with a radius of 10 mm was burned, taking the internal fistula as the center (Fig. 5). After surgery, the wound was dressed and anti-inflammatory treatment was provided. The neck wound healed uneventfully thereafter. Three months later, the fistula was confirmed to be closed by suspension laryngoscopy under general anesthesia (Fig. 6). No recurrence was detected during 9 months of follow-up.
3. Discussion

CPSF is a branchial abnormality originating from the third or fourth branchial pouch and is an important cause of anterior cervical abscess in children. The lesion mainly occurs on the left side and may be associated with asymmetric development of the original great artery and bilateral branchial apparatus or a C-cell migration disorder. A CPSF can be clinically classified as a sinus, that is, a blind-ending tract that opens to either the pyriform fossa or the skin, a cyst, which is not open to either, or a fistula, which is a tract connecting the pyriform fossa and skin. A sinus CPSF is most commonly encountered, although a fistula may develop after iatrogenic or spontaneous rupture of a cervical abscess. At least 80% of pyriform sinus fistulae occur in preschool children and the incidence is similar in both sexes. There is a general lack of awareness of this disease among non-specialist clinicians, which often leads to misdiagnosis and inappropriate treatment.

Patients with a pyriform sinus fistula usually show recurrent cervical swelling, pain, and an external opening with purulent discharge at the anterior border of the middle or lower third of the sternocleidomastoid muscle after an upper respiratory tract infection, often with symptoms of acute suppurative thyroiditis. However, because of the abundant blood supply and thick fibrous capsule in thyroid tissue, these lesions rarely become infected under normal conditions. Accordingly, the possibility of this disease should be considered in a patient with an acute suppurative thyroid inflammatory response, especially if it is on the left side and recurrent. Prompt investigation is needed to make a definitive diagnosis and to exclude other causes of a neck mass, such as a thyroglossal cyst or a second branchial cleft.

The diagnosis relies not only on clinical presentation but also on radiological findings. A barium swallow X-ray can reveal a sinus tract in the pyriform fossa, but the sensitivity of this method is only 50% to 80%. Both MRI and CT can identify a tract between the internal orifice and the neck mass or infection. MRI can detect soft tissue inflammation in the affected area and CT is considered to be more effective and requires a shorter scanning time than MRI. CT typically identifies inflammation and/or an abscess near the lesser horn of the thyroid cartilage, with air bubbles forming in the lesion or along the putative tract or may reveal a loss of high density in the thyroid gland. Further confirmation of an internal orifice in the pyriform fossa by suspension laryngoscopy or esophagoscopy is the gold standard for diagnosis of CPSF.

The external component of a pyriform sinus fistula is often absent and the internal fistula is concealed, which increases the likelihood of delayed diagnosis and treatment and leads to a relatively high rate of missed diagnosis and inappropriate treatment. For CPSF should be high in a patient with an unexplained neck abscess or recurrent acute suppurative thyroiditis, especially if it occurs on the left side. Surgical treatment, such as endoscopic cauterization or open surgery, is required as soon as the diagnosis is made.

A multicenter, prospective, randomized controlled study that includes long-term clinical follow-up is needed to investigate the advantages of surgical methods. Improvement in our understanding of this condition should lead to more effective methods of treatment.
diagnosis and treatment. In summary, CPSF should be considered in neonates with a congenital cystic mass in the neck and in older pediatric patients with this presentation, even if there is no history of recurrent neck infection. In this case, a congenital cystic mass in the neck was found in the fetus and diagnosed as CPSF after birth, as opposed to repeated neck infections previously reported.

4. Consent for publication

The need for approval to publish this report was waived by the Institutional Review Board of Shenzhen Children’s Hospital. However, written informed consent was obtained from the patient’s parents for publication of the details of the patient’s case and the accompanying images.

Author contributions

Data curation: Saihong Han.
Investigation: Shuling Huang.
Project administration: Yishu Teng.
Resources: Shuling Huang.
Supervision: Shuling Huang, Zhixiong Xian.
Visualization: Guowei Chen, Lan Li.
Writing – original draft: Yishu Teng.
Writing – review & editing: Yishu Teng.

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