Idiopathic Congenital Non-Chylous Pleural Effusion in Neonate: A Case Report

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Abstract

Congenital isolated pleural effusion is a rare cause of respiratory distress in neonates. It is usually chylous. We report a rare case of non-chylous congenital idiopathic pleural effusion.

Keywords: Idiopathic neonatal pleural effusion; Neonate; Non-chylo

Introduction

Respiratory distress is a leading cause of neonatal mortality worldwide, such as hyaline membrane disease, transient tachypnea of new born and neonatal pneumonia, congenital diaphragmatic hernia and congenital cystic adenomatous malformation, metabolic disturbances, and congenital heart diseases [1]. Pleural effusion is rare, and early diagnosis of neonatal pleural effusion is important not only for lowering mortality and morbidity, but also because it has an excellent prognosis following timely diagnosis and treatment. Pleural effusions are rare in the neonate but may be associated to several clinical conditions [2]. Chylothorax is the most common cause of pleural effusion in neonatal period [1-3]. Here, we report a rare case of non-chylous congenital idiopathic pleural effusion.

Case Report

A full-term neonate, birth weight 3.3 kg, appropriate for gestation, was born by emergency cesarean section done for maternal bleeding and abrobiplacenta. Baby did not require resuscitation and the Apgar scores were 8 and 10 at 1 and 5 min, respectively. He was admitted to the neonatal intensive care unit with a report of pleural effusion in prenatal sonography in last month of pregnancy. Respiratory rate (RR) was 66/min with mild substernal retraction and breath sounds were diminished over the right hemithorax. There were no dysmorphic features and the baby was not hydropic. Rest of the systemic examination was normal. An arterial blood gas analysis was normal with pH 7.38, HCO3 18 mmol, and PaO2 85 mm Hg. A portable chest X-ray done revealed white opacity in right side with shifting of airway and heart to the left side (Fig. 1). Chest ultrasound showed about 400 mL fluid in Rt hemithorax. A diagnostic tap was performed and 50 mL of straw colored fluid was aspirated and chest tube was inserted which drained 500 mL of straw colored fluid during 3 next days. Following pleural drainage, the infant improved remarkably. Repeat X-ray chest showed full expansion of lungs at 24 h of age (Fig. 2). Feeding was initiated on day 2. The intercostal drain was removed on day 4. The pleural fluid analysis showed transudate fluid containing protein 271 mg/dL and sugar 104 mg/dL, Cl- 90 mEq/L, triglycerides 74 mg/dL, cholesterol 16 mg/dL, and leukocyte 1-2; no microorganisms were seen on gram staining and pleural fluid culture was sterile, indicative of non-chylous pleural effusion. Computed tomography (CT) of chest was normal. The sepsis screen was normal. USG of abdomen and skull were normal and baseline renal and liver function tests were within normal limits. The karyotyping of the infant was normal, i.e. 46, XY. A diagnosis of idiopathic unilateral non-chylous pleural effusion was made. The infant was started on feeds on day 2. There was not any change in color and appearance of effusion with feeding. He recovered fully with no recurrence of pleural effusion and was discharged on day 10 of life.

Discussion

Chylothorax or the accumulation of lymph fluid in the pleural cavity is the most common cause of clinically significant pleural effusion in neonates and can be congenital or acquired. Congenital chylothorax can be seen with multiple congenital malformations that result in poor development or obstruction of the lymph system [1].

Spontaneous or idiopathic neonatal pleural effusion is defined as any effusion in a newborn of age less than 30 days,
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without any obvious etiology [2, 4]. Causes of pleural effusion in the newborn include chylothorax, immune and non-immune hydrops, Turner and Down syndromes, congenital pneumonia and congenital heart disease [5].

Idiopathic neonatal pleural effusion is frequently chylous in nature. Simple effusions are known to turn chylous after establishment of external fat feeds [3, 4]. However, in our case, the effusion remained non-chylous even after feeds were initiated. Chylous fluid is milky white in color with triglycerides more than 110 mg/dL (provided there is minimal fat intake), cholesterol between 65 and 220 mg/dL and leukocytosis with absolute cell count greater than 1,000/μL with a lymphocyte fraction greater than 80% [6].

Pleural effusion in the newborn frequently presents with respiratory distress and asphyxia ranging from mild to severe [3]. Early and active resuscitation with intubation and mechan-

Clinical ventilation are needed to establish chest wall expansion [7].

Effusion presenting antenatally acts as a space-occupying lesion and restricts the development of the fetal lungs, which, as a result, may be hypoplastic. Polyhydramnios may result from interference with normal swallowing because of increased intrathoracic pressure [8]. Polyhydramnios was present in the index case but the lungs were normal.

Conclusion

Management includes thoracocentesis followed by intercostal drain insertion. Antibiotics should be given until an infectious etiology has been excluded [7, 9]. Antenatally diagnosed pleural effusions, particularly if present prior to 32 weeks gestation, have a mortality rate as high as 55% [10, 11]. Bilateral pleural effusions are frequently associated with pulmonary hypoplasia. Postnatally, effusions persisting for more than 3 days increase the risk of chronic oxygen dependency [12].

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Conflict of Interest

The authors declare that they have no competing interests.

References

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