Case Report

Pleuropulmonary blastoma, type 1r

Abstract

A 5-year-old female admitted for pneumonia was found to have CXR concerning for right-sided congenital emphysematous lung lesion. She was initially treated with antimicrobials and pediatric pulmonology was consulted regarding normal CXR. Physical examination revealed right chest diminished air exchange with increased anteroposterior diameter. Flexible fiberoptic bronchoscopy demonstrated mucoid impaction of the right main stem bronchi causing ball-valve mechanism leading to air trapping with herniation of the right lung to left. There was also diffuse inflammation of her tracheobronchial tree, distal tracheobronchomalacia, and bronchomalacia of RML and RUL bronchi. Airway clearance was added. CTA revealed severe hyper expansion of the RML with multiple cysts that occupied the majority of her right hemithorax and left mediastinal shift. Lung V/Q scan showed minimal contribution by the right lung of only 11.3% with minimal perfusion. Pediatric general surgery recommended outpatient surgical resection. Two months later she underwent right thoracoscopy, thoracotomy with RML resection and chest tube placement. Operative findings significant for multiple adhesions from the RML to pleura, grossly distended RML which was emphysematous throughout. The patient had resolution of the right pneumothorax within a week postoperatively.

Histopathologic diagnosis of the RML lobectomy and supplying bronchi resulted pleuropulmonary blastoma, regressed type 1r. Given the rarity of this lesion, the case and pathologic specimens were sent for review at the international pleuropulmonary blastoma registry in Minnesota, USA with diagnostic consensus.

Postoperatively, pleuropulmonary blastoma tissue diagnosis raised concern for DICER1 syndrome. Medical genetics recommended DICER1 testing which revealed no identified mutations.

Her care transferred to pediatric oncology. Abdominal and thyroid ultrasonography was negative for tumors, cysts or other abnormalities. The patient was safely discharged home following her postoperative recovery. At follow up CTA revealed no tumors and CXR normalized. Outpatient PFT demonstrated mild obstructive pulmonary defect with reversibility to bronchodilators but did not reveal restriction.

Abbreviations: CXR, chest radiography; RUL, right upper lobe; RML, right middle lobe; CTA, CT with angiography; V/Q, ventilation-perfusion; RLL, right lower lobe; CPAM, congenital pulmonary airway malformation; PFT, pulmonary function testing; PIE, localized pulmonary interstitial emphysema; CDH, congenital diaphragmatic hernia

Case report

A previously healthy 5-year-old female admitted to the Miller Children’s Hospital for right-sided pneumonia was found to have chest radiography (CXR) concerning for right-sided congenital emphysematous lung lesion.1-3 Her presenting symptoms included shortness of breath with activity limitation and dyspnea with limited-to-no perfusion. Physical examination was notable for tachypnea with a maximal respiratory rate of 48 breaths per minute, tachycardia, pulse oximetry (SpO2) of 92%, diminished air exchange at right chest with asymmetric barrel chest (right-left) including increased anteroposterior diameter. Flexible fiberoptic bronchoscopy with bronchoalveolar lavage (FFB BAL) demonstrated mucoid impaction of the right mainstem bronchus causing ball-valve mechanism leading to air trapping with herniation of the right lung to the left. There was mucoid impaction within the right upper lobe (RUL) and right middle lobe (RML) bronchi, tracheobronchomalacia, particularly with the posterior distal trachea, and bronchomalacia of the right middle lobe bronchus and RUL bronchi. The mucosa of the tracheobronchial tree had diffuse inflammation. Airway clearance including beta-2 agonist were added to her inpatient treatment regimen, and further imaging obtained including chest CT with angiography (CTA) and ventilation-perfusion (V/Q) scan. The CTA revealed severe hyper expansion of the RML with multiple cysts that occupied the majority of her right hemithorax. The hyper expanded lobe was compressing atelectatic but otherwise normal appearing RUL and right lower lobe (RLL) parenchyma with severe mediastinal shift to the left. Lung V/Q scan showed minimal contribution by the right lung of only 11.3% with limited-to-no perfusion.

Received: June 20, 2017 | Published: October 13, 2017

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MOJ Anatomy & Physiology
Pleuropulmonary blastoma, type 1r

Citation: Brock J, Nussbaum E, Randhawa I. Pleuropulmonary blastoma, type 1r. MOJ Anat Physiol. 2017;4(3):315–317. DOI: 10.15406/mojap.2017.04.00138

Figure 1CXR at presentation.

Figure 2Diffuse inflammation of the tracheobronchial tree.
Flexible fiberoptic bronchoscopy with bronchoalveolar lavage. Mucoid impaction of the right main stem bronchus, causing ball-valve mechanism leading to air trapping with herniation of the right lung to the left. Mucoid impaction within right upper lobe bronchus, and right middle lobe bronchus. Tracheobronchomalacia, particularly with the posterior distal trachea, and bronchomalacia of the right middle lobe bronchus and right upper lobe bronchus.

Figure 3CT Chest coronal planes-lung and soft tissue windows.

Figure 4CT Chest lung windows.

Figure 5Gross pathology of right middle lobe.

Figure 6Histopathology-low power.

Figure 7Histopathology-high power.
Pediatric general surgery was consulted and was in agreement with highest suspicion of our differential diagnosis of type I congenital pulmonary airway malformation (CPAM). Given the patient was symptomatic with her suspected CPAM, outpatient surgical resection was recommended (lobectomy versus segmental resection, while pneumonectomy is considered in patients with extensive multilobar involvement) following resolution of the acute illness present at initial hospitalization. The patient returned two months post-discharge and underwent right thoracoscopy, thoracotomy with right middle lobe resection with right-sided chest tube placement postoperatively. Operative findings were significant for multiple adhesions from the right middle lobe to the pleura, consistent with chronic inflammation or prior infection, grossly distended right middle lobe which was emphysematous throughout. The patient had resolution of the right pneumothorax within a week of the operation.

Histopathologic diagnosis of the RML lobectomy and supplying bronchus resulted pleuropulmonary blastoma, regressed type 1r. Given the rarity of this lesion (less than 400 cases identified), the entire case and pathologic specimens were sent and gained diagnostic agreement with the international pleuropulmonary blastoma registry in Minnesota, USA.

Postoperatively, the finding of pleuropulmonary blastoma raised concern for DICER1 syndrome, also known as Pleuropulmonary Blastoma Familial Predisposition syndrome. Medical genetics was consulted and the patient underwent DICER1 testing which revealed no identified mutations.

Pleuropulmonary blastoma is a de novo tumor most commonly arising in early childhood. With tissue diagnosis, the patient was transferred to the pediatric oncology service who performed additional imaging to rule out neoplasms in other organ systems including renal and thyroid ultrasonography. No tumors, cysts or other abnormalities were identified and the patient was safely discharged home following her postoperative recovery.

At follow up, CTA has revealed no tumors and CXR has essentially normalized. Outpatient complete pulmonary function testing (PFT) has demonstrated mild obstructive pulmonary defect with air trapping and reversibility to bronchodilators with FVC 91 FEV1 73 FEF25-75% 44 (% predicted), with post bronchodilator FEV1+17% and FEF25-75% +57% change. Body plethesmography did not reveal restriction SVC 93 TLC 118 RV 187 (%predicted) RV/TLC 171%. Diffusion capacity showed no alveolar diffusion deficit DLCO 97 VA 92.

Currently, she maintains stable lung function (FEV1 73% predicted) and is controlled with medium-dose inhaled corticosteroids twice daily.

**Differential diagnosis**

- Congenital lobar emphysema
- Bronchopulmonary sequestration
- Bronchogenic cyst
- Localized pulmonary interstitial emphysema (PIE)
- Pneumatoceles
- Congenital diaphragmatic hernia (CDH)

**DICER1 syndrome associated risk**

- Pleuropulmonary blastoma
- Cystic nephroma
- Ovarian sex cord-stromal tumors
- Multinodular thyroid goiter
- Ciliary body medulloepithelioma
- Nasal chondromesenchymal hamartoma
- Embryonal rhabdomyosarcoma botryoid-type
- Pineoblastoma and pituitary blastoma
- Other rare cancers - Majority of tumors occur before age 40
- Newly discovered syndrome, data lacking on accurate risks.

**Acknowledgements**

None.

**Conflict of interest**

Author declares that there is no conflict of interest.

**References**

