

## **Unusual case of Neonatal Respiratory Distress**

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#### **#** Presented on day 43 of life with

- **#** Noisy breathing on and off since last 15 days
- **#** Difficulty in breathing x 2 days
- **#** Low grade fever x 2 days
- **#** No history of
  - **#** Sudden onset choking
  - **#** Cyanosis
  - **#** Vomiting



- Antenatal history :- was uneventful
- Natal history Birth history :- Vaginal delivery on 1-12-2021 at private hospital with birth weight of 2600 grams.
- \* Post natal history- was admitted for hyperbilirubinemia, discharged



## # Hemogram was done suggestive of

**#** Hb 12.8 mg/dl, TLC 6600 x 10<sup>3</sup> /Ul, (Polymorph 67, Lympho 30), Platelet 2.2 lac

Work up

- # CRP 4.8mg/dl
- **#** Blood culture was sterile
- **#** Baby was started on iv antibiotics



#### # Chest X ray







[Contiguous axial overlapping sections of chest taken extending from lung apex to dome of diaphragm LV. bolus contrast CT reveal the following]

FINDINGS:

Study reveals complete collapse of right lung with obliteration of right main bronchus with soft tissue density. Mediastinal structure are shifted towards right side

Left lung field is appearing normal with smooth, regular pleuroparenchymal mediastinal interface. Segmental bronchi and vessels are unremarkable.

Trachea and major brochii are normal.

Heart and large vessels are normal.

Visualized bones are unremarkable.

No e/o significant mediastinal and hilar lymphadenopathy is seen.

**CT** Report

No e/o pleural/pericardial effusion is seen.

IMPRESSION: STUDY REVEALS COMPLETE COLLAPSE OF RIGHT LUNG WITH OBLITERATION OF RIGHT MAIN BRONCHUS WITH SOFT TISSUE DENSITY S/O COLLAPSE RIGHT LUNG ?DUE TO MUCOUS PLUG/ FOREIGN BODY BRONCHUS ADVISED BRONCHOSCOPY

Please correlate with clinical findings



## **X** Rigid bronchoscopy was done to reveal cause of obstruction of right main bronchus.

**#** Baby came to blk hospital



## Clinically much better, than radiologically





- Weight 5.2kg
- # HR 141/min, RR 53/min
- Sp 02 95% (at oxygen with nasal prongs at 2l/min)
- # Respiratory distress Present (SCR+)
- # Wheeze Present
- No cyanosis / grunt /stridor
- \* No congenital abnormality (facial dysmorphism / skeletal deformity) noted



## # Chest exam

- No obvious chest deformity noted
- **#** Trachea shifted to Rt
- # B/L chest hovements symmetrical
- # AE -- Rt side
- **#** B/L chest conducted sounds

Rest of the systemic examination was unremarkable



## **\*** Right sided collapse

- **#**But why?
- #Viral Infection
- #Leuco-viscidosis (thick secretion)
  - #Primary ciliary dyskinesia
  - **#**Cystic Fibrosis
- #Congenital Lung Anomaly



After reconstruction of images











## Bronchoscopy VIDEO



- Upper trachea after 2-3 rings showed complete cartilaginous rings suggestive of right sided tracheal stenosis
- **#** Could not negotiate scope on right side of lung
- Left sided was normal



## **#** Full term/ Appropriate for gestational age/ Neonatal Hyperbilirubinemia /

**\*** Right sided aplastic lung with congenital tracheal stenosis



- **\*** Stop Antibiotics
- **#** Prognosis explained
- Only treatable Tacheoplasty



DISCUSSION

### Congenital Bronchopulmonary Foregut Malformations

FOREGUT ANOMALIES

Bronchogenic Cyst Esophageal/Neurenteric Cyst Tracheoesophageal fistula/diverticula/stenosis

Sequestration

PARENCHYMAL

Pulmonary/Agenesis/Hypoplasia CLE CCAM Pulmonary Sling Alveolar Capillary Dysplasia VASCULAR ABNORMALITIES

Pulmonary Isomerism

Tracheal Atresia Bronchial Atresia Tracheal Bronchus

**AIRWAY ANOMALIES** 

Acknowledgement: Monica Epelman, M.D.



- Pulmonary Hypoplasia
- # Primary
- Secondary
  - ☆ Intrinsic space occupying lesions
  - # External uterine compressions associated with oligohydrominos (Renal absence)
  - # Chest wall compression (kyphoscoliosis/ Skeletal dysplasias)
  - ♯ Neuromuscular or chromosomal disorders
  - Decreased pulmonary Vascular Perfusion include TOF, Pulmonary atresia, Hypoplastic right heart, Pulmonary vascular anomalies



- \* Aplasia of lung is a rare lung malformation.
- Antenatal scan may not pick up the anomaly.
- \* Antenatally diagnosed Congenital cystic adenomatoid malformation may accompany aplasia of lung.
- \* A newborn with a opaque lung warrants bronchoscopy.
- **CT** scan confirms the diagnosis.
- Constant counselling and communication are essential.
- Regular follow up, prevention of chest infection, immunization and adequate nutrition are the key factors in management



# **THANK YOU**