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fully understood, however It is characterized by the presence of large, thick, mucofibrinous plugs filling the broncho-pulmonary tree, leading to severe respiratory

n manifestations (respiratory distress) for uncommon disorder, as a first serial cases from Middle East Area according to

Case1: A-3 years old girl, who was previously healthy, presented with history of chronic cough for the last 5 months. Physical examination revealed markedly diminished chest movement and breath sound on left side. The diagnosis of foreign body in left main branchus with complete atelectasis of left lung was made. The patient underwent rigid branchoscopy by ENT Surgeon and reported there was cheesy like material occluded the left main branchus.

Case 2: 9-year-old qirl, previously healthy, she suddenly developed shortness of breath and apnea, The patient was deteriorating so she was transferred to PICU and electively intubated for 7 days. Bronchoscopy was done for the patient, and it revealed a thick secretion mainly in the right upper lobe. There was a cast which was seen in the super

then patient expectorated casts of gelatinous material for 4 days. The patient underwent Flexible Bronchoscopy which showed thick secretions but no casts.

Case4: A 3- year - old boy previously healthy presented with history of fever, cough and shortness of breath of 1-week duration. The mother gave history of expectorated whitish material few days prior to his admission. He underwent Flexible Bronchoscopy which showed a thick secretion mainly in the left upper lobe, lingual and right lower lobe. There was a cast which was seen in the left upper lobe. A few casts were expectorated during suctions.

were detected in all our case with flexible bronchoscopy except in the case number 3 his bronchoacopy showed clear airways, however he expectorated a lot of casts during his admission. Histopathology examination of specimens revealed mucus, fibrin, inflammatory exudate with ghost of inflammatory cells, charcot crystals, and eventually, a diagnosis of PB was rendered in all patients. Treatment of plastic bronchitis consists of bronchoalyeolar layage and FB aspiration. Rigid bronchoscopy was required for aspiration and extraction of the material in 3 cases. The patient with congenital heart disease did not require rigid bronchoscopy as he expectorated the all casts before the procedure. Prognosis is generally good, except in cases with congenital heart diseases, in which mortality can be as high as 29%,13 and 41% .16.

All our cases had good progress with satisfactory recovery and no death



Expectorated casts form the patient during his admission



Plastic bronchitis is an uncommon process. However, it must be taken into account in natients with recurrent/ persistent atelectasis and in case of a suspected foreign body. Patients with asthma are at great risk, other risk factors include underlying congenital heart disease, bronchopulmonary aspergillosis, sickle cell acute chest syndrome, and lower airway infections including adenovirus and mycoplasma organisms. Flexible bronchoscopy should be performed early to confirmation and clearance of airway obstructions.

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