



CLINICOPATHOLOGICAL CASE

Postoperative cytomegalovirus necrotizing pneumonia in a preschooler with tetralogy of Fallot

Neumonía necrosante por citomegalovirus en postoperatorio de cirugía cardiaca en preescolar con tetralogía de Fallot

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Summary of the clinical history (A-11-35)

We present the case of a female patient 4 years and 4 months of age referred from a second -level hospital due to complex heart disease and esophageal atresia.

Perinatal and pathological history

The patient was a product of G1 with regular prenatal care. She was born by cesarean section due to oligohydramnios with a birth weight 2400 g, length 48 cm, and Apgar 8/9. Esophageal atresia and cyanosis were noted at birth and she was transferred to a pediatric hospital where a gastrostomy and left cervical esophageal fistula were performed. Extreme tetralogy of Fallot was detected and a Blalock-Taussig fistula was created.

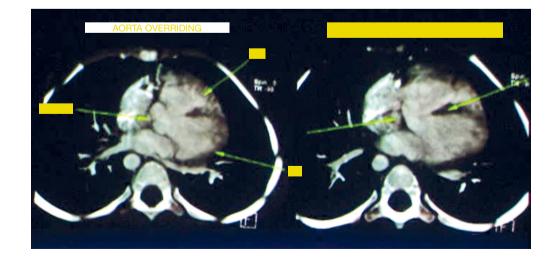
Current illness

March 4, 2009 (Admission). At 1 year and 11 months she was referred to the Hospital Infantil de Mexico Federico Gomez (HIMFG). Outpatient cardiology visit initiated approach with EKG, echocardiogram, chest computed tomography and cardiac catheterization. The case was presented in a medical-surgical meeting to decide if the patient was a candidate for total correction with placement of a valved tube.

August 9, 2010. The patient was electively admitted for the surgical procedure during which a 12-mm right ventricular Hancock tube to right pulmonary confluence was placed, 10 mm ventriculotomy, 7 mm interatrial communication, 2 mm patent ductus arteriosus and ligation. Extracorporeal circulation time was 2 h and 15 min. Clamp time was 1 h and

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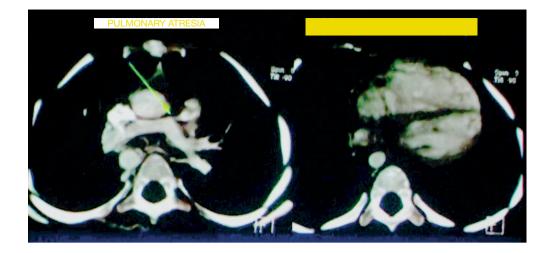




Figure 3 Right aortic arch.

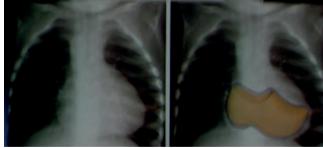
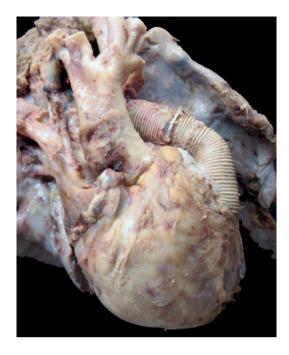


Figure 4 "Swedish boot" image characteristic of tetralogy of Fallot in the AP chest film.



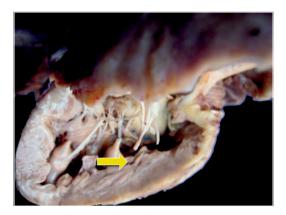


Figure 9 Repair of the 7-mm interventricular communication with a patch of pericardium (arrow).

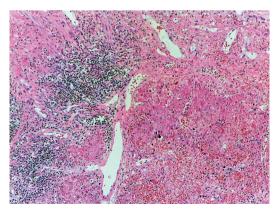


Figure 10 Necrotizing pneumonia. It was not possible to identify the pulmonary morphology due to extensive necrosis caused by the etiologic agent. In addition, areas of recent hemorrhage and inflammatory cell clusters were present.

