

Multiple Cardiac Rhabdomyomas Causing Left Ventricular Outflow Obstruction

Soham Dasgupta^{1*}and Ashraf M Aly²

¹Department of Pediatrics, University of Texas Medical Branch, USA

²Department of Pediatric Cardiology, University of Texas Medical Branch, USA

Corresponding author: Soham Dasgupta, Department of Pediatrics, University of Texas Medical Branch, USA, Tel: 832-693-1917; E-mail: sodasgup@utmb.edu

Received date: September 15, 2016; Accepted date: September 20, 2016; Published date: September 27, 2016

Copyright: © 2016 Dasgupta S, et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution and reproduction in any medium, provided the original author and source are credited.

Clinical Image

A term female infant was born via an urgent C-section for fetal arrhythmia. Postnatally, the heart rhythm was irregular and an EKG showed a normal sinus rhythm and frequent premature atrial complexes. There was cardiomegaly noted on a chest X-ray. A loud systolic murmur was appreciated. An echocardiogram showed multiple myocardial tubers (Figure 1). One tuber was attached to the septal leaflet of the tricuspid valve and another was located right below the pulmonary valve. However, the most significant one was located just below the aortic valve and measured 12×7 mm causing a moderate to severe left ventricular outflow tract obstruction (LVOTO) (50-60 mmHg peak gradient) (Figure 2). A T1 weighted brain MRI showed sub-ependymal hematomas (Figure 3) and cortical tubers consistent with tuberous sclerosis. This was further confirmed by gene sequencing showing a mutation in the TSC-2 gene. The patient is being monitored closely as an outpatient for the possible deterioration of the LVOTO or the development of arrhythmias.



Figure 1: An echocardiogram showed multiple myocardial tubers.



Figure 2: A parasternal long axis view.



Figure 3: MRI showing sub-ependymal hematomas.