Case Report

Coronary artery fistulae are communication between coronary arteries and other structures like cardiac chamber (coronary cameral fistula) or a vein (coronary arteriovenous fistula). Coronary fistulae account for 0.2 to 0.4% of the congenital cardiac abnormalities. Spontaneous closure occurs in 23% of small fistulae, primarily those arising from left coronary system in which conservative management may be appropriate [2]. Though surgery still remains the main stay of treatment, interventional closure seems to be an appropriate choice. We present a 20 day old baby with coronary cameral fistula of the right coronary ostium which was draining into the right ventricle then right atrium and into the pulmonary artery.

Examination of the cardiovascular system revealed normal first and second heart sounds with continuous murmur in the the precordium. Transthoracic Echocardiography revealed situs solitus of d loop ventricles, short axis and parasternal long axis revealed well visualized left coronary ostia but an aneurysmal dilatation of the right coronary ostia (Figures 1 & 2), dilated right atrium (RA) right ventricle (RV) and dilated pulmonary artery (Figures 3 & 4). The patient underwent coronary angiography through the right transfemoral route which revealed aneurysmal dilatation of the right coronary ostium which was draining into the right ventricle then right atrium and into the pulmonary artery (Figure 5). The patient was advised percutaneous coil closure of the fistula but the parents refused for any intervention. The patient was discharged against medical advice.

**Figure 1:** 2D Echocardiography short axis showing the presence of left coronary ostia and dilated right coronary ostia.

**Figure 2:** 2D Echocardiography parasternal long axis showing dilated right coronary cusp.

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**Abstract**

Coronary artery fistulae are communication between coronary arteries and other structures like cardiac chamber (coronary cameral fistula) or a vein (coronary arteriovenous fistula). Coronary fistulae account for 0.2 to 0.4% of the congenital cardiac abnormalities. We present a 20 day old baby with coronary cameral fistula of the right coronary ostium which was draining into the right ventricle then right atrium and into the pulmonary artery.

**Keywords:** Coronary fistulae; Congenital abnormality; Cameral fistulae
Discussion

Coronary fistulae involve communications between a coronary artery with a cardiac chamber (coronary cameral fistula) or a vein (coronary arteriovenous fistula). The coronary fistulae are rare and 60% arise from the right coronary artery. The majority (>90%) of cameral fistulae drain in the right-sided chambers of the heart and the rest into the left side of the heart or to both. The majority of these patients present in adulthood and are usually asymptomatic, most often being detected accidentally. They appear as a persistence of the sinusoidal connection between the lumen of the primitive tubular heart which supplies the myocardial blood flow in the early embryonic period. The pathophysiology of coronary fistula is myocardial stealing or decrease in myocardial blood flow distal to the site of connection [3]. The coronary vessel tries to compensate for the reduction in coronary blood flow by progressive enlargement of the ostia and the feeding artery. The management of cameral fistulae is uncertain due to the rarity of the condition. Surgery is the mainstay of treatment, in a review done [4], 57% of patients were treated surgically with a 0.5% operative mortality rate, with complications including myocardial ischaemia (3%) and recurrence (4%).

Interventional catheterization is yet another therapeutic option available for treatment. The technique uses coils or other devices and can be performed on an outpatient basis [5]. advocated intra-cardiac closure of the fistula as external plication technique had higher recurrence rates. Herein is reported a rare condition of coronary cameral fistula of right coronary artery draining into the right ventricle. The rarity is exponentiated due to the presence of this condition in a 20 day old baby. Various treatments options, interventional and surgical, were available for the management of the baby but sadly could not be undertaken due to unavailability of consent from the parents. Even though adequate management was not possible in this baby, the case report was with notifying in view of its rarity.

References
