Anomalous origin of the right coronary artery from the pulmonary artery in an asymptomatic child

Asemptomatik bir çocukta pulmoner arterden çıkan anomal sağ koroner arter

The abnormally originating right coronary artery from the pulmonary artery (ARCAPA) is a rare congenital anomaly. Although anomalous origin of the left coronary artery from the pulmonary artery (PA) presents in early infancy, symptoms in patients with the ARCAPA may rarely present in infancy. Only a few of them are discovered incidentally in childhood, most of the patients remain asymptomatic and they may be detected at the time of coronary angiography in later adult life. Therefore, the true incidence of this anomaly might be underdetermined.

A previously healthy asymptomatic 8-year-old boy was referred for the evaluation of a continuous murmur detected during school screening program. There were no diagnostic electrocardiographic or chest roentgenographic changes. Echocardiographic examination revealed spherical left ventricle shape with normal ventricular functions. Left coronary ostium was enlarged, but right coronary ostium could not be seen on echocardiographic examination. Selective left coronary arteriography showed retrograde filling of the right coronary artery from collateral vessels and opening to the PA (Video 1 and 2. See corresponding video/movie images at www.anakarder.com). Aortic root injection also showed absent right coronary artery ostium originating from the aortic root.

The surgical correction is always recommended to avoid myocardial ischemia in later life and to abolish left- to- right shunt that causing volume overload. The operative correction in our patient was performed by detaching the anomalous right coronary artery from the PA and re-implantation it to the aorta, so double-ostium coronary system is restored (Fig. 1 and 2). The patient’s postoperative course was uneventful.

Pseudoaneurysm of ascending aorta: a rare complication of mediastinitis following coronary artery bypass surgery

Assandan aortanın psödoanevrizması: Koroner arter baypas cerrahisinden sonra gelişen mediyastinitin nadir bir komplikasyonu

Postoperative mediastinal infection after open-heart surgery via median sternotomy is a devastating complication. A 58-year-old male patient had mediastinitis in the early postoperative period of coronary artery bypass graft operation because of three vessel coronary artery disease. The sternum was reopened; all necrotic and infected tissues were resected. Then sternum was closed with modified Robicsek technique and pectoralis major muscle flap. Broad-spectrum antibiotic was given according to the antibiogram results of drainage fluid. He did well and discharged 20 days after the second surgery. He delayed his control visits and two months later, he was admitted with pulsating sternum and bleeding over the incision line. Chest X ray and computed tomography (CT) of the chest showed enlargement of upper mediastinum and a huge retrosternal pseudoaneurysm originating from the ascending
aorta (Fig. 1, 2). He was taken to the operation theatre urgently and initially femoral artery cannulations were prepared. However, massive bleeding occurred at the time of sternotomy. He died although urgent cardiopulmonary bypass was tried. A huge and ruptured pseudoaneurysm originating from the proximal anastomosis site of saphenous vein graft was observed over the ascending aorta. As a result of this experience, we advice to take a control chest CT two or three weeks after a successful treatment of mediastinitis or on discharge.

Figure 1. Preoperative chest X-ray showing enlargement of the upper mediastinum

Figure 2. Chest computed tomography demonstrating the retrosternal pseudoaneurysm over the ascending aorta, which continues to the anterior side of the sternum with a pathway

RP - Retrosternal pseudoaneurysm, AA - Ascending aorta

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