Anomalous Origin of the Left Anterior Descending Coronary Artery in an Adult

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Abstract

Keywords

► coronary anomalies
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An anomalous origin of the left coronary artery from the pulmonary artery (ALCAPA) is a rare congenital heart disease. Among the variants, an anomalous origin of the left anterior descending coronary artery from the pulmonary artery (ALADPA) is extremely rare. Here, we report a case of ALADPA in an adult that was treated with coronary artery bypass grafting using the left internal thoracic artery.

Introduction

An anomalous origin of the left coronary artery from the pulmonary artery (ALCAPA) is a rare congenital heart disease. Among the variants, an anomalous origin of the left anterior descending coronary artery from the pulmonary artery (ALADPA) is extremely rare, and there have been only a few reports of ALADPA in adults. We report a case of ALADPA in an adult that was treated with coronary artery bypass grafting.

Case Description

A 43-year-old man with chest pain during exertion for several months visited our institution. Physical examination revealed no signs of congestive heart failure. Electrocardiography was normal, but X-ray showed significant cardiomegaly. Transthoracic echocardiography showed normal left ventricular function with an ejection fraction of 60% and no findings of mitral regurgitation or asynergy. Contrast-enhanced computed tomography (CT) revealed that the left anterior descending artery (LAD) originated from the pulmonary artery (PA; – Fig. 1). The left circumflex artery (Cx) originated from the aortic root, similar to the normal left main coronary artery. The right coronary artery (RCA) originated from the aortic root as usual but was abnormally dilated. Angiography showed that the LAD was dilated, tortuous and filled by collaterals from the RCA (► Fig. 2). In addition, the LAD drained to the PA. Myocardial scintigraphy showed redistribution of the anteroseptum wall, suggesting myocardial ischemia in the territory of the LAD. The patient was therefore diagnosed with ALADPA resulting in angina and underwent surgical repair with on-pump beating coronary artery bypass grafting (CABG).

The patient underwent a median sternotomy. The left internal thoracic artery (LITA) was carefully harvested. Cardiopulmonary bypass was established with aortic and right atrial cannulation. Similar to the preoperative diagnosis, the LAD was found to be dilated and originating from the anterior wall of the main pulmonary artery (► Fig. 3A). The origin was easily peeled and double ligated with silk ties in two places (► Fig. 3B). Thereafter, an arteriotomy was performed on the LAD, but there was heavy bleeding from the arteriotomy site due to collaterals from the RCA that became uncontrollable; therefore, the ascending aorta was clamped, and cardioplegic solution was administered. Under cardiac arrest, the LITA was anastomosed to the dilated LAD. The postoperative course was uneventful. The postoperative contrast-enhanced CT was performed on the seventh postoperative day and it showed that...
the bypass graft was patent. After postoperative rehabilitation for 1 month, as usual in our hospital, the patient went home on the 34th postoperative day. Two years after the operation, the patient is doing well without any chest pain and follow-up CT-angiography revealed the graft was still patent (► Fig. 4).

**Discussion**

In the present case, ALADPA was successfully treated with CABG using an LITA graft.

ALCAPA is a rare coronary anomaly, accounting for 0.24 to 0.4% of all congenital heart disease.\(^2\) In some patients with ALCAPA, the LAD independently arises from the pulmonary artery, a variant called ALADPA, it is extremely rare to diagnose ALADPA in an adult because patients with ALADPA usually become symptomatic in the first year after birth. Until now, there have been only a few case reports of ALADPA in adults.\(^1,3\)

In patients with ALADPA, the pressures in the coronary and pulmonary arteries are same in fetal life. However, after birth, the pulmonary artery pressure gradually decreases, causing ischemia of the LAD territory because of low LAD flow. Then, over time, collaterals from the other coronary arteries, normally from the RCA through the septal branches, result in LAD perfusion, depending on the collaterals. Zhang and colleagues stated that the development of collateral circulation from the RCA to the LAD determines the extent of myocardial ischemia,
suggested that the saphenous vein was more useful for the maintenance of flow than the internal thoracic artery (ITA).\(^5\) However, arterial grafting with the ITA can provide better long-term outcome than with a vein graft in general and it may be similar also in ALCAPA patients if short-term patency could be ensured against flow competition. Although ITA patency might be affected by competitive flow from collaterals, Kawasuji and colleagues stated that ITA was acceptable for a moderately stenotic coronary artery.\(^7\) In this case, the ITA graft was patent on postoperative CT, although the LAD was dilated and competitive flow was confirmed. Retrospectively, our decision to perform ITA grafting was not the false one. Additionally, Minamida and colleagues reported that a patient underwent redo CABG using ITA for postoperative vein graft stenosis with ALCAPA.\(^8\)

**Conclusion**

ALADPA in adults is an extremely rare coronary anomaly. To avoid heart failure and sudden death, surgical repair should be performed in asymptomatic patients with ALADPA. Although the surgical strategies are controversial, we recommend establishing a two-coronary system with CABG. Although ITA patency might be affected by competitive flow from collaterals, our experience suggests that ALADPA could be adequately treated with CABG using ITA in selected patients.

**Conflict of Interest**

None declared.

**References**