Percutaneous Transluminal Coronary Angioplasty for Ostial Stenosis of the Left Coronary Artery

—A case report and literature review—

Hae-Yong Lee

A 11-year-old girl developed left main coronary artery ostial stenosis after Takayasu's arteritis for which she underwent Percutaneous Transluminal Coronary Angioplasty (PTCA). The narrowing of the left coronary artery was successfully dilated by angioplasty without apparent complication.

This case suggests that PTCA may have a potential advantage as a temporary method to postpone the aortocoronary bypass surgery in a child with coronary artery stenosis due to Takayasu's arteritis.

Key Words: Takayasu's arteritis, percutaneous transluminal coronary angioplasty, coronary artery stenosis

In Takayasu's arteritis, coronary arterial involvement may be as high as 7% (Lupi-Herrera et al. 1977) by coronary arteriography or at necropsy, young female patients are most commonly affected. In patients with these obstructive lesions, aortocoronary bypass surgery may sometimes be required to preserve ventricular function (Suzuki et al. 1985; Morgan et al. 1987), however, the long term patency rate of the bypass graft remains unsatisfactory (Invert et al. 1988), especially in children. In this report, we present the case of a young girl with Takayasu's arteritis in whom left coronary ostial stenosis was demonstrated by angiography and her anginal pain was relieved by Percutaneous Transluminal Coronary Angioplasty (PTCA).

CASE REPORT

A 11-year-old girl was admitted because of syncope after exercise. She had been well until 6 months prior to admission, when she noted sharp chest pains on exertion that increased progressively. On physical examination, the patient appeared ill-looking with and no evidence of systemic disease. Her blood pressure was 100/70 mmHg and her pulse rate was 88 beats/min in the right upper extremity but the left brachial and radial pulses were not palpable. Blood pressure of lower extremities were 118/72 mmHg in the right lower extremity and 122/72 mmHg in the left lower extremity. Murmur and bruit were not audible on the abdomen. Laboratory studies revealed normal urinalysis, normal blood count, and normal serum lipid levels. Serologic tests were negative for both C-reactive protein and syphilis. The erythrocyte sedimentation rate was 46 mm/hr. All indexes of collagen disorders including LE cell, antinuclear antibody, and rheumatoid factor were negative. The resting electrocardiogram was normal, but, on
treadmill testing, she developed chest pain during mild exercise (Bruce Stage 1, 1.7 mph at a grade of 10%) associated with significant ST segment depression of 2 to 6 mm in leads II, III, AVF, and V5 to V6, together with a decrease in systolic blood pressure (Fig. 1). Selective coronary arteriograms revealed a 90% left coronary ostial stenosis (Fig. 2B) with collateral flow from the right coronary artery to the anterior descending coronary artery (Fig. 2A). The remaining coronary artery tree was normal. Aortic arch arteriogram revealed complete occlusion of the left subclavian artery with well developed collateral from the vertebral artery and ascending cervical artery (Fig. 2C). There was no abnormal angiographic finding in the thoracic and abdominal aorta. The angiographic evidence suggested a potential risk of sudden death by acute myocardial infarction and therefore, PTCA was performed. A balloon angioplasty catheter was introduced by a 7-F guiding catheter (SCIMED, Co Ltd., Maple grove, Minnesota, U.S.A.) with a 0.14" steerable guide (ACS, Co Ltd., Temecula, California, U.S.A.) wire. The balloon was inflated with diluted contrast material to 6 and 9 atm pressure. The duration of inflation was approximately 80 seconds at each dilation, and the procedure was repeated twice. Repeat angiography after PTCA showed a 40% residual left coronary ostial stenosis (Fig. 2D). Transient depression of the ST segment in lead II, III and AVF was observed during the procedure. No complications were encountered during and after the procedure. Corticosteroid therapy (prednisone, 10 to 30 mg/day for 6 months) was begun, with resultant normalization of the ESR. Follow up treadmill
testing 2 months after the procedure revealed electrocardiographic changes during moderate exercise (Bruce Stage 5, 50 mph at a grade of 18%) associated with significant ST segment depression of 2 to 4 mm in leads II, III, AVF, and V₅ to V₆ and no chest pain developed. The post PTCA clinical course was uncomplicated and the patient remains asymptomatic over a year following balloon angioplasty.

**DISCUSSION**

Takayasu’s Arteritis is an inflammatory disorder of the aorta with unknown specific etiology, and the fibrous intimal proliferation produces narrowing or obstruction of main branches of the aorta, including coronary artery ostia. Coronary artery disease is a rare involvement of Takayasu’s arteritis. The incidence of coronary artery involvement has been reported to be 8.5% in Korea (Park et al. 1992) and 9% in Japan (Matsubara et al. 1992), and is observed mainly in autopsy cases because coronary artery disease is usually not evident until the occurrence of angina pectoris or myocardial infarction, or after the onset of congestive heart failure. From 1961 to 1991, 68 patients have been reported to undergo operations for coronary artery disease resulting from Takayasu’s arteritis. Operative mortality
was eight (12.3%) and operation was repeated in two patients because of graft failure (2.8%) (Amano and Sujuki, 1991). The long term patency of coronary artery bypass graft (CABG) is 78% in Takayasu’s arteritis (Kanoh et al. 1992).

Recently, balloon angioplasty of the aorta in Takayasu’s arteritis has produced good results both immediately after and on long term follow up. Commonly, though not invariably, a small intimal tear or stretching that is observed after successful angioplasty may underlie the mechanism of angioplasty in Takayasu’s arteritis (Sanjay et al. 1992). No aneurysm developed at the site of angioplasty after balloon dilatation of the aorta in patients with Takayasu’s arteritis, in contrast to occasional aneurysms that are observed after angioplasty for coarctation of the aorta. This could be due to this markedly thickened vessel wall caused by inflammatory fibrosis of all three layers as observed in Takayasu’s arteritis (Virmamani et al. 1986). To our knowledge, this is the first case of PTCA for coronary artery stenosis in Takayasu’s arteritis. Hitherto, PTCA in Kawasaki disease has been reported. In one case who had undergone PTCA for coronary artery stenosis in Kawasaki disease, death resulted due to the obstruction of the left coronary artery immediately after balloon rupture caused by a high inflating pressure. The other two were successfully dilated (Echigo, 1988; Toshihito et al. 1991). Pathologic findings are different between Kawasaki disease and Takayasu’s arteritis. In Kawasaki disease, pathologic findings of the vessel is marked intimal thickening, so intimal tearing during angioplasty may rupture of vessel directly, but there may lesser vessel rupture in Takayasu’s arteritis because of fibrotic thickening of all three layers of the vessel.

This case suggests that PTCA may well be an effective alternative method to aortocoronary bypass surgery for coronary artery stenosis in Takayasu’s arteritis.

REFERENCES

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