Surgical angioplasty of the left main coronary artery in non-atherosclerotic lesions

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Abstract
Surgical angioplasty of the left main coronary artery confers several advantages over conventional bypass surgery: unrestricted forward flow is provided to the entire coronary bed and graft material is spared. The literature contains many reports of surgical angioplasty of atherosclerotic stenoses. The technique is described in five patients with non-atherosclerotic disease of the left main coronary artery: three children (a 7 year old girl who had undergone an arterial switch operation shortly after birth; a 9 year old boy with congenital supravalvar aortic stenosis; and a 10 year old girl with Kawasaki's disease) and two adults (a 51 year old woman with post-radiation stenosis; a 53 year old man with acute dissection). All patients had an uneventful recovery and are free from symptoms with a widely open left main trunk. Although technical difficulties are increased in these patients, excellent results can be achieved with this approach.

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The haemodynamic characteristics restored after angioplasty of a stenotic left main coronary artery are superior to those obtained after conventional aortocoronary bypass to the major branches of the left coronary arterial tree.1 2 Because ideal conditions are rarely fulfilled and the repair is technically demanding, surgical angioplasty of the left main coronary artery is not widely used.1 7 The procedure is generally restricted to myocardial revascularisation in patients with atherosclerotic lesions. We report here its application to other groups of patients with anticipated increased technical difficulties.

Patients and technique
Five patients with non-atherosclerotic disease of the coronary arteries underwent surgical angioplasty of the left main coronary artery between 1995 and 1998. The operation was performed with the aid of cardiopulmonary bypass and during a period of cold blood cardioplegia. The patients or their physicians were contacted at the end of 1998 to update follow up.

CASE 1
A newborn baby with d-transposition of the great arteries underwent an arterial switch operation. The coronary arteries originated from the corresponding facing sinuses. They were harvested and reimplanted with a button of aortic wall on the neo-ascending aorta. The Lecompte manoeuvre of the great arteries was performed, and the neo-pulmonary artery, reconstructed with autologous pericardium, was sutured without tension to the pulmonary bifurcation. The child did not have signs of myocardial ischaemia. Left ventricular ejection fraction was normal and contraction homogeneous. The child grew normally.

At age 7 years she started to complain of effort angina pectoris. Angiography revealed a stenosis at the implantation of the left coronary artery. Reoperation revealed dense pericardial adhesions. The left pulmonary artery was transected at the pulmonary bifurcation to allow mobilisation of the main pulmonary artery and access to the left coronary artery. The origin of the left coronary artery was dissected away from scar tissue that was responsible for the stenosis. A longitudinal incision was carried out in the coronary artery from its implantation on the aorta to a few millimetres distal to the stenotic segment and a patch of autologous saphenous vein used to enlarge the origin of the coronary artery. The left pulmonary artery was sutured back on the pulmonary bifurcation. Recovery was uneventful. The child, now 10 years old, is free from symptoms.

CASE 2
A 9 year old boy underwent correction of a diffuse form of congenital supravalvar aortic stenosis, that extended from the aortic root to the beginning of the descending aorta.4 Angiography also revealed a stenosis at the origin of the left coronary artery trunk. The left ventricle was hypertrophic with normal systolic function. At surgery, the beginning of the stenosis on the aorta ascendens was sharply demarcated. The aortic root was remodelled with separate patches of pericardium that were used to enlarge the sinotubular junction between all three commissures. In the
left coronary sinus, the incision was extended to the roof of the main coronary artery, beyond the ostial stenosis. In this sinus, the patch enlarged the sinotubular junction and relieved the coronary ostial stenosis. Finally, the ascending aorta was enlarged with a stripe of pericardium. Postoperative course and follow up (extending to 15 months) were uneventful. Echocardiography revealed absence of residual gradient along the left ventricular outflow tract and aorta, and a wide open left main coronary artery.

CASE 3
A 10 year old girl was admitted because of increasing angina pectoris. Eleven months earlier she had been diagnosed with Kawasaki’s disease. Coronary angiography showed giant aneurysms punctuated by severe stenosis on the left main coronary artery, proximal third of the left anterior descending artery, and proximal third of the right coronary artery. The left ventricle had a normal function without contraction abnormality. At surgery, the aneurysms were opened longitudinally and the incision was prolonged beyond the stenosis. The excess wall of the aneurysm was resected and a patch of autologous saphenous vein was used to enlarge the outlet stenoses. The arteriotomy was closed over a 3 mm probe to harmoniously calibrate the artery diameter. Because additional stenoses existed on the proximal left anterior descending artery, the left internal mammary artery was harvested and implanted on its distal half. Postoperative course was uneventful. Angiographic control at three months showed a non-restrictive left main trunk with rapid filling of the circumflex and proximal left anterior descending artery, and a patent internal mammary artery filling the distal left anterior descending artery. The child is free from symptoms 15 months after surgery.

CASE 4
A 51 year old woman presented with increasing angina pectoris 13 years after radiotherapy of the left chest for a breast carcinoma. Coronary angiography revealed a high grade stenosis of the left main coronary artery and an occlusion of the right coronary artery. The mammary arteries were hypotrophic. At surgery, the tissues were adherent to each other. After tedious dissection of the pulmonary artery, the left main coronary trunk was exposed up to its bifurcation. It was opened longitudinally and patched with a strip of autologous pericardium. A venous bypass was performed on the right coronary artery. Postoperative course and follow up (extending to 24 months) were uneventful. Coronary angiography showed a patent left main trunk and rapid opacification of the coronary arteries (fig 1).

CASE 5
A 53 year old man underwent aortic valve replacement with a mechanical prosthesis. One year later, while digging in his garden, he felt excruciating pain in the chest owing to acute type A aortic dissection. ECG showed no signs of ischaemia, and transoesophageal echocardiography showed normal ventricular function. After aortic cross clamping, the ascending aorta was transected above the aortic root. The dissection was circumferential and involved the origin of both coronary arteries. During mobilisation of the intimal flap, it appeared that a 2 cm long sequester of intimal tissue, originating from the left coronary artery trunk, was flapping in the false lumen of the aorta. The adventitial layer of the dissected coronary artery trunk was very thin with several tiny holes. The pulmonary artery was extensively dissected and retracted to permit complete exposure of the coronary artery trunk. Full thickness of the coronary artery wall was found in the middle of the trunk, about 12 mm proximal to the bifurcation. The trunk was opened longitudinally along its anterior surface up to the bifurcation. A segment of autologous saphenous vein was appropriately bevelled and end-to-end anastomosed to the coronary trunk. The aortic valve prosthesis was replaced with a composite graft. The layers of the dissected origin of the right coronary artery was fixed together with fibrin glue. Both
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coronary arteries were reimplanted in the graft, and the graft was sutured distally onto the aortic arch. Postoperative course was uneventful and the patient resumed his active life. Regular chest computed tomography showed no dilatation of the dissected aorta. Magnetic resonance imaging three years later showed unrestricted flow in the left main trunk.

**Discussion**

Restitution of normal blood flow through the left main coronary artery confers advantages over multiple and sequential bypass surgery to more distal branches. Occlusion of the main coronary trunk is prevented, antegrade perfusion to the entire coronary bed provided, competitive flow avoided, and bypass material spared. In a young patient, who may later recur with recurrent coronary artery disease, these advantages are substantial. Conventional percutaneous transluminal angioplasty may still be performed, or, if surgery is required, the risk of damaging a bypass at sternum re-entry or provoking coronary embo-lisation during dissection of a stenotic bypass is avoided. Finally, reconstruction of the main coronary trunk provides the shortest and most efficient way for blood to leave aorta (which acts as a volume capacitance) to the myocardium. The shortness and relatively large size of proximal native coronary arteries, in comparison to any bypass conduit, provide optimal conditions for myocardial perfusion under maximal stress as occurs during intense exercise.

Surgical angioplasty of the left main coronary trunk was attempted in the early days of cardiac surgery.3–7 The procedure, however, fell into disgrace because of frequent dire complications and unpredictable results, while venous and arterial aortocoronary bypasses became safe and reproducible.8 A few groups, however, showed that surgical angioplasty of the left main coronary artery provides excellent results when stringent selection criteria and rigorous surgical technique are applied.9–12 Patients with isolated, non-calcified lesions confined to the proximal half of the coronary trunk are the best candidates.

We never abandoned the procedure in our atherosclerotic patients, and we offered it to other patients who, despite enhanced technical difficulties, would particularly benefit from the ideal haemodynamic characteristics restored.13–16 Three of our patients were children with the prospect of a long life. Coronary artery bypass with the internal mammary artery has been successfully performed in children.17–19 Although the long term outlook of arterial grafting to the coronary arteries is excellent in adults, it may not be optimal in the young, especially if the child later indulges in intense physical activities.

The availability of a suitable arterial conduit was the main problem in our case 4, because the internal mammary arteries were hypertrophic after radiotherapy. A patch of autologous vein would have theoretically been superior to autologous pericardium for the angioplasty, but intrinsic fibrinolytic properties and, in this case, absence of possible radiation damage. A piece of thin, macroscopically normal pericardium found on the right side of the heart was, however, used. In case 5, surgical angioplasty of the coronary trunk avoided bypass to the coronary artery branches, which would have been difficult to find in a reoperation and without the aid of preoperative coronary angiography. Furthermore, the use of pedicled internal mammary arteries to graft coronary arteries may be dangerous in a fresh aortic dissection. The dissection frequently involves the arch arteries and may prevent adequate flow in their branches.20

The excellent clinical course of our patients validate the use of surgical angioplasty of the left main coronary artery in non-atherosclerotic lesions. Longer follow up will tell if the repair remains effective.

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