Combined percutaneous balloon dilatation of the aortic valve and coronary angioplasty

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SUMMARY A 75 year old man with severe angina caused by aortic stenosis and coronary artery disease was considered to be unsuitable for cardiac surgery after the recent removal of a bronchial carcinoma. Combined percutaneous balloon dilatation of the aortic valve and right coronary angioplasty considerably ameliorated the patient’s angina.

Case report

A 75 year old man had a two year history of exertional angina and dyspnœa that was getting worse despite treatment with propranolol and bumenidine. Left heart catheterisation in May 1985 demonstrated an aortic valve gradient of 60 mm Hg. Coronary angiography showed a discrete 90% stenosis in the mid portion of the dominant right coronary artery (fig 1) and several areas of stenosis in the left anterior descending artery with a maximum narrowing of 90%. On admission in November 1985 for aortic valve replacement and aortocoronary bypass surgery a preoperative chest x-ray demonstrated a mass in the upper lobe of the right lung. Cardiac surgery was cancelled and at lobectomy a well differentiated adenocarcinoma was removed; no local spread of tumour was found. No evidence of tumour recurrence was found in the following months but the angina became progressively worse, occurring when he walked across a warm room and at rest.

In September 1986 percutaneous coronary angioplasty and balloon dilatation of the aortic valve were attempted. The heavily calcified aortic valve was approached from the brachial artery and crossed with an exchange guide wire and progressively dilated with 10, 14, and 16 mm balloons (Meditech). Each balloon was inflated three times for 20 seconds at a pressure of 3–5 atmospheres. The femoral artery pressure was recorded continuously and fell slightly during balloon inflation. The gradient was reduced to 35 mm Hg (fig 2). Aortography demonstrated no change in the minimal aortic regurgitation. A bronchial guide catheter could not be positioned in the right coronary artery, so a femoral guide catheter was used. Narrowing of the right coronary artery was reduced from 90% to 30% with an ACS ultra-low profile 3.0 mm balloon. No attempt was made to dilate the diffusely diseased left coronary artery. The patient tolerated this 130 minute procedure and was discharged after three days.

One year later he has considerably improved effort tolerance. Apart from chest pain on walking up steep hills, he is free of angina.
Some patients with severe aortic stenosis and angina have clinically significant coronary artery disease. Our case shows that combined balloon dilatation of coronary stenosis and the aortic valve can produce worthwhile improvement in symptoms. Such palliative procedures may be appropriate in patients who are considered to be unsuitable for cardiopulmonary bypass surgery.

The duration of the combined procedure was prolonged because we could not obtain a stable position for the guide catheter in the right coronary artery from the brachial approach; but the patient remained comfortable throughout the procedure. The best sequence of balloon dilatations is uncertain. An initial coronary angioplasty would improve myocardial perfusion and might reduce the hazards of balloon dilatation of the aortic valve.

References

