Intraaortic Balloon Pump Support During Coronary Angioplasty. Initial Experience

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Objective - To evaluate the use of the intraaortic balloon (IAoB) in association with coronary angioplasty in high-risk patients.

Methods - Fourteen high-risk patients unresponsive to clinical therapy and with formal contraindication to surgical revascularization were treated by coronary angioplasty. All procedures were performed with circulatory support with the IAoB. This study reports the early results and the late findings after 12 months of follow-up. Six patients had multivessel coronary disease; of these, four had left main equivalent lesions and two had unprotected left main coronary artery disease, one of whom had severe “end-vessel” stenosis and the other was a patient with Chagas’ disease with single-vessel lesion. Eleven patients had a left ventricular ejection fraction <30%.

Results - In 100% of the patients, the procedures were initially successful. Two patients had severe bleeding during the withdrawal of the left femoral sheath. At the end of twelve months, 4 patients were asymptomatic and the others were clinically controlled. There were two late deaths in the 7th and 11th months.

Conclusion – The combined use of the intraaortic balloon pump and percutaneous coronary angioplasty in high-risk patients with acute ischemic syndromes provides the necessary hemodynamic stability to successfully perform the procedures.

Keywords: circulatory support, coronary angioplasty, intraaortic balloon pump.

Interventional cardiology has significantly progressed in the last 20 years, mainly in regard to percutaneous myocardial revascularization techniques. The advances in the development of instruments used to perform coronary angioplasty combined with the advent of new devices, such as stents, have allowed the use of this therapy in most patients

The concomitant development of techniques for circulatory support has also occurred, in particular the intraaortic balloon (IAoB), which is being increasingly used in patients with ischemic heart disease. The use of the IAoB, which required surgical placement, was initially restricted to cases of cardiogenic shock and preinfarction angina. Later, with percutaneous insertion, the complications caused by surgical handling of the femoral artery were greatly reduced, facilitating the use of this technique in intensive care units and, particularly, in catheterization laboratories.

Although widely employed, coronary angioplasty has become prohibitive in patients with severe ventricular dysfunction or in patients with lesions in coronary arteries that irrigate a significant area of viable myocardium.

Thus, the concomitant use of both methods has allowed the performance of percutaneous revascularization in high-risk patients, in which surgery is contraindicated, while achieving adequate circulatory stability.

The purpose of this study was to assess the early experience at our institution with coronary angioplasty performed with circulatory support with the IAoB in high-risk patients.

Methods

From April 1996 to August 1998, 14 patients with clinical findings of acute coronary syndromes unresponsive to clinical therapy underwent coronary angioplasty with circulatory support with the IAoB. After evaluating each individual case, surgical therapy was contraindicated mainly due to associated complicating factors, such as
significantly reduced left ventricular function (EF<30%) in 13 patients, severe pulmonary emphysema in 3 patients, and acute renal failure in 3 patients. Four patients were more than 70 years old, and 3 patients had undergone previous revascularization surgery.

In 10 patients, the procedure was complemented by the placement of 12 stents. Every patient gave written consent. All procedures were performed through the bifemoral approach. We chose the left femoral artery for the implantation of the IAoB and the right femoral artery for the placement of angioplasty catheters.

After the arteries were punctured and the sheaths inserted, the patients received 5,000U of sodium heparin. Subsequently, the IAoB was advanced through a retrograde arterial path with a flexible guide wire up to the correct position, immediately below the origin of the left subclavian artery. Then, the balloon was connected to the module, adjustments in the pressure curve were made and the electrocardiogram was monitored. Our clinic uses Datascope System 90 equipment (Datascope Corp., 15 Law Drive – CN 4011, Fairfield, New Jersey, USA). According to the manufacturer’s instructions, the balloons employed had a caliber of 9.5F and a volume of 40cc when inflated, as the patients ranged in height from 1.60m to 1.83m.

A few minutes after the beginning of the counterpulsation, coronary angioplasty was performed, according to routinely used methods. In patients undergoing stent placement, we followed our routine, which initially consists of precise definition of the lesion, mainly in relation to its extent and diameter of the vessel. A stent/vessel ratio of approximately 1.0 to 1.2 was adopted as ideal. After the liberation of the prosthesis, there must not be any residual lesion greater than 5% and it must cover all the lesion/dissection and must not restrict an important branch. Finally, control angiographies in a series of projections and the prescription of medications consisting of ticlopidine (500mg/d for 30 days) and acetylsalicylic acid (200mg/d for 6 months) are required.

Soon after the procedure, the rate of inflation of the balloon was progressively reduced, until its disconnection. The femoral sheaths were kept in place after all the instrumentation was removed.

Subsequently, the patients were referred to the intensive care unit (ICU) for 24-hours. After this period, both femoral sheaths were removed and hemostasis with manual compression, as well as concomitant control of activated clotting time (ACT), was performed.

All patients were considered high-risk patients: their left ventricular ejection fraction (EF) was <30%, the impaired vessels irrigated more than 50% of the viable myocardium, they had lesions in the left main coronary artery, with or without protection, or they had lesions equivalent to those of the left main coronary artery. Almost all patients had very poor left ventricular function (EF <30%), 6 had multivessel lesions, 4 had lesions equivalent to those of the left main coronary artery and 2 had severe lesions in an unprotected left main coronary artery. The 2 remaining patients had single-vessel lesions; 1 had a severe lesion in a final artery and the other had dilated cardiomyopathy as a result of Chagas’ disease.

The patients with multivessel disease underwent angioplasty of the culprit artery, which was confirmed by clinical and laboratory tests. These arteries also fulfilled the criterion of irrigating a critical viable myocardial mass (fig. 1). The 2 patients with severe disease of the left main coronary artery received tubular stents. (fig. 2).

The tubular stainless steel stents employed were of the Palmaz-Schatz type, by Johnson and Nir (Sci-Med).

After discharge, the patients were followed up on an outpatient basis or with telephone interviews for 12 months. Exercise testing at 3 months and control angiography at 6 months were not performed in all patients.

Results

All procedures were successful. There were no cases of hypotension or any sign of circulatory instability. None of the patients reported significant chest pain during the inflation of the angioplasty balloon.

Two patients had significant bleeding at the left inguinal region, in spite of coagulation control. This bleeding was corrected with intravenous protamine, prolonged inguinal compression and blood replacement.

The mean hospital stay was approximately 6 days. During the 6 months of follow-up, 6 patients showed recurrence of angina pectoris of slight intensity, which was easily controlled with oral drugs. In 7 patients with signs and symptoms of heart failure in addition to symptoms of chest pain before hospitalization, the symptoms recurred and were controlled by changing the oral therapy. Four patients became asymptomatic up to 6 months with the use of anti-platelet agents only. There were two late deaths as a result of sudden death after 7 and 11 months of follow-up.

After 12 months, 7 patients had controlled stable angina, 8 experienced heart failure, which was also medically controlled, and 4 were asymptomatic.

Discussion

The morphological type of coronary lesion is known to play a major role in the outcome of angioplasty. The use of the IAoB in these patients has 2 beneficial effects: the 1st occurs as a result of inflation of the balloon and consequent increase of the aortic diastolic pressure,
leading to a significant increase in coronary flow; the 2nd occurs as a result of deflation of the balloon, which creates a negative pressure in the descending aorta, significantly reducing the resistance to the left ventricle out flow. Thus, the first effect increases oxygen supply and the second effect reduces oxygen consumption.

Currently, there are several indications for the use of the IAoB in cardiac catheterization laboratories (table I).

First, it is electively used in angioplasties of high-risk patients, as in those in our series. These patients usually have decreased ventricular function and a considerable extent of viable myocardium at risk.

Second, the IAoB is employed when new devices, such as atherectomy catheters, which perform tissular ablation, are used. These devices may cause embolic phenomena during their operation, which may lead to subsequent circulatory instability, depending on the importance of the vessel. Directional atherectomy and the percutaneous extraction catheter may lead to these complications especially in the presence of coronary thrombosis, as occurs in unstable angina and acute myocardial infarction (AMI). Rotational atherectomy (rotablator), however, is the procedure that bears the strongest relation to these complications, as it is the most frequently employed. In approximately 10% of the cases in which the rotablator is used, the slow-flow and no-reflow phenomena are noted, and they usually persist for hours or days, leading to myocardial dysfunction. If the treated vessel is important and if there is previous ventricular dysfunction, serious consequences may arise. That is why prophylactic implantation of the IAoB in these selected cases is important.16,17

Acute coronary occlusion during angioplasty may also lead to circulatory failure when a major vessel is treated. Use of the IAoB for this type of complication offers the beneficial effects of hemodynamic stability and increased perfusion through the occluded vessel.18

Unstable angina, as well as AMI and its complications, are particularly improved by aortic counterpulsation. Currently, almost all cases of unstable angina resolve with the use of medications such as beta-blockers, anti-platelet agents, heparin and new substances, such as abciximab. However, in some cases, even the combined use of these drugs does not lead to stabilization of clinical findings. Refractoriness to medication and its potential risk interfere with the invasive management of these patients with the IAoB. Thus, after the patient is hemodynamically stable, elective coronary angiography is performed. The latter will show the coronary anatomy and guide the management strategy.19
In AMI, the reduction of the afterload caused by the IAoB attenuates the effects of the decreased local coronary perfusion and increases the blood flow through the occluded artery. Thus, the use of this procedure combined with thrombolytic therapy seems to significantly reduce the incidence of reocclusion of the culprit artery of the infarction. The fact that the incidence of bleeding complications is not increased is also of note.

Primary or rescue coronary angioplasty has similar effects. A series of reports have shown the benefit of the IAoB in reducing the incidence of post-angioplasty reocclusion. Many clinics use the IAoB routinely in these 2 scenarios, when the infarction is associated with signs of severe left ventricular dysfunction or when the culprit artery remains occluded in spite of all measures taken.

In cardiogenic shock, sole use of the IAoB has not lead to a significant improvement in the prognosis. Concomitant surgical or percutaneous revascularization is required.

The same is true for the mechanical complications of AMI. In addition to correction of the ventricular septal defect (VSD) or replacement of the mitral valve, revascularization is also indispensable. The IAoB temporarily provides the adequate hemodynamic stabilization required for the complete treatment of the patients.

**Table I - Intraaortic balloon in the cardiac catheterization unit**

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<th>Scenario</th>
<th>Description</th>
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<tr>
<td>1 - Elective use</td>
<td>- &quot;High-risk&quot; angioplasty</td>
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<td>- Atherectomies (rotablator)</td>
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<td>2 - Emergencies</td>
<td>- Unstable angina</td>
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<td>- Postangioplasty acute occlusion</td>
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<td>- Acute myocardial infarction with thrombolytic agents</td>
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The IAoB is also employed in nonischemic emergencies, such as mitral regurgitation caused by idiopathic rupture of chordae tendineae or in acute bacterial endocarditis. In these patients with previous ventricular function within the normal range, reduction in the afterload leads to reduction in the amplitude of the “V wave” and of the pulmonary capillary wedge pressure.

Finally, the IAoB is being increasingly used while patients wait for heart transplantation. Patients with heart failure refractory to therapy usually require circulatory support for several weeks until a heart is available for transplantation. Considering the different devices for circulatory support, the IAoB is the most adequate, as it is user-friendly and is associated with fewer side effects.

Lately, the complications associated with the use of the IAoB have been extensively examined. The overall incidence is around 10% of the cases and, currently, severe ischemia of the lower limbs occurs in less than 1% of the procedures. \(^{27,28}\)

Patients most likely to present with complications are those with severe hypertension, diabetes, females and, in particular, those with peripheral vascular disease.

Recently, a progressive decrease in the number of complications has been noted. This decrease has resulted from the reduction in the caliber of the balloons, as well as from the use of instruments with increased flexibility and percutaneous introduction without the use of sheaths.

In conclusion, the availability of the IAoB has stimulated high-risk procedures, with hemodynamic stability and a low rate of complications. Larger studies are needed to determine the cost-benefit ratio of this procedure in this group of patients.

References