Obstructed tilting disc mitral valve prosthesis associated with placenta praevia

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Successful pregnancy in women who harbour prosthetic heart valves is well documented (Bennett and Oakley, 1968; Barnard et al., 1969; Laros et al., 1970; Buxbaum et al., 1971; Reid and Barclay, 1971). Closed cardiac surgery in pregnancy was reported as long ago as 1952 (Brock, 1952; Logan and Turner, 1952). Later reports established that it was possible, in pregnancy, to undertake valve replacement with the adjunct of cardiopulmonary bypass with fetal survival. In a series of 20 cases, Zitnik et al. (1969) described a fetal mortality of 33%, while the maternal mortality was 5%, not higher than in non-pregnant patients. It is probable that the relatively non-pulsatile nature of blood flow during bypass, often with a low mean pressure, changing blood gas tensions, and disturbances of serum biochemistry, produces changes in uteroplacental perfusion and thus ultimately fetal oxygen supply, and thereby contributes to the high fetal mortality.

By a relatively simple technique of monitoring fetal heart rate with an ultrasonic transducer, Koh et al. (1975) were able to show that fetal distress, reflected by bradycardia, could in at least one instance be reversed by increasing pump output.

Case report

In 1972 a 24-year-old woman underwent mitral valve replacement with a Lillehei-Kaster prosthesis. Her mitral valve was a fixed, stenotic diaphragm unsuitable for valvotomy. When she was routinely reassessed 18 months later, as part of a study of Lillehei-Kaster valves, the prosthetic valve was thought at angiography to be functioning normally, although a mean diastolic gradient of 12 mmHg across the valve orifice was noted, which was considered to be high. The patient was, however, asymptomatic.

Eighteen months later she returned to hospital 35 weeks pregnant, with a history of having been unwell for the preceding six days with urgent dyspnoea, palpitations, cough, and ankle oedema.

On examination there was tachycardia, hypertension (85/60 mmHg), and a raised jugular venous pressure. There was a 3/6 pansystolic murmur, which radiated to the axilla, and triple rhythm. The electrocardiogram (ECG) showed sinus rhythm with an axis of +60°, a P-R interval of 0·24 sec, and depressed ST segments in leads V1 to V6. A diagnosis of an obstructed disc valve was made.

Shortly after admission vaginal bleeding was noticed and an obstetrician was consulted. Preparations were made for urgent cardiopulmonary bypass for replacement of the clotted valve. The diagnosis of placenta praevia was made. At this juncture the fetal heart was still audible and the presentation was cephalic.

As a preliminary to the induction of anaesthesia with a view to Caesarean section, arterial and central venous pressure lines were introduced under local anaesthesia, and arterial blood pressure, central venous pressure, pulse rate, and ECG were continuously monitored. The fetal heart was again thought to be audible.

Anaesthesia was then induced with the patient slightly laterally tilted, with a small dose of keta-
mine and suxamethonium, followed by rapid intubation of the trachea. When the lower uterine segment was incised, maternal blood pressure was always low, fell further and an isoprenaline infusion was started. A fresh stillborn child was delivered and, because the mother’s condition had now seriously deteriorated, an attempt was made rapidly to establish femorofemoral bypass. Cardiac arrest occurred before the groin vessels had been cannulated and attempts at resuscitation failed.

**Comment**

In an active cardiac surgical unit, painfully aware of the rapidity with which patients deteriorate when prosthetic valves become clotted and obstructed, and experienced in the rapid establishment of cardiopulmonary bypass for further valve replacement, the recognition and management of this common complication of valve surgery did not present a problem. However, when we were faced with this well recognised problem, further complicated by vaginal bleeding related to type I placenta praevia, there arose the need to establish priorities.

The pregnancy was valuable to the mother and the fetus was viable. Heparinisation was thought likely to precipitate massive bleeding and this, in an acutely ill patient, was judged likely to be fatal. Caesarean section, necessary for maternal and fetal survival, was therefore given priority.

In retrospect, a more logical course would have been to autotransfuse the patient with the blood which may have accumulated in the peritoneal cavity at Caesarean section. Autotransfusion in obstetrics is well documented. Highmore (1874) described its use in postpartum haemorrhage, while Miller (1973) discussed autotransfusion in the management of ectopic pregnancy.

Stehling et al. (1975) described the indications for and contraindications to the use of this technique. The indications listed were blunt or penetrating trauma to the chest or abdomen, ruptured ectopic pregnancy, elective surgery associated with large blood loss, and rare blood types or antibody peculiarities complicating the cross-matching of blood. The contraindications to the use of autotransfusion were gastrointestinal tract contamination, concomitant distant fractures or cerebral trauma in respect of the use of intravenous heparin, and the presence of cancer cells in the operative field.

As a precaution against microemboli-induced pulmonary changes, a 20–40 micron filter should be included when using autotransfusion. Prolonged use of suction should also be avoided, since further destruction of the blood occurs. In units not accustomed to the use of cardiopulmonary bypass, the stringent precautions necessary to avoid air embolism should be carefully observed.

Faced with this problem again, we would establish femorofemoral bypass with its attendant necessity for heparinisation. The fetus would be carefully monitored by recording fetal heart rate, ECG, and the acid-base status of the mother. In our case, the excessive bleeding would have been managed by means of autotransfusion until Caesarean section was performed, followed by replacement of the obstructed mitral prosthesis.

**References**


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