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Bilateral congenital coronary artery fistula

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Bilateral congenital coronary artery fistula may be defined as a condition in which more than one coronary artery gives rise to a fistulous tract. Twenty such cases have been reported.¹⁻⁶ Only two of them were diagnosed angiocardiographically and corrected surgically.⁶ A case studied by selective coronary arteriography and operated upon successfully is presented here.

Case report

A 38-year-old man was admitted for investigation of precordial pain on exertion, of one year's duration, which had increased in the previous month. On admission the heart rate was 72, blood pressure 160/100 mmHg, there were no signs of congestive heart failure, and findings on physical examination were normal. The electrocardiogram and thoracic radiograph were normal. Selective coronary arteriography demonstrated aneurysmal dilatation of the left anterior descending coronary artery at its origin, with complete obstruction distal to it. The distal segment of this artery was filled via a wide collateral vessel from the left circumflex artery which showed multiple narrowings. Multiple vessels appearing late after the injection filled a fistulous tract draining into the main pulmonary artery (fig 1).

The right coronary artery showed severe obstructive lesions, and filled, via multiple branches, a fistulous tract draining into the main pulmonary artery (fig 2). At operation, both fistulous tracts were ligated and aorto-coronary bypass grafts to the left anterior descending, left circumflex, and right coronary arteries were inserted. The patient was released from hospital after an uneventful postoperative course of 21 days.

Discussion

Congenital coronary artery fistula may be defined as a communication of a long coronary artery with one of the atria or ventricles, the coronary sinus, superior vena cava, or the pulmonary trunk. It is an uncommon congenital cardiac lesion, about 400 cases having been reported since the first description by Krause in 1865.8 Its estimated incidence is 0.4% of the cases with congenital heart disease.

Reviewing 172 cases of congenital coronary artery fistula, McNamara and Gross⁹ found four cases of Address for reprint requests: Professor HN Neufeld, Heart Institute, Chaim Sheba Medical Center, Tel Hashomer, Israel.

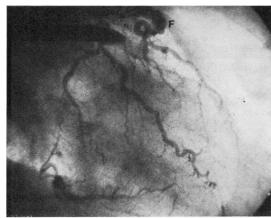


Fig 1 Selective left coronary arteriography in the right anterior oblique projection. The left anterior descending artery is completely occluded, and the circumflex artery shows obstructive lesions. Multiple vessels, appearing late, fill a fistulous tract (F) which drains into the main pulmonary artery.

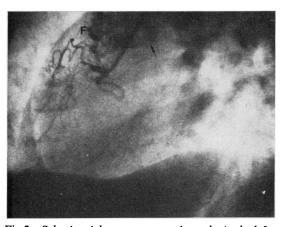


Fig 2 Selective right coronary arteriography in the left anterior oblique projection. The right coronary artery shows severe obstructive lesions, and fills via multiple branches a fistulous tract (F) which drains into the main pulmonary artery (arrow).

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bilateral fistula, involving both coronary arteries. In 1968 Ogden¹ found 11 such cases. Another case was found at necropsy by Rose.⁴ Two other cases were diagnosed angiocardiographically but not corrected surgically.² ³ Two others were corrected surgically.⁶ In our case bilateral fistulae, involving tracts from the right and left coronary arteries were present, and were probably a contributing factor to the patient's chest pain. It is possible that the increased coronary flow predisposed to the relatively early development of severe coronary atherosclerosis in this patient. The fistulous tracts were ligated and the patient has remained asymptomatic for six months since. A similar case also diagnosed angiocardiographically and treated surgically was reported by Gupte et al⁵ in 1979.

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