Pericarditis due to *Bacteroides melaninogenicus* secondary to a teratoma

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Over the past 50 years the aetiology of purulent pericarditis has changed. Most cases used to be caused by Gram positive bacteria, but now there is a higher incidence of cases due to Gram negative bacteria and fungi.1,2 Anaerobes are rarely found.3 In the following case *Bacteroides melaninogenicus*, an anaerobic Gram negative bacillus, caused a purulent pericarditis secondary to a perforating teratoma.

**Case report**

A 17 year old girl was admitted with signs and symptoms of acute pericarditis with tamponade. Apart from an upper respiratory tract infection 8–10 days earlier she had been well.

The chest radiograph showed a small pleural effusion and an enlarged cardiac silhouette. An echocardiogram showed a pericardial effusion and about 250 ml fluid was subsequently removed at pericardiocentesis. Culture of the fluid yielded an anaerobic Gram negative bacillus after three days' incubation. This was later identified as *B melaninogenicus* and found to be sensitive to metronidazole but resistant to penicillin.

The patient was treated with intravenous metronidazole, 500 mg thrice daily, for 12 days and her condition gradually improved. Nineteen days after admission, however, signs of constrictive pericarditis developed and a total pericardiectomy was performed.

At operation a cystic tumour measuring 8 × 6 × 5 cm was found in the right mediastinum. It was attached to the thymus and communicated with the pericardium. The tumour, thymus, and pericardium were excised. On histological examination the tumour was found to be a teratoma; there was no evidence of malignancy. The postoperative period was uneventful.

**Discussion**

This is the first report of isolation of *B melaninogenicus* from a case of purulent pericarditis. This organism is a normal inhabitant of the upper respiratory tract. It is slow growing, requiring 72 hours' incubation for visible growth, and will be missed if cultures are incubated for only 40–48 hours.

The pathogenesis of the present case, in which a mediastinal teratoma perforated into the pericardial sac, is very unusual. Mediastinal teratomas, accounting for 10–20% of all tumours of the mediastinum in children and young adults,4 have been estimated to perforate into the pericardium in less than 1% of cases.5 Formerly such events were diagnosed at necropsy.

The first report of a successfully treated case was published in 1966.6 Since then three rather similar cases have been described,7,8 though in none of these were bacteria or fungi identified in the contents of the pericardial sac.

**References**