“The plug and the bug” – a case of coronary embolism in bacterial endocarditis

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Summary

Systemic embolism is a classic complication of infective endocarditis. Coronary involvement and acute myocardial infarction (MI) are rare and increase mortality significantly. Recognising this unusual entity is crucial to provide adequate care. Percutaneous coronary intervention and thrombus aspiration is preferred to thrombolysis, which classically increases intracerebral haemorrhage risk. The present article describes the case of an acute inferior ST-elevated MI due to a \textit{Streptococcus salivarius} endocarditis in a patient with known bicuspid aortic valve.

Key words: bacterial endocarditis; septic embolism; coronary embolism; myocardial infarct

Case report

A 79 year-old female with prior history of hypertension and moderate aortic insufficiency due to bicuspid aortic valve presented to the emergency department with ST-elevation myocardial infarction (MI). She complained of acute retro-sternal chest pain and left arm radiation of 2 hours duration. The presentation followed a one-month period of fatigue, weight-loss and progressive dyspnea. Physical examination revealed an acutely ill lady. Non-invasive blood pressure was 150/60 mm Hg on both sides; pulse regular at 100 beats per minute; respiratory rate at 20/minute; saturation 97% under 2L/min oxygen and temperature was 37.9 °C. Significant signs were a rapid upstroke followed by quick collapse of peripheral pulses, an enlarged hyperdynamic cardiac apical impulse, a blowing pandiastolic decrescendo murmur in the left upper sternal border, a midsystolic ejection murmur with carotid radiation and no extra heart sounds. Pulmonary auscultation was consistent with bilateral basal crackles. Chest radiograph revealed a marked cardiomegaly and dilation of the aortic knob and root but no signs of widened mediastinum or pulmonary congestion. Troponin Ic was 0.35 \textmu g/L, white blood cell count of 13.5 G/L with normal differential and a C-reactive protein of 60 mg/L, and the rest of the laboratory work-up was unremarkable.

On arrival, the 12-lead electrocardiogram (ECG) showed normal sinus rhythm, first-degree atrioventricular block and a 2 mm ST-elevation in leads II, III avF, V5 and V6 (fig. 1A). Therapy was initiated with aspirin 500 mg, heparin 5000 U and clopidogrel 600 mg. An urgent coronary angiography revealed the distal occlusion of a marginal branch of the circumflex artery but otherwise normal coronary arteries (fig. 1B). The aortogram showed significant ascending aorta dilatation (fig. 1C) as well as severe aortic valve regurgitation. The left ventricular ejection fraction was deemed normal with inferolateral hypokinesia. Considering the history and presentation, a septic embolus was thought to be the cause of the occlusion and, due to vessel tortuosity, distal lesion and spontaneous symptom alleviation no intravascular treatment was attempted. Peak troponin Ic was 12.3 \textmu g/L and total CK 545 U/L. Four different peripheral blood cultures grew penicillin sensitive \textit{Streptococcus salivarius} (minimum inhibitory concentration =\textless 0.125 mg/L). A trans-oesophageal echocardiogram demonstrated a calcified and moderately remodelled bicuspid aortic valve with no evidence of vegetation or abscess and a normal mitral valve. All other findings were in line with the angiographic assessment. The abdominal scan showed a superior splenic infarction and the brain MRI was consistent with multiple, bilateral, embolic lesions with secondary haemorrhage (fig. 1D). The retina, skin and other organs were free from embolic complications. The patient was treated with i.v. ceftriaxone 2 g daily for 4 weeks, all subsequent blood cultures were negative and she remained symptom-free. The severe aortic regurgitation warranted a close cardiological follow-up and to date, the patient has had no further complaints.
CASE REPORT

Discussion

In 1856, Rudolf Virchow was the first to describe coronary embolism as a complication of bacterial endocarditis. Autopsy studies in the mid 20th century revealed that it was found in up to 40% of patients with fatal bacterial endocarditis [1]. Coronary involvement leading to MI is nevertheless a rare condition with retrospective studies accounting for incidences of 0.3 to 3% [2, 3]. The prognosis is difficult to quantify, but there is...
a clear trend towards excess mortality when compared to those without septic coronary embolism [4]. Virtually any described microorganism causing endocarditis may be involved but Streptococci are thought to be the most commonly isolated pathogen [5].

Certain angiographic features such as normal coronary arteries, abrupt termination of blood flow or absence of collateral vessels may suggest embolism [6]. The left-anterior descending (LAD) coronary artery is by far the most commonly involved. It is noteworthy to remember that not all MIs are due to coronary embolism. Other conditions have been described such as direct obstruction of coronary ostia due to aortic cusp vegetation, severe aortic insufficiency with low diastolic pressures or para-aortic abscess compression of the left main coronary artery [7].

In 1972, Pfeifer was the first to describe a successful surgical reperfusion of the LAD in a patient with MI due to septic coronary embolus [8]. Thrombolysis was later attempted with, over the years, periodic case reports of fatal intracerebral haemorrhages probably due to undetected cerebral mycotic aneurysms, such that it is now considered a relative contraindication [9, 10].

The most convincing management seems to be the percutaneous coronary approach with, when possible, embolus aspiration. However, this is not without potential adverse consequences, as angioplasty, with or without stent deployment, may lead to mycotic aneurysm or pannus migration [9, 11, 12]. Moreover, stenting in an infected area could lead to early stent thrombosis [13]. The optimal treatment strategy has yet to be determined but it is unlikely, given the small amount of cases, that any will be based on solid evidence. We would advise coronary angiography to be considered as the first approach.

This case illustrates that when confronted with myocardial infarction in unusual settings, one should think of endocarditis as an embolic source. In doing so, the patient benefits from prompt and vital antibiotics but caution should be drawn when attempting reperfusion, as adverse consequences lie behind the perilous path of each chosen strategy.

References