We report the case of a 36 years old man who presented with an incidental echocardiographic finding of a 1.1 × 0.8 cm mobile structure on the mitral valve associated with mild to moderate eccentric jet of mitral regurgitation. Surgical excision confirmed the diagnosis of nonbacterial thrombotic endocarditis (NBTE), formerly known as marantic endocarditis. The cardiolipin IgG and IgM antibodies and the lupus anticoagulant were strongly positive confirming the diagnosis of antiphospholipid syndrome. The patient was promptly anticoagulated.

CASE PRESENTATION

A 36 year old asymptomatic man was referred to our clinic for further assessment and management of hypertension. He denied fevers, weight loss, rigor, or sweating. He was on treatment for anxiety and depression, and chronic pain resulting from accidental division of the digital nerve of the right index finger and also the radial nerve. He was on treatment with Diclofenac, Gabapentin, Tramadol, Lansoprazole. Olanzapine, Venlafaxine and Perindopril 4 mg od. On examination he was apyrexial, the pulse was 66 bpm and the supine blood pressure was 159/109 mm Hg. The serum sodium was 141 mmol/L, potassium 5.0 mmol/L and creatinine was 86 umol/L. Furthermore, the Hb was 13.0g/dl and the CRP 9 mg/L. The resting ECG confirmed sinus rhythm with rare unifocal ventricular ectopic beats. The transthoracic echocardiogram confirmed a 1.1 × 0.8 cm mobile and echogenic structure on the atrial surface of the posterior mitral valve leaflet with mild to moderate eccentric jet of mitral regurgitation. There was mild left ventricular hypertrophy and the other valves were structurally normal. The above findings were also confirmed on transoesophageal echocardiography. In view of the diagnostic dilemma the patient underwent a surgical excision of the above structure, and the mitral valve was repaired with a bovine pericardial patch. He made an uneventful recovery. Histology, however, revealed a mass composed of fibrin and thrombus with occasional endothelial cells suggestive of non-bacterial thrombotic endocarditis (NBTE) or marantic endocarditis (Figure 1).

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DISCUSSION

Non-infective thrombotic endocarditis (formerly known as marantic endocarditis) is a condition characterized by sterile vegetations on the endocardium and mainly affects the cardiac valves. It consists largely of fibrin and platelet aggregates and without an inflammatory response and is associated with numerous diseases including autoimmune disorders and in particular hypercoagulable states like SLE or in patients possessing the antiphospholipid antibody (aPL). Other conditions associated with NBTE include mucin producing adenocarcinomas, haematological malignancies, acquired immune deficiency syndrome, disseminated intravascular coagulation (DIC) and chronic conditions like tuberculosis, trauma from indwelling right sided cardiac catheters or central lines, snake bites, burns, overdose and late effect of thoracic radiation treatment (1-3). There is no literature associating NBTE with administration of specific drugs (Figure 2).

Hypercoagulability with the associated venous or arterial thromboembolism was first described in 1865 by Trouseau as a presenting sign of malignancy (4). Adenocarcinomas of the lungs and the ovary constitute 50% of the cases and haematological malignancies account for approximately 25% (5). Zeiger in 1888 considered the lesions to be bland thrombi deposited on cardiac valves, and coined the word thromboendocarditis (6). This was subsequently described as marantic endocarditis due to its association with malignant and wasting diseases (7). At autopsy Libman and Sacks in 1924 described the above findings in a large number of patients with systemic lupus erythematosus(up to 40%). Gross and Friedberg in 1936 subsequently used the word nonbacterial thrombotic endocarditis (8). Angrist and Marquiss described the association between systemic emboli and this condition likely to be the result of either the superficial nature of these lesions or the lack of cellular organization (9). The diagnosis its mostly reached at post-mortem with an incidence of between 0.3% to 9.3% in the adult population (10,11).

Nonbacterial thrombotic endocarditis - Not forgotten

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Figure 1) Transesophageal echocardiogram illustrating the intracardiac vegetation identified by a circle: it shows a large, nodular echodensity on the atrial aspect of the posterior mitral valve leaflet. AV Aortic valve; LA Left atrium; LV Left ventricle; RV Right ventricle
Furthermore, the incidence of visceral embolism (the majority being cerebral) is between 14-90% (12).

NBTE commonly affects those between the 4th and 8th decades of life and the diagnosis can be confused with culture negative endocarditis or the presence of fibroelastoma on the cardiac valves. With the advent of echocardiography more asymptomatic lesions are being diagnosed during life. The vegetations are mostly located on the left-sided heart valves (majority on the mitral valve) and cases of NBTE involving the right-sided heart valves are rare. The mural endocarditis, chordae tendineae and papillary muscles may also be involved. Atrial myxomas or papillomas rarely arise from the cardiac valves. Fibroelastomas on the other hand usually arise from the surface of the valves and represent <10% of primary cardiac tumours.

There are no ante mortem clinical findings or pathognomonic signs and symptoms that allow for NBTE and the diagnosis is made either at post-mortem (13) or as an incidental finding on echocardiography. Negative blood cultures, the absence of clinical signs of infection and a normal thrombophilia screen are also important clues. Spontaneous venous thromboembolism is also more likely to occur in the presence of NBTE. However, the presence of a disease process known to be associated with NBTE, the presence of a heart murmur or the presence multiple systemic emboli (in several cerebral territories on diffusion-weighted MRI) may point towards the above diagnosis. On the other hand single or focal lesions and limited infarction are more characteristic of infective endocarditis. Both thrombotic and infective endocarditis may occur without the above diagnoses and pose a diagnostic dilemma. We propose that in patients with high likelihood of NBTE a transthoracic echocardiogram should be considered to assess for the above diagnosis. Surgical intervention may expedite the diagnosis and significantly reducing the risk of life threatening embolic events.

The management of NBTE relies on the correction of the underlying disease process. Anticoagulation with vitamin K antagonists remains the mainstay treatment for NBTE in patients with SLE and positive antiphospholipid antibodies. In patients with malignancy and associated DIC, with the use of intravenous heparin is supported by the available evidence. In this group of patients the use of warfarin is not as effective (14-16). Furthermore, in patients with malignancy (17,18) and asymptomatic valvar vegetations, but without DIC, the role of anticoagulation has not been proven, and the use of either warfarin or heparin should be balanced against the bleeding risks. Surgical excision of the vegetations may be considered in those patients with potentially curable cancers (19-21) in those with severe regurgitant valves, recurrent thromboembolic events (22) despite adequate anticoagulation and, as a clinical case demonstrates, in those patients where the diagnosis is uncertain. Surgical preservation of the affected valve should be the preferred surgical approach.

NBTE is a rare condition associated with advanced mucin secreting malignant tumours and hypercoagulable states. It is associated with high incidence of embolic events. The diagnosis is usually reached at post-mortum. Our case clearly demonstrates that malignant vegetations on the cardiac valves may occur without the above diagnoses and pose a diagnostic dilemma. We propose that in patients with high likelihood of NBTE a transthoracic echocardiogram should be considered to assess for the above diagnosis. Surgical intervention may expedite the diagnosis and significantly reducing the risk of life threatening embolic events.

REFERENCES