Background

Since the seminal observation by Armand Trousseau in 1865, numerous studies have confirmed the common association between cancer and thromboembolism. The etiology of the hypercoagulable state in cancer is multifactorial. Contributing factors include procoagulant alterations associated with the malignancy (tissue factor and cancer procoagulant expression by tumor cells, vascular compression by tumor masses) and the host’s inflammatory response (monocyte tissue factor expression, increased levels of factor VIII, fibrinogen, and von Willebrand factor). Although venous thromboembolism is the most common manifestation, cancer-associated thrombophilia may also present as a migratory superficial thrombophlebitis (the original syndrome described by Trousseau), arterial thrombosis, disseminated intravascular coagulation (DIC), a thrombotic microangiopathy or nonbacterial thrombotic endocarditis (marantic endocarditis).

Case presentation

A 48-year-old woman with no significant past medical history, previously asymptomatic, who suddenly showed a lowered level of consciousness, was admitted to the department of intensive care unit for stroke. She had no history of tobacco use, consumed an occasional alcoholic beverage, and did not have any recent nonsteroidal anti-inflammatory drug use. The computer tomography (CT) of the brain confirmed multiple embolic infarctions throughout both cerebral hemispheres and cerebellum. The distribution of the signal abnormalities was compatible with multiple embolic infarcts. Cardiovascular examination revealed intact carotid and distal pulses without any bruits, regular rhythm with no cardiac murmurs. ECG showed normal sinus rhythm with no ischemic changes. Chest x-ray was normal. Transthoracic and transesophageal echocardiography were significant for multiple vegetations on a trileaflet aortic valve associated with moderate aortic regurgitation (Fig. 1 and Fig. 2). During hospitalization, the patient had worsening neurological and repeated CT of brain control with evidence of new ischemic lesions with embolic origin, despite he was treated with IV heparin. Findings were consistent with a new embolization resulting in multiple infarcts. Given the worsening of the clinical condition the patient underwent CT of the abdomen/pelvis (Fig. 3) that showed presence of a mass in the tail of the pancreas. Initial laboratory data were notable for a complete blood count, basic metabolic panel, liver function test, coagulation studies, amylase, lipase, cardiac enzymes, triglycerides, and erythrocyte sedimentation rate: all within normal limits.

Nonbacterial Thrombotic Endocarditis in Pancreatic Cancer

Endocardite trombotica non batterica in associazione con una neoplasia pancreatica

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Nonbacterial thrombotic endocarditis (NBTE), known as marantic endocarditis, is a phenomenon due to hypercoagulability with a complex pathogenesis. Originally described by Ziegler, the lesions of NBTE were considered to be fibrin thrombi deposited on normal or superficially degenerated cardiac valves [1]. Numerous reports have identified the relationship between NBTE and a variety of different inflammatory states, including chronic diseases like malignancy and autoimmune disease [2, 3]. NBTE is a serious manifestation of prothrombotic state that is characterized by the deposition of thrombi on previously undamaged heart valves in the absence of a bloodstream bacterial infection and by the increased frequency of arterial embolic events in patients with chronic debilitating diseases. Although hypercoagulability is often seen in patients with pancreatic cancer, NBTE has rarely been reported antemortem. We report a case of marantic endocarditis in patient with pancreatic cancer, in which neurological symptoms preceded the diagnosis of pancreatic cancer.

Keywords: Nonbacterial thrombotic endocarditis, pancreatic cancer, echocardiography.

Workup for bacterial endocarditis was negative, including multiple sterile blood cultures. In suspected endocarditis associated with antiphospholipid syndrome, the patient has performed immunological tests, but they were negative for antinuclear, anticardiolipin, antimitochondrial, and antismooth muscle antibodies. Additional laboratory findings showed CEA of 660 ng/mL, CA 19.9 of 47 U/mL and CA 125 level of 721 U/mL. Pancreatic enzymes, erythrocyte sedimentation rate, C-reactive protein, remained within normal limits. The diagnosis of marantic endocarditis was made.

The patient required admission to the intensive care unit for stroke. She was treated with aspirin, beta blocker therapy, and unfractionated heparin. Definitive therapy includes antitumor therapy directed at the underlying malignancy and indefinite anticoagulation with unfractionated or low molecular weight heparin. Vitamin K antagonists such as warfarin are ineffective and should not be employed in management. In our experience, prompt diagnosis and treatment with heparin and treatment of the patient’s pancreatic cancer may prevent catastrophic neurological events and decrease morbidity.

The patient continued on heparin iv and neurological symptoms worsened. Unfortunately, she continued to deteriorate from the pancreatic cancer and died 6 weeks after this hypercoagulable state was diagnosed. This case demonstrates a rare presentation of advanced pancreatic adenocarcinoma. Marantic endocarditis is likely underreported and underdiagnosed. Cancer is associated with a significant hypercoagulable state that increases the risk of thromboembolism fivefold. NBTE is an uncommon but devastating manifestation of malignancy-associated thrombophilia. NBTE should be considered in any cancer patient suffering an episode of arterial thromboembolism. TEE is useful imaging modality in the diagnosis of NBTE. The long-term outcome for most patients with malignancy-associated NBTE is poor, because it is typically associated with disseminated and incurable malignancies. Health care professionals should be alert to the possibility of marantic endocarditis in any patient with cancer, especially pancreatic cancer, who presents with symptoms of neurological dysfunction or an arterial thrombotic event.

Discussion

NBTE is a phenomenon due to hypercoagulability from an underlying cause. Clinical presentation is most often a sudden neurological deficit. It has been associated with a variety of inflammatory states including malignancy. It is the result of a prothrombotic condition characterized by deposition of sterile fibrin and platelet-rich thrombi on previously undamaged heart valves. Cardiac function is generally unaffected. Associated morbidity and mortality are due to increased frequency of arterial embolic events, specifically in the central nervous system [5]. NBTE is characterized by the presence of an amorphous mixture of platelets and fibrin on a valve leaflet in the absence of inflammation or micro-organisms, or destruction of the underlying cardiac valve [6]. It is most often diagnosed only at postmortem examination because
the small, friable vegetations frequently embolize, leaving remnants too small to be identified by echocardiography. Vegetations caused by NBTE are found in approximately 1.2% of patients during autopsy. Although the reported incidence of systemic emboli varies widely, on average 42% of patients embolize [7]. Arterial embolization most frequently involves the spleen, kidney, brain, and heart, respectively [8]. It is also the major cause of morbidity and mortality in these patients. The most common sites of vegetations are on the mitral and aortic valves along the coaptation line [9]. However, the tricuspid and pulmonary valves as well as bivalvular involvement have been identified [10, 11]. The great majority of underlying neoplasms associated with NBTE are adenocarcinomas of the pancreas, colon, lung, and prostate. The pathogenesis of NBTE is incompletely understood. The lesions of NBTE are classically found in areas of high flow on valvular leaflets; therefore, blood flow likely contributes to the location if not the initiation of these valvular lesions. Elevated levels of circulating cytokines associated with cancers, such as tumor necrosis factor or interleukin-1 may also result in local tissue damage that instigates vegetation formation. In addition, the interaction between these cells activates the coagulation cascade leading to the generation of thrombin and thrombosis [12]. While NBTE has been reported in every age group, it most commonly affects patients between the fourth and eighth decades of life. No sex predilection has been reported [13]. Therefore, if a diagnosis of endocarditis is made, but the cultures and serology are negative, and there is no response to antibiotic treatment, then NBTE should be strongly considered. Patients with newly acquired murmurs should be evaluated with two-dimensional echocardiography to elucidate the presence of valvular vegetations. The preferred diagnostic test for NBTE is a TEE, which is more sensitive in the detection of valvular vegetations than the transthoracic approach [14]. Although no established treatment strategies exist for tumor related NBTE and DIC, there are limited data supporting the use of anticoagulation in these patients. The most effective anticoagulant appears to be unfractionated heparin, which has been shown to be effective in reducing the incidence of recurrent episodes of thromboembolism [15, 16]. When delivered in therapeutic doses, both i.v. and s.c. heparin therapy has been effective. In contrast to heparins, vitamin K antagonists such as warfarin should not be used in patients with malignancy-associated NBTE, as recurrent thromboembolic events while on warfarin are common. A multicenter, randomized trial by Lee et al. [17] demonstrated low-molecular weight heparin was more effective than warfarin in reducing the risk of recurrent thromboembolism in patients with cancer without increasing the risk of bleeding. Although the exact reason for warfarin’s inability to control the coagulopathy associated with NBTE is not precisely known, the presence of non-vitamin K dependent agents that induce the thrombotic coagulopathy has been suggested by some investigators [18]. Anticoagulation must be continued indefinitely in patients suffering from NBTE, because recurrent thromboembolism has occurred in patients following discontinuation of heparin therapy. Although no trial has compared unfractionated heparin with low-molecular weight heparin in patients with tumor associated NBTE, a multicenter double-blind trial published by Sakuragawa et al. [19] demonstrated that low-molecular weight heparin can be used as an alternative to unfractionated heparin in patients with DIC. Although most patients do not require surgery to address NBTE valvular lesions, cardiac surgery is a reasonable intervention in selected circumstances where the risk-benefit balance is favorable.

References


