Case report

Acute paraparesis due to acute aortic thrombosis in young adult a mysterious case

Gaurav Singhal¹, Vijay Pathak², Sunil Sampley³, Pradeep Meena⁴, Manish Ruhela⁵, Rahul Choudhary⁶

¹⁵Senior Resident (Dept of cardiology,SMS hospital,jaipur)
²Professor (Dept of Cardiology,SMS hospital,jaipur)
³Senior Resident (Dept of Cardiothoracic and Vascular Surgery,SMS hospital,jaipur)

ARTICLE INFO

Keywords:
- Aortoiliac
- Paraplegia
- Young adults
- Thrombosis

ABSTRACT

The occlusion of the Aorta is a rare event, which is potentially life threatening. Acute onset paraplegia is rare event due to sudden aortic thrombosis. We present the case of a 38-year-old man who was referred to the emergency department of our hospital because of sudden onset bilateral lower limb weakness and abdominal pain. He had no history of any disease. He was a non-smoker, non-alcoholic and had no family history. A computerized tomography angiography of abdominal aorta and bilateral lower limb revealed the thrombosis of the abdominal aorta and of both iliac arteries. After 18 hour of admission he died due to ventricular tachycardia and ventricular fibrillation.

Introduction

Acute abdominal aortic occlusion is an infrequent, vascular condition that requires emergency surgery. It should be managed promptly due to its fatal outcome. Thrombosis of abdominal aorta is usually insidious onset. Most cases of aortic thrombosis occur in fifth to sixth decades of life but the condition has been reported in patients as young as 29 years.¹ In majority of cases thrombosis is secondary to atherosclerosis or embolism. Less commonly, thrombosis may occur in the sac of an aneurysm and after syphilitic aortitis,² pressure by tumors,³ pelvic peritonitis, and irradiation to the abdomen and/or cardiovascular procedure and trauma. The following description of a case of aortic thrombosis occurring in a young adult man is of interest because the thrombosis without atherosclerosis and any predisposing condition, is rarely reported.

CASE REPORT

A 38-year-old male who was admitted to our hospital with sudden onset paralysis of the bilateral lower extremities and abdominal pain. His physical examination revealed that his femoral pulses were both nonpalpable, the bilateral lower extremities were cold and pallor. There was history of pain abdomen radiating to the back. There was no history of bladder and bowel involvement. There was no history of any similar complaint in the past. There was no history of diabetes mellitus, hypertension, tuberculosis, coronary artery disease, trauma, or surgical intervention in the past. The patient was a non-smoker and non-alcoholic.

On examination, his pulse rate was 100/min, regular, and equal in both upper limbs. His bilateral lower limb pulses, dorsalis pedis, popliteal and femoral were absent. His respiratory rate was 20/min at the time of examination. His blood pressure was 170/100 mmHg in the right arm in supine position. Systemic examination was not significant.

Routine investigations revealed hemoglobin was 11.5 gm/dl, total leucocyte counts were 22000 per mm³, platelets counts 3.59 lakh per ml, and his blood urea was 16.86 mg/dl, creatinine 1.08 mg/dl, and serum sodium was 141.2 mEq/L, serum potassium was 5.19 mEq/L. Serum cholesterol was 246 mg%, random blood glucose was 112mg%, Urine analysis showed frequent red blood cells without glucose and albumin. ECG was normal. CT angiography of abdominal aorta and bilateral lower limb showed thrombus in abdominal aorta with iliac arteries extension. Figure 1. Figure 2 shows aortic thrombosis with right renal infraction. Other investigations including 2-D Echo, protein C and S, were all negative. Homocysteine levels were normal. PT and aPTT were normal.

After 6 hour of admission patient developed difficulty in breathing. ECG was showing hyperacute T waves. Serum potassium was done which revealed 9.6 mEq/L. Anti-hyperkalemic treatment started immediately. Initial urine output which was 300 ml was not increased. Nephrology consultation was taken for dialysis but patient died due to ventricular tachycardia and fibrillation.

* Corresponding Author: Gaurav Singhal
Senior Resident (Dept of cardiology, SMS hospital, Jaipur

Copyright 2010 BioMedSciDirect Publications. All rights reserved.
Discussion

Acute abdominal aortic occlusion is a condition that requires emergent surgery. High morbidity and mortality rates occur even when treated in a timely manner. Embolic occlusion and abdominal aortic thrombosis (in situ thrombosis) of the previously diseased aorta are the main pathophysiological mechanisms which underlie this condition. Dossa et al. reported that heart disease and female gender were risk factors for embolization while smoking, diabetes, and a history of intermittent claudication were risk factors for thrombosis. In our case, the patient was male with no risk factors and predisposing conditions.

Although occlusion may occur in any segment of the aorta, the terminal part of the abdominal aorta distal to the renal arteries (infrarenal), especially the iliac bifurcation portion, is the most frequent site of the occlusion. In our patient total thrombotic occlusion was found in the infrarenal abdominal aorta and iliac arteries.

Most emboli originate from the heart, especially from the left atrium, in the setting of atrial fibrillation. Atrial myxoma, bacterial or fungal endocarditis, and prosthetic valve thrombus are infrequent causes of embolization. In our case patient was in sinus rhythm and 2D ECHO was normal. The patient had no other possibilities to account for the embolization. In situ thrombosis occurs in the setting of severe atherosclerotic occlusive disease in approximately 75% of the cases and is precipitated by conditions such as dehydration, diabetic ketoacidosis, and heart failure that slow the circulation at the vicinity of the stenosis.

Different clinical presentations may be observed in patients with involvement of arterial branches of the abdominal aorta. A full-blown case is characterized by the sudden onset of sharp pain located mainly in the lumbar area, abdomen, perineum, and bilateral lower extremities. Later in the course, weakness, numbness, paresthesia, and dysesthesia occur and in the end, paralysis of the lower limbs may dominate the clinical picture. In the literature, only a few cases have been found that present with the sudden onset of paraplegia or paralysis of the lower extremities without pain due to the occlusion of the Adamkiewicz artery, the major artery supplying blood to the inferior part of the spinal cord. Our patient was admitted to the hospital complaining of the development of paralysis of the bilateral lower extremities, thus revealing that he might have Adamkiewicz artery involvement. Although the major determinant of the mortality rate has been stated as the time elapsed until revascularization at the previous studies, Dossa et al. reported that the neurological state of the extremities had a more significant effect on the mortality rate than the ischemia time in their study.

Our patient’s physical examination revealed cold, pale, cyanotic lower extremities with an absence of femoral and distal pulses. Muscle weakness, absence of deep tendon reflexes, and loss of sensation are some of the manifestations of ischemic neuropathy associated with significant mortality.

Although aortography is the gold standard test, the use of Doppler USG, contrast-enhanced CT and magnetic resonance...
angiography (MRA) can also provide additional information to confirm the diagnosis. It is suggested that patients should have this procedure done preoperatively, especially for those with abdominal pain, hypertension, or anuria. In our case we could not do this test due to early death of the patient.

Since inadequate collateral arterial development occurs, immediate revascularization is necessary for both survival and limb salvage. An embolus can be removed by using balloon-tipped Fogarty catheters through transverse arteriotomies made on common femoral arteries under local anesthesia. In patients with thrombotic occlusion, decisions should be made based on the patient's clinical status and comorbidities. Aortic reconstruction or femoral revascularization via aortofemoral bypass may be undertaken. Several reports have advocated aortic reconstruction for all patients with infrarenal aortic occlusion due to the potential risk of propagation of thrombosis at the distal aorta up to the renal and mesenteric arteries; in patients with renal artery involvement, revascularization by reconstruction or an aortofemoral bypass should be performed promptly.

In order to maintain sufficient circulation to the distal extremities, relieve the symptoms, and preserve the limbs, we planned urgent surgery, but patient develop acute breathlessness with tachypnea with hypotension and anuria. Blood urea, serum creatine and serum electrolytes were normal but after 6 hour of admission serum potassium was 9.6mMol/l blood urea was 54.3mg/dl and serum creatinine was 2.3 mg/dl. Patient died due to refractory ventricular fibrillation.

In conclusion, patients promptly admitted to the hospital because of paralysis of the bilateral lower extremities should be examined there for pulses. If pulselessness or acute aortic occlusion is suspected, additional imaging procedures, including angiography, should be undertaken. In order to prevent the higher rates of morbidity and mortality in cases who develop paralysis in their lower extremities, immediate removal of the obstruction via a percutaneous thrombectomy should be performed using a balloon-tipped Fogarty catheter, and the thrombus material should be retrieved from the aorta. In patients with renal artery involvement, revascularization by reconstruction or aortofemoral bypass should be performed as quickly as possible.

REFERENCES