Case Report

Tubercular Aortitis - A Pictorial Review

Chetana R Ratnaparkhi¹, Sushupti M Chaukhande²

ABSTRACT

Tuberculous aortitis is a rare disease entity indicative of disseminated tuberculosis. We report a case who presented with newly diagnosed hypertension along with tuberculous lymphadenitis. Tuberculous aortitis should be included in the list of differential diagnosis for secondary hypertension, especially if the patient has a recent disease history of pulmonary tuberculosis.

Key words: Tubercular, aortitis, CT angiography.

Introduction:

Aortitis is a general term that refers to a broad category of infectious or noninfectious conditions in which there is abnormal inflammation of the aortic wall¹. Tubercular aortitis is a rare entity that is invariably indicative of disseminated tuberculosis². The prevalence of tuberculous aortitis is expected to increase with the rise in M. tuberculosis infection due to the rise in co-infection with HIV and multiple drug resistant tuberculosis¹. It occurs in less than 1% of patients with latent tuberculosis, with mortality rates as high as 60%³. The clinical manifestations are usually vague and nonspecific¹. As a result, aortitis is often overlooked during the initial work-up of patients with constitutional symptoms and systemic disorders¹. Noninvasive cross-sectional imaging modalities such as magnetic resonance (MR) angiography, and computed tomographic (CT) angiography play a critical role in initial evaluation and further assessment of aortitis¹. Here, we present a rare case of tubercular aortitis which was suspected on clinical examination and diagnosed on CT angiography. Multidetector computed tomography (CT) has largely replaced conventional angiography¹. CT angiography has excellent spatial resolution and is commonly the initial imaging study performed. In this article, the imaging manifestations of a case of tubercular aortitis have

¹Associate Professor, ²Junior Resident NKPSIMS & RC, Nagpur

Address for Correspondence -Dr. Sushupti M. Chaukhande E-mail: sushupti90@gmail.com been discussed.

Case Report:

A 13 years old boy, presented with the complaints of generalized body ache, fever and swelling in the neck since 4-5 days. Patient There was no past history of tuberculosis. Onexamination Patient was mildly febrile, Pulse was 96/min & BP (130/90) which falls in hypertensive range for that age. He also had bilateral cervical and axillary lymphadenopathy. Rest of general and systemic examination was unremarkable. USG abdomen revealed multiple pre and para-aortic lymph nodes with loss of fatty hilum. FNAC from the neck swelling was suggestive of chronic granulomatous inflammation. Cytological features favoured tubercular lymphadenitis. Her CT aortic angiography revealed thickening of wall of aorta (figure 1), reduction in caliber of a rta just below origin of renal arteries (figure 2), Narrowing of caliber of bilateral renal arteries just after their origin (figure 3), Narrowing of caliber of SMA just after its origin (figure 4, 5). These imaging features along with the history of tubercular lymphadenitis were consistent with the diagnosis of tubercular aortitis.

Figure 1: Axial CT Angiography image showing thickening of the wall of the abdominal aorta



Figure 2:



Figure 3: Axial and coronal CT angiography image shows Narrowing of caliber of bilateral renal arteries just after their origin.

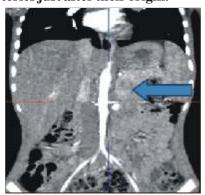


Figure 4: Sagittal - CT angiography

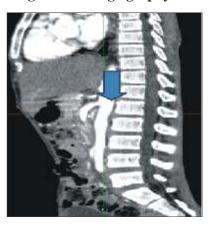
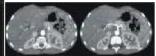


Figure 5: Axial - CT angiography





Discussion:

The most common causes of aortitis are the large vessel vasculitides - giant cell arteritis (GCA) and Takayasu arteritis⁴. Takayasu arteritis is an autoimmune disorder which typically affects the descending abdominal aorta and presents as pulseless disease due to occlusion of the vessels. Tubercular aortitis is a rare entity that is invariably indicative of disseminated tuberculosis². The prevalence of tuberculous aortitis is expected to increase with the rise in M. tuberculosis infection due to the rise in co-infection with HIV and multiple drug resistant tuberculosis¹. Tuberculous aortitis usually involves the distal aortic arch and descending aorta as seen in our case¹. It is generally due to direct extension from contiguous mediastinal lymph nodes, empyema, or pericarditis or to hematogenous or lymphatic spread of distant infection¹. It occurs in less than 1% of patients with latent tuberculosis, with mortality rates as high as 60%³. In a patient with tuberculosis and thickening of the aortic wall, the possibility of tubercular aortitis is likely.

The risk of perforation increases when a cold abscess is present⁵. In a study of 39 cases of tuberculousmycotic aneurysms by Long et al⁶, 75% of the aneurysms appeared to originate from aortic wall erosion by a contiguous focus; most of the aneurysms (90%) were saccular and false, and disseminated tuberculosis was present in almost one-half of cases. Involves the distal aortic arch and descending aorta. Prevalence is on increase with rise in co-infection with HIV and multiple drugresistant tuberculosis

Conclusion:

We present a case of Tubercular aortitis which is is a rare entity that is invariably indicative of disseminated tuberculosis. Tubercular aortitis occurs in less than 1% of patients with latent tuberculosis, with mortality rates as high as 60%. Multidetector computed tomography (CT) has largely replaced conventional angiography. CT angiography has excellent spatial resolution and is commonly the initial imaging study performed.

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