Case Report

False Positive ¹⁸F-FDG PET/CT in Cardiac Sarcoidosis

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Abstract

Cardiac sarcoidosis is an underdiagnosed disease that may be present in as many as 25% of patients with systemic sarcoidosis. ¹⁸F-FDG PET/CT is playing an increasing role in the management of sarcoidosis patient with suspected cardiac involvement. Physiologic FDG uptake by myocardium needs to be suppressed and differential diagnosis must be considered. Here we present a case of false positive ¹⁸F-FDG PET/CT due to myocardial ischemia.

Keywords: Cardiac sarcoidosis; PET/CT; False positive; ¹⁸F-FDG; Myocardial ischemia

Case Presentation

A 46-year-old non-smoker, non-diabetic woman with newly diagnosed sarcoidosis, presented with atypical chest pain at rest and was referred for a ¹⁸FDG PET-CT exam looking for cardiac involvement. She had no prior history of heart disease. Rest electrocardiogram and dobutamine stress echocardiogram were normal. PET-CT revealed hypermetabolic lymph nodes on both sides of the diaphragm, thickening of the peribronchovascular interstitium and large sized antero-septo-apical myocardial uptake (Figure 1). Because of these observations, the patient was diagnosed as having cardiac sarcoidosis. Also, because of the persistent chest pain and the atypical clinical presentation, a coronary angiography was performed. This showed a proximal left anterior descending artery sub occlusion, which lead to myocardial ischemia at rest and was responsible for PET-CT findings (Figure 2).

Discussion

In our case cardiac catheterization (right anterior oblique angulated view) was performed because of persistent chest pain, which is an atypical clinical presentation for cardiac sarcoidosis as well as palpitations, syncope, bradycardia, peripheral edema and dyspnea [1,2]. Patient has not reported any chest pain symptoms since angioplasty was performed. Cardiac involvement in patients with systemic sarcoidosis is probably underestimated. Autopsy studies show a prevalence of cardiac involvement in sarcoidosis in more than 25% of cases [3], although clinical evidence of myocardial involvement is found in only 5% of patients with systemic sarcoidosis. Clinical manifestations include conduction disturbances, arrhythmia, congestive cardiac failure, and sudden cardiac death [4]. Early diagnosis is challenging and requires different imaging techniques [5]. Revised guidelines of the "Japanese society of sarcoidosis and other granulomatous disorders" provide recommendations for the diagnosis of this potentially life-threatening disease [6]. These



Figure 1: PET/CT fusion images from attenuation-corrected ¹⁸F-FDG PET/ CT scans (**A**, anterior maximum intensity projection PET image; **B**, horizontal long axis; **C**, vertical long axis; and **D**, short axis) showing hyper metabolic lymph nodes on both sides of the diaphragm (arrows; SUV_{max} 12.7) and large sized antero-septo-apical myocardial uptake (arrowhead; SUV_{max} 5.8).



Figure 2: Cardiac catheterization (right anterior oblique angulated view) showing a proximal left anterior descending artery sub occlusion (arrowhead).

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guidelines do not include recent emerging techniques such as ¹⁸F-FDG PET/CT even if this is a promising tool for early disease detection as well as for therapy monitoring and image-guided biopsy [7,8]. However, ¹⁸F-FDG PET/CT must be performed following specific patient preparation protocols [4,7,9] (long fasting [10], highfat low-carbohydrate diet [11], use of unfractionated heparin [12]) to suppress cardiac physiological ¹⁸F-FDG uptake and avoid false positive results. Focal or patchy ¹⁸F-FDG uptakes are consistent with a diagnosis of cardiac sarcoidosis whereas the diffuse myocardial uptake pattern or the focally increased lateral-wall uptake pattern are likely to represent normal variations [13]. The present clinical case shows another pitfall in interpreting cardiac ¹⁸F-FDG PET/CT exam in patient with sarcoidosis. In spite of a well-done cardiac preparation, intense myocardial uptake can be seen in ischemic tissue because of the preferential use of glycolysis under anaerobic conditions [14,15]. In our case the imaging protocol did not include myocardial perfusion imaging since dobutamine stress echocardiogram had already been done in order to exclude coronary macroangiopathy or to detect other cardiomyopathies [16].

Conclusion

We would like to emphasize the importance of ruling out coronary artery disease before interpreting cardiac ¹⁸F-FDG PET/CT exam in patients with sarcoidosis, especially in case of atypical chest pain.

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