A case of multisystem sarcoidosis on 18F-FDG PET/CT

Seval Erhamamci¹, Saime Ramadan², Bülent Yaşar³

¹Department of Nuclear Medicine, Baskent University Faculty of Medicine, Baskent University Istanbul Hospital, Istanbul, Turkey
²Department of Pathology, Baskent University Faculty of Medicine, Baskent University Istanbul Hospital, Istanbul, Turkey
³Department of Gastroenterology, Istanbul Medipol University, Çamlıca S.U.A.M. Hospital, Istanbul, Turkey

ABSTRACT

We described a 67-year-old woman with uncommon distribution of multisystem sarcoidosis to disseminated lymph node, lung, liver and sinonasal region on 18F-FDG PET/CT imaging, which showed confounding scintigraphic features of mimicking widespread malignancy due to high metabolic uptake. The present case emphasizes the importance of whole-body assessment by PET/CT in multisystem sarcoidosis associated with rare sinonasal involvement and also the selection of the biopsy site by PET-guidance.

Keywords: Sinonasal sarcoidosis, extrapulmonary sarcoidosis, 18F-FDG PET/CT

INTRODUCTION

Sarcoidosis, a chronic inflammatory disease of unknown etiology, characterized by noncaseating granulomas, most commonly affects pulmonary and mediastinal involvement (1-5). Extrapulmonary sarcoidosis is seen in 25-50% of cases, usually associated with thoracic involvement (1). Extrapulmonary sarcoidosis can be involved multiple organ systems, such as the peripheral lymph nodes, liver, spleen, bones, heart, kidneys, skin, eyes, muscles and central nervous system (2-7).

18F-FDG PET is a non-invasive diagnostic imaging technique widely used in oncological clinical practice, and also provides a valuable assessment of infectious and inflammatory diseases (1-5). Sarcoidosis may exhibit characteristic thoracic PET findings related to its predominant pulmonary and mediastinal involvement (1-5). Disseminated lymph node and extrathoracic lesions may mimic widespread high metabolic metastases or diffuse lymphomatous disease on 18F-FDG PET imaging (4-7). This case report illustrates an example of uncommon distribution of multisystem sarcoidosis to disseminated lymph node, lung, liver and sinonasal region on 18F-FDG PET/CT imaging, which showed confusing scintigraphic features of mimicking widespread malignancy due to high metabolic uptake.
CASE REPORT

A 67-year-old woman without a significant medical history, non-smoker admitted with fatigue and dyspnea. Physical examination and initial laboratory results were unremarkable. Abdominal CT revealed hypodense lesion in the left lobe of liver and wall thickening in the caecum localization. However, no malignancies were determined by colonoscopy. Chest CT showed bilateral multiple pulmonary nodules, and enlarged mediastinal and hilar adenopathy, prompting referral for PET/CT to assess for malignancy of unknown origin. PET imaging revealed widespread foci of intense 18F-FDG uptake in multiple lymph nodes, lung, liver, and sinonasal region (Figure 1). No primary malignancy was identified, findings were interpreted as highly concerning for granulomatous disease, and recommended transbronchial or inguinal lymph node biopsy for differential diagnosis from malignancy. Histopathological examination of excisional biopsies from the right inguinal lymph node led to a diagnosis of nonnecrotizing granuloma with asteroid body which is highly suggestive of sarcoidosis (Figure 2). The patient was started on a regimen of systemic steroids.

Figure 1. Maximum intensity projection anterior (A) and lateral (B), transaxial PET and fusion PET/CT (C-E) images revealed widespread foci of high 18F-FDG uptake in multiple lymph nodes including bilateral cervical, supraclavicular, mediastinal, hilar, abdominal and right inguinal nodes (largest was 2.0 cm in diameter with SUVmax=18.5), in left maxillary sinus and left nasal concha (SUVmax=31.4), in numerous peribronchovascular region (SUVmax=11.8) and parenchymal nodules in both lungs, in the liver (SUVmax=11.7). No primary malignancy was identified.

Figure 2. (A) Hematoxylin–eosin stain (original magnification × 100): Nodal effacement by numerous well-defined, small nonnecrotizing granulomas composed of epithelioid cells with Langhans giant cells and lymphocytes. (B) Hematoxylin–eosin (original magnification × 400): Star-like asteroid body within giant cell not specific but compatible with sarcoidosis.
DISCUSSION

18F-FDG PET/CT can significantly contribute to the management of multisystem sarcoidosis because it can detect active inflammatory granulomatous disease (1-3). 18F-FDG PET/CT can increase the accuracy of the diagnosis of extrathoracic involvement due to whole-body functional mapping of active inflammatory sites (4). However, active sarcoidosis lesions are known to be hypermetabolic like malignancies on 18F-FDG PET/CT, and may cause to misinterpretation on imaging (5-7). Malignancies cannot be ruled out based on hypermetabolic lesions; therefore, a biopsy was required to confirm the diagnosis. 18F-FDG PET/CT may guide the selection of the biopsy site.

Extrathoracic sarcoidosis is most often associated with thoracic involvement (1-3). Detecting extrathoracic lesions is sometimes difficult by physical examination, standard thoracic radiography or CT. 18F-FDG PET/CT may be helpful in patients with sarcoidosis for determining the intrathoracic and extrathoracic extensity of disease, detecting active disease and accessing the response to treatment (2-4). In the current case, thoracic involvement and also extrathoracic disseminated lymph node, liver and sinonasal involvement was successfully revealed by 18F-FDG PET/CT imaging. The most common extrathoracic involvement was reported the abdomen, which includes liver, spleen, biliary tree, peritoneum, and lymphatic sarcoidosis (2-7). Hepatic involvement is encountered in 50-80% of cases in autopsy specimens (3). Hepatic involvement usually asymptomatic until advanced stage of the disease, is characterized by enlargement, mottled pattern of FDG accumulation, or scattered small nodular granulomatous lesions on cross-sectional imaging (1-8). This may be associated with an enlargement of abdominal lymph nodes close to the liver hilus or in celiac regions, as in our case.

Sinonasal involvement in sarcoidosis is uncommon, account for less than 5% of cases, and is usually seen with advanced stage of the disease (9). Involvement of sinonasal mucosa could be isolated, or a part of multisystem involvement. Sinonasal involvement is most likely underdiagnosed type of sarcoidosis because it is usually occult lesion. Even if a biopsy is required to confirm sarcoidosis, PET/CT imaging may play an important role in diagnosis and treatment follow-up (9). Unfortunately, biopsy can not be performed on newly detected sinonasal areas due to technical and ethical reasons. The changes in FDG uptake intensity on 18F-FDG PET/CT may reflect the effectiveness of medical treatment and provide a change in therapeutic strategy.

The present case emphasizes the importance of whole-body assessment by 18F-FDG PET/CT in multisystem sarcoidosis associated with rare sinonasal involvement and also the selection of the biopsy site by PET-guidance. In addition, PET/CT provides prognostic information and guides the therapeutic management.

ETHICAL CONSIDERATIONS

Informed Consent: Written informed consent was obtained from all participants who participated in this study.

Referee Evaluation Process: Externally peer-reviewed.

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