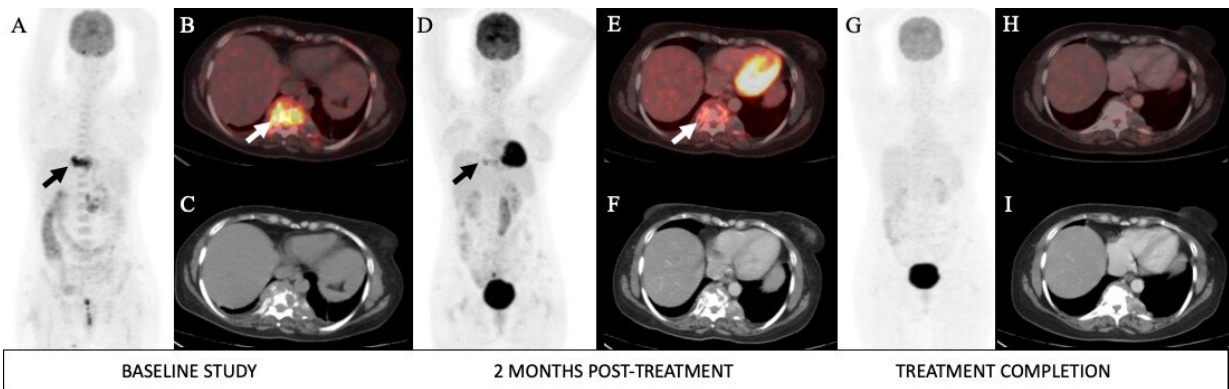


# Earlier diagnosis and treatment assessment of tuberculosis achieved with pet/ct

June 24 2019



A 38-year-old woman diagnosed to have with Pott's spine was subjected to F-18 FDG PET/CT scan. Baseline study--maximum intensity projection image (A), fused PET/CT (B) and CT image (C) showing increased FDG uptake in a lytic lesion with significant soft tissue component in the D9 vertebra with associated paravertebral soft tissue density. Study done at 2 months post-ATT (D, E, F) showing favourable response to therapy. Study done at treatment completion (G, H, I) showing complete metabolic response to therapy. Credit: Mittal BR, Parihar AS, Sood A, et al.

Research presented at the Society of Nuclear Medicine and Molecular Imaging's 2019 Annual Meeting shows that molecular imaging with 18F-FDG positron emission tomography/computed tomography (PET/CT) can evaluate tuberculosis at the molecular level, effectively identifying diseased areas and guiding treatment for patients.

According to the World Health Association, tuberculosis is one of the top 10 causes of death worldwide. Although the [disease](#) is curable and preventable, in areas with high rates of tuberculosis, it contributes to significant morbidity, mortality and an increased risk of transmission from infected individuals. Tuberculosis most frequently involves the lungs; however involvement of tissues and organs other than the lungs is referred to as extra-pulmonary tuberculosis.

"Extra-pulmonary tuberculosis presents a particular challenge as the disease site is often not accessible for performing an invasive diagnosis. The physician thus relies on the clinical diagnosis for initiating treatment as well as deciding the duration of therapy, which can be difficult," said Bhagwant R. Mittal, MD, DNB, professor and department head of nuclear medicine at the Postgraduate Institute of Medical Education and Research in Chandigarh, India. "In our study, we aimed to evaluate the utility of 18F-FDG PET/CT in the initial diagnosis and response assessment of patients with extra-pulmonary tuberculosis."

Ninety-three patients with extra-pulmonary tuberculosis were prospectively enrolled in the study prior to the initiation of treatment. The patients underwent 18F-FDG PET/CT imaging as a baseline, and then received follow-up imaging after two months and after treatment completion. Follow-up scans were categorized into three groups: complete metabolic response (no abnormal lesions), [residual disease](#) (persistent lesions, but no new lesions) and disease progression (new lesions compared to the baseline scan).

In the baseline scans, 176 lesion sites were detected among the 93 study participants. The most common sites included the lymph nodes and central nervous system. Two month follow-up scans were performed on 47 patients, and 21.2 percent were classified as having complete metabolic response, 72.3 percent had residual disease and 6.4 percent were characterized as having disease progression. A final scan was

conducted post-treatment and, of the 28 patients imaged, 28.6 percent had complete metabolic response, 53.6 percent of patients showed residual disease and 17.8 percent had disease progression. During the course of the study, 12.9 percent of patients died. Of these deaths, the patients who fell in the [disease progression](#) category had the highest mortality rate—60 percent.

"This study has the potential to change the way we manage [tuberculosis patients](#). Our results show that 18F-FDG PET/CT provides a whole-body survey and identifies the disease sites in various organs and tissues in a single study. This helps to provide an early estimation of disease extent, and in suspected cases, helps to identify accessible biopsy sites for obtaining tissue diagnosis," Mittal said. "Further, follow-up scans can point towards response to treatment and thus suggest predict a more accurate outcome."

**More information:** Abstract 227: "18F-FDG PET/CT in Extra-Pulmonary Tuberculosis - Role in Initial Evaluation and Response Assessment"

Provided by Society of Nuclear Medicine and Molecular Imaging

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