

TECHNOLOGIST PRESENTATION

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Initial experience of imaging cardiac sarcoidosis using hybrid PET-MR - a technologist's case study

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Background

Cardiac imaging has been identified as a potential use of hybrid PET-MRI. This new technology allows simultaneous Positron Emission Tomography (PET) and MR scanning to occur.

This case study features a 34 year old male with known pulmonary sarcoidosis which was diagnosed in 2006 and initially treated with immunosuppressants. He presented with NYHA functional class 2 symptoms. ECHO showed asymmetrical LV anterior and lateral wall hypertrophy with a maximum wall thickness of 18mm in the basal and mid anterior and lateral wall.

The patient was referred for a cardiac¹⁸F Fluorodeoxyglucose (FDG) PET-MR study to determine the cause of the hypertrophy. The possible differential diagnoses included hypertrophic cardiac myopathy with pulmonary sarcoidosis or cardiac sarcoidosis.

Methods

Prior to the scan, the patient followed a high protein, low carbohydrate diet for one day. This was then followed by a 12 hour fast to suppress physiologic glucose metabolism by the heart.

The patient was administered with 355mBq of ¹⁸F FDG 180 minutes prior to the PET-MR study.

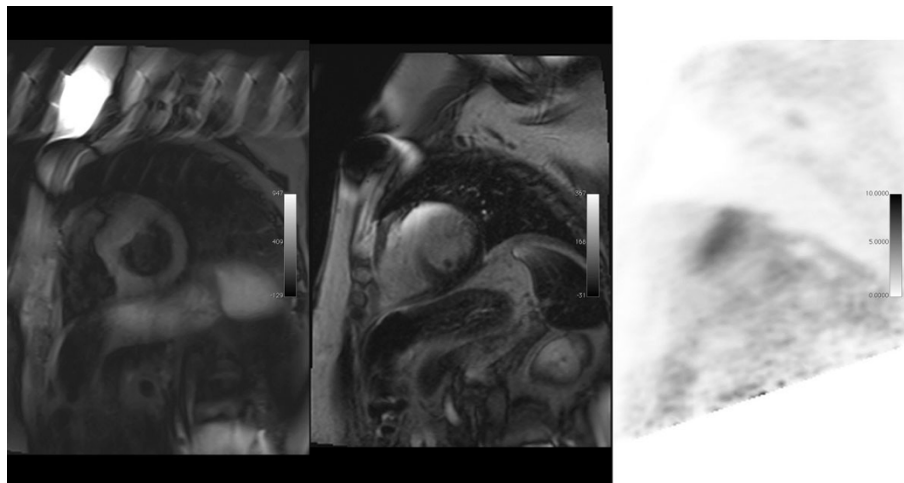


Figure 1 From left to right: 1) SA STIR demonstrating shows homogeneous signal from the LV myocardium. 2) SA LGE demonstrating focal mid myocardial wall LGE in the antero-septal segments. 3) SA ¹⁸F FDG PET showing focal uptake in the antero-septal segments Note is made of a Reveal device visualised on images 1 & 2.

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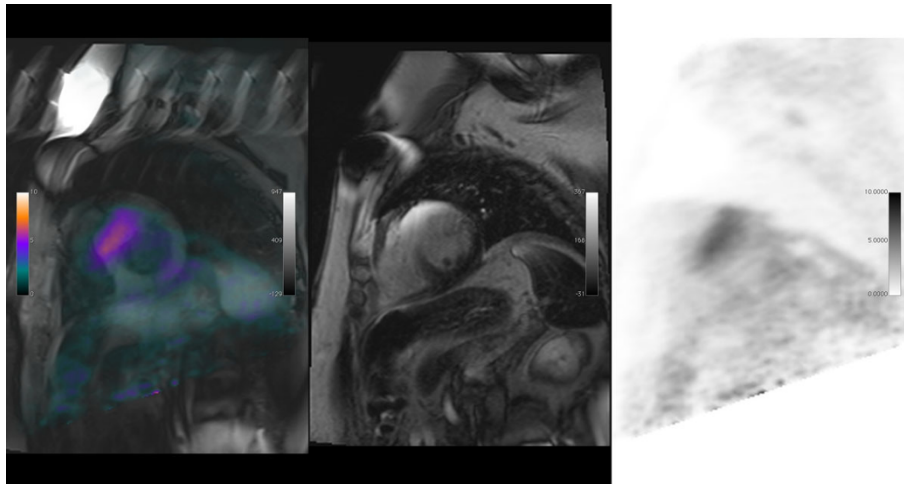


Figure 2 These images show the fusion of the PET onto the STIR sequence, demonstrating the area of active inflammation despite there being no oedema.

A simultaneous PET-MR study was acquired. A standard CMR protocol was undertaken in conjunction with a simultaneous 10 minute single PET bed using a Siemens Biograph mMR hybrid PET-MR scanner. This is a 3T magnet containing a 260mm field of view PET detector located at the isocentre.

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Results

The scan demonstrated mediastinal ¹⁸F FDG avid lymphadenopathy with antero-lateral myocardial thickening with mid-myocardial LGE and focal ¹⁸F FDG uptake compatible with active myocardial inflammation within an area of "scar" in keeping with active cardiac sarcoidosis.

The oedema sensitive STIR sequence did not differentiate between the hyper-metabolic active disease and the chronic "burnt out" disease.

Conclusions

This patient is one of the first examples of imaging active cardiac sarcoidosis using a hybrid PET-MR technique. It demonstrates the potential for differentiating between active and chronic cardiac sarcoidosis during one scan. This new imaging technique could have a substantial impact on the diagnosis and management of cardiac sarcoidosis

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