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Case Report: Brown Fat Accumulation of Tc-99m Macroaggregated Albumin in a Lung Perfusion Study in a Patient With Multiple Lung Arteriovenous Malformations and Right-to-Left Shunting

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Abstract: An 18-year-old man was preoperatively assessed for a varicocele and found to be hypoxemic. A Tc-99m macroaggregated albumin lung perfusion scan showed right-to-left shunting, evidenced by increased radiotracer uptake in the brain, kidneys, thyroid gland, and bilateral supraclavicular areas, a typical location for brown adipose tissue. Chest computerized tomography angiogram study showed supraclavicular fat density areas and multiple pulmonary arteriovenous malformations.

The authors report a rare case of brown fat visualization on a lung perfusion scan in a patient with right-to-left shunting, likely because of increased perfusion to activated brown adipose tissue.

Abbreviations: AVM = arteriovenous malformation, BAT = brown adipose tissue, CTA = computerized tomography angiogram, FDG = fluorodeoxyglucose, MAA = macroaggregated albumin, SPECT = single photon emission tomography.

INTRODUCTION

Brown adipose tissue (BAT) is type of adipose tissue that has a role in thermogenesis, especially in neonates and young children during cold exposure, by aging BAT becomes white adipose tissue; however, a lesser amount of persistent brown fat is present in adults, which also has a some role in energy metabolism, in the last decade due to common use of F-18 fluorodeoxyglucose (FDG) BAT is commonly seen on these scans, especially in the supraclavicular, axilla, and paravertebral regions; however it can also be seen adjacent to visceral organs in the mediastinum and abdomen. In cases of cold exposure, BAT becomes activated and ultimately becomes more apparent on F18 FDG scans.

METHODS

No Institutional Review Board approval was requested because such approval is not required for single case reports at the Johns-Hopkins Medical Institution. No consent was obtained because this publication contains no identifiable information.

This publication methodology included review of the electronic medical record and review of the available images for our case.

Patient Information

An 18-year-old patient was preoperatively assessed and found to be hypoxemic. The patient was investigated by a chest computerized tomography angiogram (CTA) and a Tc-99m macroaggregated albumin (MAA) lung perfusion study to evaluate for possibility of arteriovenous malformations (AVMs) within the lung.

Diagnostic Findings

The findings on an anterior perfusion view of the head and upper chest using Tc-99m MAA shows increased radiotracer uptake in the brain, thyroid, and bilateral supra-clavicular regions (Fig. 1). It has been demonstrated previously that in the presence of shunting in the lungs, a portion of the Tc-99m MAA particles bypass the lung capillary circulation and appear in the systemic circulation.

The uptake in the supra-clavicular region is presumed to be brown fat because of its location and anatomic correlation with the expected location of BAT.

A chest CTA image shows an AVM in the right lower lobe (Fig. 2). Other axial slices showed multiple AVMs. The multiplicity of these lesions suggested a diagnosis of hereditary hemorrhagic telangiectasia, which was subsequently treated successfully by vascular embolization, which improved the patient’s hypoxemia.

A coronal chest CTA image shows structures that have fat density in the supraclavicular regions as shown in (Figure 3); these areas of well-perfused fat correspond to the areas of Tc-99m MAA accumulation seen on the perfusion scan and likely represent areas of BAT perfusion because of brown fat activation.

DISCUSSION

Brown adipose tissue uptake has been demonstrated previously with multiple radiotracers, such as Tl-201 chloride, I-123 metaiodobenzylguanidine, Tc-99m sestamib, F-18 and H-3 FDG, H-3 methionine, and H-3 thymidine. The regions of
BAT uptake in these radiotracers are usually the supraclavicular and paravertebral adipose tissues. The available images we have, however, show only Tc-99m MAA accumulation in the supraclavicular adipose tissue with no definite evidence of accumulation in the paravertebral adipose tissues, which if present may be obscured by the lung activity.

Although we do not have a single-photon emission computed tomography image to show the accumulation of Tc-99m MAA in the supraclavicular brown fat, it can be considered that the supraclavicular region of uptake on the perfusion images is a site of brown fat accumulation of Tc-99m MAA because of the typical location and the presence of fat tissue in the supraclavicular region as shown on the chest CTA coronal image provided.

The presence of intrapulmonary shunting causing uptake in highly perfused organs including activated brown fat may be an important factor that allowed visualization of the activated brown fat in the supraclavicular regions in this patient. Given the fact that the examination was performed in a summer month (August) and although we do not have information regarding the environment surrounding the patient during the examination day, we believe the brown fat uptake is attributed to the presence of metabolically active brown fat with increased vascularity, which is mainly because of young age (18 years old) with the possibility of being because of a surrounding cold environment less likely.

Although the shunt fraction should have been estimated for this study, it was not calculated because the clinical concern was to evaluate for the presence of pulmonary embolism and the images were sufficient to suggest the diagnosis of intrapulmonary shunting.

We report a rare case of Tc-99m MAA accumulation in BAT in the supraclavicular region, which will help resolve diagnostic dilemma’s in the future if a similar case is encountered.

REFERENCES


