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(G*) (POS-45) Applying the Hybrid PET/MRI Technique PMRO_x in Healthy Humans to Measure Cerebral Metabolic Rate of Oxygen

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Introduction

Positron Emission Tomography (PET) is the gold standard for imaging CMRO₂, the cerebral metabolic rate of oxygen (Fan et al., NeuroImage, 2020). However, the procedure requires up to three radiotracers and invasive arterial sampling. Incorporating MRI techniques can simplify the procedure (Ssali et al., JNM, 2018; Narciso et al., Phys Med Biol, 2021). PET/MR imaging of oxidative metabolism (PMRO_x) uses whole-brain (WB) CMRO₂ measured by MRI to calibrate simultaneously acquired [15O]O₂-PET data, eliminating arterial sampling. Arterial spin labeling (ASL) can be used to replace [15O]H₂O-PET, reducing the requisite number of radiotracers to one. PMRO_x is non-invasive yet maintains the ability of PET to quantify the oxygen extraction fraction (OEF). CMRO₂ can be imaged under different metabolic states (e.g., rest and during a functional task) in a single imaging session as the acquisition time is ~5 min. The accuracy of PMRO_x was previously demonstrated in a porcine model (Narciso et al., JNM, 2021). The aim of the current work was to present initial data translating PMRO_x to human participants.

Methods

Data were acquired from n=13 healthy subjects on a Siemens 3T Biograph mMR scanner. Five minutes of list-mode PET data were acquired after inhalation of ~2000 MBq of [15O]O₂ while measuring WB CMRO₂ by MRI (Jain et al., J Cereb Blood Flow Metab, 2010) at rest and during a finger-tapping task. PET images were reconstructed using MR-based attenuation correction maps, motion-corrected, and smoothed by a 4mm Gaussian filter. Pseudo-continuous ASL (TR/TE: 4210/37.86 ms, post-labeling delay: 1.7s, labeling duration: 1.5s) was collected during the PET acquisition. ASL images were motion-corrected and smoothed by a 6mm filter. All images were pre-processed in SPM12 and calculations were completed with in-house MATLAB scripts.

Results

Results were normalized to Montreal Neurological Institute (MNI) atlas space. Average resting CMRO₂ across subjects was 4.5 and 3.5 mL/100g/min for grey and white matter, respectively. CMRO₂ was observed to increase ~25% in the motor cortex during tapping.

Discussion

This preliminary study demonstrates the feasibility of imaging CMRO₂ in a span of 5 minutes by combining [15O]O₂-PET with MRI, reducing the requisite number of radiotracers to one and eliminating arterial sampling. CMRO₂ values were in good agreement with literature values.

Keyword-1

PET/MRI

Keyword-2

Cerebral Metabolic Rate of O₂

Keyword-3

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