

Multi-slice PO₂-weighted ³He imaging in a rabbit model of regionally impaired perfusion

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Introduction: Oxygen mapping with hyperpolarized ³He MRI yields quantitatively accurate PO₂ measurements in phantoms when spatially non-selective RF excitations are used [1]. It is generally recognized that to accurately depict steady-state P_AO₂ values in the lung, it is also necessary to ensure that the inhaled gas contains 21% oxygen evenly mixed with the dose of hyperpolarized gas. But just as the goal of conventional ¹H MRI is rarely to obtain absolute quantification of T1 or T2, absolute quantification of P_AO₂ may not be necessary to obtain useful information from oxygen maps generated using ³He MRI. By striving simply for PO₂ weighting, the sequence programmer is free to use slice-selective excitations to improve SNR and/or spatial resolution, and a much less cumbersome inhalation procedure can be used. The purpose of the present work is to demonstrate the ability of multi-slice oxygen-weighted ³He imaging to depict regional P_AO₂ variations, by applying this technique in an animal model of regionally impaired perfusion.

Methods: Ventilation-perfusion mismatch was created in four anesthetized rabbits using a model of reversible pulmonary embolism. Under fluoroscopic guidance, a balloon angiocatheter was inserted through the right femoral vein and positioned in the left pulmonary artery. Upon inflation, the balloon obstructed perfusion to the lower lobes of the left lung. PO₂-weighted ³He MR images were acquired in each rabbit with the balloon alternately inflated and deflated. Each ³He scan was performed at forced breath hold, immediately following inhalation of 30 ml hyperpolarized ³He, using a 1.5T whole-body scanner (Siemens Sonata) equipped with a homebuilt ³He birdcage RF coil. ³He gas was polarized using a prototype commercial system (Magnetic Imaging Technologies, Inc.)

Two different PO₂-weighted spoiled-gradient-echo pulse sequences were used [1,2]. Both pulse sequences were implemented so as to require acquisition of the fewest possible number of images per scan, and multi-slice capability was implemented by interleaving 2D slice acquisition during the oxygen-sensitization time. Imaging parameters included: TR/TE, 10/3.1 ms; bandwidth, 200 Hz/pixel; matrix, 32×64; pixel size 4×4 mm, oxygen sensitization time, 5s. Slice thickness was 7mm for axial images and 9mm for sagittal images. Pixel-by-pixel P_AO₂ maps were calculated from each set of images using the standard formulas [1,2], but the values served only to provide a quantitative basis for regional contrast and were not interpreted to represent absolute measurements of P_AO₂.

Results: Imaging results were qualitatively similar for all rabbits using both pulse sequences. The fluoroscopic image in Fig.1 shows the position of the inflated balloon (red arrow), and the DSA (digital subtraction angiography) image shows the lack of perfusion to the lower left lung. The coronal-projection oxygen-weighted images show elevated P_AO₂ in the lower left lung with the balloon inflated, in contrast with the small perfused region in the upper right lung (yellow arrows). The axial multi-slice images in Fig.2 also show elevated P_AO₂ elevated in the lower left lung with the balloon inflated (yellow arrows), while the sagittal images show that the boundary of the non-perfused region occurs along a fissure between upper and lower lung lobes (red arrows).

Conclusions: PO₂-weighted ³He imaging is sensitive to regionally obstructed perfusion, and 2D multi-slice application is capable of resolving the location and extent of the affected lung region. The resulting P_AO₂ maps are not absolutely quantitative, but they have good spatial resolution and contrast-to-noise ratio. While ventilation anomalies remain difficult to characterize using oxygen-weighted methods, this study suggests that PO₂-weighted ³He MRI may be useful for identifying perfusion anomalies in the lung. Although the region studied here occupies a large fraction of the rabbit lung, the absolute size of the lesion is relatively small (a few inches), suggesting that respectable spatial resolution is possible in human applications.

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References: [1] Miller GW et al. ESMRMB 22 (2005). [2] Fischer MC et al. MRM 52:766-773 (2004).

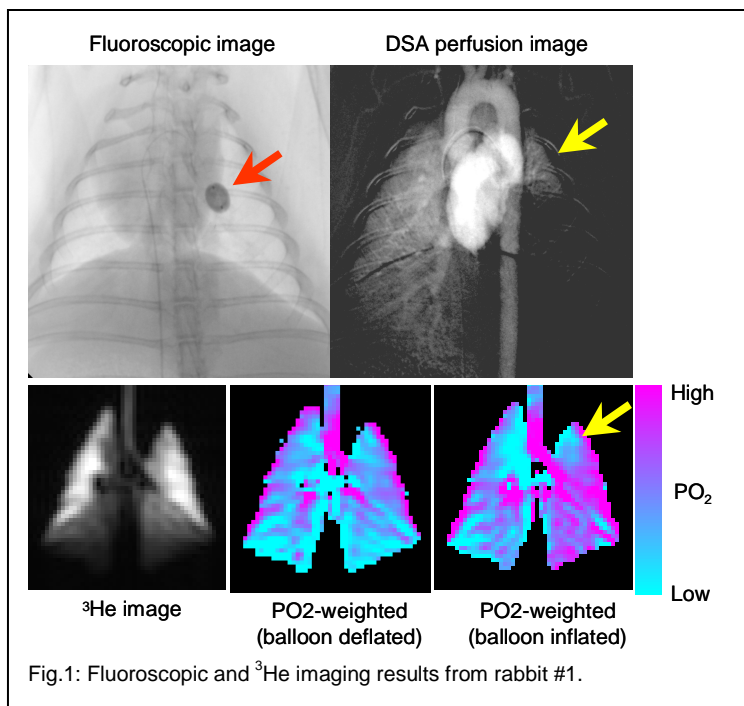


Fig.1: Fluoroscopic and ³He imaging results from rabbit #1.

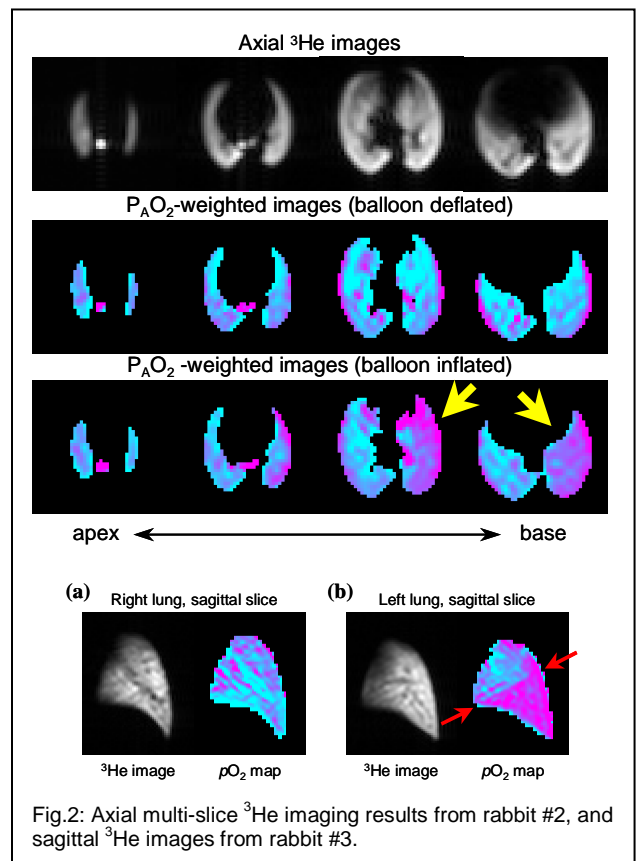


Fig.2: Axial multi-slice ³He imaging results from rabbit #2, and sagittal ³He images from rabbit #3.