## Dynamical <sup>17</sup>O imaging in tumor bearing mice at 7T

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**Introduction:** Oxygen consumption rate and blood flow are important parameters for the physiological and pathological evaluation of brain, myocardium and tumors<sup>1,2</sup>, where <sup>15</sup>O PET has been utilized. <sup>17</sup>O MRI will be another tool for the direct observation of tumor oxygenation. Measurements of the blood flow and oxygen consumption rate by *in vivo* <sup>17</sup>O NMR has been reported recently<sup>3,4</sup>. We have developed <sup>17</sup>O imaging by FISP and succeeded in the visualization of natural abundance  $H_2^{17}O$  distribution in the mouse with 10 min data acquisition<sup>5</sup>. In this study, we will report the dynamical study of <sup>17</sup>O imaging using <sup>17</sup>O enriched saline in the tumor bearing mice. Phantom experiments demonstrated the feature of <sup>17</sup>O *in vivo* NMR signals.

**Methods:** MRS/MRI was performed on 7T/400mm/SS system (NIRS/KOBELCO/Bruker) with 40 mm  ${}^{1}$ H/ ${}^{17}$ O Litz coil (Doty Scientific Inc.). Water, ethanol or 25-100 % ethyleneglycol -water was used in phantom experiments.  ${}^{17}$ O images of healthy and tumor bearing C3H/He mice (20 – 25 g) were obtained under Ketamine:Xylazine anesthesia by true FISP with a TR/TE = 4.3/2.15 ms. A saline (0.5ml) containing 5%  ${}^{17}$ O prepared from 10%  ${}^{17}$ O water (Cambridge Isotope Laboratories, Inc) was i.v. injected to tumor bearing mice. After imaging experiment, organs were excised for  ${}^{17}$ O spectral measurements.

**Results:** Phantom studies: The <sup>17</sup>O signal of ethyleneglycol or water in 25% ethyleneglycol-water phantom was detected by fid acquisition mode but not by echo mode within our experimental condition. Animal studies: The S/N in the <sup>17</sup>O images of a mouse obtained by FISP with 10 min data acquisition was dramatically improved from 3.8 to 13.4 after the injection of a saline containing 5% <sup>17</sup>O. The result of 10 sec dynamical imaging of  $H_2^{17}O$  with under the spatial resolution of 2.5 mm without slicing was shown in Fig.1. The process of initial accumulation of the injected  $H_2^{17}O$  in the heart and re-distribution to whole body was demonstrated. Spatial resolution of 1.25 mm was attained with the 10 min data acquisition.

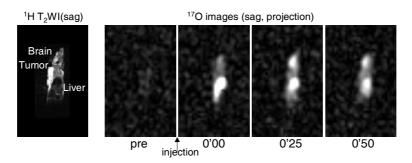


Fig.1. <sup>17</sup>O images of a tumor bearing mouse measured by FISP in 10 sec before and after i.v. injection of 5% <sup>17</sup>O saline

**Discussion:** The dynamical study of  $H_2^{17}O$  was achieved in the tumor bearing mice with a temporal resolution of 10 sec. The temporal and spatial resolutions attained in this study will lead this imaging method to the evaluation of blood flow or oxygen consumption rate using <sup>17</sup>O gas and water. FISP is shown to be an effective method for imaging *in vivo* <sup>17</sup>O signals from free water. The phantom experiments with ethyleneglycol-water strongly suggest that the *in vivo* <sup>17</sup>O images obtained here is from the mobile  $H_2^{17}O$  and not from other molecular species or immobilized water. The echo image of <sup>17</sup>O detecting solely the signal from free water, not from the body constituents contributing as background should have an advantage over the other NMR method with FID detection in the <sup>17</sup>O<sub>2</sub> gas study.

**Reference:** (1) Secomb TW. et al, 34 :313 (1995), (2) Ando K, et al, Int J Radiat Biol 75 :505 (1999), (3) Zhu XH. et al, MRM 45:543 (2001), (4) Fiat D. et al, Neurol Res 26:803 (2004), (5) Narazaki M. et al, 14th ISMRM 3113 (2006)