

# Fast 4D Metabolite Mapping of Hyperpolarized $^{13}\text{C}$ Compounds using Multiple Echo 3D FIESTA

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**OBJECTIVE:** The objective of this work is to develop a technique which will allow repeated measurements of spatial (3D imaging) and spectral (1D chemical-shift) distributions following administration of a hyperpolarized  $^{13}\text{C}$  substrate in order to determine spatially localized metabolic kinetics (e.g. spatial distribution of tumor metabolism of  $^{13}\text{C}$ -pyruvate to  $^{13}\text{C}$ -lactate).

**BACKGROUND:** Reeder et. al [1] first demonstrated the ability to derive high spatial resolution chemical-shift images from multiple sequential gradient echoes acquisitions at echo times shifted by  $\Delta$ . Recently Wieben et. al.[2] proposed a much faster method for acquiring both the spatial and spectral information in one scan using a multiple-echo 2D balanced SSFP (FIESTA) imaging technique. The Nyquist frequency ( $N_f$ ) for this technique is determined by the echo spacing,  $\Delta$ , where  $N_f=1/(2\Delta)$ , and the spectral resolution ( $\Delta$ ) is determined by  $\Delta = 1/(N \Delta)$ , where N is the number of acquired echoes. The multiple-echo 2D FIESTA technique is fast, has relatively high spatial resolution, and is able to adjust the number of echoes (N) and the echo spacing ( $\Delta$ ) to provide the desired spectral resolution. We have implemented a multiple gradient echo 3D FIESTA technique in order to provide the increased SNR necessary for imaging hyperpolarized  $^{13}\text{C}$  labeled compounds (products and substrates) at low concentration.

**MATERIALS and METHODS:** All  $^{13}\text{C}$  imaging was performed in a transmit/receive  $^{13}\text{C}$  surface coil designed and built in our laboratory. All imaging was performed on a 1.5 T General Electric Signa MR scanner operating with version 9.1 software. The manufacturer's standard 3D FIESTA pulse sequence was modified to allow multi-nuclear and multiple-echo imaging (ME-3DFIESTA). A phantom consisting of a sphere containing  $^{13}\text{C}$ -acetate and a vial containing  $^{13}\text{C}$ -2-hydroxyethylacrylate (280 Hz chemical shift) were placed on the  $^{13}\text{C}$  surface coil. ME-3DFIESTA imaging was performed with a 16 x 64 x 64 matrix giving isotropic 7 mm spatial resolution, BW=62.5 MHz, 12 echoes, 1.68 ms echo spacing,  $\Delta=49.6$  Hz,  $N_f=297$  Hz, and TR=21.2 ms for a 25 second scan time. In a second experiment multiple ME-3DFIESTA acquisitions were also obtained on a syringe containing 50 mM of both  $^{13}\text{C}$ -hydroxyethylpropionate and  $^{13}\text{C}$ -cis-fumarate simultaneously hyperpolarized using the PASADENA technique[3,4]. The parameters for this acquisition were the same as above except a TR of 14.5 ms, acquisition of 8 echoes with echo spacing of 1.34 ms, and an imaging time of 17 seconds. The spectral data was reconstructed with the technique reported by Leupold et. al.[5]

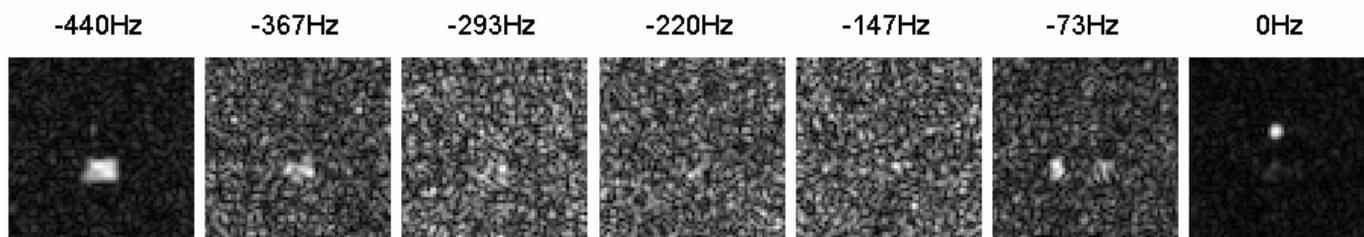


FIGURE 1: Selected chemical shift images of the  $^{13}\text{C}$ -acetate and  $^{13}\text{C}$ -2-hydroxyethylacrylate phantom.

**RESULTS:** Selected chemical shift images calculated from imaging the  $^{13}\text{C}$ -acetate and  $^{13}\text{C}$ -2-hydroxyethylacrylate phantom using the ME-3DFIESTA pulse sequence are shown in Figure 1. The chemical shift images in Figure 1 demonstrate the ability of the ME-3DFIESTA to provide metabolite images with the correct chemical shifts under static conditions. The first echo spatial image of the syringe containing simultaneously hyperpolarized  $^{13}\text{C}$ -hydroxyethylpropionate and  $^{13}\text{C}$ -cis-fumarate is shown in Figure 2. Selected chemical shift images of this phantom are shown in Figure 3 demonstrating the ability of the ME-3DFIESTA technique to provide spatial and spectral information of hyperpolarized  $^{13}\text{C}$ -labeled substrates and products in a timely manner.

**CONCLUSION:** We have demonstrated the feasibility of implementing a fast multiple gradient echo 3D FIESTA technique for the rapid measurement of spatial location and chemical shift of hyperpolarized  $^{13}\text{C}$ -labeled substrates and products.

**REFERENCES:** [1]Reeder et al., MRM 51, 35-45, 2004. [2] Wieben et al. Proceedings of the ISMRM, p. 2386, 2005 [3]Bhattacharya et al., Proceedings of the ISMRM, p 171, 2005, [4]Bhattacharya et. al. 2005) MAGMA, 18.5 [5] Leupold et al., Proceedings of the ISMRM, p 102, 2005.

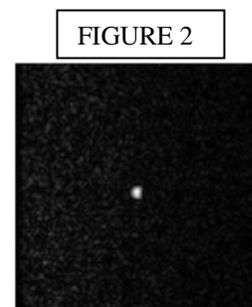


FIGURE 2

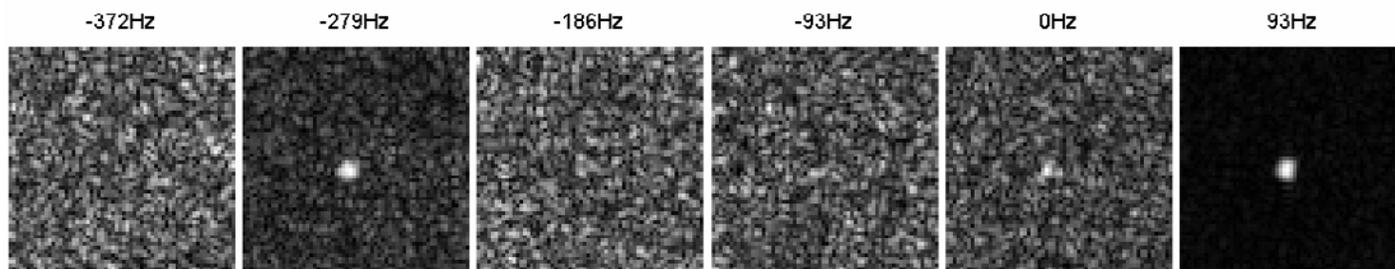


FIGURE 3: Selected chemical shift images of simultaneously hyperpolarized  $^{13}\text{C}$ -hydroxyethylpropionate and  $^{13}\text{C}$ -cis-fumarate.