

# Inner volume BOLD-fMRI at ultra-high spatial resolution

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## Introduction:

Multi-Echo FLASH imaging in combination with echo train shifting [1,2] offers BOLD sensitivity for matrix sizes up to 256 x 256 leading to in-plane resolutions of about 1 mm<sup>2</sup>. Long measurement times and low Signal-to-Noise-Ratio (SNR) preclude BOLD-fMRI going to higher resolutions. Hence, only a reduction in the Field-of-View (FOV) further increases spatial resolution. Latter, however, is limited by aliasing artifacts due to signals from outside of the FOV. 2D-selective excitation [3] in slice- and phase encoding direction, as shown in Fig. 1, allows for small inner FOV sizes resulting in increased spatial resolution. Recently, applications with two-dimensional pulse profiles such as spiral [4] or blipped planar [5] excitation trajectories were reported. In this study, latter one was chosen because the limiting side excitations only occur in one single dimension.

## Methods:

The design of the 2D-selective excitation pulse was based on a duration of 27.5 ms, spatial dimensions of 50 mm in read direction and 4 mm in slice direction and a flip angle of 50°.  $k_{\max,x}$  was set to 2.5 cm<sup>-1</sup>,  $k_{\max,y}$  to 0.84 cm<sup>-1</sup> and  $Dk_y$  to 0.04 cm<sup>-1</sup>.

For data acquisition, the Double Contrast Echo Train Shifted Multi-Echo FLASH sequence [2] was chosen with the following sequence parameters:  $T_E=38$  ms,  $T_R=126$  ms. The matrix size was 192 x 114 leading to an in-plane resolution of 0.5 x 0.5 mm<sup>2</sup>.

For data acquisition, the Double Contrast Echo Train Shifted Multi-Echo FLASH sequence [2] was performed on a 3 T (Trio, Siemens, Erlangen, Germany) using a matrix head coil with the following sequence parameters:  $T_E=38$  ms,  $T_R=126$  ms, flip angle  $\alpha=50^\circ$ . The matrix size was set to 192 x 114 leading to an in-plane resolution of 0.5 x 0.5 mm<sup>2</sup>. To demonstrate BOLD-sensitivity, an initial BOLD-experiment was performed by acquiring 200 images in 2.9 s/image. Functional activation was achieved by a visual stimulation task showing a checkerboard alternating with a gray screen (5 images in 14.5 s each).

Functional results were processed and analyzed with Statistical Parametric Mapping 5 (SPM5). The images were realigned to the average image (Fig. 3) and subsequently smoothed with a 2 mm Gaussian Kernel. The alpha error was set to  $\alpha=0.01$  and minimal shown cluster size was  $k=10$  pixels.

## Results:

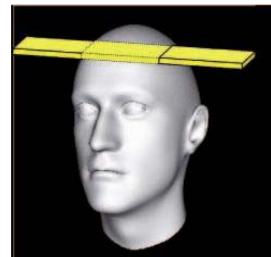
Fig. 3 shows the average image of 200 acquisitions during the fMRI experiment. It should be noted that the image is free of noticeable artifacts at an in-plane resolution of 0.5 x 0.5 mm<sup>2</sup>. Activation maps of the human brain (Fig. 4) show BOLD-signal only close to the venules.

## Discussion:

We presented a high resolution T2\*-weighted acquisition sequence in combination with a two-dimensional excitation pulse suitable for functional experiments. Its feasibility includes both, high resolution morphological T2\*-weighted imaging with high contrast to the venule system (Fig. 3) and local high resolution functional studies (Fig. 4). Latter showed activation only in areas close to venules instead of a blurred activation-signal in the whole cortex, as achieved with generic EPI-BOLD sequences.

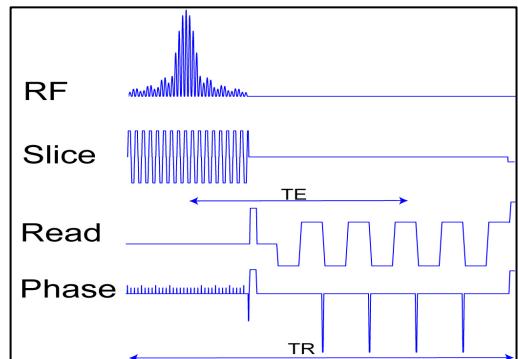
## References:

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- [2] Heiler P et al, Proceedings of the ISMRM 15, 1940 (2007).
- [3] Haase A et al, J. Magn Reson 67, 258-266 (1986)
- [4] Pauly J et al, J. Magn Reson 81, 43-56 (1989)
- [5] Rieseberg S et al, Magn. Reson. Med. 47, 1186-1193 (2002)



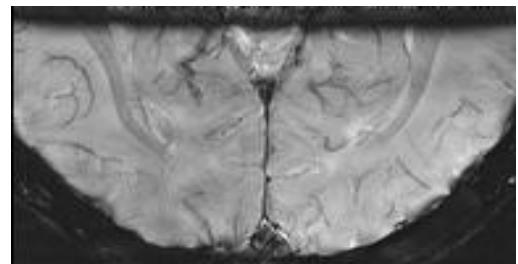
**Figure 1:**

The excitation profile of the 2D selective blipped planar excitation pulse trajectory allows a reduction of the FOV to inner volume sizes.



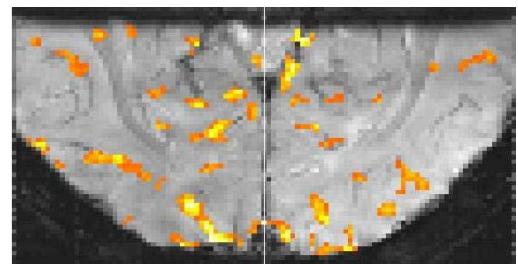
**Figure 2:**

Sequence Scheme for Multi-Echo FLASH with two dimensional RF excitation based on a blipped planar trajectory



**Figure 3:**

Average of 200 Multi-Echo FLASH images with 2D-RF excitation.  $T_E$  was 38 ms and  $TR$  126 ms at a total scan time of 2.9 s/image.



**Figure 4:**

Activation map of the visual cortex of a healthy volunteer. Activation is located close to the venules