B1-dependence of single-voxel MRS sequences: STEAM, PRESS and MEGA-PRESS

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The signal detected by an MRS experiment depends on accurate calibration of transmit B_1 power; it is B_1 -dependent. Progress to higher field strengths has brought increasing issues of B_1 inhomogeneity (see Figure 1) and increasing application of exotic RF pulse shapes to deliver increased slice-selection bandwidth. At 7T in particular, B_1 is sufficiently inhomogeneous that the peak B_1 achievable in one region of the brain may be only half the global peak. It is often assumed that the B1-dependence of a 90° pulse goes as $\sin(\pi/2*B_1/B_{1nom})$, and a 180° pulse as $\sin^2(\pi/2*B1/B1nom)$, where B_1 and B_{1nom} are the actual and intended B_1 field. However, these expressions are based upon the on-resonance case for rectangular, non-selective pulses, , when shaped slice selective pulses are being used, it is the signal integral across the slice that determines

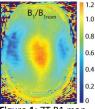


Figure 1: 7T B1 map

output signal intensity. In this abstract we present simulated and phantom data describing the B₁-dependence of three common MRS experiments: STEAM, PRESS and MEGA-PRESS.

Method

Simulations Slice profiles were simulated at a range of B_1/B_{1nom} from 0 to 1 for 'Spredrex', an asymmetric excitation pulse and 'GTST', a high-bandwidth refocusing pulse (see Fig 2). The STEAM sequence localizes with three 90° pulses and PRESS with one 90° pulse and two 180° pulses. The B1-dependence of these sequences is calculated assuming that the three orthogonal slice profiles independently impact signals.

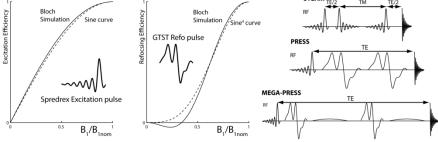


Figure 2: B1-dependence of excitation (left) and refocusing pulse (center). Pulse sequences (right).

Experimental A 10 mM solution of GABA in phosphate-buffered saline was scanned using a Philips Achieva 7T scanner. The B1-sensitivity profiles were measured in a region of maximum B1-field homogeneity. STEAM (TE=14ms) and PRESS (TE=30ms) experiments were performed without water suppression using the the following parameters: TR=3s; $(2 \text{ cm})^3$ volume; 8k datapoints at 5 kHz spectral width (SW); 4 averages; sampling B_1/B_{1nom} factors from 0.1 to 1 in increments of 0.05. MEGA-PRESS experiments were performed with excitation water suppression and the following parameters: TR/TE=3s/70ms; $(3 \text{ cm})^3$ volume; 8k datapoints at 5 kHz SW; 64 averages; sampling B_1/B_{1nom} from 0.65 to 1 in mean increments of 0.025. Water supression pulses were not scaled by B_1/B_{1nom} to maintain good water suppression over all experiments.

Results

Single-pulse Bloch simulations (shown in Figure 2 left) show that the excitation pulse has sine-like behavior, as expected, whereas the refocusing pulse has sin⁴ rather than the expected sin² behavior. Experimental data shown in Figure 3 are best matched by sin⁴ (STEAM), sin¹¹ (PRESS) and sin¹⁵ (MEGA-PRESS) functions.

Discussion

Although simple ('naïve') theory expects the B₁ dependence for PRESS to be sin⁵, the experimental dependence measured GTST pulses would predict that PRESS will have a sin⁹ dependence (based on factors of sin¹ for the spredrex pulse and sin⁴ for each GTST pulse). However, the B₁-sensitivity found in the experimental data was even greater than that predicted by simulation - sin⁴ for STEAM, and sin¹¹ for PRESS. – the progressively worsening B₁-sensitivity from naïve, to simulated, to

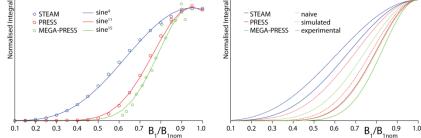


Figure 3: Experimental B1-dependence of sequences (left), comparison of naïve and simulated expectations to experimental curves (right).

experimental is shown above right. The increased experimental sensitivity may be due to B_1 inhomogeneity within the MRS voxel, which was not accounted for, or the interaction of the flip angle variation across the slice profile in the 3 different directions used for localization.

Use of lower bandwidth sinc-Gaussian-like refocusing pulses with \sin^2 behavior may result in improved signal, especially for MRSI applications, however chemical shift dispersion effects will be greater. The different B_1 sensitivity of PRESS and MEGA-PRESS presents issues for the quantification of MEGA-PRESS, as PRESS measurements of creatine or water are typically used as reference signals. The severity of this issue is underlined in the MEGA-PRESS case, for which a 20% miscalibration of the B1 power will result in over 50% loss in signal. This is especially pertinent at 7T when B_1 field homogeneity is a major challenge. For MRS experiments, careful localized flip angle calibration is critical for optimum SNR, and the use of parallel transmit techniques for improved B_1 field homogeneity will be especially important for MRSI experiments with wider spatial coverage.

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