

Accurate temperature measurements in the presence of field inhomogeneities

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Target audience: Clinical researchers involved in MR-guided thermal ablation work or, more generally, in field mapping.

Purpose: The goal of this work is to improve the measurement of temperature in the presence of B_0 inhomogeneities. Transcranial MRgFUS can be used to treat essential tremor^{1,2} near the thalamus. However, B_0 inhomogeneities near the treatment target often prevent accurate measurement³ of the therapy heating at the target site. With proton resonance frequency (PRF) thermometry, phase changes are converted into temperature changes by mapping the echo time, TE , to temperature. In the presence of field inhomogeneities, MR echoes may be translated and/or spread in k-space, potentially making the 'true' echo time vary spatially within the imaged object and take on values that differ from the nominal user-input TE value. Errors in TE lead to errors in temperature, and the purpose of this work is to describe and correct for these errors.

Methods: Consider a dual-echo gradient-echo sequence with imaging bandwidth BW_1 and BW_2 and nominal user-input echo time TE_1 and TE_2 for the two echoes, respectively. A relative temperature measurement could be obtained independently from each echo, and combined through a weighted sum with weights w_1 and w_2 : $\Delta T = (w_1\Delta T_1 + w_2\Delta T_2)/(w_1 + w_2)$. It can be derived, when selecting weights such that $w_2 = -w_1 \times BW_2/BW_1$, errors in ΔT_1 precisely cancel out errors in ΔT_2 , giving rise to a temperature measurement immune to field inhomogeneities. Ideally, however, weights should be chosen to optimize the temperature-to-noise-ratio (TNR), based on expected signal strength and/or TE value, and constraining the choice of these weights would have a TNR penalty. Alternately, inhomogeneity-induced errors in TE could be explicitly calculated, allowing any weights w_1 and w_2 to be subsequently selected for optimum TNR. With $\Delta\phi$ the measured temperature-induced phase shift, and τ the evolution time available for phase accumulation (in most sequences τ is simply equal to TE but in more exotic cases it may take on different and even negative values⁴), one can obtain:

$$\Delta\tau_1 = -\frac{\tau_1 \times (\Delta\phi_1 \times \tau_2 - \Delta\phi_2 \times \tau_1)}{\Delta\phi_1 \times \tau_2 \times (BW_1/BW_2) - \Delta\phi_2 \times \tau_1}, \quad \frac{\Delta\tau_1}{\Delta\tau_2} = \frac{BW_2}{BW_1} \times \frac{\tau_1}{\tau_2}.$$

However, these equations require $BW_1 \neq BW_2$. Different bandwidths lead to different spatial distortions in the presence of field inhomogeneities, making it difficult to perform pixel-by-pixel processing. A more general but more computationally-involved method is introduced below that remains valid even for a single-echo sequence or for a dual-echo sequence with $BW_1 = BW_2$.

Given a field map $B_0(\vec{r})$, spatial gradients in $B_0(\vec{r})$ are responsible for shifting signal in k-space.

A pulse-sequence specific function f_s is introduced that returns the time at which any given k-space point is sampled. The error in the echo time is given by:

$$\Delta TE_{j,i}(\vec{r}) = f_s(\Delta\vec{k}_{j,i}) - TE, \quad \text{with } \Delta\vec{k}_{j,i}(\vec{r}) = 2\pi \times \gamma \times \hat{v}_{j,i} \times \nabla B_{0,i}(\vec{r}), \quad \text{and } \hat{v}_{j,i} = TE_j + \Delta TE_{j,i-1},$$

where j is the echo number, i the iteration number, $\Delta TE_{j,0} = 0$. Corrected temperature measurements are obtained from corrected τ values. The field map also needs to be corrected in the process, hence the ' i ' dependence in $B_{0,i}$. For example, with a dual-echo sequence, the field map would be obtained by comparing the phase of the two echoes and dividing by $(TE_{2,i} - TE_{1,i})$, and thus is corrected from one iteration to the next. A very small number of iteration (about 3) appeared to be sufficient in the present work.

Results: A series of thermometry experiments were performed by heating a gel phantom using an ultrasound transducer inside a 3 T MR system (55 W for 20 s, $1.5 \times 1.5 \times 5$ mm³, 128×128 , $TE/TR=10/17$ ms, 30° flip angle). Temperature measurements were made in the presence of field inhomogeneities (introduced by intentionally de-adjusting shim parameters) as well as with carefully-shimmed scans and using a number of different settings for the sampling bandwidth, BW . Figure 1 shows that as expected the presence of field inhomogeneities cause errors in the measured temperature values. These errors take the form of a multiplicative factor that increases linearly with $1/BW$, with a slope that depends on local field gradients at the transducer's focus. Through careful shimming, local field gradients could be removed, leading to accurate measurements at all sampled $1/BW$ settings ('o' symbols in Fig. 1). The proposed work aims at enabling accurate measurements despite field inhomogeneities.

Results in Fig. 2a were reconstructed with weights w_1 and w_2 chosen as described above ($BW_1 = -BW_2$, $w_1 = w_2$, 37 W for 30 s, $TE_1/TE_2/TR = 7.2/17.8/25$ ms), and reference results were obtained from carefully-shimmed scans. For Fig. 2b, a sequence with $BW_1=BW_2$ was used instead, and data were reconstructed with the iterative method described above (57 W for 30 s, $TE_1/TE_2/TR = 6.2/16.7/24$ ms). In both cases please note that corrected results obtained in bad shim conditions, once reconstructed as proposed, were in good agreement with reference good-shim results.

Discussion: PRF thermometry measures temperature-induced field changes, and is in nature a field-mapping method. The type of errors described and corrected for here would presumably be present in any field-mapping MR application, not just thermometry. These errors can be quite significant, especially when a narrow imaging bandwidth is chosen to help boost SNR, and errors by as much as 26% were readily observed and corrected for.

Conclusion: Field inhomogeneities create errors in temperature measurements, which can be detected and corrected using the proposed strategies.

References: 1. Elias et al, N Engl J Med 369:7 (2013). 2. Lipsman et al, Lancet neurology. 12:5 (2013). 3. Sprinkhuizen, Chapter 7, Ph.D. thesis (2010). 4. Madore et al, MRM 66:658(2011). Support from grants R01CA149342, P41EB015898 and R01EB010195 is acknowledged.

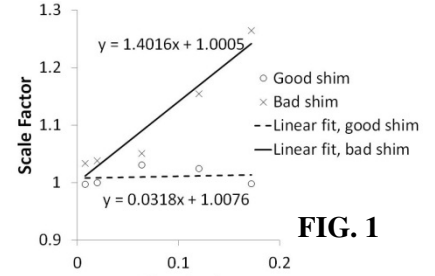


FIG. 1

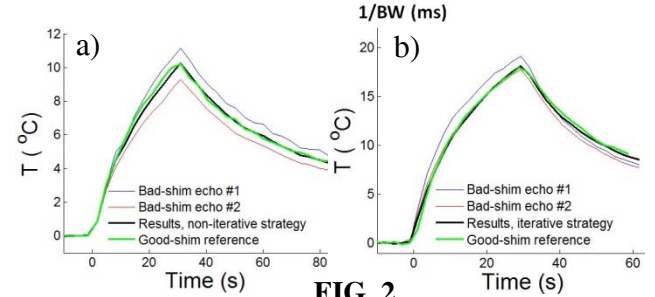


FIG. 2