## Blipped mGESEPI (bmGESEPI) for Fast and Accurate T2\* Measurements with B0 Inhomogeneity Compensation

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**Introduction**: Knowledge of  $T_2^*$  relaxation times at ultra-high field strength ( $\geq 7$  T) is needed for optimizing acquisition parameters for  $T_2^*$ -weighted imaging and understanding relaxation mechanisms. However, standard  $T_2^*$  measurements (*e.g.*, using multi-echo gradient echo (GE)) are affected by static magnetic field (B<sub>0</sub>) inhomogeneity, which is particularly severe at ultra-high field strength. The multi Gradient Echo Slice Excitation Profile Imaging (mGESEPI) method was developed for  $T_2^*$ -weighted imaging and  $T_2^*$  measurement with B<sub>0</sub> inhomogeneity compensation [1,2]. We implemented this method at ultra-high field strength, but realized that it requires excessive acquisition times to provide accurate  $T_2^*$  measurements. We therefore developed a more efficient method that allows significantly faster and accurate measurements, and demonstrate these advantages in a phantom as well as *in vivo* and postmortem human brains.

**Theory:** In the mGESEPI method, a compensation gradient  $G_c$  is combined with the slice rephaser gradient, a train of M GE images are acquired at different TEs, and this is repeated for N equidistant  $G_c$  values. For each echo, an image is reconstructed by 3D Fourier transform followed by partial summation along the slice direction. A  $T_2^*$  map is then computed by fitting a monoexponential decay pixel-by-pixel to the M images. The susceptibility-induced gradient in the slice direction  $G_{z,susc}$  that can be compensated for by a given  $G_c$  at a time TE is given by  $\int G_c(t) dt = G_{z,susc}$  TE. Since the  $G_{z,susc}$  that that be compensated for <u>decreases</u> as TE increases, the  $T_2^*$  measurements are accurate only if the largest  $G_c$  ( $G_{c,max}$ ) is able to compensate for the largest  $G_{z,susc}$  at the <u>last</u> echo. However, satisfying this condition at ultra-high field strength requires a large  $G_{c,max}$  and N, resulting in excessive acquisition times for *in vivo* studies.

We developed a new method based on the mGESEPI method in which the compensation gradient  $G_c$  is also added as a blipped gradient in the slice direction between each echo acquisition (Figure 1). As such,  $\int G_c(t) dt$  increases linearly with TE, so that the <u>same</u>  $G_{z,susc}$  is compensated for at each echo. For the  $T_2^*$  measurements to be accurate, the  $G_{c,max}$  required to compensate for the largest  $G_{z,susc}$  is a factor M smaller than in the mGESEPI method, thus allowing a reduction of N as well as the acquisition time by the same factor. This so-called blipped mGESEPI (bmGESEPI) method is therefore more efficient and allows significantly faster and accurate  $T_2^*$  measurements.



Figure 1: bmGESEPI pulse sequence.

**Methods**: The studies were performed on an ultra-high field human whole-body MRI system using transverse electromagnetic RF coils.  $T_2^*$  maps were acquired in a phantom (air-filled tube orthogonal to B<sub>0</sub> surrounded by a CuSO<sub>4</sub> solution) using the multi-echo GE, mGESEPI, and bmGESEPI methods with equivalent parameters, namely TR 100 ms, interecho spacing  $\Delta$ TE 3.5 ms, M 10, field-of-view (FOV) (8 cm)<sup>2</sup>, matrix (MTX) 128<sup>2</sup>, one 3 mm thick slice,  $G_{c,max}$  160% (of the slice rephaser gradient) and N 80 (mGESEPI) or  $G_{c,max}$  16% and N 8 (bmGESEPI). In addition,  $T_2^*$  maps were reconstructed using only partial data from the original mGESEPI data set, thus simulating mGESEPI acquisitions with a range of smaller N values and either the same  $G_{c,max} / N$  ratio or  $G_{c,max}$  as the original data set. We also studied 2 healthy volunteers (2 male, age 32–34) who gave informed consent and 4 postmortem unembalmed human subjects (1 male, 3 female, age 57–84). For the *in vivo* study shown below,  $T_2^*$  maps were acquired using the GE and bmGESEPI methods with TR 500 and 100 ms respectively,  $\Delta$ TE 4.4 ms, M 10, FOV (18 cm)<sup>2</sup>, MTX 256<sup>2</sup>, one 3 mm thick slice,  $G_{c,max}$  20% and N 16 (bmGESEPI).

**Results and Discussion**: Figure 2 shows the results of the phantom study. The GE  $T_2^*$  map clearly shows artifacts due to  $B_0$  inhomogeneity. These artifacts can be largely corrected for by the mGESEPI method with  $G_{c,max}$  160% and N 80, as shown by the more homogeneous  $T_2^*$  map, however at the cost of a long acquisition time. As expected, decreasing both  $G_{c,max}$  and N progressively reduces the amount of correction, as shown by the artifacts appearing around the tube on the mGESEPI  $T_2^*$  map with  $G_{c,max}$  16% and N 8. Reducing only N while keeping the same  $G_{c,max}$  also results in artifacts, as seen on the mGESEPI  $T_2^*$  map with  $G_{c,max}$  160% and N 8, because the spacing between the  $G_c$  values becomes too large (*i.e.*, the oversampling becomes insufficient) for an accurate reconstruction of the images. On the other hand, the bmGESEPI method can provide a  $B_0$  inhomogeneity compensation that is even better than that achieved by the mGESEPI method with  $G_{c,max}$  160% and N 80, as shown by the very homogeneous  $T_2^*$  map, while requiring only 10% of its acquisition time. These results clearly show the significant advantages of the bmGESEPI method.

Figure 3 shows the results of an *in vivo* study. The GE  $T_2^*$  map shows very low values with no clear depiction of the anatomy due  $B_0$  inhomogeneity, whereas the bmGESEPI  $T_2^*$  map shows higher values with a clear delineation of anatomical structures such as the lateral ventricles, putamen, and globus pallidus. The high  $T_2^*$  values in the right posterior cortex are due to  $B_1$  inhomogeneity (*i.e.*, low flip angle and/or receive sensitivity), as typically observed on axial images of the human brain acquired at ultra-high field strength. While the parameters still need to be optimized in further studies, these results clearly show the effectiveness of the bmGESEPI method for *in vivo*  $T_2^*$  measurements.



**Conclusion**: We developed a new method for  $T_2^*$ -weighted imaging and  $T_2^*$  measurement with  $B_0$  inhomogeneity compensation that is more efficient than the existing mGESEPI method and allows significantly faster and accurate  $T_2^*$  measurements. This method will be particularly useful at ultrahigh field strength, which is affected by severe  $B_0$  inhomogeneity.

References: [1] Yang QX. Proc 6<sup>th</sup> ISMRM 1998. p. 578 [2] Liu H. Proc 9<sup>th</sup> ISMRM 2001. p. 1352

Figure 3: Results of an in vivo study (34-year-old male).