

A Comparison of Breast Tissue T₁ Mapping Using Conventional Multi-flip Angle and 2-point Dixon Techniques

Reem Bedair¹, Mary McLean², Andrew Patterson³, Roie Manavaki¹, John Griffiths², Fiona Gilbert¹, and Martin Graves³

¹University of Cambridge, Department of Radiology, Cambridge, Cambridgeshire, United Kingdom, ²Cancer Research UK Cambridge Research Institute, Cambridge, Cambridgeshire, United Kingdom, ³Department of Radiology, Cambridge University Hospitals NHS Foundation Trust, Cambridge, Cambridgeshire, United Kingdom

Target Audience: Clinical scientists' interested in quantitative mapping in the breast.

Introduction: Dynamic contrast enhanced (DCE) MRI applications particularly for breast imaging, utilize high spatiotemporal resolution acquisitions to assess enhancement kinetics. Quantitative pre-contrast T₁₀ mapping is required in order to convert signal intensity changes to gadolinium concentration in order to apply quantitative pharmacokinetic models. Whilst DCE-MRI acquisitions typically use some form of fat-suppression/water excitation, T₁₀ mapping is generally performed without fat suppression. Recently DCE studies have used 2-point Dixon (2PD) acquisitions as a time-effective method of obtaining robust water-only images. In this work, we compare B₁-corrected pre-contrast T₁₀ relaxation times obtained using a conventional non-fat suppressed variable flip angle (VFA) approach with a VFA 2PD method in a cohort of patients with locally advanced breast cancer at 3T.

Methods: Data Acquisition: Following IRB approval and informed consent, six patients (range 51- 67, mean age = 56 years) with proven breast malignancy (5 invasive ductal, and 1 mucinous carcinoma) referred for DCE-MRI were recruited into the study. Imaging was performed on a 3T MR system (MR750, GE Healthcare, Waukesha, WI) using an 8 channel-breast coil. Following the localizer scans, standard three-dimensional (3D) axial, non-fat suppressed, SPGR T₁₀ images were acquired at three flip angles (2°, 3°, 5°) TE/TR= 2.3/5.3ms, 256x256 matrix with slice thickness of 1.4mm, slices=112, parallel acceleration = 2. The acquisition time at each flip angle was 41s. Subsequently a dedicated breast 3D 2PD sequence was acquired with the same geometrical parameters and TE/TR = 1.2, 2.2/5.3ms, parallel acceleration = 2 generating water-only images. The acquisition time for each flip angle was 32s. B₁-mapping was performed using a multi-slice 2D Bloch-Seigert shift method (TR/TE/α= 30/13.5ms/20°, 128x128 matrix). DCE data was then acquired for 8 minutes with a 9s temporal resolution using a 3D segmented k-space SPGR sequence with fat suppression and an in-plane resolution of 1x1x1.4mm. Contrast agent (Magnevist, BayerSchering, Berlin, Germany) at 0.1mmol/kg body weight was administered using a power injection. The total scan time was 20 minutes.

Image Analysis: Both the conventional and water-only T₁₀ maps were calculated and corrected for B₁ transmit non-uniformity using DCETool² in Osirix (Pixmeo, Geneva). Representative regions of interest (ROIs) of the malignant breast tumour and the fibroglandular tissue (FGT) in the ipsilateral and contralateral breasts were drawn on the maximally enhancing DCE images and copied to both T₁₀ maps. The ROIs were manually adjusted in the case of misregistration between the T₁₀ maps and the DCE data.

Results: Figure 1 shows the acquired DCE series, standard and thresholded (10 %) water-only T₁₀ maps corrected for B₁ non-uniformity. The mean T₁₀ values for FGT tissue measured in both ipsilateral (T₁₀ = 1200 ± 240 ms) and contralateral (T₁₀ = 1230 ± 160 ms) breasts using the standard sequences were lower when compared with the 2PD technique in the ipsilateral (T₁₀ = 1450 ± 230 ms) and contralateral (T₁₀ = 1340 ± 330 ms) breasts. Whereas, the mean T₁₀ values of the malignant breast lesions were lower using the 2PD technique (T₁₀ = 1450 ± 230 ms) compared to the standard method (T₁₀ = 2070 ± 360ms) (Figure 2).

Discussion: Fat is a possible confounder for DCE-MRI of the breast. As far as we are aware this is the first report of B₁-corrected T₁₀ measurements in both FGT and breast tumour at 3T. Rakow-Penner et al have previously reported native T₁ values in the right breast of five normal volunteers at 3T using an IR-FSE sequence with and without IDEAL processing¹. They reported mean values for FGT of 1324.42±167.63ms and 1444.83±92.7ms using the non-IDEAL and IDEAL methods respectively, which are consistent with our results. The mean difference in T₁₀ values between the 2PD and the standard method of approximately 200ms in tumors is a little surprising given the relative paucity of fat in tumors but this may be attributed to either partial volume effects or increased uncertainty of the techniques at the longer T₁₀ values observed in tumors compared to FGT. Further studies will optimize the choice of flip angles to improve the accuracy and reproducibility of these measurements.

Conclusion: This pilot study demonstrates the application of a rapid VFA 2PD technique for the measurement of water-only T₁₀ values suitable for subsequent quantitative DCE-MRI analysis without the confounding influence of fat.

References: [1] Rakow-Penner *et al.* JMRI 2006 23:87-91 [2]Sung K *et al.* RSNA 2011

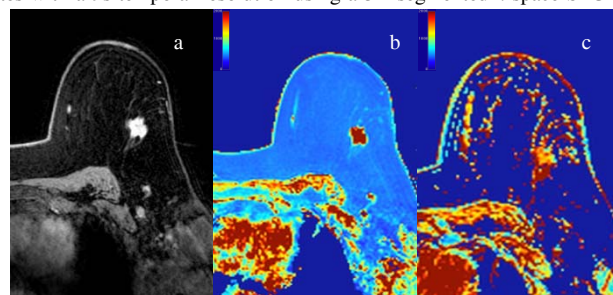


Figure 1 shows a) T₁W CE image b) standard non fat-suppressed c) water-only T₁₀ maps.

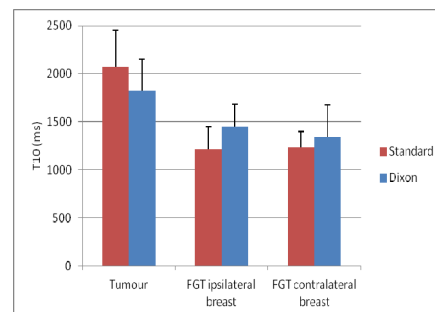


Figure 2 shows the mean T₁ relaxation values measured in tumor, ipsilateral FGT and contralateral FGT.