

A Random Regressor Model for T1-Correction of SPGR Variable Flip-Angle Acquisition without Experimental B1-Inhomogeneity Correction

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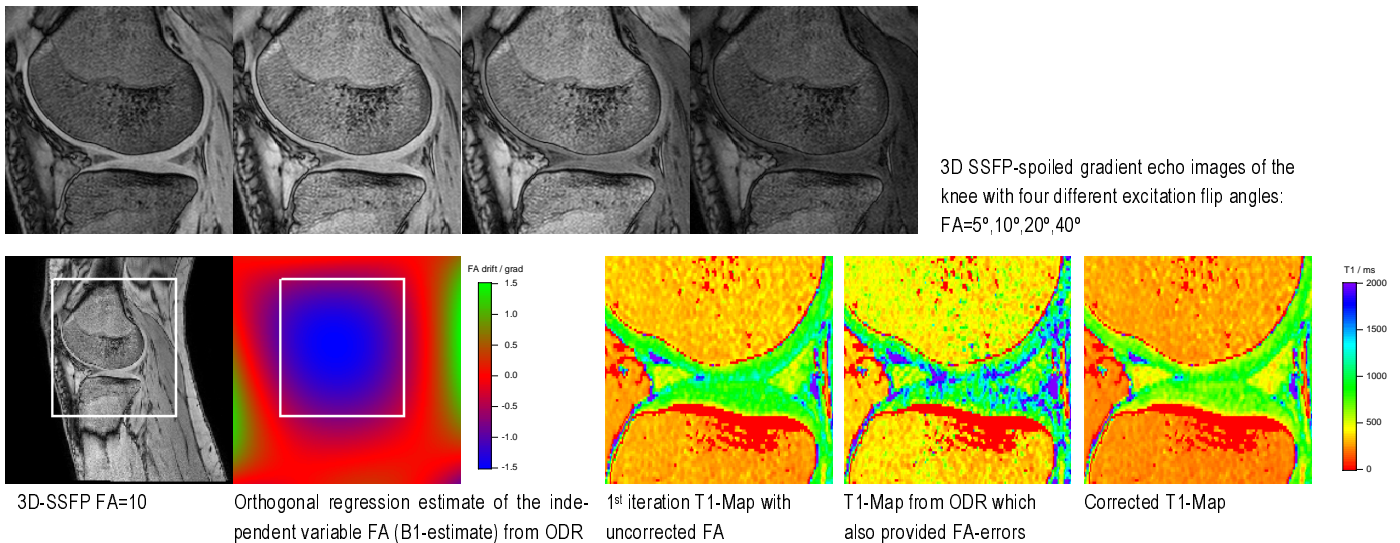
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Introduction: The methods of quantitative T₁-mapping from spoiled gradient recalled echo (SPGR) image acquisition with varying flip angles under steady state conditions allow fast and high resolution T₁-quantification in a clinical feasible timeframe [1]. The accuracy of the computed T₁-maps from the measured SPGR signal depends strongly on the SNR and number of measurement points (number of various flip angles FA) but even more critical is the correct knowledge of FA and its variation across the field of view. Especially at higher field strength (>1.5T) a calibration of the true FA across the VOI becomes inevitable. Previously introduced methods have been described that either provide the necessary FA-information by acquiring separate RF-transmit homogeneity maps (B₁-maps) [2] or incorporating B₁-determination approaches into the same sequence [3]. Nevertheless, both strategies require additional scan time and if the B₁-information was not provided for already performed scans the estimated T₁-values might differentiate significantly from the true values. We introduce a novel method of numerically estimating the error of the presumably known FA from the behaviour of the SPGR-signal with varying FA by means of a "Trust Region" fitting method also known as orthogonal distance regression ODR or measurement error model [4]. The improved algorithm (ODRPACK95) provides estimates for the error in the independent variable FA and even allows implementation of bound constrains very efficiently [5]. The primer application is for T₁-quantification in dGemric (delayed Gd-enhancement of MRI in cartilage) of the knee and hip joints.

Methods: Measurements were performed on a Philips Achieva 3.0T using a phased-array eight-channel knee coil and a 3D-SPGR sequence with the following parameters: FOV=160mm, acquisition matrix 320x320 leading to an in-plane resolution of 0.5mm at a slice thickness of 2mm. 40 slices in sagittal orientation were required to cover the entire region of the medial and lateral condyle of the femur as well as the patella cartilage; TR=9ms, TE=3.5ms (out-of-phase); four different flip angles were used: 5,10,20,40° leading to a total scan time of 8:00min. The ODRPACK95 code was implemented in IGOR 6.02 (Wavemetrics, Boulder, CO). The corrected T₁-values for articular cartilage, bone marrow and muscle were compared to values obtained from other already verified quantitative T1-mapping approaches.

Analysis: Common Least-Squares (LS) fitting of an even non-linear function y_i with an error implies that there are no errors in the independent measurements of x_i : $y_i + \varepsilon_i = f(x_i, \beta)$. Under this conditions the parameters β can be estimated by solving the classical LS-problem: $\min \frac{1}{2} \sum_{i=1}^n [f(x_i, \beta) - y_i]^2$. If, however the error in x_i cannot be ignored LS modifies to: $y_i + \varepsilon_i = f(x_i + \delta_i, \beta)$ with δ_i being the error in x_i . Solving this equation for the parameter vector β becomes known as the ODR: $\min_{\beta, \delta} \frac{1}{2} \sum_{i=1}^n [(f(x_i + \delta_i, \beta) - y_i)^2 + \delta_i^2]$ and provides error estimates δ_i for the independent variable x_i (FA).

Results: We tested the robustness and reliability of this technique for providing estimates of the FA-variation on a pixel-by-pixel base. The first iteration using ODR estimates T₁ and provides error for the flip angles α_i from the signal equation of a SPGR-sequence: $S(\alpha_i) = (M_0 \cdot (1-A) \sin \alpha_i) / (1 - A \cos \alpha_i)$, with $A = \exp(-TR/T_1)$. The resulting FA-error maps $\Delta \alpha_i(x, y)$ are then fitted with a 4th order polynomial to produce smoothly varying noiseless FA-maps. During the second iteration which uses the classical LS-Levenberg-Marquardt algorithm, the corrected T₁ are estimated with prior knowledge of the FA-maps. The resulting FA-corrected T₁-maps obtained similar values for cartilage, bone marrow and muscle as previously established techniques [6].



References: [1] S.C.Deoni et al., MRM 46 (2003): 515-526; [2] S.C.Deoni, Proc ISMRM 15 (2007): 42; [3] R.D.Newbould, Proc ISMRM 15 (2007): 38; [4] P.T. Boggs et al, SIAM J.Sci.Stat.Comput. 8, 6 (1987): 1052-1078; [5] J.W.Zwolak et al. ACM Trans. Math.Soft., 33,4 (2007); [6] D. Burstein, et al. MRM. 45 (2001): 36-41.