

A New 3D Isotropic $T_{1\rho}$ Mapping Technique for *In Vivo* Human Knee Cartilage at 7T MRI

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Introduction: Osteoarthritis (OA) is a degenerative disease affecting Articular Cartilage (AC) of synovial joints in shoulders, knee, hip, etc. It is reported that ~5% of the population over 25 years of age and ~12% over 60 years of age are diagnosed with knee OA¹. Spin-lattice relaxation in the rotating frame ($T_{1\rho}$) has been a promising quantitative MRI imaging technique to diagnose OA at an early stage by detecting changes in the proteoglycan and collagen matrix in the cartilage². High resolution MRI is desired across the thickness of AC (which varies from 1-6 mm) particularly in OA patients to identify the extent of the cartilage degeneration. With current methods, due to the practical scan time considerations, two 2D/3D scans in axial and coronal/sagittal orientations are performed for patellar and femoral/tibial cartilages respectively with <1mm in-plane resolution and 3-5 mm slice thickness^{3,4}. Here we propose a new 3D-GRE based time efficient method to acquire $T_{1\rho}$ weighted images of the whole knee at an isotropic resolution (0.5 mm³) within 12 min scan time that allows reformatting of the isotropic images of knee to any desired orientation with $T_{1\rho}$ maps calculated for patellar, femoral and tibial cartilages and validated the proposed method with existing technique.

Materials and Methods: All the human studies were conducted under an approved Institutional Review Board protocol of the University of Pennsylvania. Seven healthy volunteers of age 32.8 ± 14 years underwent MRI at whole body 7T scanner (Siemens Healthcare, Erlangen, Germany) using a 28-channel Knee Coil (Quality Electrodynamics, OH, USA). A new 3D $T_{1\rho}$ sequence has been developed as shown in Fig 1. Readout is done with very low flip (~5°) angle spoiled gradient echo segments (~600 echo segments per shot with a GRAPPA factor of 2 along phase encoding direction) to obtain rapid 3D images with 30 shots of TR 6s each. Using this 'burst mode' shot strategy, a high throughput rate (>80%) for readout is achieved. Spin-lock pulse amplitude was kept constant at $B_1 = 500\text{Hz}$ for various spin lock times, $TSL = 0, 14, 28$ and 42 ms with $TR/TE = 8.7/3.32$ ms and $NEX = 1$, $FOV = 160 \times 160 \times 80$ mm³, matrix size = $320 \times 320 \times 160$ with 0.5 mm³ isotropic resolution in axial orientation. Total acquisition time was 12 min. In addition, $T_{1\rho}$ weighted images in axial and coronal orientations were acquired in $0.5 \times 0.5 \times 3$ mm³ resolution for the comparison of $T_{1\rho}$ maps with the proposed technique. The data corresponding to different TSLs were fitted to a mono-exponential decay expression $S(TSL) = (M_0 - M_{ZSS}) \times e^{(-TSL/T_{1\rho})} + M_{ZSS}$ to generate $T_{1\rho}$ maps, where ' TSL ' is the spin lock time in 'ms', ' M_0 ' is the thermal equilibrium and ' M_{ZSS} ' is the steady-state Z component of magnetization. The images of isotropic resolution were reformatted to $0.5 \times 0.5 \times 3$ mm³ slices in axial and coronal orientations to compare the results with non-isotropic acquisitions.

Results: Axial $T_{1\rho}$ maps of a healthy volunteer obtained from the proposed technique and reformatted to coronal and sagittal orientations are shown in Fig 2. $T_{1\rho}$ values of knee cartilages calculated using the proposed method in isotropic resolution (center slice) and reformatted to 3 mm thick slices are shown in Fig 3A (patellar, 0.5 mm center slice / 3mm reformatted: 34.36 ± 4.0 / 32.66 ± 4.0 ms; femoral: 44.6 ± 2.0 / 43.38 ± 1.3 ms and tibial: 39.71 ± 4.1 / 38.28 ± 3.3 ms). Accuracy of the proposed approach was validated by reformatting the isotropic images to axial and coronal orientations to 3mm thick slices and the $T_{1\rho}$ values of each cartilage are compared with the $T_{1\rho}$ values obtained

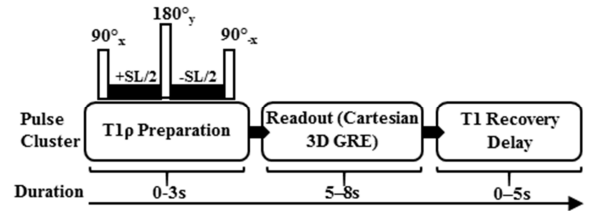


Fig 1 The preparation block consists of a Spin lock preparation⁵ for $T_{1\rho}$ measurements followed by 3D GRE readout to spatially encode the signal and a delay block to allow T_1 recovery

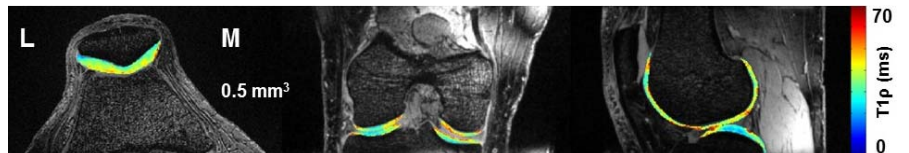


Fig 2 Axial, reformatted - coronal and sagittal sections of the healthy knee showing $T_{1\rho}$ maps acquired using 3D isotropic MRI.

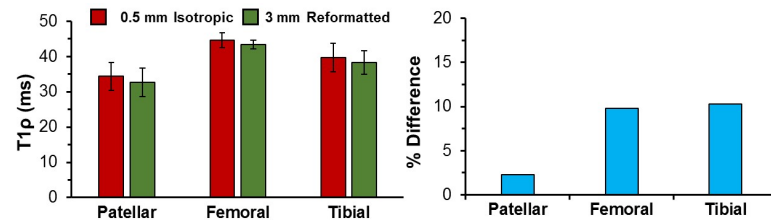


Fig 3 A. $T_{1\rho}$ values of cartilages calculated from images acquired at isotropic resolution (center slice) and reformatted to 3mm thick slices. B. % change in $T_{1\rho}$ values calculated from isotropic acquisition with non-isotropic acquisition.

and rapid isotropic sequence to acquire $T_{1\rho}$ -weighted images and quantitatively validated the accuracy of the $T_{1\rho}$ values calculated. Since AC of knee is a curved surface, the proposed isotropic $T_{1\rho}$ imaging may improve the diagnostic accuracy particularly in OA patients at a practically achievable scan time. Work is in progress to evaluate the reason for the deviation of $T_{1\rho}$ values in femoral and tibial cartilages from the conventional method and validate this technique with additional healthy volunteers of different age groups and subjects with knee pain.

References: 1. Lawrence et al (2008) *Arthritis Rheum* 58:26-35; 2. Wheaton et al (2005) *MRM* 54:1087-1093; 3. Singh et al (2014) *PLoS ONE* 9(5): e97486; 4. Borthakur et al (2003) *JMRI* 17:730-736; 5. Witschey et al (2007) *JMR* 186:75-85.

Acknowledgments: This work is supported by the National Institute of Biomedical Imaging and Bioengineering under award number P41-EB015893.

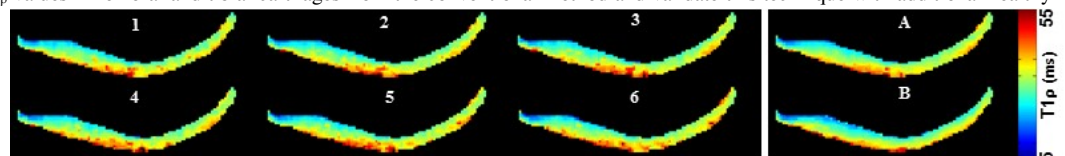


Fig 4 $T_{1\rho}$ map of segmented patellar cartilage in axial view, 1 to 6 shows slices with isotropic resolution, A reformatted slice by combining 1 to 6 and B $T_{1\rho}$ map calculated from $T_{1\rho}$ weighted image acquired with 3mm slice thickness