3D distribution of T1p relaxation times in the human knee joint

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Introduction: Previous studies have shown the MR relaxation time T1p is a surrogate biomarker for changes in the extracellular matrix of articular cartilage and may be sensitive to osteoarthritis (OA) progression (1). T1p increases in controlled degradation studies in ex vivo bovine plugs (2), porcine (3) and guinea pig (4) models of arthritis. In this study, we present the first 3D assessment of T1p among all regions of the knee joint articular cartilage in a subset of asymptomatic subjects (N=17) of the early OA T1p clinical trial.

Methods: Each subject was scanned on a 1.5 T clinical MRI system equipped with 40 μT/mm gradients (Siemens AG; Erlangen, Germany; Sonata Model) and an 8-channel transmit/receive 1H RF coil (Philips; Amsterdam, Netherlands; previously Invivo; Latham, NJ). The patella was centered with the leg flexed in the RF coil and scanned using the imaging protocol listed in Table 1. T1p images were acquired with a previously described T1p prepared balanced gradient echo sequence. Cartilage segmentation was performed using a LiveWire algorithm in

3DViewnix (MIPG; University of Pennsylvania; Philadelphia, PA) and imported into a custom image processing software constructed in Matlab (Natick, MS) to extract T1p by cartilage layer (superficial, middle, deep zones), side (medial, lateral), and location (patellar, femoral and tibial articular cartilage). A musculoskeletal

Sequence	Orientation	Contrast	Acquisition Type	TE/TR/flip (ms/ms/degrees)	Resolution (mm)	Imaging Matrix	Duratio n (min)	Slices	Thickness (mm)
b-SSFP	Triplane	T1	2D	2.56/5.12/70	1.17x1.17	256x256	0.5	10*3=30	3.0
b-SSFP	Sagittal	T1	2D	2.56/5.12/70	0.55x0.55	256x256	0.5	30	3.0
FSE	Axial	T2	2D	25/2850/90-180	0.55x0.55	256x256	2.0	25	3.0
FSE	Coronal	T2	2D	25/2850/90-180	0.55x0.55	256x256	2.0	25	3.0
MP-RAGE	Sagittal	T1	3D	18.6/37.1/90	0.55x0.55z0.55	256x256x208	8.2	208	0.55
SLIPS	Axial	T1p	3D	3.7/7.4/20	0.55x0.55x3.0	256x128x30	15.0	30	3.0
SLIPS	Coronal	Tlρ	3D	3.7/7.4/20	0.55x0.55x3.0	256x128x30	15.0	30	3.0

radiologist evaluated all images for bony cysts, meniscal tears, cartilage thinning, loss and other irregularities.

Results: Among the 17 asymptomatic subjects ages 30-60, T1p was significantly different between patellar, femoral and tibial regions (p<0.001) and between the superficial, middle and deep zones of the cartilage (p<0.01). Fig.1 depicts mean T1p for all subjects, medial and lateral sides of the knee and shows patellar cartilage T1p is elevated, but with greater variance than the middle and deep zones. Medial and lateral mean T1p were 44.6 ± 14.0 and 45.3 ± 14.0 ms, respectively. Superficial, middle and deep zone T1p were 47.9 ± 15.1, 43.7 ± 13.4, and 42.3 ± 13.2 ms, respectively. Patellar, femoral and tibial T1p were 55.7 ± 16.9, 43.2 ± 8.5, and 36.0 ± 6.2 ms, respectively. No significant difference in T1p was detected by side or age. Elevated T1p was confirmed in one subject corresponding with focal loss of the articular cartilage on the medial patellar facet with an underlying bony cystic change as shown in Fig. 2.

Discussion: T1p is lower in tibial and femoral cartilage compared to the patellar cartilage likely reflecting differences in loading and associated changes in molecular content. The T1p gradient from superficial to deep zones likely reflects the increasing concentration of proteoglycan from the joint space surface to the subchondral bone. The complete characterization of T1p relaxation times among all regions of the knee articular cartilage will aid diagnosis of cartilage lesions and progression of early OA symptoms as shown in Fig. 2.

References: (1) Borthakur, et al. NMR Biomed, 2006. (2) Wheaton AJ, et al. Magn. Res. Med. 2005. (3) Wheaton, AJ, et al. Acad. Radiol. 2004. Witschey II W.R.T., et al. Rapid 3D T1p-weighted Imaging (JMRI: accepted for publication).







Fig. 2: 3D crossections of the patellar cartilage in a subject with focal cartilage loss and bony cyst (arrow) and corresponding 2D T2 slice.