Quantitative T2*-Mapping of the Achilles Tendon using a Multi-Echo VTE SPGR-sequence at 3 Tesla: preliminary results. Sebastian Apprich ${ }^{1}$, Vladimir Juras ${ }^{1}$, Oliver Bieri ${ }^{2}$, Stefan Zbyn $^{1}$, and Siegfried Trattnig ${ }^{1}$
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## Target audience

Musculoskeletal radiologists, orthopedic surgeons

## Purpose

A new multi-echo (me) spoiled gradient echo sequence (SPGR) with an variable echo time (vTE) scheme (me-vTE-SPGR) enables quantitative MR imaging of tissues with very short T2 relaxation times (1) such as the Achilles tendon (AT). The purpose of this study was to compare the T2*-relaxation times of patients with an injured AT with intact AT in volunteers using this noval multi-echo VTE sequence at 3 T .

## Methods

Institutional Review Board approval and written, informed consent were obtained.
Ten patients (mean age $43.9 \pm 13.4$ years) suffering from a painful AT and 10 age-matched, healthy volunteers (mean age $43.7 \pm 11.2$ years) were examined with a 3T whole-body system, using an 8channel knee coil. Morphological MRI included a sagittal T2-w fast spin echo sequence. T2* maps were calculated from an isotropic 3D me-vTE-SPGRsequence ( 20 TE's from 0.8 to 20.1 ms ) using a monoexponential fit least square analysis. T2* values were manually assessed using a ROI analysis for the most severe pathology within patients and for the distal twothirds of the AT in volunteers. All subjects completed the Achilles tendon Total Rupture Score (ATRS; 0-100 points). Statistical measures included an analysis of variance and Pearson-Correlation.

## Results

Mean T2*-values in patients ( $8.0 \pm 4.7 \mathrm{~ms}, 95 \% \mathrm{Cl} 2.5-$ 18.3 ms ) differed statistically significant $(P=0.002)$ from mean T2*-relaxation times in healthy volunteers ( $2.6 \pm 0.6 \mathrm{~ms}, 95 \% \mathrm{Cl} 1.5-3.3 \mathrm{~ms}$ ). Furthermore, a strong correlation was found between the clinical ATRS and mean T2*-values (Pearson coefficient $0.846, R^{2}=0.72, \mathrm{P}<0.001$ ).
Morphological observation revealed 3 cases of achillotendinitis (thickening of AT, but no increase in signal intensity (SI)), 3 cases of thickened ATs with increase in SI (but not isointense to fluid), and 4 partially ruptured ATs (SI within the tendon isointense to fluid). Mean T2*-values for these cases were 3.6 $\pm 0.5 \mathrm{~ms}, 7.1 \pm 0.4 \mathrm{~ms}$, and $12.3 \pm 4.1 \mathrm{~ms}$ (Figure 1.), respectively.

## Discussion

Conventional MRI sequences are not able to detect MR signal from tissues with short T2 relaxation times. The me-vTE-SPGR-sequence enables quantitative imaging of healthy ATs with short T2 relaxation as well as of injured ATs in which T2* relaxation times are prolonged due to increased water content and anisotropy of collagen fibers.
This new sequence might be in future a non-invasive predictive marker for the probability of a rupture of an


## Conclusion

T2*-Mapping using a multi-echo VTE sequence is able to quantify changes in the water content and collagen fibre organisation of the injured AT.

## References

1. Deligianni et al, High resolution FourierEncoded Sub-Millisecond Echo Time Musculoskeletal Imaging at 3T and 7T, MRM

Figure 1: Patient with partial rupture of Achilles tendon at the insertion zone. Note the increased T2*relaxation times in the distal part of the AT.


