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### Ultrahigh-Field Multimodal MRI Assessment of Muscle Damage

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#### To The Editor:

Magnetic resonance imaging (MRI) is a powerful tool to assess skeletal muscle damage both in injured athletes and patients with neuromuscular diseases. Alterations in muscle have been assessed with  $T_2$ -weighted,<sup>1</sup> diffusion tensor,<sup>2</sup> and sodium MRI.<sup>3</sup> However, these sequences require a high signal-to-noise ratio. Considering these limitations, one could expect that measurements performed at ultrahigh-field strength (eg, 7T) would provide a more accurate assessment of skeletal muscle diffusivity properties<sup>4</sup> and sodium concentration<sup>5</sup> so that relevant information regarding changes in biochemical and structural tissues properties related to muscle damage may be obtained.

A 33-year-old man (188 cm, 82 kg) volunteered to participate in the study after providing informed consent. Calf muscles were injured during a high-intensity electrostimulation exercise performed using a commercial device (Compex Performance, Djo global, France). Muscle MRI was performed using a whole-body scanner (Magnetom 7T, Siemens Healthcare, Erlangen, Germany). Five days after the damaging exercise, a multimodal imaging session was performed. Highresolution T1-weighted images (a gradient-recalled echo [GRE] with parameters previously detailed<sup>4</sup> and a turbo spin-echo, repetition time [TR] = 3500 msec, echo time [TE] = 34 msec, in-plane field of view  $[FoV] = 180 \times 180 \text{ mm}^2$ , matrix = 640 × 640, 15 slices, ST = 6 between slice = 6 mm, bandwidth = 230 Hz/px, mm, gap TA = 240 sec) were acquired and the corresponding images were used as anatomical reference images. A multiecho GRE sequence (TR = 43 msec, TEs = 3.57/8.51/13.60/18.69/23.78/28.87/33.96/39.05 msec, flip angle =  $10^{\circ}$ , in-plane FoV =  $180 \times 180 \text{ mm}^2$ , matrix =  $448 \times 10^{\circ}$ 448, 60 slices, ST = 3 mm, bandwidth = 370 Hz/px, TA = 456 sec) was used to produce  $T_2^*$  maps, while  $T_2$  maps were generated from T<sub>2</sub>-weighted images acquired with a segmented (15 segments) echo planar imaging sequence with TEs = 8/18/28/38/48/58 msec. Other acquisition parameters were as follows: FoV =  $180 \times 180 \text{ mm}^2$ ; matrix = 192 × 192; TR = 4800 msec; 20 slices; ST = 6 mm; gap between slices = 6 mm, short-tau inversion-recovery for fat saturation; TA for each TE = 132 sec. Axial multidirectional diffusionweighted images with fat saturation were acquired to characterize diffusion tensor imaging derived parameters as previously described.<sup>4</sup> A 3D radial sequence was used to obtain sodium images using the following parameters: TR = 110 msec, TE = 0.2 msec, FoV = 180 ×  $180 \times 180 \text{ mm}^3$ , voxel resolution =  $2 \times 2 \times 2 \text{ mm}^3$ ,  $10^4$  radial

spokes, bandwidth = 276 Hz/px, TA = 1102 sec. High-resolution T<sub>1</sub>-weighted images were used as reference images (GRE, TR = 13 msec, TE = 5 msec, in-plane FoV = 180 × 180 mm<sup>2</sup>, matrix = 512 × 512, 88 slices, ST = 2 mm, bandwidth = 260 Hz/px, TA = 587 sec). Sodium images were acquired in the presence of six calibration phantoms with different sodium concentrations (10, 30, and 50 mM) and placed within the FoV. Images were then corrected from B<sub>1</sub> inhomogeneities using precalibrated B<sub>1</sub> sensitivity maps from a phantom acquisition measured using the phase sensitive method.<sup>6</sup>

Additional investigations were performed at 1.5T (Avanto, Siemens Healthcare) in order to quantify muscle  $T_2$  values averaged on all slices 3 (D3), 5 (D5), 7 (D7), and 14 (D14) days after the damaging exercise, as previously described.<sup>7</sup> Control muscle  $T_2$  values were determined 6 months (M6) after the damaging exercise.

As illustrated in Fig. 1a, 1.5T measurements showed an increased  $T_2$  value in gastrocnemius medialis reaching the highest intensity 5 days after the damaging exercise and being resolved after a 6-month period. A slight  $T_2$  increase was observed in the soleus at D3 but no other changes could be seen in antagonist muscles. Ultrahigh-field MRI measurements confirmed the  $T_2$  increase (Fig. 1b) and extended our knowledge, showing concomitant local changes in  $T_2^*$  and sodium concentration (in comparison to unaffected muscles, eg, lateral compartment of the lower leg) in addition to transverse diffusivity ( $\lambda_3$ ) and fractional anisotropy (FA) as compared to published control values.<sup>4</sup> In addition, diffusion parameters were increased in other areas, ie, FA and sodium concentration in the gastrocnemius lateralis, eigenvalues ( $\lambda_1$ ,  $\lambda_2$ ,  $\lambda_3$ ), mean diffusivity (MD),  $T_2^*$ , and sodium concentration in the anterior compartment of the lower leg muscles (Fig. 1c).

The 7T MRI measurements showed additional changes in the same muscle regarding diffusion properties and sodium concentration together with changes in agonist and antagonist muscles that were not visible from the 1.5T measurements.

Studies have previously reported  $T_2$  changes resulting from damaging exercise and assigned them to inflammatory/edematous processes.<sup>1</sup> Changes in DTI metrics have been reported in the early phase of muscle damage,<sup>8</sup> illustrating subtle alterations of muscle diffusion properties. Sodium MRI can provide further information regarding biochemical changes in the damaged muscle, as recently shown in Achilles tendon<sup>9</sup> and knee cartilage.<sup>10</sup>

From 1.5T MRI measurements, a large  $T_2$  increase was quantified in the gastrocnemius medialis muscle as a result of the damaging muscle stimulation. Interestingly, measurements performed at higher field (7T) confirmed the  $T_2$  changes but also illustrated changes in agonist and antagonist muscles that were not detected at 1.5T. Larger diffusion parameters values were measured in the anterior compartment of the lower leg muscles, thereby suggesting subtle changes in the diffusion properties. These changes could be

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FIGURE 1: (a)  $T_2$  maps overlaid on anatomic images recorded at 1.5T for a proximal and a distal slice of the lower leg. Mean  $T_2$  values for the triceps surae muscles (gastrocnemius medialis [GM], gastrocnemius lateralis [GL], and soleus [SO]) and the muscles from the lateral and the anterior compartments of the lower leg (LCLL and ACLL, respectively), and maximal voluntary isometric plantarflexion force (MVC) determined 3 days (D3), 5 days (D5), 7 days (D7), 14 days (D14), and 6 months (M6) after the neuromuscular electrical stimulation exercise inducing muscle damage are specified below. (b) Quantitative maps of  $T_2$ ,  $T_2^*$ , diffusion tensor-derived parameters ( $\lambda_1$ ,  $\lambda_2$ ,  $\lambda_3$ , mean diffusivity [MD], fractional anisotropy [FA]) and sodium concentration ([Na]) overlaid on transverse anatomical images of the lower leg muscles recorded 5 days after the damaging exercise at 7T. Arrows indicate local changes in  $T_2$ ,  $T_2^*$ ,  $\lambda_3$ , FA, and [Na] (yellow arrow: decrease, white arrow: increase). Specific localized changes in FA and [Na] are displayed, suggesting a specific characterization of muscle structure and biochemical environment, respectively. The sodium concentration and  $T_2^*$  averaged on all slices were 63 ± 16 mM and 22.5 ± 5.2 msec for the GM, 27 ± 7 mM and 13.9 ± 4.9 msec for the GL, 32 ± 7 mM and 15.2 ± 1.9 msec for the soleus, 15 ± 6 mM and 15.1 ± 2.3 msec for the LCLL, and 27 ± 11 mM and 16.3 ± 1.8 msec for the ACLL. (c) Diffusion tensor imaging-derived parameters (green marker) including eigenvalues ( $\lambda_1$ ,  $\lambda_2$ ,  $\lambda_3$ , mean diffusivity (MD) and fractional anisotropy (FA) of the triceps surae muscles (gastrocnemius medialis [GM], gastrocnemius lateralis [GL] and soleus [SO]) and the muscles from the lateral and the anterior compartments of the lower leg (LCLL and ACLL, respectively) for the case subject superimposed on boxplots of values obtained in 14 healthy controls from a previous study.<sup>4</sup>

explained by muscle co-contraction of agonist and antagonist muscles during the damaging exercise and/or muscle balance modification leading to overcompensation of the undamaged muscles. In addition, an increased FA was observed in the neighboring soleus muscle area, which could be due to the stress mediated by the increased muscle volume of the gastrocnemius medialis associated with the edema/inflammation processes.

In conclusion, multimodal 7T MR of skeletal muscle shows potential. Performed in a larger number of subjects, the corresponding longitudinal changes could provide a better understanding of the physiopathological mechanisms underpinning muscle tissue alteration and regeneration and of the intramuscular changes that are driving  $T_2$  changes. This ultrahigh-field MRI multimodal approach could be helpful in athletes and patients with neuromuscular disorders for the assessment of biochemical and/or microstructural changes occurring in skeletal muscle after injuries, during sport reconditioning, and in case of surgical intervention.

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