Diffusion Tensor Imaging of Liver Fibrosis in an Experimental Model

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Introduction

Though percutaneous liver biopsy is considered the gold standard for assessing liver fibrosis, its utility as a tool for longitudinal monitoring has been limited because repeated biopsies of liver are not practicable clinically. Early diagnosis of liver fibrosis could facilitate early interventions and thus alleviate its progression to cirrhosis and/or hepatocellular carcinoma. Therefore, noninvasive imaging technique is needed to detect liver fibrosis at early stage and monitor the disease progression or treatment effects. Several studies have shown that measurement of water diffusivity by diffusion-weighted imaging (DWI) was useful in evaluation of liver fibrosis and cirrhosis. The aim of this study was to characterize longitudinal changes in diffusion properties of liver using diffusion tensor imaging (DTI) in an experimental model of liver fibrosis.

Methods

Animal Procedures: Liver fibrosis was induced in male Sprague-Dawley (SD) rats (220-260 g; N = 8) by subcutaneous injection of 1:1 mixture of CCl4 in olive oil at a dose of 0.2 mL/100g of body weight twice a week for 4 weeks. Animals were scanned before, 2 and 4 weeks after CCl4 insult.

In Vivo MRI: MRI experiments were performed on a 7T Bruker MRI scanner with a 60 mm RF Tx/Rx quadrature resonator. Animals were fasted before scanning to reduce peristalsis. Each animal was anesthetized with ~1.5% isoflurane and maintained at about 36.5°C with respiratory monitoring. DTI was performed on one axial slice passing through the liver with respiratory-gated single-shot spin-echo-echo-planar imaging (SE-EPI) sequence using TR = 2 respiratory cycles (~2.0-2.5 s), TE = 32 ms, two b-values were used (0 and 1000 s/mm2) for 6 diffusion gradient directions, FOV = 5.12×5.12 cm², slice thickness = 2 mm, acquisition matrix = 64×64, voxel size = 0.8×0.8×2 mm³, NEX = 10, and total scan time of ~3 min. The DTI acquisition was repeated twice, and measurements of parameters were averaged.

Data Analysis: DW images were first co-registered using AIR5.2.5. Apparent diffusion coefficient (ADC), fractional anisotropy (FA) and directional diffusivity (ADC⊥ and ADC∥) maps were generated using DTIStrain2. A large region of interest (ROI) was defined in the anatomical scout image to encompass a large homogeneous region of liver parenchyma for DTI measurements. One-way ANOVA was employed to compare differences in parameters among different time points, with P values less than 0.05 considered statistically significant.

Histology: Of the 8 animals scanned at 2 weeks after CCl4 insult, 2 were sacrificed after MRI for histological analysis. 4 weeks after CCl4 insult, 2 of the remaining 6 animals were sacrificed after MRI. Furthermore, one normal animal was sacrificed as a control. The livers were embedded in paraffin, sectioned, and examined by light microscopy after standard hematoxylin-eosin (H&E) staining.

Results

Fig. 1 shows the representative anatomical scout images, ADC maps and FA maps of liver before, 2 and 4 weeks after CCl4 insult. ADC, FA, ADC⊥, and ADC∥ values of different time points of CCl4 insult are shown in Fig. 2. Significant decrease was found for ADC 2 weeks (P < 0.05) and 4 weeks (P < 0.01) after CCl4 insult. FA at 2 weeks after CCl4 insult was significantly lower than that before (P < 0.01) and 4 weeks after (P < 0.05) the insult. Compared with normal liver, collagen depositions and widespread fat vacuoles were consistently observed in livers after CCl4 insult, as shown in Fig. 3. Cell necrosis/apoptosis was evident in liver after 2-week CCl4 insult (Fig. 3b), while collagen depositions were more pronounced in liver after 4 week of CCl4 insult.

Discussions

The histological observations confirmed established liver fibrosis in the animals studied. Despite the potential water diffusivity increase caused by cell necrosis/apoptosis, ADC was observed to be decreasing gradually after CCl4 insult, probably due to the increased extracellular collagen deposition and intracellular fat droplets during the progression of liver fibrosis (Fig. 3b and 3c). FA decrease at 2 weeks after CCl4 insult resulted from the significant decrease of ADC (P < 0.01). This was likely due to the prominent cell necrosis/apoptosis occurred (Fig. 3b), perturbing water diffusion along the radially oriented hepatic architecture. The subsequent FA increase at 4 weeks after CCl4 insult arose from the significant decrease of ADC (P < 0.01) thereafter. This might be caused by the pronounced extracellular collagen depositions (Fig. 3c), leading to decreased water diffusivity in the more isotropic extracellular compartment.

Conclusion

The experimental results in this study demonstrated that DTI could detect longitudinal changes in diffusion properties of liver in an experimental model of liver fibrosis. Therefore, DTI may be valuable in detecting liver fibrosis at early phase and monitoring its progression.

References