

## Rapid evaluation of liver fat content using in-out-phase imaging in nonalcoholic fatty liver disease

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### Introduction

The aim of the current study was to examine the possibilities for quantification of liver fat content (LFC) in patients with nonalcoholic fatty liver disease (NAFLD) by in-out-phase MR-imaging. Signal intensities (SI) measured from in-out-phase imaging were compared to liver fat measurements by proton magnetic resonance spectroscopy (<sup>1</sup>H MRS), which was used as a reference method.

### Materials and methods

Sixty-one in-out-phase and <sup>1</sup>H MRS examinations of patients with type 2 diabetes at high risk for NAFLD were included in the study. Written informed consent was obtained from all subjects and the study protocol was approved by the local ethics committee. A 1.5 T MR imager (Signa Horizon LX, GE Medical Systems, USA) was used for transverse T1W dual-echo FSPGR (in-out-phase) imaging (TR = 150 ms, TE = 2.1 ms and 4.4 ms, FA = 75°) and a single voxel <sup>1</sup>H MRS (PRESS, TR= 3000, TE = 25 ms, figure 1) examination of the liver. Liver SI differences between in-phase and out-phase images were measured at the same location as the <sup>1</sup>H MRS examination. Pearson's correlation was used to study the relationship between measured SI differences from in-out-phase images and liver fat content measured by <sup>1</sup>H MRS.

### Results

A highly significant linear correlation was observed between LFC measured by <sup>1</sup>H MRS and SI differences calculated from the in-out-phase images (P<0.001, r=0.94, figure 2). Using the simple difference in SI between in-phase images and out-phase images an intercept of the regression line with the x-axis was observed at 5.1%, comparable to the upper limit of normal LFC. A discrimination with high sensitivity (95%) and specificity (98%) between normal and elevated LFC was observed.

### Conclusion

The findings of this study suggest that in-out-phase imaging can be used for estimation of LFC in patients with NAFLD. The cut-off value of 5.1% allows for objective rapid and reliable discrimination between normal or elevated LFC.

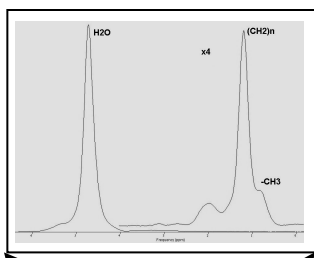


Figure 1: Typical voxel location and spectrum

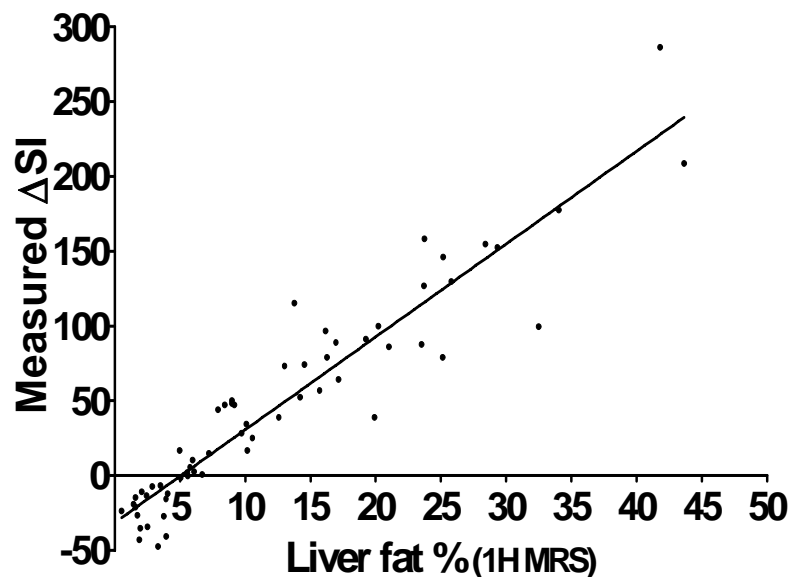


Figure 2: Linear correlation between in-out-phase SI differences and liver fat content measured by <sup>1</sup>H MRS