

# Influence of Magnetisation Transfer on established T1 mapping methods

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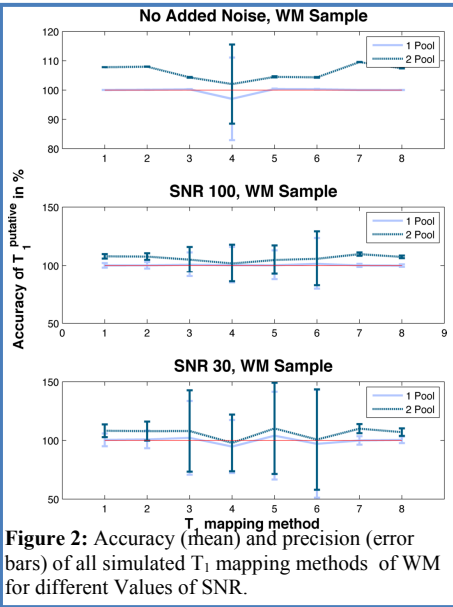
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## Introduction

T<sub>1</sub> mapping has proven to be useful for many purposes, e.g. in characterisation of different diseases or for quantitative water mapping [1,2]. Proper T<sub>1</sub> quantification mandates consideration of MT effects, as suggested e.g. by Ou [3]. However, consideration of MT introduces a large number of additional free parameters in T<sub>1</sub> fitting models. Thus, the stability of the fitting routine is decreased and high SNR is required in order to obtain robust fitting results. This limits the application of such methods to *in vitro*, phantom or post mortem studies. However, MT becomes important especially *in vivo*. To account for MT, the Bloch Equations need to be extended. This is done by the Bloch McConnell Equations [4], which describe the dynamics of a system in the chemical exchange regime. There, the observed or putative T<sub>1</sub> relaxation time is given as:

$$\frac{1}{T_1^{\text{putative}}} = \frac{1}{2} \left( \frac{1}{T_1^A} + k_{AB} + \frac{1}{T_1^B} + \frac{k_{AB} M_0^A}{M_0^B} \right) - \sqrt{\left( \frac{1}{T_1^A} + k_{AB} + \frac{1}{T_1^B} + \frac{k_{AB} M_0^A}{M_0^B} \right)^2 - 4 \left( \frac{1}{T_1^A T_1^B} + \frac{k_{AB}}{M_0^B T_1^A} + \frac{k_{AB} M_0^A}{M_0^B T_1^B} \right)}$$

As a consequence, the standard fitting models and signal equations used for T<sub>1</sub> fitting fail and compromise the accuracy and precision of the obtained T<sub>1</sub> values. This study aims at quantifying changes in the accuracy of T<sub>1</sub> in presence of MT if the signal equations used for T<sub>1</sub> determination specifically neglect the MT effect.



**Figure 2:** Accuracy (mean) and precision (error bars) of all simulated T<sub>1</sub> mapping methods of WM for different Values of SNR.

T<sub>1</sub> fitting, corresponding to different levels of SNR. T<sub>1</sub> maps were calculated neglecting possible effects on the signal equation due to MT. The accuracy [%] =  $100 - \frac{\text{mean}(T_1^{\text{sequence}_i}) - T_1^{\text{putative}}}{T_1^{\text{putative}}} \cdot 100$

and precision =  $\frac{\text{std}(T_1^{\text{sequence}_i})}{T_1^{\text{putative}}} \cdot 100$  of the method were obtained for each sample consisting of 576 voxels.

## Results

Figure 1 exemplarily shows the precision and accuracy of the T<sub>1</sub> of GM for different SNR obtained through variation of the flip angles in SpGE (3). For all the discussed methods, the accuracy and precision of conventional T<sub>1</sub> quantification without added noise, SNR 100 and SNR 30 in WM and GM are displayed in Figures 2 and 3, respectively.

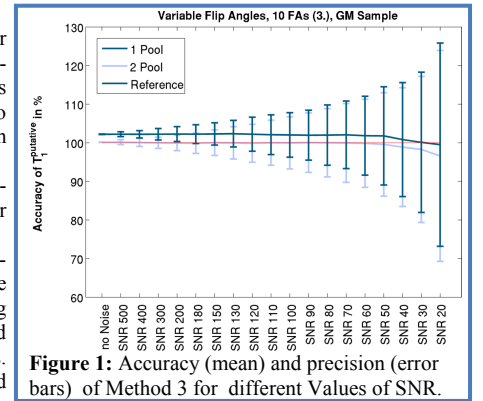
## Conclusion/ Discussion

It was shown that all methods give accurate T<sub>1</sub> estimations, with an absolute error below 10% for both samples with or without MT. All methods but one perform more accurately without chemical exchange. Method 4 is the exception where the accuracy actually worsens without MT. This comes at no surprise, as its fitting model is only valid in presence of the effect. Even without added noise, the precision of method 4 is comparatively low; this can be attributed to the larger number of fitting parameters (4) and as a consequence the less stable fit. Furthermore, the Look Locker method exhibits high sensitivity to the right choice of the flip angles.

Adding noise (SNR ≥ 20) to the simulated images decreases the precision while leaving the accuracy essentially unchanged. Consequently, the T<sub>1</sub> values of the two pool samples cannot be separated from the one-pool samples any more. One can conclude that the effect of MT is below the error of the method itself, if noise is present. The methods based on SpGE with different FAs are less prone to deviations due to MT, while the precision of the Look Locker based methods is higher, despite the fact that it is the approach with the shortest measurement time.

## References

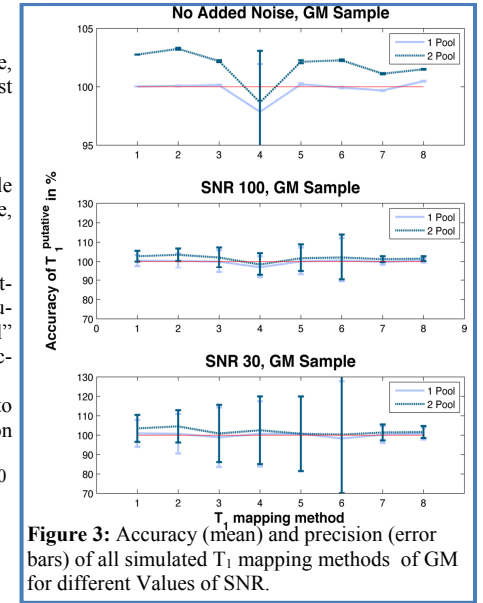
- [1] Shah N.J. et al. Hepatology 2003; 38: 1219 [2] Gowland P. et al. MRM 1992; 26:241 [3] Ou X. et al. MRM 2008; 59: 835 [4] McConnell. J. Chem. Phys: 1958; 28: 430  
[5] Kubach M.R. et al. 18th ISMRM, 2010 [6] Steinhoff S. et al. MRM 2001;46:131 [7] Stanisiz. et al. MRM 2005; 54: 507



**Figure 1:** Accuracy (mean) and precision (error bars) of Method 3 for different Values of SNR.

Sample Parameter	White Matter (WM)			Gray Matter (GM)		
	No MT	Pool A	Pool B	No MT	Pool A	Pool B
M <sub>0</sub>	1	0.861	0.139	1	0.95	0.005
T <sub>1</sub> / ms	1071	1084	1000	1748	1820	1000
T <sub>2</sub> / ms	69	69	0.01	99	99	0.0091
k <sub>AB</sub> / Hz	-	3.197	-	-	-	2
T <sub>1</sub> <sup>putative</sup> / ms	1071			1748		

**Table 1:** Sample parameters used in the simulations.



**Figure 3:** Accuracy (mean) and precision (error bars) of all simulated T<sub>1</sub> mapping methods of GM for different Values of SNR.