

# Combining Model Driven and Data Driven Approaches for Inverse Problems in Parameter Estimation and Image Reconstruction: From Modelling to Validation

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Thomas Yu  
Ecole Polytechnique Fédérale de Lausanne, LTS5



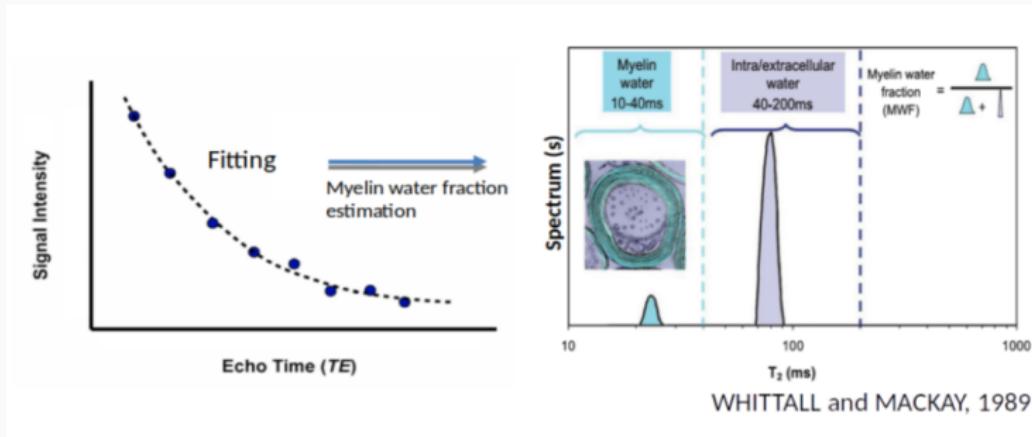
Realistic Modelling

Multicomponent  $T_2$  Relaxometry

# Realistic Modelling

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Multicomponent  $T_2$  Relaxometry



Can we integrate realistic models in a **supervised** machine learning approach to solve the highly ill-posed problem of recovering the  $T_2$  distribution?

*The following slides are based on the postprint version of the article: "Model-informed machine learning for multi-component  $T_2$  relaxometry" published in Medical Image Analysis [Yu et al., 2021a]. DOI: 10.1016/j.media.2020.101940.*

Within a single voxel, there are different tissues/materials each with different  $T_2$ 's We can recover these properties through MR spin-echo/gradient echo acquisitions,

$$\mathbf{s}(TE_i) = \int EPG(TE_i, T_1, T_2, \alpha)p(T_2)dT_2. \quad (1)$$

Traditional approaches include doing a non-parametric, non-negative least squares fitting using a dictionary of EPG signals:

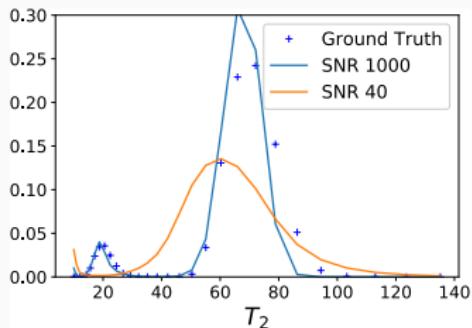
$$\arg \min_{\mathbf{p} \geq 0} \|D_\alpha \mathbf{p} - \mathbf{s}\|_2^2 + \lambda \Phi(\mathbf{p}) \quad (2)$$

or modelling  $p(T_2)$  as a mixture of distributions:

$$p(T_2) = \sum_{i=1}^n v_i F_i(\mathbf{m}_i, T_2) \quad (3)$$

and solving a nonlinear optimization problem.

1. State of the art non-parametric approaches tend to be highly sensitive to noise, with heuristic setting of regularization parameter. This leads to oversmoothing.
2. Gaussian mixture fitting requires long computation time and can be unstable without fixing parameters to set values. Furthermore, number of compartments need to be fixed.



**Figure 1:** Reconstructions from two different SNRs (40,1000) using a non-parametric approach( NNLS with Laplacian regularization)

In brain tissue, several assumptions are traditionally made about the  $T_2$  distributions:

1. In white matter, there are well-separated lobes corresponding to the myelin water and the intra/extraxonal space.
2. Overall, lobes of distinct tissue are well-separated.

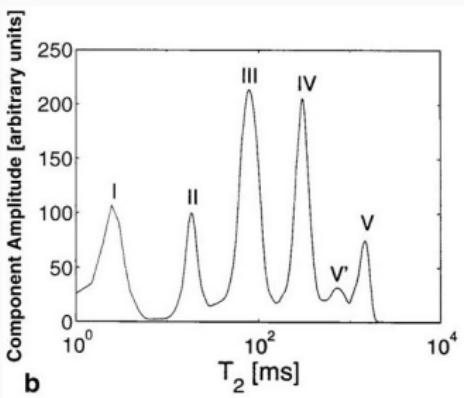
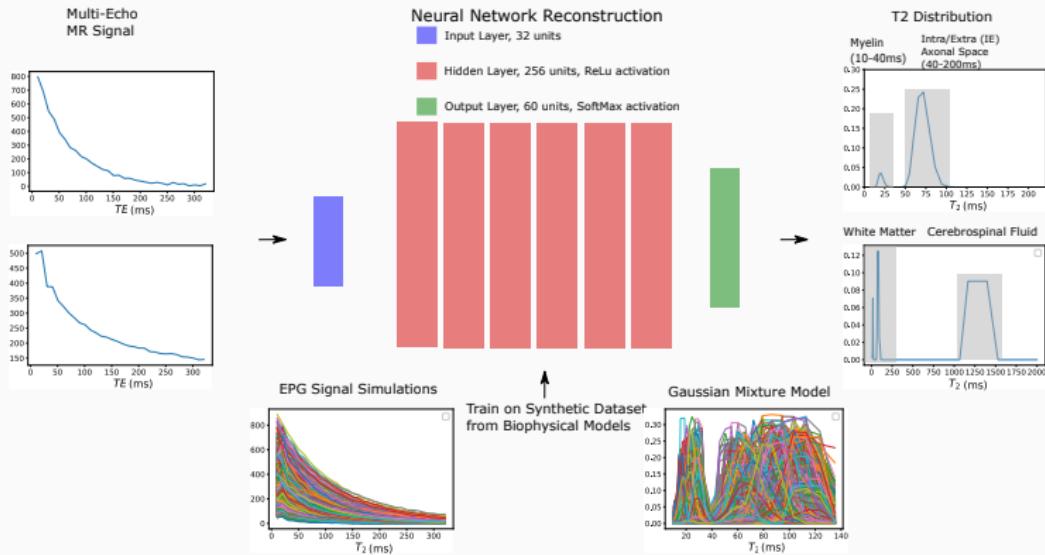


Figure 2: Multicomponent  $T_2$  of frog nervous tissue (SNR 20,000)  
[Peled et al., 1999]

## Model-Informed Machine Learning (MIML)



**Figure 3:** An overview of our proposed method for recovering  $T_2$  distributions [Yu et al., 2021b].

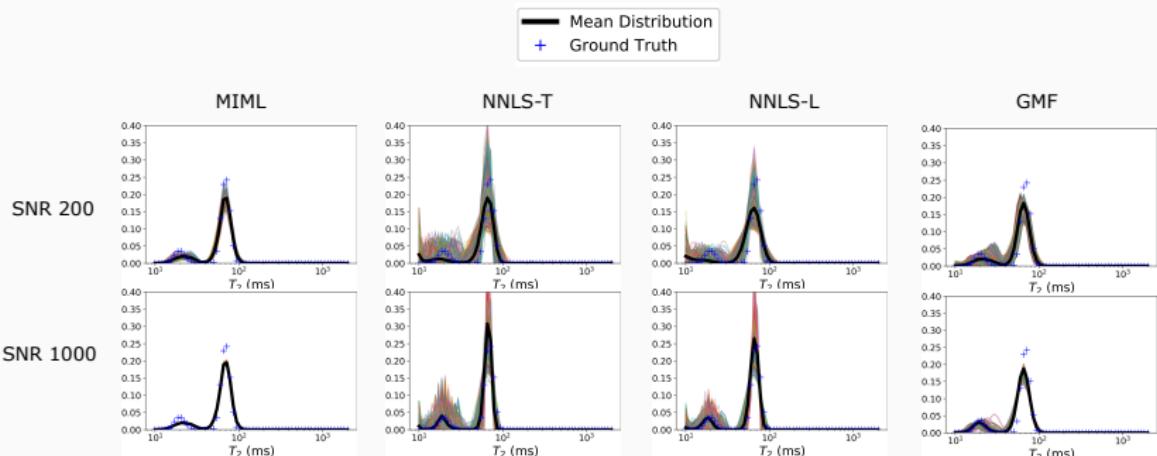


Figure 4: Results over a realistic, synthetic  $T_2$  distribution

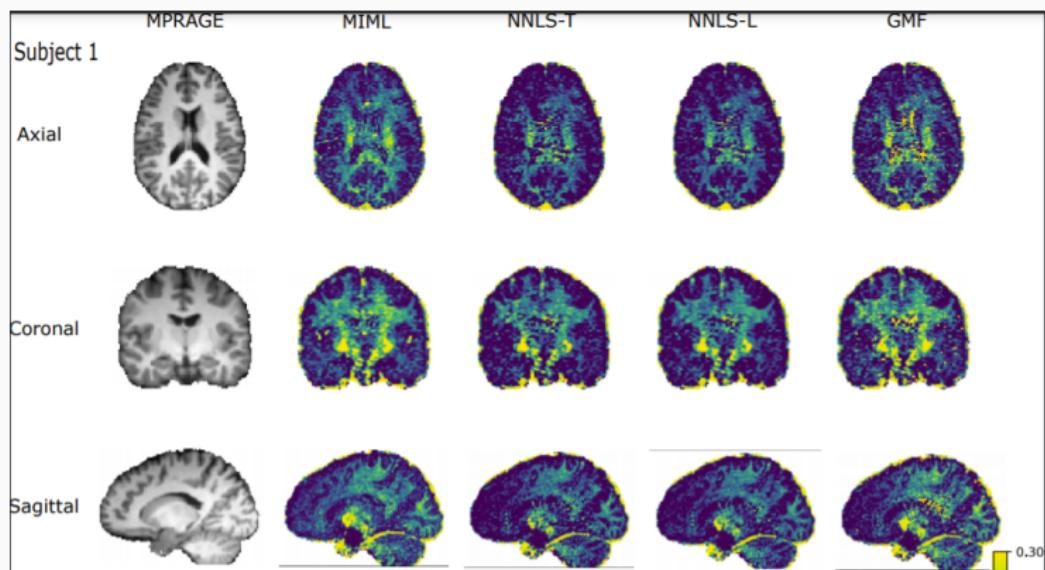


Figure 5: Myelin water fraction (MWF) maps from a healthy subject

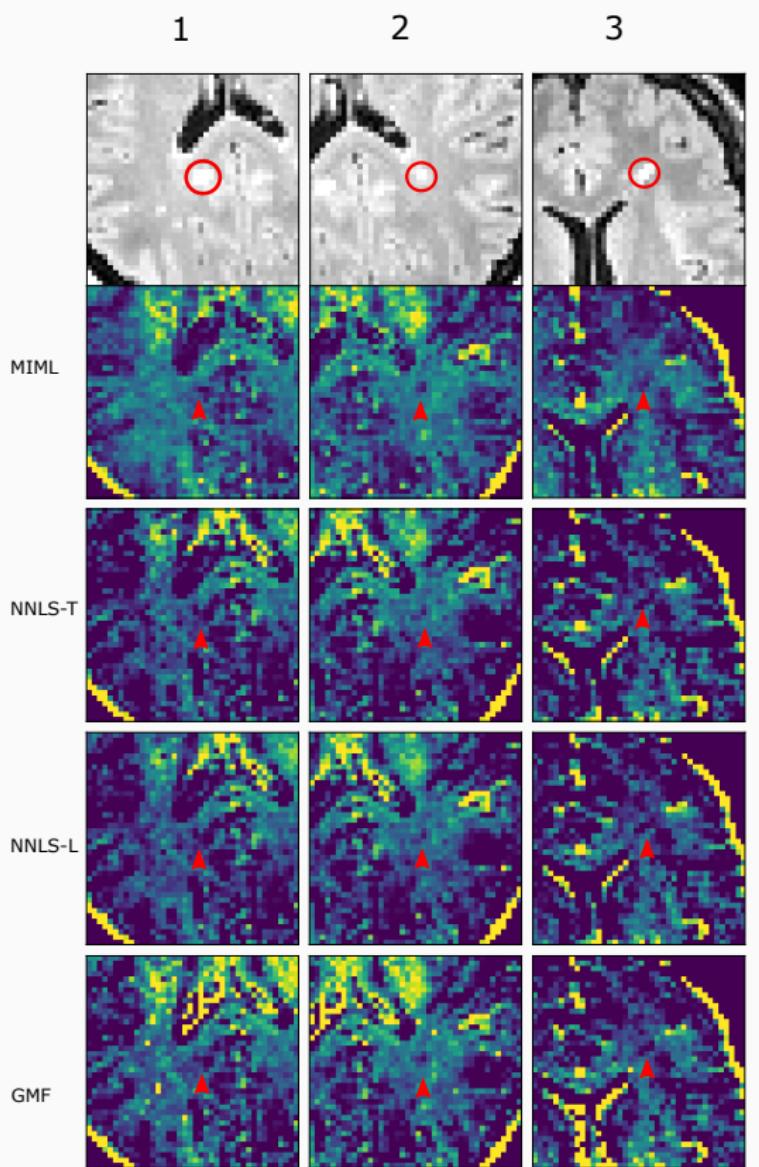


Figure 6: Closeup on MS lesions

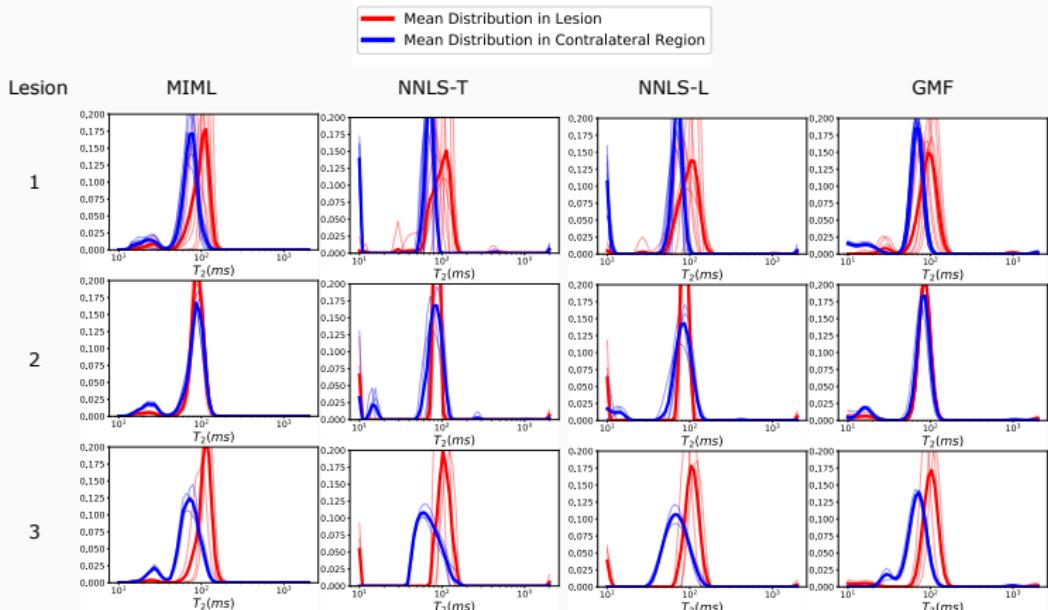


Figure 7: Mean  $T_2$  distributions in MS lesions.

1. Supervised machine learning trained solely on synthetic data which encodes realistic modelling/common assumptions on the structure of solutions can robustly and accurately solve the ill-posed problem of multi-component  $T_2$  estimation.
2. Furthermore, MIML is 22-4980 times faster than non-parametric or Gaussian mixture fitting.

-  Peled, S., Cory, D. G., Raymond, S. A., Kirschner, D. A., and Jolesz, F. A. (1999).  
**Water diffusion,  $t_2$ , and compartmentation in frog sciatic nerve.**  
*Magnetic Resonance in Medicine: An Official Journal of the International Society for Magnetic Resonance in Medicine*, 42(5):911–918.
-  Yu, T., Canales-Rodriguez, E. J., Pizzolato, M., Piredda, G. F., Hilbert, T., Fischi-Gomez, E., Weigel, M., Barakovic, M., Bach Cuadra, M., Granziera, C., Kober, T., and Thiran, J.-P. (2021a).  
**Model-informed machine learning for multi-component  $t_2$  relaxometry.**  
*Medical Image Analysis*, 69:101940.

-  Yu, T., Canales-Rodríguez, E. J., Pizzolato, M., Piredda, G. F., Hilbert, T., Fischi-Gomez, E., Weigel, M., Barakovic, M., Cuadra, M. B., Granziera, C., et al. (2021b).  
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