Contribution of quantitative MRI in the preclinical detection of early neurodegeneration of fronto-temporal dementia in patients with amyotrophic lateral sclerosis.

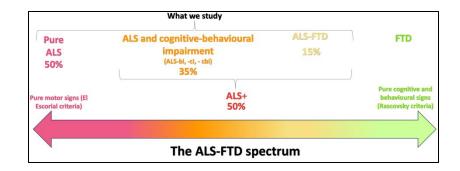
Geron, C.^{1,2}, Lamalle, L.¹, Philips, C.^{1a,1b}, Vandewalle, G.¹, Collette, F.^{1,3} & Maguet, P.^{1,2}

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Predicted changes in

ALS + vs. pure motor ALS & controls :

introduction



Currently: no validated biomarkers for early detection of FTD and ALS+

- ⇒ Misdiagnosis and delays
- ⇒ Impact disease management and prognosis

⇒ Shortened survival

Urgent need for validated biomarkers

Aim of the study

Identify early microstructural changes in ALS+ patients using quantitative MRI (qMRI) and use the derived parameters as imaging biomarkers for FTD.



Hypothesis

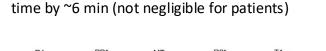
Early neurodegenerative microstructural changes can be identified before clinical symptoms

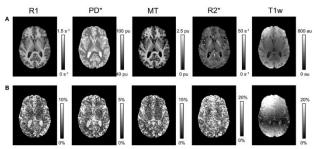
- ⇒ Precise quantification and localisation of the neurodegenerative processes
- \Rightarrow Early diagnosis in FTD and ALS+
- ⇒ Presymptomatic diagnosis (in familial forms)

Study Design

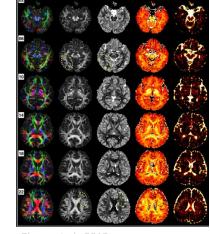
3T- quantitative MRI Protocol

- Multiparameter mapping: R1, R2*, MTsat, PD, QSM for microstructural tissue characterization
- Diffusion imaging (NOD DI) to estimate axonal diameter & neurite density
- Optimization: Spatial resolution adjusted (0.80 mm \rightarrow 1.07 mm), reducing scan





Weiskopf et al., 2013.



Zhang et al., 2012

Stastistical analysis

- PARTICIPANTS: 5 groups of patients and Healthy controls.
- MANOVA model in MSPM toolbox, with F-test to assess the overall multivariate effect.

Values	Peptide accumulation	Inflammation	Neuronal ⇒ myelin loss
R1			\
R2*		↑	\
QSM		↑	\
MT sat	Focal 个		\
Neurite density + orientation			\