

POSTER PRESENTATION

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Improving the classification of brain tumors in mice with perturbation enhanced (PE)-MRSI

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Classifiers based on pattern recognition analysis of MRS(I) data are becoming important tools for the non-invasive diagnosis of human brain tumors [1,2]. Here we investigate the potential interest of perturbation-enhanced MRSI (PE-MRSI), in this case acute hyperglycemia, during moderate brain hypothermia [3,4], for improving the discrimination between mouse brain MRS patterns of glioblastoma multiforme (GBM), low grade oligodendro-glioma (ODG2), and non-tumor brain parenchyma (NT). Six GBM-bearing mice and two ODG2-bearing mice were scanned at 7 Tesla by PRESS-MRSI with 12 and 136 ms echo-time, during euglycemia (Eug) and also during induced acute hyperglycemia (Hyp), generating altogether four datasets per animal (echo time + glycemic condition): 12Eug, 136Eug, 12Hyp, and 136Hyp. For classifier development, using in-house built software *SpectraClassifier 2.0* [5,6], all spectral vectors (spv) selected were unit length normalized (UL2) and used either as training set (76 GBM spv, four mice; 70 ODG2 spv, 2 mice; 54 NT spv, 6 mice) or as independent test set (2 mice, 61 GBM spv and 17 NT spv). All Fisher's LDA classifiers obtained had very good descriptive performance when extracting at least 10 features from the training sets as evaluated by Bootstrapping: correctly classified cases $\geq 99\%$. Evaluation of predictive performance with the independent test set clearly revealed that 12Hyp MRSI-based classifiers with at least 5 features provided the best robustness: balanced error rate (BER) for spv prediction $< 0.9\%$. This highlights the potential interest of perturbation-enhanced MRSI protocols for improving the non-invasive characterization of brain tumors at a preclinical level.

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