

Interleaved $^1\text{H}/^{31}\text{P}$ STEAM Spectroscopy of Exercising Human Gastrocnemius Muscle at 3 Tesla

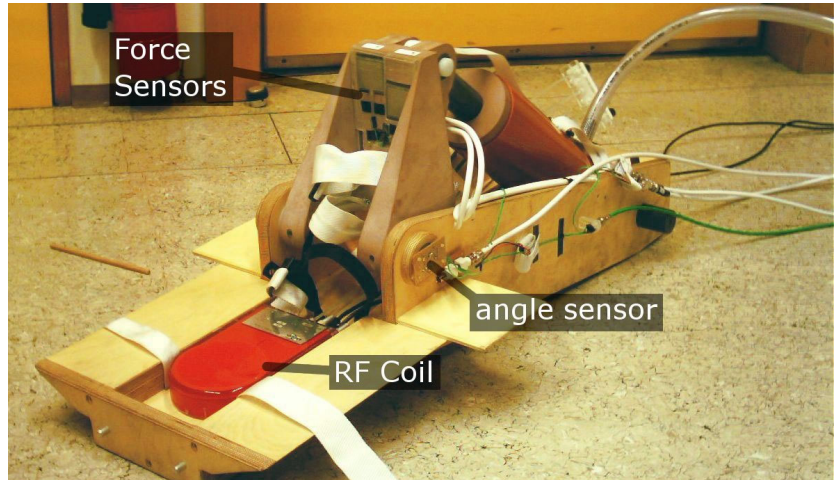
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Purpose/Introduction Proton and phosphorus magnetic resonance spectra can provide complementary information on tissue metabolism. Acquiring two sets of spectra in consecutive experiments is time-consuming and furthermore may result in undesirably modified test conditions, e.g. due to fatigue during studies of exercising muscle. Whilst ^{31}P MRS has long been used for investigation of exercising muscle, interest in ^1H spectroscopy of muscle tissue has increased over recent years [1]. In particular, the possibility to quantify changes in lactate concentration and compare them with changes in pH, PCr and P_i during exercise will remove a long-standing technical limitation to the study of the regulation of glycolysis, and also of cellular acid-base buffering mechanisms [2]. An interleaved STEAM sequence was developed to acquire localised ^1H and ^{31}P spectra from two independently positioned voxels in a single experiment, which has the additional benefit of NOE enhancement of the ^{31}P spectra. This work demonstrates the feasibility of interleaved acquisition of ^1H and ^{31}P spectra during plantar flexion in a pneumatic exercise rig.

Methods Interleaved acquisition of ^1H - and ^{31}P spectra with STEAM localisation was implemented on a 3T Bruker Medspec whole-body scanner via the MultiScanControl tool. The size, shape and position of the two voxels and other sequence parameters can be chosen independently. The integer number of excitations/acquisitions of ^1H spectra per ^{31}P excitation can be chosen to optimise T_R for the respective metabolites. T_1 of ^1H metabolites is typically on the order of 1s [3] while ^{31}P metabolites have significantly longer T_1 s, ranging from 3s to 6s at 3 Tesla [4]. A reasonable acquisition ratio is therefore $^1\text{H} : ^{31}\text{P} = 4 : 1$, yielding T_R s of e.g. 2s:8s and a time resolution of 16s (for minimum phase cycling). CHESS was used for efficient water suppression.

A non-magnetic rig (Fig. 1) was constructed for plantar flexion exercise with defined force during dynamic NMR studies. Whilst pushing a pedal against a pneumatic piston ($V=1.7\text{ l}$) the calf of the extended leg lies on a double tuned transmit-receive surface coil ($d=10\text{ cm}$) which is counter-sunk in a wooden frame. By varying the pneumatic pressure, pedal force can be adjusted arbitrarily e.g. to match the subject's MVC – even remotely, during the NMR measurement, and can be modified dynamically. Exercise is monitored continuously by force and position sensors on the pedal.



Results The good performance of the interleaved $^1\text{H}/^{31}\text{P}$ STEAM was verified using a two-compartment test object [4]. In vivo ^1H spectra of resting human calf muscle (5 subjects) were equivalent with standard STEAM experiments and with the interleaved $^1\text{H}/^{31}\text{P}$ STEAM sequence, while ^{31}P SNR was increased due to NOE by a factor of 1.34 ± 0.06 . Repetition time was $T_R=2\text{ s}$ for ^1H spectra (left) and $T_R=8\text{ s}$ for ^{31}P spectra (right). $T_E=8\text{ ms}$, $T_M=30\text{ ms}$, $\text{BW}=2.5\text{ kHz}$, 1024 data points. VOIs were 1.7 cm^3 (^1H) and 34 cm^3 (^{31}P).

Spectral quality was sufficient for quantification of creatine (Cr), choline (TMA) and extra- and intramyocellular lipid (EMCL & IMCL) from ^1H spectra and P_i and PCr in ^{31}P spectra, even during exercise. Significant PCr/ P_i changes could be achieved using the pneumatic exercise rig without deterioration of spectra e.g. by motion (Fig. 2).

Discussion/Conclusion Interleaved acquisition of STEAM spectra of ^1H and ^{31}P at 3 Tesla was demonstrated to be feasible during plantar flexion exercise. Interleaved acquisition ^1H spectra are equivalent to standard acquisition, and there is an SNR benefit in ^{31}P spectra. Future developments will focus on implementation of multiple T_E s for protons during one ^{31}P MRS acquisition, lactate editing and adiabatic water suppression.

References

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