Dynamic susceptibility induced asymmetry of MR proton resonance frequency (PRF) thermometry maps during simultaneous RF ablation causing large and spatially dependent temperature errors

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Introduction
MR thermometry based on the proton resonance frequency (PRF) method (1) has gained good acceptance for guiding RF ablation of liver tumors (2). Thermal dose model (TD) is a reliable predictor of cell death in thermal ablation technique (3). To derive a correct TD thus a correct cut-off point for the application of energy, temperature evolution has to be accurately measured throughout the treatment. Previous works focused on proving the accuracy of MR thermometry with the PRF method: ex-vivo using optical temperature probes to provide reference temperature measurements during simultaneous RF ablation or in animals (4) after in-vivo procedure by comparison of the effectively coagulated lesion size defined by histopathology and the measured TD size (5). None of these experiments reported spatially related errors in temperature maps and TD during power application, while using single slice 2D experimental imaging setup. We described here dynamic susceptibility induced 3D asymmetry effect causing errors in the TD estimates.

Materials and Methods
A bipolar internally-cooled electrode was inserted in a homogeneous sample of fresh pork muscle. The experiment was repeated twice: once with the RF electrode positioned along the Bo field to minimize the susceptibility artifact of the needle, and second orthogonally to Bo to maximize the susceptibility effect. RF power (15W, duration of heating 166 s, energy of 2.5 kJ) was applied simultaneously to the MR imaging with an RF generator (Colon AG, Teltow, Germany) working at 475 kHz. The generator was placed outside the Faraday cage, and the transmission line traversed a wave guide of the cage. Three low pass LC stopband filters (fc = 63.5 MHz) and 6 ferrite cores were added to the transmission line to provide an 80dB attenuation of harmonic at the resonant frequency. A low power setting was chosen to minimize disrupting effects such as vaporization along the muscle fibers (4KJ over 160sec i.e. 25W/35W average/peak power). The sample was imaged in a 1.5T MR system (Espree, Siemens AG, Germany). 3 orthogonal slices were chosen: a transverse plane orthogonal to the RF electrode and cutting it midway between its anode and cathode active zones, and both a sagittal and coronal plane parallel to the electrode. All were determined from a high resolution T1 3D gradient-echo (Vibe) acquisition. A dynamic series of 60 images during the power application (21-25, 36-40 and 51-60 pausing power application). Main Imaging parameters were: slices = 3, dynamics = 60, acquisition time/dynamic = 2.55s (total imaging time was 178.5 s) repeated every 5 sec (simulated respiratory cycle), voxel = 1.6 x 1.6 x 6.0 mm, matrix =128 x 128, TR/TE/FA = 200/40 and 51-60 pausing power application. Temperature levels increase from blue (37°C) to white (65°C and above) for a qualitative appreciation of the shape of the heating pattern. During heating, the expected isothermal surface is a 3D ellipsoid with the long axis coinciding with the axis of the RF electrode (Top left images). Bottom images show that the temperature pattern change from the symmetric expected 3D ellipsoid pattern to a pattern varying depending on the orientation between the electrode and Bo, suggesting susceptibility change over time and/or T°.

Discussion and Conclusion
The direction of the effect (apparent T° increase/decrease) is dependent on the slice orientation and electrode orientation and can cause either overestimation (worst case) or underestimation of the thermal dose. This artifact in the temperature maps is likely to be caused by gas bubbles forming during heating, which can be approximated as a spherical object with a different susceptibility compared to tissue and causing a recognizable dipole magnetic field disturbance. Control experiment with passive electrode in water bath at 20°C and 90°C confirms that the observed effects cannot be attributed to odd needle susceptibility changes with temperature. For future in-vivo studies, the MR method will require correction of the thermal dose estimate for this effect, since such inaccuracy will prevent using the actual TD for treatment cut-off. Further theoretical analysis and simultaneous experimental comparison with optical temperature probes are needed to solve this problem in order to reliably base clinical procedure end point on TD estimates.

References: