

RF coils for high resolution imaging of the human visual cortex at ultra-high fields

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Propositions of the thesis

RF coils for high resolution imaging of the human visual cortex at ultra-high fields

Shubharthi Sengupta, Maastricht, November 28th, 2018

- 1. Anatomical, functional and quantitative MR imaging methods at ultra-high fields have become essential tools for studying and understanding the functional and structural organization in the human cortex.
- 2. A densely populated, small diameter, conformal phased array coil will show increased SNR gains (over a volume coil), through a combination of a tight-fitting former and small coil loops.
- 3. Electromagnetic simulations are indispensable for RF coil development.
- 4. Cable traps are an underrated component in the RF chain.
- 5. Ultra-high resolution (<60μm) images can be acquired on a wide-bore, 9.4T system by using parallel-transmit (pTx) methods in conjunction with optimized receiver arrays on large, ex-vivo samples that would otherwise not fit in a pre-clinical, animal scanner.
- 6. A combination of homogeneous transmit excitations and superior receive signal SNR can help acquire $100\mu m$ isotropic GRE T_2^* weighted data over the entire post mortem human brain sample, allowing clear definition and contrast of anatomical structures in the deep brain nuclei and especially in the cortical grey matter.
- 7. The ability to rotate the sample container on top of a coil surface and tilting the coil surface, thereby providing 2 full degrees of rotational freedom, enables advanced phase contrast and quantitative susceptibility imaging while also allowing diffusion imaging with up to $\sqrt{3}$ more diffusion gradient amplitudes and hence up to 3 times higher b-values.
- 8. Dipole elements will generally outperform traditional loops when imaging deeplying anatomical structures.
- 9. The electric things have their life too. Paltry as those lives are. *Rick Deckard, Do Androids Dream of Electric Sheep?*
- 10. The future is already here it's just not very evenly distributed. William F. Gibson