

Double tunable TxRx $^1\text{H}/^{19}\text{F}$ Helmholtz pair for MR imaging and spectroscopy at 11.7T

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Introduction.

For non-invasive imaging of targeted tissues, cells and biological tissue substitutes in living species the application of fluorinated agents are becoming more popular as ^{19}F is a relative sensitive nucleus and images can be obtained without disturbing background signals as seen in ^1H MRI. However, compared to ^1H nuclei the number of available ^{19}F nuclei is usually limited and thus optimal signal to noise detection is required. Apart from designing contrast agents with a large number of ^{19}F nuclei, this can be aimed for by using the highest possible magnetic field, optimal RF coils and the shortest possible echo time and repetition time in acquisition. Furthermore, for anatomical matching a background ^1H MR image is needed. The aim of this study was to realize these MR conditions to enable ^1H and ^{19}F MRI of rat leg bone and bone substitutes containing ^{19}F contrast, by: 1. designing and building a Helmholtz pair RF probe that is capable of generating a homogeneous RF-field for ^1H and ^{19}F at 11.7T magnetic field 2. performing zero echo time (ZTE) ^1H and ^{19}F imaging to capture short T2 species.

Materials and Methods

MR was performed on a 11.7T MR-system (Bruker Biospin, Germany). The RF-coil consists of a Helmholtz pair with two separate, slightly bended, elements with a size of 25 x 35mm. The average distance between the elements is 25mm. Both elements are separately tuned and matched (balanced) and are combined with a home-built lumped element Wilkinson power splitter/divider [1]. Since the elements couple strongly a split in the resonant curve is observed. The elements must be tuned to the lowest frequency [2]. For best performance both nuclei have a separate power/divider. The circuit diagram of the complete coil can be found in figure 1. The circuit in the red box with the solid line is the RF-coil with the 2 similar elements. In the red box with the dashed line the circuit for the power splitter/ divider can be found. This circuit splits the Tx signal into two outputs with half the power and the same phase. During receive the splitter/ divider combines the received signals. Note the mirrored cable connections at the elements to achieve similar phases for both elements. Otherwise the transmit and receive RF-field will partly cancel out due to the 180° phase difference between the elements.

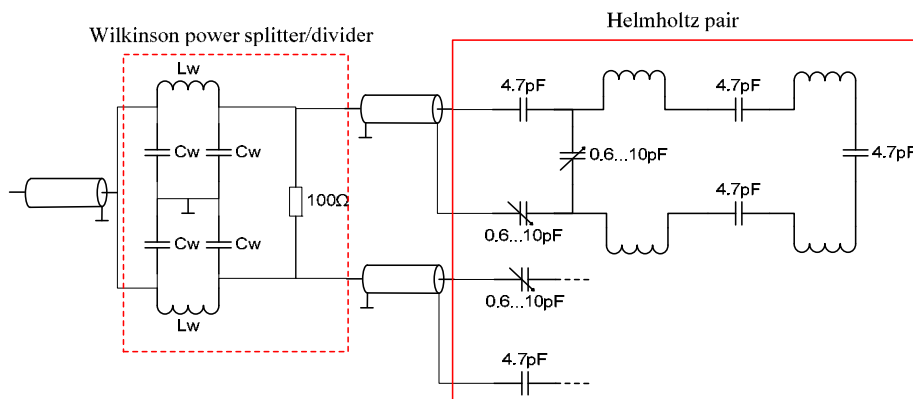


Figure 1: The circuit diagram of the coil and Wilkinson power divider/ splitter. The component values for the divider/ splitter at ^1H and ^{19}F are respectively for 22nH and 23.9nH for L_w and 4.5pF and 4.78pF for C_w .

Zero echo time imaging was done with 200kHz bandwidth, TE/TR=0ms/4 ms, flip angle= $3-6^\circ$, FOV=45*45*45mm.

Results and Discussion

The new RF-coil was first tested on a bone sample that was put in a 15ml phantom containing a salt solution. In ^1H gradient echo imaging with conventional pulses and small flip angles very little signal for the bone is observed, but this image clearly demonstrates the homogeneity of the coil over 3 cm (Fig 2, left image). When ZTE (zero echo time) ^1H MRI of the same sample is performed, the bone species with differential tissue contrast becomes visible (Fig 2, middle image). Finally we tested the ^1H MRI performance of the coil in an in vivo experiment with the hind limb of a rat in the coil. The ZTE ^1H image of an in vivo experiment of a rat leg placed in the new coil also reveals structural details in the bone (Fig 2, right image). Good ^{19}F imaging performance of the probe was also demonstrated by a gradient echo image with conventional pulses and small flip angles, obtained of a 2ml phantom containing a TFA (trifluoroacetic acid) solution (Fig 3, left image). Finally, we obtained a first ZTE ^{19}F image of the same sample (Fig 3, right).

Conclusion

We developed an RF-coil that is capable of generating a homogeneous RF-field large enough to cover a rat hind leg. The coil can be tuned to ^1H and ^{19}F for (spectroscopic) imaging of both nuclei. This setup combined with zero echo time imaging can be used to visualize bone and bone substitutes labeled with ^{19}F compounds. Thus regeneration of bone and degradation of these substitutes present at defects can be monitored by imaging both nuclei with overlay of the images.

Reference: 1) Jon B. Hagen, Radio-frequency Electronics, Circuits and Applications. [2] Mihaela Lupu, NMR Probeheads: For Biophysical and Biomedical Experiments. [3] Weiger et al. Ann Meeting ISMRM-ESMRMB Stockholm, 2010

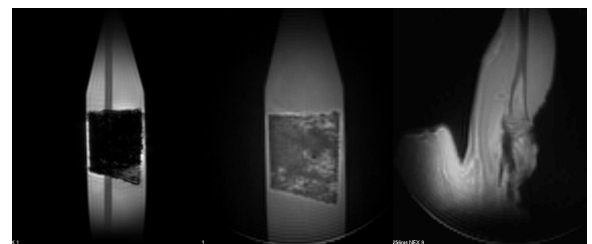


Figure 2: ^1H MR images of bone sample in a aqueous salt solution (left: GRE image, middle: ZTE image) and of in vivo rat hind limb (right: ZTE image).

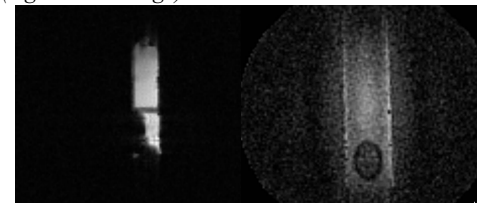


Figure 3: ^{19}F imaging of a 2ml TFA phantom (left: GRE, right: ZTE). The phantom has an air bubble at the bottom.