



An open multi-B0-multi-B1-CEST dataset of the healthy human brain

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## Introduction:

- CEST quantitatively described by Bloch-McConnell (BMC) equations
- First in-vivo multi-B0-multi-B1 dataset of a healthy human brain ready for BMC fitting and full quantification

## **Methods:**

- 1 healthy subject on 2 scanners (MAGNETOM Prisma 3T and Terra.X 7T)
- Conventional Spin-Lock pulse train
  (10 pulses, DC = 50%,

tp = 100 ms, td = 100 ms)

 Benefits: fast simulation similar to CW, without artifacts at decent B0/B1

- B1 levels of 0.3, 0.6, 0.9, 1.5, 2, 2.7, 4 μT
- Image readout: 3D snapshot-CEST GRE [1]
- Offsets for B1 < 4 μT equidistantly between -100 and 100 ppm with -6:0.25:6 ppm,B1 = 4 μT -6:0.5:6 ppm
- Offsets were interpolated for evaluation
- WASABI B1 and B0 mapping [2]
- Z-spectra from GM and WM with decent B0/B1 (B0: ±0.1 ppm, B1: ±5%)
- Within these regions, grey and white matter ROI were defined

## Results:

- Identification of suitable B0/B1 area (figure 2a,b)
- B1 and B0 maps from 7T (figure 2c,d)
- Z-spectra in grey and white matter for all B1 levels at 3T (figure 2e,f) and 7T (figure 2g,h)
- BMC fitting for GM and WM data are currently under investigation (figure 3)

## Outlook

- Publish data with exact definition of preparation and acquisition
- Invite other research groups to do BMC fitting
- Improve models and acquisition for deeper understanding of in vivo CEST



MultiB0\_B1\_qCEST\_brain

Create library for prediction of fitting parameter with Deep Learning





