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Pitfalls and perspectives

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On the reconstruction of magnetic resonance current density images of the human brain: Pitfalls and perspectives



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ABSTRACT

Magnetic resonance current density imaging (MRCDI) of the human brain aims to reconstruct the current density distribution caused by transcranial electric stimulation from MR-based measurements of the current-induced magnetic fields. So far, the MRCDI data acquisition achieves only a low signal-to-noise ratio, does not provide a full volume coverage and lacks data from the scalp and skull regions. In addition, it is only sensitive to the component of the current-induced magnetic field parallel to the scanner field. The reconstruction problem thus involves coping with noisy and incomplete data, which makes it mathematically challenging. Most existing reconstruction methods have been validated using simulation studies and measurements in phantoms with simplified geometries. Only one reconstruction method, the projected current density algorithm, has been applied to human in-vivo data so far, however resulting in blurred current density estimates even when applied to noise-free simulated data.

We analyze the underlying causes for the limited performance of the projected current density algorithm when applied to human brain data. In addition, we compare it with an approach that relies on the optimization of the conductivities of a small number of tissue compartments of anatomically detailed head models reconstructed from structural MR data. Both for simulated ground truth data and human in-vivo MRCDI data, our results indicate that the estimation of current densities benefits more from using a personalized volume conductor model than from applying the projected current density algorithm. In particular, we introduce a hierarchical statistical testing approach as a principled way to test and compare the quality of reconstructed current density images that accounts for the limited signal-to-noise ratio of the human in-vivo MRCDI data and the fact that the ground truth of the current density is unknown for measured data. Our results indicate that the statistical testing approach constitutes a valuable framework for the further development of accurate volume conductor models of the head. Our findings also highlight the importance of tailoring the reconstruction approaches to the quality and specific properties of the available data.

1. Introduction

Knowledge of electrical current density (J) and conductivity (σ) distributions in the human brain is important in many neuroscience applications. It enables the control and optimization of the brain areas targeted by transcranial brain stimulation (TBS) and is needed for localizing neural sources from electro- and magnetoencephalography data. Also, it might be useful for characterizing malignant tissue

(Holdefer et al., 2006; Miranda et al., 2014; Nathan et al., 1993). TBS techniques either inject weak electrical currents via scalp electrodes (transcranial direct or alternating current stimulation; tDCS, tACS) or use electromagnetic induction (transcranial magnetic stimulation, TMS) to modulate neuronal activity (Barker et al., 1985; Nitsche and Paulus, 2000). Stimulation targeting and dose control can be improved using electric field *simulations* in individualized head models constructed from structural magnetic resonance (MR) head scans of a subject (e.g.,

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Received 29 April 2021; Received in revised form 21 July 2021; Accepted 25 August 2021 Available online 1 September 2021. 1053-8119/© 2021 The Authors. Published by Elsevier Inc. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/) Thielscher et al., 2015). Ensuring the accuracy of the simulations is important to prevent false conclusions about the applied stimulation pattern and strength. However, this is a challenging problem that requires measurements of the current density distribution in the human brain to serve as ground truth, ideally using non-invasive measurement approaches that can be broadly applied both in healthy persons and patients.

Magnetic resonance current density imaging (MRCDI) and magnetic resonance electrical impedance tomography (MREIT) are two modalities using transcranial current injections and MR imaging (MRI) to reconstruct current density and conductivity distributions in the brain (Scott et al., 1991). In both methods, the currents are injected in synchrony with an MRI pulse sequence to create a magnetic flux density distribution that changes the magnetic field of the MR scanner. As a result, the phase of the measured MR signal is modulated by the z-component of the current-induced magnetic flux density (B_z), i.e. the part that is parallel to the static scanner field. The MR phase information thus can be used to obtain cross-sectional images of the current-induced magnetic flux density B_z .

Several methods have been developed to reconstruct the current density distribution from the measured B_{τ} images. Considering Ampere's law, a unique and unambiguous reconstruction requires knowledge of all three components of the current-induced magnetic flux density. As only the B_{τ} component parallel to the main magnetic field of the MRI scanner can be measured during an MRCDI experiment, accurate and complete current density mapping requires rotating the measured object inside the scanner (Eyüboğlu, 2006; Scott et al., 1991; Woo and Seo, 2008), which is impractical for human in-vivo brain imaging. Alternative methods thus aim to reconstruct the two components of the electrical current density that are orthogonal to the main scanner field (i.e., J_x and J_y) from a single B_z image (Ider et al., 2010; Jeong et al., 2014; Park et al., 2007). In particular, the "projected current density" method introduced by Jeong et al (2014) was used in recent studies on in-vivo human brain MRCDI (Chauhan et al., 2018; Göksu et al., 2018a; Kasinadhuni et al., 2017). Combined with data from diffusion tensor imaging, projected current densities estimated with this method for two current injection directions (left-right and anterior-posterior) have also been used to reconstruct the ohmic conductivity distribution in the human brain (Chauhan et al., 2018). The accuracy of the reconstructed current density distribution depends on both the quality of the B_{z} images and the properties of the reconstruction methods. Usually, their performances are good for simple phantoms with one or more conductivity inhomogeneities inside a homogenous background composed of a saline solution or an agar gel with a conductivity value in the order of 1 S/m (Kwon et al., 2007; Oh et al., 2003; Park et al., 2007). Also the reconstruction method used so far for in-vivo human brain MRCDI studies was validated in this way (Jeong et al., 2014). However, this situation is strongly simplified compared to MRCDI measurements of the human head, which has a complex anatomy and where the brain is surrounded by the highly resistive skull and conductive scalp.

In an earlier study (Göksu et al., 2018b), we observed that the reconstructed current densities were only coarse estimates of the current flow in the brain even when applied to noise-free simulated data. Here, we explore the underlying causes and reveal fundamental limitations that occur when applying the projected current density method of Jeong et al (2014) to B_z image of the human brain. We then explore whether the method can be modified to achieve better reconstruction performance. We also test an alternative approach to estimate the current density distribution that is based on the optimization of the conductivities of an anatomically detailed head model to fit the measured B_z image. Using both simulated B_z data serving as ground truth and B_z data from human in-vivo measurements, we show that the alternative approach performs better than the projected current density method. In particular, we advocate for systematic statistical evaluations of the goodness of fit between measured and simulated B_z data as a principled way to draw conclusions on the accuracy of simulations. A preprint of this paper was published on biorxiv.

2. Material and methods

2.1. Human brain imaging

Two healthy volunteers, who had no previous neurological and psychiatric disorders, were included in this study. Written informed consent was obtained from the participants prior to the scans and they were screened for contraindications to MRI and transcranial electric stimulation (TES). The study complied with the Helsinki declaration on human experimentation and was approved by the Ethics Committee of the Capital Region of Denmark. We injected electrical current with ±1 mA magnitude in the left-right (LR) and anterior-posterior (AP) directions via surface electrodes attached to the head. The currents were applied in synchrony with a multi-gradient-echo MR pulse sequence (Göksu et al., 2018a; Göksu et al., 2018b) on a 3T MRI scanner equipped with a 64-channel head coil (Magnetom PRISMA, Siemens). We measured B_z distributions for a single slice covering the bottom part of the brain by post processing the acquired MR phase images (Göksu et al., 2018a) and corrected unwanted effects of the magnetic stray field caused by the cable currents on the B_{τ} measurements (Göksu et al., 2019). The B_7 measurements were only available for white matter (WM), gray matter (GM), and cerebrospinal fluid (CSF) and lack the data in the scalp and skull. The B_{z} measurements were denoised with a Gaussian filter with a full width at half maximum (FWHM) value of 3 pixels.

We generated three dimensional (3D) individualized head models of the subjects by utilizing prior structural T1- and T2-weighted measurements (Nielsen et al., 2018) to simulate the current density distribution in the head and the resulting magnetic field distribution caused by the currents. As described in more detail below, we used different head models of increasing anatomical complexity (single compartment, three compartments, four compartments, five compartments, see Fig. 1A) for the simulations.

2.2. Forward simulations

Current densities were calculated using the Finite Element Method (FEM) implemented in SimNIBS 3.1.0 (Thielscher et al., 2015). The initial part of the study relied on the head model *ernie* included in the example dataset. This volume conductor model has a high resolution and is composed of 4.58×10^6 tetrahedral elements with an average volume of 1 mm³. For the later part, individual head models of the subjects with similarly high resolutions were reconstructed from T1-weighted and T2-weighted structural MR images. Detailed explanations on the construction of the head model are given in (Nielsen et al., 2018).

The full models with all tissue types include five compartments (5c). The ohmic conductivities of the 5c models were assigned as $\sigma_{scalp} =$ 0.465, $\sigma_{skull} = 0.01$, $\sigma_{WM} = 0.126$, $\sigma_{GM} = 0.275$, and $\sigma_{CSF} = 1.654$ S/m (Fig. 1A) (Thielscher et al., 2011). In addition, we simulated anatomically simplified models by combining tissue types. A single compartment (1c, i.e. homogenous) model was used by setting all conductivities to $\sigma_0 = 1$ S/m. A model with three compartments (3c) was created by assigning the same conductivity $\sigma_{inside} = 0.34$ S/m to CSF, GM and WM volumes in the 3D head model ($\sigma_{scalp} = 0.465$ S/m, $\sigma_{skull} = 0.01$ S/m). The conductivity σ_{inside} was calculated by taking the average conductivity of the region composed of WM, GM, and CSF volumes, weighted according to their relative volumes, in the slice being imaged. Finally, a model with four compartments (4c) was created by combining GM and WM volumes in the 3D head model with a conductivity of $\sigma_{brain} = 0.18$ S/m that was determined as the average of σ_{WM} and σ_{GM} in the imaged slice ($\sigma_{scalp} = 0.465$, $\sigma_{skull} = 0.01$ S/m, $\sigma_{CSF} = 1.654$). In all models, we injected electrical current with 1 mA magnitude in LR and AP directions to calculate J distributions in the brain using FEM for numerically



Fig. 1. Current density reconstruction for a LR current injection. (A) Volume conductor models of the head with different levels of anatomical detail (1c, 3c, 4c, and full models). The results are shown for an axial slice of these models. (B) The conductivity distribution σ of the full model used as ground truth and the corresponding simulated $|J_{xy}^{true}|$ and B_z^{true} distributions are shown in columns 1-3. The current density distribution $|J_{xy}^{rec}|$ that was reconstructed from B_z^{true} by the projected current density algorithm is shown in column 4. (C) Conductivity, current density and magnetic field distributions for the homogenous (1c) head model that was used in the projected current density algorithm. The current injection scheme is shown in column 4. The electrodes have a rectangular shape with a size of $4 \times 7 \text{ mm}^2$ and they are centered at positions C6 (right, visible in the image) and C5 (mirrored position to C6 on the left side) according to the EEG 10-20 system. (D) Visualization of the terms that are neglected and included, respectively, in the projected current density algorithm (see Eq. 3) for the 1c head model.

solving the Laplace equation for the electric potential (Saturnino et al., 2019a). The current-induced **B** distributions were determined from the current density by applying the Biot–Savart law using the Fast Fourier Transform (Yazdanian et al., 2020). In particular, the z-component of the current-induced magnetic flux density B_z is given by

$$B_{z}(p) = \frac{\mu_{o}}{4\pi} \int_{V} \frac{J_{x}(y-y') - J_{y}(x-x')}{|p-p'|^{3}} dp',$$
(1)

where p = [x, y, z] is the position at which B_z is evaluated, V denotes the head model and p' = [x', y', z'] are positions inside V. $J = [J_x, J_y, J_z]$ is

the current density in the head model and constant μ_o is the magnetic permeability of free space. For all models, we first calculated the J and B_z distributions in the 3D head volumes, then we coregistrated them to the imaged slice.

In addition, we aimed to test how much the projected current density J^{rec} , reconstructed in the imaging slice using the projected current density algorithm (Eq. (2) below), changes the simulated current-induced B_z distributions. For that, we substituted the calculated J distributions by J^{rec} at the positions corresponding to CSF and brain in the imaging slice before determining the current-induced B distribution by using (1).

2.3. Projected current density algorithm

The projected current density algorithm introduced by Jeong et al. (2014) reconstructs a current density distribution J^{rec} from a B_r image

$$\boldsymbol{J}^{rec} = \boldsymbol{J}^o + \frac{1}{\mu_o} \left[\frac{\partial (\boldsymbol{B}_z - \boldsymbol{B}_z^o)}{\partial y}, \frac{-\partial (\boldsymbol{B}_z - \boldsymbol{B}_z^o)}{\partial x}, 0 \right],$$
(2)

using simulated current densities $J^o = [J_x^o, J_y^o, J_z^o]$ and magnetic flux densities B_z^o that are based on a volume conductor model of the subject. In the original form of the algorithm, J^o and B_z^o are obtained for a model with a homogeneous conductivity σ_o (1c) and the outer shape of the imaged object (Fig. 1B). The reconstructed current density distribution J^{rec} is then composed of a smooth and curl-free component J^o that is superimposed onto a high contrast component based on directional derivatives of $B_z - B_z^o$. The latter scales with the local difference between measured and simulated magnetic flux densities.

The algorithm can be easily derived from Ampère's law. The unknown true current density J that gives rise to the magnetic field B can be written as:

$$J = J^{o} + (J - J^{o}) = J^{o} + \frac{1}{\mu_{o}} (\nabla \times B - \nabla \times B^{o})$$
$$= J^{o} + \frac{1}{\mu_{o}} \left[\frac{\frac{\partial (B_{z} - B_{z}^{o})}{\partial y_{z}} - \frac{\partial (B_{y} - B_{y}^{o})}{\partial z}}{\frac{\partial (B_{z} - B_{z}^{o})}{\partial x}} - \frac{\partial (B_{z} - B_{z}^{o})}{\partial x}}{\frac{\partial (B_{y} - B_{y}^{o})}{\partial x}} - \frac{\partial (B_{x} - B_{x}^{o})}{\partial y}}{\partial y} \right].$$
(3)

As we only measure B_z , our best possible assumption about the spatial derivatives of the two unknown components of **B** are $\partial B_{x/\partial z} \approx \partial B_{x/\partial z}^o \approx \partial B_{y/\partial z}^o \approx \partial B_{y/\partial z}^o$. In this case, Eq. (3) simplifies to Eq. (2). For example, this assumption will approximately hold in phantoms that is uniform along the z-direction and in which the z-component of the injected current can be kept small.

The algorithm is straightforward to implement and can be applied to single cross-sectional simulated B₇ images in case of limited volume coverage. In contrast to the original projected current density method that employs a second-order Laplacian operator (Park et al., 2007), the modified variant proposed by Jeong et al. (2014) relies only on first-order spatial derivatives of the B_z image data. This reduces the amplification of high-frequency noise by the derivative operations and potentially improves the quality of the reconstructed current density images from B_{z} data with lower signal-to-noise ratios (SNR). In addition, the method introduced by Jeong et al. (2014) only requires local derivative operations, making it applicable to data from objects that contain large regions with low or no signal such as the human head with the skull and air cavities. On the downside, an error analysis by Jeong et al. (2014) suggests that the simplified method will likely suffer from higher reconstruction errors for high SNR B_z images compared to the original approach suggested in (Park et al., 2007).

2.4. Error metrics

We evaluated the difference between a current density J_{xy} and a reference current density J_{xy}^{ref} using the relative root mean square (RMS)

differences:

$$\delta_{J_{xy}} = \sqrt{\frac{\sum_{i=1}^{N} \left[\left(J_x(i) - J_x^{ref}(i) \right)^2 + \left(J_y(i) - J_y^{ref}(i) \right)^2 \right]}{\sum_{i=1}^{N} \left(J_x^{ref}(i)^2 + J_y^{ref}(i)^2 \right)}} \times 100\%$$
(4)

Here, *i* is the pixel index and *N* is the number of pixels inside the measured region composed of WM, GM and CSF. Please note that we restricted the comparison to the x- and y-components of the current flow, as only those influence the z-component of the magnetic flux density B_z measured by MRI and as only the x- and y-components of the current density are changed by the projected current density algorithm. For completeness, we also evaluated the difference between J_z and the z-component of a reference current density J_z^{ref} in selected cases as

$$\delta_{Jz} = \sqrt{\frac{\sum_{i=1}^{N} \left(J_{z}(i) - J_{z}^{ref}(i)\right)^{2}}{\sum_{i=1}^{N} \left(J_{z}^{ref}(i)\right)^{2}} \times 100\%.}$$
(5)

In addition, we quantified the differences between the z-component of the current induced magnetic flux density B_z and a reference case B_z^{ref} as

$$\delta_{B_{z}} = \sqrt{\frac{\sum_{i=1}^{N} \left(B_{z}(i) - B_{z}^{ref}(i)\right)^{2}}{\sum_{i=1}^{N} \left(B_{z}^{ref}(i)\right)^{2}} \times 100\%.}$$
(6)

Finally, we calculated relative noise floors (nf, the inverse SNR) for the experimentally measured current-induced B_z images as

$$af = \sqrt{\frac{\sum_{i=1}^{N} \left(B_z^{m@0mA}(i) \right)^2}{\sum_{i=1}^{N} \left(B_z^m(i) \right)^2} \times 100\%,}$$
(7)

where B_z^m and $B_z^{m@0mA}$ are the measured B_z fields with and without current injection, respectively.

2.5. Optimization of conductivities

As alternative approach for estimating current densities from magnetic flux densities, we tested optimizing the tissue-specific conductivities $\sigma = [\sigma_{WM}, \sigma_{GM}, \sigma_{CSF}, \sigma_{scalp}, \sigma_{skull}]$ with the goal to minimize the error δ_{B_z} between the simulated magnetic flux density B_z given by the head models and the reference B_z^{ref} . We formulated a constrained optimization problem as

$$\begin{aligned} \text{Minimize } \delta_{B_z} &= \sqrt{\frac{\sum_{i=1}^{N} \left(B_z(\sigma, i) - B_z^{ref}(i)\right)^2}{\sum_{i=1}^{N} \left(B_z^{ref}(i)\right)^2} \times 100\%} \\ &\text{subject to} \\ \sigma_{low} &< \sigma < \sigma_{high} \end{aligned}$$
(8)

where vectors σ_{low} and σ_{high} contain the lower and higher bounds for the to-be-optimized tissue conductivities. The choices of these bounds are detailed below. We solved (8) for σ using the sequential least squares programming (SLSQP) algorithm (Kraft, 1988), implemented in the *optimize* module of *SciPy v1.7.0* (Virtanen et al., 2020). SLSQP is an iterative algorithm for solving non-linear constrained optimization problems. At each iteration of the algorithm, a quadratic approximation of the constraints. The approximated sub-problem is solved for an updated search direction and the optimal step-size is found by substituting the updated search direction are updated by using the optimized step-size and search directions.

Initially, we used the simulated magnetic flux density of the full model with all tissue types as reference to confirm the stability of the optimization approach. Subsequently, measured B_{τ} images from the human imaging data served as reference. We implemented constraints on the conducitivities to reduce the likelihood of overfitting that could occur, e.g. in case of systematic differences of the volume conductor models to the true head anatomy. For the 3c model, we constrained σ_{scalp} , σ_{skull} and σ_{inside} to the ranges [0.2 < σ_{scalp} < 1, 0.003 < σ_{skull} < 0.03, 0.2 < σ_{inside} < 0.95] S/m. For the 4c model, we used the ranges $[0.2 < \sigma_{scalp} < 1, \ 0.003 < \sigma_{skull} < 0.03, \ 1.2 < \sigma_{CSF} < 1.9, \ 0.05 < \sigma_{brain} < 0.05 < \sigma_{brain} < 0.05 < \sigma_{brain} < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0.05 < 0$ 0.95] S/m. We also fitted the full model with five conductivities (5c) to human imaging data by using the ranges [0.2 < σ_{scalp} < 1, 0.003 < $\sigma_{skull} < 0.03, \ 1.2 < \sigma_{CSF} < 1.9, \ 0.05 < \sigma_{WM} < 0.95, \ 0.05 < \sigma_{GM} < 0.95]$ S/m. For the 5c model, we imposed $\sigma_{GM} > \sigma_{WM}$ as an additional constraint. For the 5c model, the ranges were chosen to well include the conductivities reported in studies that measured relatively fresh or live tissue, preferably human at low frequencies (0-100 kHz) near body temperature (Table 1 in Saturnino et al., 2019b). The ranges of the combined tissue regions of the 3c and 4c models were chosen correspondingly using volume-weighted averages of the ranges for the tissues of the 5c model. We initialized the optimization algorithm with $[\sigma_{scalp} = 0.465,$ $\sigma_{skull} = 0.01$, $\sigma_{inside} = 0.34$] S/m for the 3c model and $[\sigma_{scalp} = 0.465$, $\sigma_{skull} = 0.01, \ \sigma_{brain} = 0.18$] S/m for the 4c model and $[\sigma_{scalp} = 0.465,$ $\sigma_{skull} = 0.01, \ \sigma_{CSF} = 1.654, \ \sigma_{WM} = 0.126, \ \sigma_{GM} = 0.275$] S/m for the 5c model.

It is worth noting that without additional knowledge of the total resistance between the two stimulation electrodes, the tissue conductivities are only determined up to a common scaling factor. That is, for a head model with N tissue compartments, only N-1 conductivity *ratios* are uniquely determined via the optimization algorithm. We could have directly optimized N-1 conductivity ratios instead. However, this would have required the choice of an arbitrary "reference" tissue and would have made the use of biologically plausible conductivity ranges as constraints in the optimization difficult.

Applying the optimization algorithm requires the repeated evaluation of B_z for varying conductivities of the head models. This is computationally expensive when it is based on FEM simulations for obtaining the current densitiy J and subsequent solving of the Biot-Savart law using the Fast-Fourier Transform. Therefore, we employed a non-intrusive generalized polynomial chaos (gPC) expansion for the fast evaluation of B_{τ} as a function of the conductivities (scalp, skull, CSF, GM, WM) (Codecasa et al., 2015; Ghanem et al., 2017; Saturnino et al., 2019b; Wiener, 1938). For the gPC expansions, we used a Jacobian polynomial basis set with a maximum order of 10 and sampled the conductivities by using uniform probability density functions. We selected the conductivity ranges of the gPC expansions slightly larger than the corresponding bounds of the optimization algorithm in order to ensure sufficient accuracy of the B_{π} fields across the tested conductivity range. We used Kfold cross validation with K = 10 for the evaluation of the relative error between the B_z fields obtained from the forward simulations and gPC expansions, and set the relative error of the gPC method to 1%. Training of the gPC converged after ~250 FEM simulations. Using the gPC model, we were then able to reevaluate the B_z fields in the complete head model as a function of the conductivity parameters with a computation time in the order of a few seconds. The relative error of 1% that was ensured by the training procedure using cross validation corresponds well to the actual error when applying the gPC later on to reevaluate B_z (Fig. S11 exemplarily compares the results obtained with the gPC expansion versus the forward simulations using FEM and Biot Savart Law for a few selected conductivity combinations). For further details on our gPC implementation, please refer to (Saturnino et al., 2019b).

2.6. Statistical testing

Taking the measured current-induced B_z images from the human imaging data as reference, we assessed the differences δ_{B_z} (Eq. 6) between the simulated and measured B_z data as an indication of the quality of the simulations. First, we evaluated the quality of models with

fixed pre-determined conductivities. For that, we compared two simulations obtained from a simple and a more complex model (e.g., 1c and 3c) by calculating $\Delta \delta_{B_z} = \delta_{B_z^1} - \delta_{B_z^2}$, where $\delta_{B_z^1}$ denotes the error of the simpler model and $\delta_{B_2^2}$ denotes the error of the more complex model. Our hypothesis was that the simpler model has a higher error compared to the more complex model, i.e., that the difference $\Delta \delta_{B_z}$ is positive. To test this hypothesis formally, we first generated a null distribution H_{a} of error differences that could occur by random chance (i.e., if there was no systematic difference between the errors of simpler and more complex models) and then compared our observed error difference to that distribution. This null distribution H_{a} was generated by taking the two simulated B_r images, one from the simpler and one from the more complex model, switching voxels in between the two randomly, and calculating the error difference $\Delta \delta_{B_z}$. The random switching of voxels establishes the difference between the errors we would observe if we assumed that the simpler and more complex models can reconstruct the data equally well. We repeated the random switching of voxels between the two simulated B_z images 10,000 times which generates 10,000 samples of the error difference $\Delta \delta_{B_z}$. As we hypothesized that the error difference should be positive, we then calculated a one-sided p-value (right tail) by computing the fraction of the random samples where the difference was greater than the observed true difference between the errors. Significance was then assessed at p = 0.01, Bonferroni corrected for multiple comparisons.

Furthermore, in order to also systematically compare head models with optimized conductivities, we adapted the statistical approach slightly. Fitting the conductivities requires the measurement data so that the error defined in Eq. (8) can be minimized. To perform unbiased testing, we split the data randomly to training (80%) and testing (20%) parts. We again repeated this random splitting 10,000 times, and for each split first optimized the conductivities on 80% of the data and then calculated $\delta_{B_{\perp}}$ on the remaining 20%. This gave us 10,000 $\delta_{B_{\perp}}$ values for the simpler and more complex models, from which we calculated the error differences $\Delta \delta_{B_z} = \delta_{B_z^1} - \delta_{B_z^2}$. Here we again hypothesized that the simpler model would have a higher error compared to more complex one, as the more complex model has more degrees-of-freedom to fit the data. Thus, similar to the strategy outlined in the previous paragraph, we calculated a one-sided (left tail) p-value by assessing the proportion of the samples where the difference was smaller than zero and tested for significance at p = 0.01, Bonferroni corrected for multiple comparisons. We used the same procedure to compare the conductivity-optimized 3c head model with the 1c head model, however keeping the conductivity of the 1c model fixed and only repeatedly optimizing the 3c model.

2.7. Data and code availability

The original MR data cannot be made publicly available due to privacy restrictions. The largest part of the methods is publicly available via our project homepage www.simnibs.org. This includes the methods for building the head volume conductors from structural MR images, forward simulations of the current densities and current-induced magnetic fields and non-intrusive generalized polynomial chaos expansions for fast re-evaluation of the current-induced magnetic fields in case of changing conductivities. The scripts for statistical testing were developed to address our immediate research needs, but not as easy-to-use software for dissemination and will therefore not be made publicly available at the current time point.

3. Results

In the following, we start by characterizing the performance of the "standard" projected current density algorithm, using simulated ground truth data as reference. We then assess whether the reconstruction accuracy of the algorithm can be improved by using more realistic volume conductor models, and test how sensitive the reconstructed current density is to errors of the head models when the latter become more

detailed. We continue by evaluating the performance of an alternative approach to estimate the current density that is based on optimizing the conductivities of detailed volume conductor models to maximize the fit of the simulated and the reference B_z data. Complementing our initial assessments based on simulated data, we test the two approaches for B_z data obtained from MR imaging of the human brain. In particular, we introduce a hierarchical statistical testing approach as a principled way to test and compare the quality of reconstructed current density images that accounts for the limited signal-to-noise ratio of measured B_z data and the fact that the ground truth of the current density is unknown in reality.

3.1. Projected current density algorithm based on a homogenous (1c) head model

Fig. 1B shows the conductivity distribution σ in an axial slice through a realistic head model (first column) and the simulated current density distribution $|J_{xy}^{true}|$ for a left-to-right (LR) current injection (second column). The z-component B_z^{true} of the magnetic field arising from the injected current flow is shown in the third column. In practice, MRI measures B_z only inside CSF and the brain, which is considered here by applying a corresponding mask. Applying the standard projected current density algorithm to the B_z distribution results in the reconstructed current distribution $|J_{xy}^{rec}|$ shown in the fourth column of Fig. 1B. It is obvious that the reconstructed current density differs strongly from the true simulated current density distribution inside the cranial cavity both in terms of its spatial pattern and average strength (R²=0.22 between $|J_{xy}^{true}|$ and $|J_{xy}^{rec}|$; average strengths: $|J_{xy}^{true}|$ 0.039 A/m², $|J_{xy}^{rec}|$ 0.058 A/m²). In particular, $|J_{xy}^{rec}|$ fails to capture the spatial details of the current flow, even those that originate from large anatomical structures such as the ventricles. In contrast, the spatial pattern of $|J_{\gamma\gamma}^{rec}|$ is more similar to the current flow $|J_{xy}^o|$ inside a homogeneous conductor (Fig. 1C, second column; R²=0.68 between $|J_{xy}^{o}|$ and $|J_{xy}^{rec}|$, average strength of $|J_{xy}^{o}|$: 0.068 A/m²). This shows that the results of the projected current density algorithm are dominated by the simulated current density J^{o} (Eq. 2), and are less influenced by the actually measured B_z data. The reason for this low performance gets apparent when visualizing the terms that are neglected in the projected current density algorithm (see Eq. 3) as they involve the non-measurable B_x and B_y components of the magnetic field (first and second column of Fig. 1D). For the simulated head anatomy, the neglected terms are as strong and even stronger than the included terms that depend on the measured B_z component (third and fourth column of Fig. 1D), which prevents a reasonable reconstruction of the true current density distribution.

The projected current density algorithm shows similar performance for an anterior-to-posterior (AP) current direction (rows one and two of Fig. S1A). Very similar results were also obtained when using a head model as ground truth that incorporated anisotropic brain conductivity, derived from diffusion MR data (Fig. S2A-C). These control results confirm the limited applicability of the algorithm for realistic head anatomies. In contrast, the algorithm performs better for a volume conductor that is uniform along the z-axis, as the neglected terms are close to zero in this case (row three in Fig. S3). For this simplified "head" model, $|J_{xy}^{rec}|$ captures the spatial details of $|J_{xy}^{rue}|$, especially throughout the ventricles (R^2 =0.63 between $|J_{xy}^{true}|$ and $|J_{xy}^{rec}|$; average strengths: $|J_{xy}^{true}|$ 0.035 A/m², $|J_{xy}^{rec}|$ 0.032 A/m²). The remaining differences show mostly up as blurring, caused by a limited numerical accuracy of the gradient calculations in Eq. (2). This effect has also been observed in previous studies using simplified phantoms (Sajib et al., 2012).

3.2. Projected current density algorithm based on inhomogeneous head models

Using more realistic volume conductor models of the head can help to make the simulated J^o more similar to the true current density. By that, the strength of the terms that are neglected in the projected current density algorithm will also be reduced. In the following, we tested whether this helps to improve the performance of the algorithm. For that, the reconstructed current densities were compared when using three different head models to calculate J^o : 1) A model with a single compartment (1c) as used in the original algorithm. 2) A model with three compartments corresponding to scalp, skull and the cranial cavity (3c in Fig. 2B). 3) A model with four compartments corresponding to scalp, skull, CSF and the brain (4c in Fig. 2B). Please see the Methods section for the choice of the conductivities for the 3c and 4c models.

Expectedly, the reconstructed current density (4th column in Fig. 2B) approaches the true current density distribution (Fig. 2A) when the head models get closer to the full model used as reference. For example, the current density in the brain is clearly overestimated close to the electrodes when using the 1c model that does not account for the "shielding" effects of the low-conductive skull. This effect is mostly corrected by using the 3c model, but only the 4c model also achieves a reasonable estimation of the current density around the ventricles and in the sulci. Evaluation of the RMS errors $\delta_{J_{XY}}$ (Eq. 4) for the 1c, 3c and 4c models (Fig. 3A) reveals that most of the improvement resulted from the better models used to calculate J^o , while the projected current density algorithm causes only little additional improvement. The underlying reason is that the neglected terms are still very strong even for the more detailed models (Fig. 2C shows all terms for the 3c model as example). For the 3c model, the algorithm reconstructs some of the details of the current flow in the brain that are missing in the underlying J_{xy}^{o} distribution (second row of Fig. 2B). However, this results only in a small overall improvement in $\delta_{J_{XY}}$ (Fig. 3A). A visual inspection of the spatial distribution of the reconstruction error revealed that the projected current density algorithm decreased the error in many parts of the brain, but also resulted in localized increases at several positions, in particular around the sulci (data not shown). For confirmation, an anteriorto-posterior current direction was also tested (Figs. S1 and S4), giving similar results.

While using more realistic head models seems straightforward to reduce the reconstruction error, it is important to note that these models still include errors, e.g. due to inaccurate segmentations of the structural MR images or incorrectly chosen ohmic tissue conductivities which deviate from their unknown ground truth. That is, more detailed head models will result in current density distributions appearing realistic with more spatial high frequency content, but which are not necessarily closer to the real current density distribution. In the following, it is therefore tested how robustly improvements can be obtained when the head model for the J^o simulations differs from the real volume conductor and whether the projected current density algorithm is helpful to correct errors introduced by deviations of the head models from the ground truth.

Fig. 3A shows the RMS errors δ_{Jxy} for selected variations of the anatomy of the 4c model, namely overall thinner and thicker skulls (affecting the amount of current entering the cranial cavity) and opened and closed sulci (affecting the amount of well-conductive CSF in the sulci and by that the local current flow distribution). While varying the skull thickness changed the current flow only little, modifying the sulci increased the error almost to the level of the 3c model (Fig. S5 shows the corresponding $|J_{xy}^{o}|$ and $|J_{xy}^{rec}|$). Fig. 3B depicts the RMS errors for the homogenous, 3c and 4c models when varying selected conductivities of the full model that was used as ground truth. The scalp conductivity of the full model was set to half and double of the standard value, and the conductivities of GM and WM were halved and doubled. In all cases, the 3c and 4c models maintained a better performance than the 1c model. Both for the tested variations of anatomy and conductivity, the projected current density algorithm achieved no or only moderate improvements. In particular, changing to anatomically more accurate head models resulted in stronger improvements than those that could be obtained by using the algorithm. It is also worth noting that the differences between the results of the 1c, 3c and 4c models were larger



Fig. 2. Current density reconstructions based on different head models for a LR current injection. (A) The conductivity distribution σ of the full model used as ground truth and the corresponding simulated $|J_{xy}^{rue}|$ and B_z^{true} distributions. (B) Results for the 1c, 3c and 4c head models. Columns 1-3 show the conductivity distributions and the simulated $|J_{xy}^{e}|$ and B_z^{e} distributions. Column 4 depicts the current densities $|J_{xy}^{rec}|$ that were reconstructed by the projected current density algorithm with the corresponding head model. (C) Visualization of the terms that are neglected and included, respectively, in the projected current density algorithm based on the 3c head model.

than the effects caused by modeling brain conductivity as anisotropic vs isotropic (Fig. S2D). Thus, isotropic conductivities were used for all tissue compartments.

3.3. Optimization of the tissue conductivities of the inhomogeneous head models

The above results indicate that the fit between the estimated and true current densities can be improved more by choosing an appropriate volume conductor model than by applying the projected current density algorithm. A straightforward approach to estimate the current density distribution is thus to use the measured B_z data to optimize the tissue conductivities of anatomically detailed volume conductor models that are determined via the segmentation of structural MR data. The following section provides a basic proof-of-concept that this optimization approach can successfully account for conductivity changes of the full model used as ground truth. The performance for volume conductor models that differ in the amount of anatomical detail is also tested



Fig. 3. RMS errors $\delta_{J_{xy}}$ of different head models for a LR current injection, with and without applying the projected current density algorithm. Solid and dashed lines represent the $\delta_{J_{xy}}$ values for the J_{xy}^{rec} distributions, respectively. (A) Dependence of $\delta_{J_{xy}}$ on the anatomical level of detail of the head model (1c, 3c and 4c models) and on segmentation errors of the 4c model. (B) Dependence of $\delta_{J_{xy}}$ on variations of the conductivity of the full model that was used as ground truth.

and it is assessed how the presence of inaccurate segmentations impacts reconstruction performance.

It can be exemplarily seen in Fig. 2A and B that the similarity of the estimated and true current density distributions J_{xy}^o and J_{xy}^{true} co-vary with the similarity between B_z^o and B_z^{true} when changing between the 1c, 3c and 4c models. This is also revealed in the corresponding co-variation between the RMS errors $\delta_{J_{XY}}$ and $\delta_{B_{\tau}}$ in Fig. 4A and C (solid lines – 1c, 3c and 4c). In addition, when varying the conductivities of the full model that serves as ground truth (solid lines in Fig. 4B and D), both errors show similar dependences on the conductivity and the model (R²=0.86 between $\delta_{J_{XY}}$ and δ_{B_z} for the non-optimized cases). That is, the errors in J_{xy} relative to its ground truth are well reflected in the errors in B_z . This favorable behavior of B_z occurs as, by the Biot-Savart law, B_z is fully determined by the three-dimensional distribution of J_{XY} , with the impact of the current density scaling with distance to the measured B_{z} (Eq. 1). It is interesting to note that, by Ampère's law, J_{xy} is determined by the spatial changes of all components of the magnetic field rather than only B_z (Eq. 3). This explains why the projected current density algorithm can have low performance despite a close relation between J_{xy} and B_z .

The demonstrated relation between $\delta_{J_{xy}}$ and δ_{B_z} suggests that minimizing the RMS errors for B_z by optimizing the conductivities of the head models might be a useful approach to improve the similarity between the simulated J_{xy} and the ground truth. Indeed, optimizing the tissue conductivities of the 3c and 4c models to minimize δ_{B_z} consistently improves the fit for J_{xy} (dashed lines in Fig. 4B and D, R²=0.86 between $\delta_{J_{xy}}$ and δ_{B_z} for the optimized cases). Expectedly, the achievable improvement scales with the level of detail of the head model. Rather trivially, the errors approach zero when the full head model is used for the optimizations (data not shown). In general, optimization reduces δ_{B_z} more than $\delta_{J_{xy}}$ when compared to the RMS errors of the non-optimized cases. Still, also after optimization, the remaining δ_{B_z} reflect well the relative differences between the $\delta_{J_{xy}}$ of the different head models, making the former a useful parameter for model comparison.

Optimizing the tissue conductivities in the presence of anatomical inaccuracies hardly changes $\delta_{J_{XY}}$ (dashed line in Fig. 4A) although it reduces δ_{B_z} for all cases (R²=0.61 between $\delta_{J_{XY}}$ and δ_{B_z} for the op-

timized cases). This indicates that constraining the optimization to a few conductivities which affect the current flow globally reduces the risk of overfitting. However, it also shows that accurate segmentation of the brain anatomy from the MR images is required to ensure accurate estimations of local details of the current density distribution via this approach. In particular, also for the non-optimized cases, changing the sulcal structure of the 4c model affects J_{xy} , but hardly changes B_z (solid lines in Fig. 4A and C). This indicates that B_z is not a sensitive marker of localized errors in the estimated current density distribution.

For confirmation, an anterior-to-posterior current direction was also tested (Fig. S6), giving similar results. The error for J_z is shown in Fig. 4E and F for completeness. Even though the relationship between J_z and B_z is only indirect, it follows the same general dependence on the head model and conductivities as observed for J_{xy} and B_z . As an alternative to optimizing the model conductivities, Kwon and colleagues (2016) introduced an iterative version of the projected current density algorithm that, combined with an anatomically detailed head model, achieved good reconstruction accuracy for simulated ground truth data. However, our initial tests suggest that stable convergence depends on the availability of B_z data for the complete head volume including scalp and skull (Supplementary Material B).

3.4. Hierarchical model selection based on statistical testing

The results so far indicate that the tested projected current density algorithm has limited capabilities to reconstruct the current density distribution for the complex anatomy of the human head. Instead, a better estimation of the true current density distribution is achievable by using detailed volume conductor models that are reconstructed from structural MR scans and optimizing their conductivities based on the measured B_z data. In contrast to the above tests that used simulated B_z data also as ground truth, measured B_z data contains noise. Therefore, the difference between the simulated and measured B_z fields would not fully disappear even if the head model was perfect. In order to quantify the quality of current density distributions calculated via head models in practice, the difference between the simulated and measured B_z field thus has to be compared to the expected noise level of the B_z mea-



Fig. 4. RMS errors $\delta_{J_{xy}}$, δ_{Bz} and δ_{Jz} of different head models for a LR current injection, with and without optimized conductivities. Solid and dashed lines correspond to the error values obtained for the models with literature and optimized conductivities, respectively. The first column (A, C & E) shows the dependence of $\delta_{J_{xy}}$, δ_{Bz} and δ_{Jz} on the anatomical level of detail of the head model (1c, 3c and 4c models) and on segmentation errors of the 4c model. The second column (B, D & F) shows the dependence of $\delta_{J_{xy}}$, δ_{Bz} and δ_{Jz} on variations of the conductivity of the full model that was used as ground truth.

surements, for example obtained in independent control measurements without current injection. Specifically, when optimizing model parameters such as the conductivity values, this comparison can help to identify cases of overfitting in which the remaining difference can be lower than the expected noise level. This is particularly relevant for models with many free parameters. Finally, as the ground truth with regards to the conductivity distribution calculated via a detailed volume conductor model results in B_z data that is closer to the measured B_z data when compared to the current density distribution obtained for an anatomically simpler model. This ensures that increasing the complexity of the head model indeed helps to improve the approximation of the unknown true current density distribution.

In the following, rigorous statistical testing is employed to assess the quality of current density estimates for measured B_z data for two current injection directions in two participants. Initially focusing on the current density estimates obtained for head models with pre-defined conductivities taken from literature, permutation testing is used to assess the difference between the models (see the Methods section for details). Fig. 5A shows the measured B_z data of two electrode montages (LR and AP) for one participant, and Fig. 5B and 5C depict the simulated data and the remaining difference. The results also confirm that the current density estimated via the 4c model explains the measured B_z significantly better than that of simpler models (Fig. 5D) ($\Delta=77.0$ % and $\Delta=10.7$ % between 4c and 3c models for LR and AP, $p \le 0.01$ corrected). This also holds for the current density estimated via the standard projected current density algorithm ("1c+rec." in Fig. 5), which achieves moderate improvements in δ_{B_z} compared to the 1c head model ($\delta = 25.3 \%$ and $\delta =$ 8.8% for LR and AP, $p \le 0.01$ corrected). As the projected current density algorithm changes the current density only within the brain and CSF area of the imaged slice, the moderate differences in $\delta_{B_{\perp}}$ are partly caused by the need to use J^{o} of the 1c model in the remaining parts of the head volume for calculating B_z . In order to ensure a fair comparison to the 3c model, we use the same approach when determining B_z of the 3c model (indicated by the squares in Fig. 5D), still revealing statistically significant differences of 11.8 % and 3.1 % for the LR and AP injections, respectively. The results also show that the current density estimated using the 5c model explains the measured B_z significantly better than the 4c model for the LR injection ($\delta = 14.6\%$, $p \le 0.01$ corrected), whereas we do not observe a statistically significant difference for the AP injection (Fig. 5D). For the AP montage, the impact of the ventricles on the B_z distribution in the image center are visible in both the measured B_7 data and the results of the 4c and 5c models, but are absent for the simpler models. However, even for the 4c and 5c models, the remaining differences are above the expected noise floor seen for the control measurement for 0 mA. In addition, they exhibit clear spatial patterns, suggesting that the 4c and 5c models do not fully explain all aspects of the underlying current density distribution.

Fig. 7A shows the corresponding current density estimates for the different models. At the SNR of the measured B_z , the projected current density algorithm (1c model) does not recover details of the inhomogeneous current flow pattern that are expected to occur due to the different conductivities between the brain and CSF. Also, the 3c model does not reconstruct any details of the inhomogeneous current flow pattern, but still explains the measured B_z field better because it corrects for the impact of the low-conductive skull on the average strength of the current flow in the cranial cavity. The 4c and 5c models improve the fit to the measured B_z field by accounting for the spatially varying conductivities in the cranial cavity. The results for the second subject are shown in Figs. S7 and S9A, confirming the above observations.

3.5. Hierarchical model selection for models with optimized conductivities

The B_z fields of the 3c, 4c and 5c models with optimized conductivities are shown in Fig. 6B for the first subject, and the remaining differences to the measured fields are shown in Fig. 6C. Compared to the models using literature values for the conductivities, optimization reduced the average strength of the simulated B_z fields. This decrease is a main cause of the largely improved δ_{Bz} of the 3c, 4c and 5c models compared to the 1c case (Fig. 6D vs 5D), and is correspondingly reflected in the on average lower current densities of the optimized 3c, 4c and 5c models in Fig. 7B compared to their counterparts in Fig. 7A. In addition, the optimized B_z fields in Fig. 6B also better reflect the measured spatial variations of the simulated B_z fields close to the electrodes in particular for the LR injections.

The improvement of δ_{Bz} for the 4c versus 3c model is statistically significant for both LR and AP injections. Increasing the level of detail of the model further from 4c to 5c results in statistically significant improvements for the LR injection, but not in case of the AP injection (Fig. 6D). The B_z fields obtained for the 3c, 4c and 5c models result in residuals with similar spatial patterns (Fig. 6C). However, visual inspection of the 4c and 5c models with the 3c model for the AP injection reveals localized variations of the simulated B_7 fields in the middle of the brain due to the CSF-filled ventricles. The peak absolute differences in this region are in the order of 0.3 nT, which is similar to the differences of the simulated B_2 field of the 3c model and the measured B_2 field in that region. These findings indicate the minimal measurement sensitivity that is required to reliably resolve B_{z} variations caused by large brain structures for a current injection of 1 mA baseline-to-peak. Similar effects of the CSF-filled ventricles are also visible for the AP injection for the second subject in Fig. S8B (Fig. S9B shows the corresponding current densities).

While optimization strongly reduces the differences between simulated and measured B_z fields, the remaining differences still show spatial patterns that are above the strength of the noise floor for the 0 mA results. This indicates that also the 4c and 5c models with optimized conductivities do not account for all aspects of the measured B_z field. The optimized conductivities are shown in Fig. S10 for completeness.

4. Discussion

Using simulated and measured data, we demonstrated that the tested variant of the projected current density algorithm (Jeong et al., 2014) achieves only coarse reconstructions of the current density distribution in the human brain, while working as expected for artificial geometries that are uniform along the z-direction. The algorithm neglects the components of Ampère's law (Eq. 3) that depend on the B_x and B_y components of the current-induced magnetic field. These components are weak in geometries that are uniform along the z-direction, but can be dominant in case of the human head. This might also explain why the projected current density algorithm achieved little additional improvements when we tested it in combination with more detailed head models instead of a single compartment model. In contrast, increasing the anatomical detail of the head model alone already had a large effect on the accuracy of the estimated current densities. In our tests with simulated ground truth data, this still held when the head model suffered from uncertainties of the ohmic tissue conductivities or had a limited segmentation accuracy. We confirmed that these conclusions derived from simulated ground truth data also apply to measured data: Current densities calculated with a 3-compartment model and tissue conductivities taken from literature explained measured B_z images significantly better than current densities reconstructed using the standard projected current density algorithm.

Our results demonstrate the importance of formal statistical testing for systematic comparisons of volume conductor models that have different levels of anatomical detail. Statistical testing within a hierarchical model selection framework ensured that more detailed models did indeed provide more accurate estimations of the unknown true current density. Expanding the framework beyond models with fixed conductivities, we also demonstrated that the framework allows for the systematic assessment of current density estimates derived from volume conductor models that use optimized tissue conductivities based on the

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Fig. 5. Experimental B_z measurements and the corresponding simulated B_{z} distributions for the first subject. The conductivity values of the 3c, 4c and 5c models were chosen from literature values, using the procedure described in Methods section. (A) Measured B_z distributions with and without 1 mA current injections in LR and AP directions. (B) Simulated B_r distributions obtained for the 1c, 3c, 4c and 5c models and for the projected current density reconstruction based on the 1c model (1c+rec.). For the latter, B_z was calculated by using the reconstructed current density J^{rec} in the imaged slice and J^{o} of the homogenous head model in the rest of the head volume. (C) Differences between the measured and simulated B_z distributions. (D) Dependence of δ_{Bz} on the model. Asterisks (*) indicate significant differences between the models at $p \le 0.01$, Bonferroni corrected for 8 comparisons. The squares () represent δ_{Bz} values obtained for B_z distributions reconstructed using J^o of the 3c model in the imaged slice and J^o of the homogenous model in the rest of the head volume. This was done for a fair comparison of the 1c+rec. results with the 3c model, as the reconstructed current density algorithm only changes the current distribution in the imaged slice and the change in B_z is correspondingly lower. The dashed green and purple lines represent the noise floors *nf* (Eq. 7). The δ_{Bz} difference between the 4c and 5c models of the AP case is non-significant (ns).

measured B_z images. Complemented by results for simulated ground truth data that showed a stable convergence towards the true current densities with varying tissue conductivities for the employed optimization approach, these findings suggest that using the measured B_z images to fit the tissue conductivities of personalized volume conductor models

might be a valuable approach in order to estimate the unknown true current density.

It is worth noting that the possibility to distinguish between the quality of different volume conductor models using measured B_z data and statistical testing also scales with the measurement SNR. Specifically,



Fig. 6. Results for the 3c, 4c and 5c models with optimized conductivities for the first subject, determined by minimizing the difference to the measured B_z distributions. (A) Measured B_z distributions with and without 1 mA current injections. (B) Optimized B_z distributions for the 3c, 4c and 5c models, corresponding to the median δ_{Bz} shown in D. (C) Difference between the measured and optimized B_z distributions. (D) Dependence of δ_{Bz} on the model. The red lines indicate the median, the boxes mark the 25th and 75th percentiles and the whisker lengths is set to 1.5 times the interquartile range. Values outside that range are marked as outliers. Blue asterisks (*) indicate significant differences between the models at $p \le 0.01$, Bonferroni corrected for 6 comparisons. The dashed green lines indicate the noise floors.



Fig. 7. Estimated current density distributions $|J_{xy}|$ for the first subject. (A) Results for the 1c, 3c, 4c and 5c models based on literature conductivities. Also, the results for the projected current density reconstruction based on the 1c model are shown (1c+rec.). The corresponding B_z fields are shown in Fig. 5. (B) Results for the 3c, 4c and 5c models based on optimized conductivities. The corresponding B_z fields are shown in Fig. 6.

differences between models will occur as non-significant when the corresponding changes of the B_z data are below the noise levels. While a systematic evaluation of the required SNR levels is outside the scope of this study, our results indicate that a sensitivity of around 0.3 nT is required for a current injection of 1 mA in order to reveal the conductivity contrast caused between the ventricles and surrounding brain areas.

Increasing the accuracy of the anatomical model, both in terms of spatial accuracy and better conductivity estimates, will directly impact on transcranial electrical stimulation approaches by providing better individualized targeting and dose control. Measuring the current-induced B_z fields for specific stimulation montages can serve as an important validation step. In particular, it might provide helpful information for identifying the reasons behind inaccurate model predictions in cases where the measurements and simulations do not match.

4.1. Relation to prior studies

Our findings show that the blurriness and lack of detail of current density reconstructions for human in-vivo data do not merely result from a low SNR of the B_z images (Chauhan et al., 2018; Göksu et al., 2018b; Kasinadhuni et al., 2017) and a lack of accounting for the magnetic stray fields cause by the electrode cables in two of the studies (Göksu et al., 2019). Rather, they are also due to a limited applicability of the reconstruction algorithm to the human head anatomy. As the reconstructed current density is biased towards the smooth current flow occurring in a volume conductor with homogenous conductivity (Fig. 1A&B), it is likely that the similarity between the current density distributions reconstructed from measured B_r data of different persons will be artificially increased, camouflaging interindividual differences. This effect also explains why in our prior study the fit between current densities reconstructed from simulated and measured B_{τ} data were consistently better than the fit between the simulated and measured B_{τ} data itself (Göksu et al., 2018a). It is likely that also the performance of methods such as DT-MREIT (Jeong et al., 2017), which uses the reconstructed current densities in combination with diffusion tensor images for estimating the conductivities of brain tissues, will be markedly affected by the low accuracy of the spatial patterns and strengths of the reconstructed current densities, leading to biased conductivity estimates. We only assessed a specific variant of the projected current density method (Jeong et al., 2014) so that our findings will not necessarily generalize to other current density or conductivity reconstruction methods. However, they point towards the need to carefully ensure the desired behavior of reconstruction algorithms when applying them to B_z data of the human head. The lacking uniformity along the z-direction and the lack of data from the scalp and skull regions render the reconstruction problem fundamentally more difficult than for typical MREIT phantoms.

Kwon and colleagues (2016) introduced a method that combines an anatomically detailed head model with an iterative updating procedure to improve the accuracy of the reconstructed current densities by minimizing the difference between calculated and measured B_z data. It showed promising performance on a simulated case in which B_z data were available for the complete head volume. However, our initial tests indicate that it requires further development to be applicable to measured B_z data with limited volume coverage and lack of data from the scalp and skull (Supplementary Material B).

Our approach to optimize conductivities shares similarities with the algorithm proposed in (Gao et al., 2006). Both methods optimize the conductivity values of a few tissue types instead of aiming at a voxelwise conductivity reconstruction, and both methods use polynomial expansions (here: generalized polynomial chaos expansion; Gao et al.: response surface method) for representing the functional dependence between simulated B_z and conductivity in order to avoid the costly reevaluation of the forward model during conductivity optimization. They differ in the level of detail of the head models and the optimized cost functions. While we chose to minimize the relative root mean square differences between measured and simulated B_{zy} Gao and colleagues choose a more complex cost function that additionally incorporates the correlation coefficient. While beyond the scope of this study, the impact of the choice of the cost function on the stability and accuracy of the optimization procedure for low-SNR B_z data might be a relevant topic to develop the overall approach further. The largest difference concerns our use of statistical testing for model selection in order to ensure the quality of the estimated current density distribution despite the lack of a ground truth in case of measured B_z data.

4.2. Limitations and future steps

We used the optimization of model conductivities as a means to estimate the current density distribution, and systematically ensured the quality of the calculated current density distribution by statistical testing. However, this does not imply that the optimized conductivities (or their ratios) are necessarily robust and trustworthy, as conductivity variations of different tissues can have quite similar effects on the current flow patterns and the measured current-induced magnetic fields. For example, increasing scalp conductivity and decreasing skull conductivity both have the same main effect of increasing the amount of current shunted through the scalp and decreasing the current that enters the intracranial cavity. Alternatively, also decreasing the overall conductivity of the intracranial tissues will have a similar impact on the current flow pattern. In other words, various combinations of the tissue conductivities can give similar current flow patterns, rendering a stable estimation of the conductivities in the presence of measurement noise challenging. This limitation does not affect the stability of the estimated current flow distributions. The conductivities are fitted to best explain the magnetic field in the imaged slice that is strongly depending on the current flow in the surrounding region. The simulated current flows in distant regions are thus not necessarily more accurate than simulations based on standard conductivities taken from literature. Similarly, they do not necessarily generalize to other electrode positions, which becomes obvious when comparing the scalp conductivities for the LR and AP injections (Fig. S10). Interestingly, both skull thickness and composition underneath the electrode positions are different for the two injection schemes, which might be reflected in the change of the conductivities.

The main aims of this study were to analyze how well the projected current density algorithm can be applied to in-vivo B_z data from the human intracranial region and to compare it with an approach that relies on the optimization of a low number of parameters (the compartmental tissue conductivities) of anatomically detailed head models. While voxel-wise reconstructions of the current densities or conductivities remain the ultimate aim of MREIT and MRCDI, our findings highlight the importance of tailoring the reconstruction approaches to the quality and specific properties of the measurements. We based our conclusions also on hierarchical statistical testing and in-vivo B_z data, as we feel that this best ensured the validity of our findings. In contrast, for pure simulation studies, it is more challenging to ensure transferability of the results to the envisioned application and to properly account for factors such as deviations of the noise floor from spatially independent and identically distributed Gaussian noise.

Our results suggest that MRCDI data might be very valuable for the further development of personalized volume conductor models of the head. A relevant question to address would be to examine which aspects of the head modeling need to be improved to ensure that the same conductivity values for the modelled tissue compartments are reached independent of electrode montage. Considering that the residuals still contain spatial patterns above the noise floor also for the 4c and 5c head models with optimized conductivities, it seems promising to test the impact of new segmentation methods (Puonti et al., 2020) on data from a larger group of subjects. In addition, as the simulations assume homogenous conductivities of the brain tissues, it might be interesting to explore whether localized changes of the tissue conductivities due to pathologies such as tumors are detectable as outlier regions in the residual images. This effect would be similar to the local increases in the residual images around the ventricles for 3c models and the AP injections seen here. Increased coverage of the imaged region beyond a single slice would be beneficial for these developments. However, this requires a careful tradeoff against maintaining and further increasing the SNR and robustness to physiological noise.

Credit authorship contribution statement

Hasan H. Eroğlu: Conceptualization, Methodology, Software, Validation, Formal analysis, Investigation, Interpretation of results, Writing - Original Draft, Visualization

Oula Puonti: Conceptualization, Methodology, Software, Investigation, Interpretation of results, Writing - Review & Editing

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Declaration of Competing Interest

HRS has received honoraria as speaker from Sanofi Genzyme, Denmark and Novartis, Denmark, as consultant from Sanofi Genzyme, Denmark, Lophora, Denmark, and Lundbeck AS, Denmark, and as editor-inchief (Neuroimage Clinical) and senior editor (NeuroImage) from Elsevier Publishers, Amsterdam, The Netherlands. He has received royalties as book editor from Springer Publishers, Stuttgart, Germany and from Gyldendal Publishers, Copenhagen, Denmark. The other authors report no conflicts of interests.

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Supplementary materials

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