RESEARCH ARTICLE

Magnetic Resonance in Medicine

Rapid estimation of 2D relative B_1^+ -maps from localizers in the human heart at 7T using deep learning

Felix Krueger^{1,2} | Christoph Stefan Aigner¹ | Kerstin Hammernik^{3,4} | Sebastian Dietrich¹ | Max Lutz¹ | Jeanette Schulz-Menger^{5,6,7} | Tobias Schaeffter^{1,2,8} | Sebastian Schmitter^{1,9,10}

Correspondence

Felix Krüger, Physikalisch-Technische Bundesanstalt, Braunschweig and Berlin, Germany.

Email: felix.krueger@ptb.de

Funding information

German Research Foundation, Grant/Award Numbers: SCHM 2677/2-1, SCHM 2677/4-1, GRK2260-BIOQIC **Purpose:** Subject-tailored parallel transmission pulses for ultra-high fields body applications are typically calculated based on subject-specific B_1^+ -maps of all transmit channels, which require lengthy adjustment times. This study investigates the feasibility of using deep learning to estimate complex, channel-wise, relative 2D B_1^+ -maps from a single gradient echo localizer to overcome long calibration times.

Methods: 126 channel-wise, complex, relative 2D B_1^+ -maps of the human heart from 44 subjects were acquired at 7T using a Cartesian, cardiac gradient-echo sequence obtained under breath-hold to create a library for network training and cross-validation. The deep learning predicted maps were qualitatively compared to the ground truth. Phase-only B_1^+ -shimming was subsequently performed on the estimated B_1^+ -maps for a region of interest covering the heart. The proposed network was applied at 7T to 3 unseen test subjects.

Results: The deep learning-based B_1^+ -maps, derived in approximately 0.2 seconds, match the ground truth for the magnitude and phase. The static, phase-only pulse design performs best when maximizing the mean transmission efficiency. In-vivo application of the proposed network to unseen subjects demonstrates the feasibility of this approach: the network yields predicted B_1^+ -maps comparable to the acquired ground truth and anatomical scans reflect the resulting B_1^+ -pattern using the deep learning-based maps.

Conclusion: The feasibility of estimating 2D relative B_1^+ -maps from initial localizer scans of the human heart at 7T using deep learning is successfully demonstrated. Because the technique requires only sub-seconds to derive channel-wise B_1^+ -maps, it offers high potential for advancing clinical body imaging at ultra-high fields.

KEYWORDS

7 Tesla, B₁⁺-mapping, body MRI, deep learning, heart, parallel transmission

Parts of this work have been presented at the 2022 Annual Meeting of the International Society for Magnetic Resonance in Medicine.

This is an open access article under the terms of the Creative Commons Attribution License, which permits use, distribution and reproduction in any medium, provided the original work is properly cited.

.....

© 2022 Physikalisch-Technische Bundesanstalt (PTB) and The Authors.. Magnetic Resonance in Medicine published by Wiley Periodicals LLC on behalf of International Society for Magnetic Resonance in Medicine.

Magn Reson Med. 2022;1–14. wileyonlinelibrary.com/journal/mrm | 1

¹Physikalisch-Technische Bundesanstalt, Braunschweig and Berlin, Germany

²Technische Universität Berlin, Biomedical Engineering, Berlin, Germany

³Technical University of Munich, Munich, Germany

⁴Imperial College London, London, United Kingdom

⁵Charité—Universitätsmedizin Berlin, corporate member of Freie Universität Berlin and Humboldt-Universität Zu Berlin, Experimental Clinical Research Center, Berlin, Germany

⁶DZHK (German Centre for Cardiovascular Research), Partner Site Berlin, Berlin, Germany

⁷Department of Cardiology and Nephrology, HELIOS Hospital Berlin-Buch, Berlin, Germany

⁸Division of Imaging Sciences and Biomedical Engineering, King's College London, London, United Kingdom

⁹Center for Magnetic Resonance Research, University of Minnesota, Minneapolis, Minnesota USA

¹⁰Medical Physics in Radiology, German Cancer Research Center (DKFZ), Heidelberg, Germany

INTRODUCTION

Magnetic resonance imaging (MRI) at ultra-high fields (UHF) provides ample opportunities for imaging the human body because of an intrinsic gain in signal-to-noise ratio (SNR), a higher chemical shift, and in many cases, a stronger contrast. However, the shortened radiofrequency (RF) wavelength within the tissue at 7T and the resulting constructive and destructive interference of the transmit (Tx) magnetic fields (B₁⁺) can lead to spatially nonuniform distributions of the flip angle (FA) and contrast, as well as to local signal dropouts. Such effects reduce the diagnostic value of UHF MRI³ when relying on conventional single-channel transmission imaging, especially when targeting the human body.

Parallel transmission (pTx) in combination with multi-Tx coils allows to spatially shape the amplitude and phase of B_1^+ -distributions in the human body^{2,4–7} using dedicated RF pulses. Such pTx methods can be divided into (1) calibration-free methods, which do not require additional calibration before the actual measurement, and (2) subject-tailored approaches.

Subject-tailored pTx pulses are typically calculated based on subject-specific B₁⁺-maps of all Tx channels, which require potentially lengthy adjustment times at the beginning of the study. Although rapid 3D B₁⁺-mapping of the entire brain at 7T can be achieved within only 40 seconds or less for 8 Tx channels, 13 the acquisition in the body is challenging. Scan times in the thorax can be substantially longer because of the larger field of view (FOV) and different sources of motion.¹⁴ Furthermore, some of the faster techniques applied to the brain, such as DREAM¹⁵ or Bloch-Siegert based mapping, ¹⁶ cannot be transferred straightforwardly to the body because of the sensitivity to blood flow or high RF power levels required at 7T, therefore, leaving only a few options for B₁⁺-mapping in the human body. Several works have used B₁-mapping methods in a slice-selective manner to limit scan times to acquire the data within a breath-hold of fewer than 20 seconds. 10,15,17,18 However, when FA optimization over a 3D volume is desired, multi-slice 2D or 3D B_1^+ -maps with scan times of more than 3 minutes 19 need to be acquired, requiring multiple breath-hold scans or free respiratory navigation. Recent preliminary work showed fast B_1^+ -mapping based on pre-saturated TurboFLASH sequences $^{20-22}$ in under a minute. However, reconstruction times of at least 8 minutes were needed. In addition to the time required for mapping and reconstructing B_1^+ , further calibration time is necessary for the RF pulse calculation, which depends on the complexity of the desired FA pattern. In total, an overall calibration time of more than 10 minutes is not uncommon for the human body in practice, which is one of the major drawbacks of UHF body MRI in clinical applications.

In recent works, deep learning (DL) techniques have shown promising results in reducing the RF pulse calculation times. 23-25 Vinding et al. 23 proposed a DL-based RF pulse design for high-FA 2D RF pulses at 7T with a pulse prediction time of under 10 milliseconds. Furthermore, different neural networks have been suggested to approximate channel-combined or channel-wise B₁⁺-distributions in various application scenarios.²⁵⁻²⁹ Abbasi-Rad et al.²⁸ reduced the specific absorption rate (SAR) in T₂-FLAIR imaging at UHF in the human head by determining a scaling factor for adiabatic RF pulses from a predicted channel-combined B₁⁺-magnitude. Plumley et al.²⁹ presented a system of conditional generative adversarial networks to predict how rigid motion changes complex, channel-wise B_1^+ -distributions in the head. This framework is highly suited for real-time pTx pulse re-design, but requires the initial channel-wise B₁⁺-maps. Eberhardt et al.²⁵ used B₁⁺-maps of only some of the 16 Tx channels of a 7T head coil and applied image-to-image translation networks to augment the data to a complete set of channel-wise B₁⁺-maps in the head. This approach achieved comparable FA homogenization results to those obtained from a full calibration data set if half of the Tx-channel-wise B₁⁺-maps were acquired.²⁵ Nevertheless, the need to substantially reduce the calibration time in the human body at 7T to a duration comparable to typical adjustment times of clinical scans of a few seconds remains unanswered. Yet, all recent DL techniques focused on the human brain or still needed a significant amount of input information.

In this proof-of-concept study, we reduce the time to obtain subject-specific B₁⁺-maps for subject-tailored cardiac pTx at 7T to a minimum. This is achieved by deriving complex, relative, channel-wise 2D B₁⁺-maps of 8 Tx channels from a single gradient-echo (GRE) localizer scan obtained in a combined 1-Tx mode with 32 receive (Rx) channels using DL. Because such localizer scans are acquired anyway for planning at the beginning of the session, this approach does not add any additional scan

time for B_1^+ -mapping. B_1^+ -phase shimming vectors are calculated based on the estimated B₁⁺-maps for a region of interest (ROI) covering the heart, and the predicted (PR) channel-wise B₁⁺-maps and resulting shimmed B₁⁺-fields are tested against the ground truth (GT). Finally, the feasibility of this approach is demonstrated in vivo.

2 **METHODS**

The proposed method of this work is to replace the B₁⁺-mapping step for subject-tailored pTx at 7T by a DL estimation approach. The neural network was trained on acquired localizer data serving as input and the corresponding B₁⁺-maps as GT. The performance of the network was assessed by calculating B₁+-phase shimming vectors based on the PR data and by an in-vivo application.

2.1 Hardware and setups

All data were acquired on a 7T whole-body MRI scanner (Magnetom 7T, Siemens, Erlangen, Germany) with an 8-channel transmit array and a whole-body gradient system using a commercial 32-element (8 dipoles and 24 loops) body coil array (MRI.TOOLS, Berlin, Germany). The coil comprised of two halves, a posterior and an anterior, each containing 16 elements. In the transmit case, the 32 elements were driven in an 8 Tx mode, in which four elements (1 dipole and 3 loops arranged along the dipole) were combined with hardware-fixed RF phases. In the receive case, all 32 elements operated independently using the 32 Rx channels of the system. The subjects were scanned in a supine position with the heart in the isocenter.

Reconstruction of the complex localizer data, manual selection of 2D heart and thorax ROIs, and static pulse design was carried out on a separate workstation (12 cores with 2.1 GHz, 128 GB RAM) using a custom-built toolbox written in MATLAB (Version 2020a, The MathWorks, Natick, MA, USA). The prediction of the relative B₁⁺-maps using the suggested neural network was executed within the Google Colab environment (Google LLC, Mountain View, CA, USA).

2.2 Data acquisition and processing

For the proposed neural network, 2 libraries containing complex localizers as input and channel-wise B₁+-maps as GT (Figure 1A) of healthy subjects were included in this study following approval of an internal review board and after written and informed consent. A performance library used for cross-validation comprised of 44 subjects (age: 34 ± 11 years, max/min: 66/21 years; body mass index (BMI): $23.79 \text{ kg/m}^2 \pm 3.10 \text{ kg/m}^2$, max/min: $34/19 \text{ kg/m}^2$, skewness: 0.96; male/female: 27/17), whereas an application library was used for in-vivo application.

For the input of the neural network, a GRE localizer was acquired with all Tx channels transmitting with the default phase setting (i.e., an RF phase optimized by the vendor based on electromagnetic simulations for the heart and aorta). However, the performance of this default setting depends on the size and shape of the human body and may yield destructive interferences in the heart.6 The parameters for this protocol were as follows: nominal FA = 15° , $T_E = 2.87$ ms, $T_R = 5.17$ ms, $T_A = 496.32$ ms, FOV = (384×384) mm², resolution = (4×4) mm², slice thickness = 4mm, slice distances = 20mm, flow compensation along slice/readout direction, no parallel imaging, no partial Fourier applied.

Subsequently, the same acquisition was repeated eight times using identical parameters with only a single Tx channel active per measurement. The active channel was incremented across scans. Following the approach by van de Moortele et al., 19,30 channel-wise, relative 2D B₁+-maps (i.e., an absolute scaling factor of the maps was unknown) were calculated using these eight measurements. These maps served as the neural network's GT. Although this fast technique provides only relative B₁⁺-maps, it has been demonstrated to be highly suited for multiple body applications in UHF imaging, including static⁷ and dynamic pTx for cardiac MRI.^{6,31,32} Unlike in previous works,^{6,32,33} the phase distributions were not calculated relative to one channel, but instead relative to the phase of the superimposed complex data of transmit channels 2, 3, and 4, which are located posterior and anterior to the subject. This resulted in a sufficient B_1^+ -magnitude in the heart for the reference signal, ensuring smooth, complex-valued data. The nine GRE acquisitions (one localizer and eight for B₁⁺-mapping) were obtained transversal in all subjects for 3 slice locations with 20 mm distance. In some subjects, the top or bottom slice contained only a small section of the apex or atria. For four subjects, one affected slice was not included in the final data set. Only one slice was available in one case. As a result, the performance library for cross-validation of the neural network comprised 126 slices, including the complex, 2D localizer data from 32 Rx channels and relative, channel-wise, 2D B₁⁺-maps from 8 Tx channels (Figure 1A).

The data were pre-processed as follows: real and imaginary parts of the 32 Rx-channel-wise localizers and 8 calculated Tx-channel-wise B₁⁺-maps were separated. Normalization by the maximum absolute value of the complex localizer for the input and the maximum absolute value of the complex B₁⁺-distribution for the GT was conducted to ensure a comparable dynamic range in the complex data across different subjects between 0 and 1. The

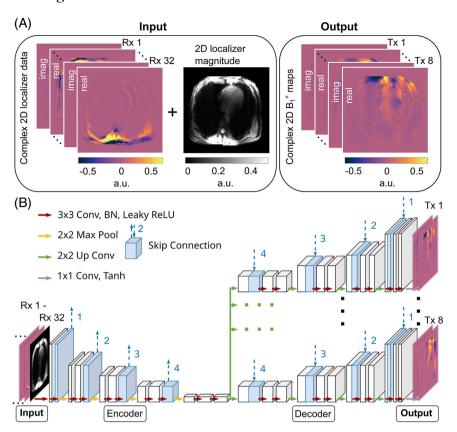


FIGURE 1 (A) Data structure for the input and output of the proposed neural network. The recieve (Rx) -channel-wise localizer data is split up into real and imaginary part, and the corresponding localizer magnitude image is used as the input. This results in an overall input size of $96 \times 96 \times 65$. Similarly, the ground truth transmit (Tx) -channel-wise B₁+-maps are split up into real and imaginary data and arranged to a size of $96 \times 96 \times 16$. (B) The architectural structure of the proposed deep convolutional neural network. Each stage in the encoder consists of a max pool layer, two 3×3 convolutions with a batchnorm (BN), and a LeakyReLU activation function. The decoder is split into eight decoder pipelines for each Tx channel of the B₁⁺-output. The decoder pipelines are similar to the encoder, the max pool is replaced by a 2×2 up convolution. Skip connections and dropouts are included.

32 Rx-channel-wise localizer slices in the image domain were used as input for the network leading to an overall input size of $96 \times 96 \times 65$, i.e., the 32 real, 32 imaginary Rx-channel-wise localizers, and one magnitude image using the root-sum-of-squares across all Rx channels.

Correspondingly, the 8 Tx-channel-wise B_1^+ -maps resulted in a size of $96\times96\times16$ serving as GT. The sum-of-magnitudes (SOM) of the B_1^+ -maps were used to manually select a binary mask (ROI1) covering the thorax for every slice and subject. This mask was then multiplied pixel-wise with the input and output data to suppress the noise signal outside the thorax. This step was repeated for another mask (ROI2) covering the heart to be used for static pulse design and quantitative evaluation.

2.3 Network architecture and model evaluation

The proposed neural network was based on a modified version of the established 2D UNet architecture^{34,35} to predict complex, Tx-channel-wise, 2D B₁⁺-maps from complex, Rx-channel-wise 2D localizer data (Figure 1B). This architecture is suitable for a plethora of applications, for example, image reconstruction,^{36,37} quantitative parameter mapping,^{38,39} and artifact correction.^{37,40} The chosen architecture comprised four encoding stages,

each including a 2×2 max pooling layer (downsampling) and 2 repetitions of 3 × 3 convolutions, batch normalizations, and LeakyReLU activation functions. For decoding, the structure was split into eight parallel pipelines, with one output B₁⁺-map for each Tx channel connecting the 32-Rx-channel localizer input data with an overall input size of 96 × 96 × 65 with the corresponding 8 Tx channel B_1^+ -maps with an overall size of $96 \times 96 \times 16$. Each decoder stage included a 2×2 up convolution layer (upsampling), 2 repetitions of 3×3 convolutions, batch normalizations, and LeakyReLU activation functions. The number of output filters was set to 32 in the first and last layer and was doubled/halved per encoding/decoding stage, up to a maximum of 512 filters. The network used skip connections between corresponding stages to reduce spatial resolution losses and avoid vanishing gradients.41 To minimize overfitting, dropout layers were used for regularization with an increasing dropout rate per stage.

The neural network was implemented in Python 3.8.3 using Tensorflow $2.2.0^{42}$ and trained on a 24 GB NVIDIA Titan RTX on-premise. Encoder-decoder models typically rely on minimizing an L^{1-} or L^{2-} loss between PR and GT image, which introduces an asymmetry in the loss function regarding magnitude and phase. In contrast, the proposed model relies on a symmetric loss to achieve better performance for the training process. ⁴³ This \bot – loss is defined as:

with $P(x, y) = \frac{|\text{Re}(x)Im(y) - Im(x)\text{Re}(y)|}{|x|}$ for two vectors x and y.

The minimization was conducted using the ADAM optimizer. For better convergence, a decaying learning rate⁴⁴ was used: LR_i = $1 \times 10^{-4} e^{-0.99944(j-1)}$ for the j-th epoch to achieve a decrease in the learning rate of one magnitude after 4000 epochs. The model was trained for 4000 epochs with a batch size of 2, leading to a training time of approximately 240 min. The code for implementing the neural network, the network weights, the data libraries, and pulse design and analysis algorithms can be downloaded from https://github.com/felixkrueger90/

The quality of the PR B₁⁺-maps (i.e., its resemblance to the GT) was evaluated via image quality metrics assessed through the root-mean-squared error (RMSE) and structural similarity index measure (SSIM) performed on the ROI1-masked thorax data. A 5-fold cross-validation was carried out on the entirety of the performance library (Figure 2A) to evaluate the model's generalization performance on all thorax geometries. First, all data were randomly shuffled at the subject level and split into 5 subsets containing 8/9 subjects each. Therefore, every subject is included in only one of the subsets. One subset is kept as unseen test data, and the other 4 are used for network training. This process was repeated five times, with each subset being the test data once. For every data split, approximately 20% of all 44 subjects (8/9 subjects) were used for testing and 80% (35/36 subjects) for training.

Pulse design and analysis

To assess the suitability of the network for calculating optimized excitations in the body at 7T, B₁⁺ -phase shimming was carried out based on the DL predicted maps. Shimming was performed by determining a shim vector $\mathbf{b}_{\mathrm{Tx}} = \left(e^{i\overline{\varphi_1}}, \cdots, e^{i\varphi_8}\right)^{\mathrm{T}}$ that is applied to the individual PR B_1^+ -maps with φ_i being the transmit offset phase for channel i. The optimization was executed regarding three cost functions based on the Tx-channel-wise B₁⁺ -maps, superimposed by ROI2 covering the heart. The cost functions were optimized using the fmincon constraint optimization function in MATLAB with the constraint that the amplitudes of **b** equal 1. The first cost function was based on the coefficient of variation (CV),⁶ which is a surrogate for the spatial homogeneity of the combined B_1^+ -field:

$$CV = \frac{SD\left(\left|\sum_{Tx=1}^{N_{Tx}} \widehat{B}_{1,Tx}^{+} \boldsymbol{b}_{Tx}\right|\right)}{\operatorname{mean}\left(\left|\sum_{Tx=1}^{N_{Tx}} \widehat{B}_{1,Tx}^{+} \boldsymbol{b}_{Tx}\right|\right)}\Big|_{ROI}, \qquad (2)$$

where SD is the spatial standard deviation and N_{Tx} is the number of independent Tx channels (i.e., 8 for our case). When minimizing the CV, the resulting homogeneous B₁⁺ -shim vector is denoted by \mathbf{b}_{Hom} . The second cost function aimed to maximize the mean transmit efficiency $\eta^{6,7,19}$ to obtain the phase shim setting \mathbf{b}_{Eff} with highly constructive interference:

$$\eta = \text{mean} \left(\frac{\left| \sum_{Tx=1}^{N_{Tx}} \hat{B}_{1,Tx}^{+} \boldsymbol{b}_{Tx} \right|}{\sum_{Tx=1}^{N_{Tx}} \left| \hat{B}_{1,Tx}^{+} \right|} \right) \Big|_{ROI}.$$
 (3)

In some cases, maximizing the transmit efficiency yielded highly localized signal dropouts, particularly when the ROI was large. A modified cost function was used to overcome this effect that aimed to fix the mean efficiency within the ROI to a user-defined target value η_{tar} :

$$\eta_{\text{fix}} = \text{mean} \left(\left| \frac{\left| \sum_{\text{Tx}=1}^{N_{\text{Tx}}} \widehat{B}_{1,\text{Tx}}^{+} \boldsymbol{b}_{\text{Tx}} \right|}{\sum_{\text{Tx}=1}^{N_{\text{Tx}}} \left| \widehat{B}_{1,\text{Tx}}^{+} \right|} - \eta_{\text{tar}} \right| \right)_{\text{ROI}}.$$
 (4)

This strategy yielded a trade-off between a homogeneous and an efficient B₁⁺ -field, typically avoiding dropouts. According to our experience, it is beneficial for cardiac MRI applications. Therefore, as a third optimization problem, the shim setting \mathbf{b}_{Enf} was applied with $\eta_{tar} = 0.5$.

To access the quality of the PR B₁⁺-maps and the feasibility of the approach in context of a subject-specific calibration, the B_1^+ -shim vectors \mathbf{b}_{Hom} , \mathbf{b}_{Eff} , and \mathbf{b}_{Enf} were calculated based on the PR B₁⁺-maps. The shim vectors were then applied to both the PR and GT B₁⁺-maps to generate the combined B₁⁺-map when all Tx channels transmit together. These combined maps based on the GT and PR were quantitatively compared for all three settings and the default shim \mathbf{b}_{Def} in terms of CV and mean efficiency optimized for ROI2 covering the heart.

2.5 **Experimental application**

The application library was acquired for in-vivo application of the proposed approach. The library contained three subjects (1 female/ 2 male, 28/32/40 years, $19/23/26 \text{ kg/m}^2$) who underwent the same protocol as the volunteers of the performance library. For B₁⁺-prediction, a separate neural network was trained on all thorax geometries from all subjects of the performance library. Therefore, a second 5-fold cross-validation was performed (Figure 2B), where the 126 slices from all 44 subjects were split randomly into 5 subsets at the slice level. In contrast to the first cross-validation, not all slices from a single subject were contained in one subset. Therefore, the same subject

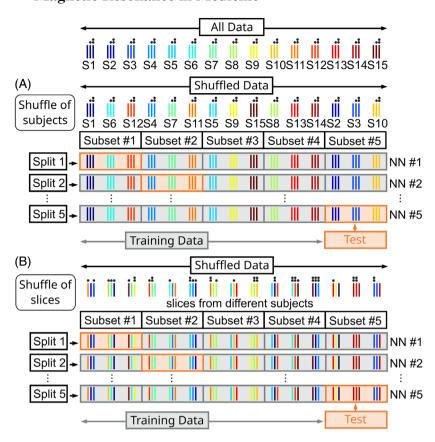


FIGURE 2 Workflow of the 5-fold cross-validation. Here, the initial data sample consists of 15 subjects (S1-S15) with three slices each. Slices from the same subject are tagged with the same color, whereas the central slice within one subject is marked with one and the upper slice with two dots. (A) For performance evaluation, the data is randomly shuffled at the subject level. The shuffled data is partitioned into five equal subsets. For each data split, one subset is used as the test data, whereas the remaining subsets are used for training the model. This is repeated five times leading to five different neural networks (NN). (B) For the in-vivo application, the process is adapted. The shuffling is carried out at the slice level, meaning slices from one subject can simultaneously be used for testing and training. This ensures that the network is trained on all thorax geometries.

(but different slices) may be used for testing and training simultaneously. This approach ensured that the resulting networks were always trained on all thorax geometries, except for the subject, which contained only 1 slice. Subsequently, the network leading to the highest SSIM when evaluating the corresponding test data is applied to the three subjects of the *application library*. In addition, B_1^+ -shimming was performed based on the PR B_1^+ -maps using the same optimization functions described above.

3 | RESULTS

This section demonstrates the prediction of channel-wise 2D B_1^+ -maps from an initial localizer for the heart at 7T using DL. The process takes under 1 second to derive the PR maps. The cross-validation results for unseen test cases are provided. The validity of the approach is exemplified by different phase-only B_1^+ -shim vectors calculated for the PR and successive in-vivo acquisition using the introduced calibration pipeline.

3.1 | Model evaluation and selection

Table 1 contains the quantitative differences between the PR and the GT channel-wise B_1^+ -maps. Five neural

networks were assessed for the corresponding subsets of the 5-fold cross-validation concerning the RMSE, SSIM, and symmetric loss. For example, the PR maps for unseen test subset 2 generated by network 2 yield a mean symmetric loss value of 0.0172, a mean RMSE value of 0.0438%, and a mean SSIM value of 0.7638 (Table 1). Similar results are observed for all five networks of the cross-validation. Network 2 yields the lowest mean value in the symmetric loss, the second lowest mean value in the RMSE, and the highest in the SSIM. Therefore, network 2 is selected for the performance analysis. The boxplot in Figure 3 depicts the performance of network 2 on all unseen slices from subset 2 regarding the RMSE (Figure 3A) and SSIM (Figure 3B). The performance of the network varies among slices ranging from 0.0314% to 0.0615% for the RMSE and from 0.7024 to 0.8195 for the SSIM. As representative examples, three of the 26 unseen test cases are visualized in the following with a high SSIM value of 0.7825 (example 1), a medium value of 0.7552 (example 2), and a low value of 0.7218 (example 3).

3.2 | Evaluation of image quality

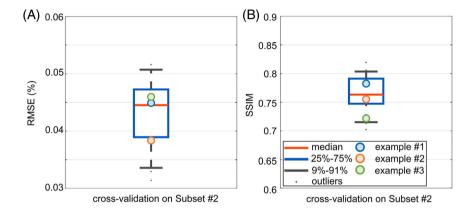
Figure 4 shows the channel-wise PR compared to the GT B_1^+ -maps for unseen example 1. The results for examples 2 and 3 are provided in the Supporting Information

TABLE 1 RMSE, SSIM, and symmetric loss values obtained from the evaluation of the five NN applied to the corresponding five test subsets when performing a 5-fold cross-validation. The BMI distribution is given for every test set and the corresponding training data. Network 2 evaluated on subset 2 (highlighted in bold) is used for further assessment because it results in the highest mean SSIM.

NN	Training BMI mean ± SD (kg/m²)	Tested on	Test BMI mean ± SD (kg/m²)	RMSE mean ± SD (%)	SSIM mean ± SD	Symmetric loss mean ± SD
1	23.57 ± 3.19	Subset 1	23.99 ± 2.60	0.0440 ± 0.0064	0.729 ± 0.049	0.019 ± 0.003
2	24.06 ± 3.11	Subset 2	22.12 ± 2.43	0.0438 ± 0.0067	$\boldsymbol{0.764 \pm 0.033}$	$\boldsymbol{0.017 \pm 0.002}$
3	23.42 ± 2.72	Subset 3	24.53 ± 4.08	0.0435 ± 0.0096	0.727 ± 0.051	0.019 ± 0.004
4	23.48 ± 3.21	Subset 4	24.37 ± 2.43	0.0449 ± 0.0063	0.734 ± 0.030	0.019 ± 0.002
5	23.76 ± 3.10	Subset 5	23.15 ± 3.01	0.0518 ± 0.0084	0.728 ± 0.030	0.020 ± 0.004

Abbreviations: BMI, body mass index; NN, neural network, RMSE, root-mean-square error; SD, standard deviation; SSIM, structural similarity index measure.

FIGURE 3 Boxplots summarizing the root-mean-squared error (RMSE) (A) and the structural similarity index measure (SSIM) (B) for all unseen test cases of subset 2 predicted by network 2. Three representative examples with a high SSIM value of 0.7825 (example 1), a medium value of 0.7552 (example 2) and a low value of 0.7218 (example 3) are marked separately and used for further assessment.



(Figure S1 and Figure S2). Qualitatively, the channel-wise PR matches the GT data not only for the magnitude, but especially for the phase distributions, which is a key requirement for subsequent pTx applications. A more detailed inspection of the magnitude difference ΔB_1^+ (last columns in Figure 4) reveals larger differences in the chest region closer to the coil at Tx channels 1 and 8 compared to the other channels. This observation is consistent among subjects. A similar observation is not made with elements closer to the spine.

The observed match is further supported in Figure 5, showing a detailed view (Figure 5A) and a quantitative comparison (Figure 5B,C) between the PR and GT data for the third and fourth Tx channel for example 1. Line plots for cross-sections through the heart for magnitude and phase of the DL predicted (black) and measured (red) B_1^+ -maps are displayed. Although the magnitude and phase of the PR distributions follow the GT, sudden changes or edges (e.g., in the phase) are reflected less accurately by the prediction. This is not unexpected because neural networks minimizing distance metrics tend to smooth the data. The mean error for the magnitude (nRMSE, definition Plumley et al. $^{29} \pm SD$) averaged over the cross-section amounts to $5.90\% \pm 8.87\%$ for Tx

channel 3 and $0.41\% \pm 1.01\%$ for Tx channel 4, whereas the mean phase differences are $0.37 \, \text{rad} \pm 0.38 \, \text{rad}$ for Tx channel 3 and $0.43 \, \text{rad} \pm 0.50 \, \text{rad}$ for Tx channel 4.

Similar results can be observed when the eight B_1^+ -maps are combined for the three unseen test cases (Figures 6 and Supporting Information Figure S3). The network yields a prediction for example 1 with a mean error for the SOM over the heart of 0.69%, for the magnitude of the complex sum (MOS) of 3.86%, and with a mean difference in the phase of the complex sum (POS) of 0.015 rad. The pattern, i.e., the local signal dropouts and the corresponding phase wraps in the heart are approximated accordingly (see yellow arrows).

3.3 \mid B_1^+ -shimming performance

Figure 7 depicts the B_1^+ -shimming results for the three unseen test cases (examples 1–3) after calculating three B_1^+ -shim vectors based on the PR B_1^+ -maps that are applied to the PR and GT B_1^+ -maps. The results are shown for the default shim setting \mathbf{b}_{Def} (Figure 6), homogenous shim \mathbf{b}_{Hom} , efficiency setting \mathbf{b}_{Eff} , and the shim setting \mathbf{b}_{Enf} , when enforcing the mean efficiency to 50%.

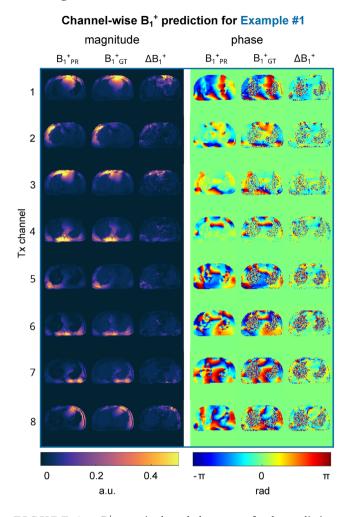


FIGURE 4 B_1^+ -magnitude and phase maps for the prediction of the neural network $B_{1\,PR}^+$ compared to the GT $B_{1\,GT}^+$ for the unseen example 1 with a high structural similarity index measure value. The absolute error ΔB_1^+ between the prediction and ground truth shows a higher residual error for the first transmit (Tx) channel and the eighth channel as compared to channels 2–7. Overall, the prediction qualitatively matches the GT for both the magnitude and the phase.

The default setting \mathbf{b}_{Def} (left column of Figure 7) calculated based on electromagnetic simulations yields different magnitude distributions among subjects, demonstrating the need for optimized pTx excitations. However, both PR and GT B₁⁺-maps show visually consistent results for all three subjects. Although **b**_{Hom}, tends to improve the homogeneity to the cost of lower magnitude values, the efficiency shim $\mathbf{b}_{\mathrm{Eff}}$ leads to higher amplitudes but at the expense of stronger spatial B₁⁺-variation. In contrast, the enforced shim \boldsymbol{b}_{Enf} causes lower mean efficiency but also less pronounced dropouts, as seen in the $b_{\rm Eff}$ results. Importantly, all such features are always reflected by both the GT and PR maps. This high level of visual agreement among PR and GT maps is observed independently of the example and applied shim.

To investigate this agreement in more detail, a quantitative evaluation for the settings \mathbf{b}_{Hom} and \mathbf{b}_{Eff} are shown in Figure 8 for all 26 test cases of subset 2. When calculating a homogenous shim based on the PR maps (Figure 8A) the mean CV decreases from 43.5% (range = 18.7%–55.9%) to a value of 17.2% (range = 7.4%-29.0%). The mean decreases from 39.6% (range = 25.0%-54.6%) to 34.2% (range = 24.9%–54.6%) when applying \mathbf{b}_{Hom} to the GT. Those CV values are overall higher for the GT compared to the PR case, but the CV values after shimming for both are lower than compared to \mathbf{b}_{Def} . For the efficient shim \mathbf{b}_{Eff} , the mean efficiency increases from a mean value of 47.1% (range = 37.9%-57.2%) to a value of 68.3% (range = 63.0%–88.4%) for the PR maps. When applying the same vector to the GT, the mean efficiency increases from 46.4% (range = 37.2%-53.3%) to 62.5%(range = 48.3%–78.6%), leading to similar results for the GT and PR data.

3.4 Performance of the neural network in unseen test cases

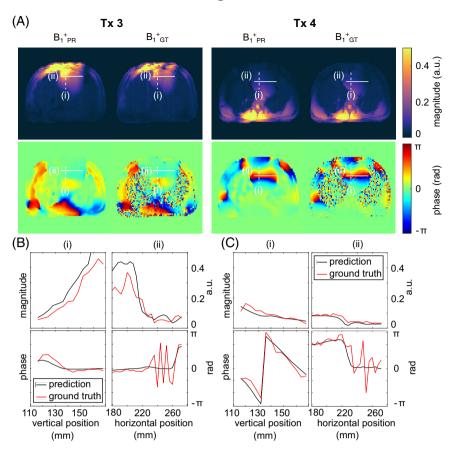
Figure 9 illustrates the FA prediction for the measured channel-combined B₁⁺-maps, the reconstructed 2D GRE image, and cardiac 2D cine GRE for the unseen test case 1 acquired with the corresponding pulses designed on the PR B₁⁺-maps. Qualitatively, a close match between FA predictions, the 2D GRE images, and cardiac cine images is observed, demonstrating the feasibility of the DL-based calibration approach. Similar results are obtained for two other subjects, illustrated in Supporting Information Figure S4. The results of the 5-fold cross-validation using all thorax geometries are provided in Supporting Information Table S1.

4 | DISCUSSION

This work presents a novel, fast DL-based approach for obtaining relative B_1^+ -maps in the human heart at 7T, addressing the urgent need for reducing B_1^+ -calibration times in UHF body imaging. The data-driven approach predicts the 2D B_1^+ -maps in 0.2 seconds per slice from 2D localizer images typically acquired at the beginning of the session for planning. Therefore, this approach could make an additional B_1^+ -mapping scan obsolete. Although this proof-of-principle is applied to estimate B_1^+ only in a 2D slice, it may be extended toward multi-slice application and potentially to 3D. Thereby, it may save several minutes of calibration time and potentially several breath-holds.

Some related DL approaches in UHF imaging have been recently reported for subject-specific calibration of the transmit B₁⁺-fields.^{25–29} Abbasi-Rad et al.²⁸ predicted

FIGURE 5 (A) Magnitude and phase for the predicted $(B_{1\,PR}^+)$ and the ground truth $(B_{1\,GT}^+)$ relative B_{1}^+ -maps for transmission (Tx) channel 3 (anterior channel) and 4 (posterior channel) of example 1. The two channels have been chosen because of their high combined magnitude within the heart. Vertical (dotted) and horizontal (solid) profiles of the phase and magnitude for the deep learning predicted (black) and ground truth (red) 2D B_1^+ -maps are provided for Tx 3 (B) and Tx 4 (C).



channel-combined B_1^+ -magnitudes from a standard localizer to determine a FA scaling factor for adiabatic RF pulses. Plumley et al.²⁹ used DL to predict the complex, channel-wise B_1^+ -maps of an 8 Tx channel head coil after motion based on an initially acquired set of B_1^+ -maps. Eberhardt et al.²⁵ used a 2-to-16-fold under-sampled set of B_1^+ -maps for a 16 Tx channel head coil and used neural networks to augment a full set of 16 complex, channel-wise B_1^+ -maps. The presented work, in contrast, derives a complete set of complex, channel-wise B_1^+ -maps of an 8 Tx channel body coil from an initial localizer image and applies the technique to the human body at 7T.

The proposed network is based on a standard UNet architecture and translates complex, 32 Rx-channel-wise localizers (B_1^-) as input into complex, relative 8 Tx-channel-wise B_1^+ -maps. Early results of this study showed that the magnitude data could be retrieved with reasonable accuracy from the localizer. All features of the magnitude B_1^+ -maps were approximated accordingly. Only localized deviations were visible. The maps appear smoothed, likely introduced by the loss function during network training. The effect seems not to affect results or hinder shimming applications because the spatial variation of B_1^+ is smooth. Estimating the Tx phase turned out to be more difficult, which was also reported by Plumley et al., 29 and particularly the loss function and reference RF

phase impacted the result. Although in a few cases very localized residual phase prediction errors are still observed at locations with rapidly changing phases or at singularities, an overall high similarity is observed between PR and GT phase patterns. This property is a prerequisite for any static and dynamic pTx applications.

The DL models were all trained on a considerably larger set of maps from 44 healthy subjects compared to previous work for the brain^{25,28,29} to account for stronger inter-subject variations of the thorax geometry. 12 However, a shift in the training data regarding different property distributions (BMI, age, and gender) can lead to a bias in the performance of the network. For example, the performance was higher for subjects with smaller BMI due to a skewed distribution of BMIs in the training library toward lower values. Because the BMI distribution in Germany is shifted toward higher values $(26.85 \pm 4.95 \text{ kg/m}^2)$, 46 simply increasing the data may further amplify the bias. A selection from a more representative sample of the general population is needed for the training data. Furthermore, maps from patients may need to be included to account for anomalies in the thorax geometry.

When evaluating the pulse design with a phase-only B_1^+ -shim setting optimized on the PR, all features in the magnitude and phase match when applying the shim

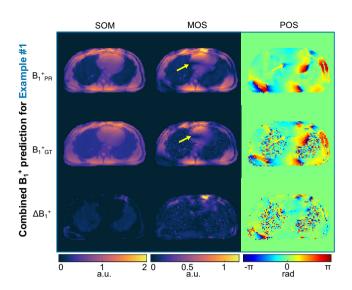


FIGURE 6 Combined B_1^+ -maps for the unseen test case example 1. The predicted $(B_{1\,PR}^+)$ and the ground truth $(B_{1\,GT}^+)$ data are shown for the sum of magnitudes (SOM) over the 8 transmit (Tx) channel, the magnitude of sum (MOS), and the phase of the summed-up data (POS). When evaluating the region of interest over the heart the average error is 0.69% regarding the SOM, 3.86% for the MOS, as well as a mean difference for the POS of 0.015 rad. The absolute error ΔB_1^+ between the prediction and the ground truth is presented. The signal dropouts and associated phase-wraps are marked by the yellow arrows.

vectors. For the homogeneity shim setting \mathbf{b}_{Hom} , the CV on the GT is higher compared to the optimized PR for all test cases, which is likely because of the smoothing effect on the magnitude data for the PR. Nevertheless, the resulting CV values for the GT are lower for b_{Hom} than for the default setting **b**_{Def}. For some test cases (Figure 7, example 1), optimizing the homogeneity still leads to localized signal dropouts in the heart. This observation, however, is consistent with our experience with previous studies that apply B₁ phase shimming to the human heart. The same observation is made in this work when relying on GT data used for calibration. If comparing the channel-combined PR and GT B_1^+ -maps after applying \mathbf{b}_{Eff} or \mathbf{b}_{Hom} , the match was higher for the efficient than for the homogeneous shim. The mean values for examples 1-3 regarding the SSIM, when applying \mathbf{b}_{Hom} , were lower (0.7412) than compared to $\mathbf{b}_{\rm Eff}$ (0.7569). Small phase deviations may explain this because an error in the phase is expected to impact a homogeneous shim more than an efficient shim.

Although this study successfully demonstrates the feasibility of deriving complex multi-Tx-channel B_1^+ -maps from localizer scans, the work is still subject to a few limitations. In this proof-of-concept study, only a single RF coil, a commercial body array coil with 32 elements (8 dipoles and 24 loop elements), has been used because this is the only available multi-Tx-channel coil at our center that can be applied in vivo. The same 32 elements are used for transmission and reception when operating this

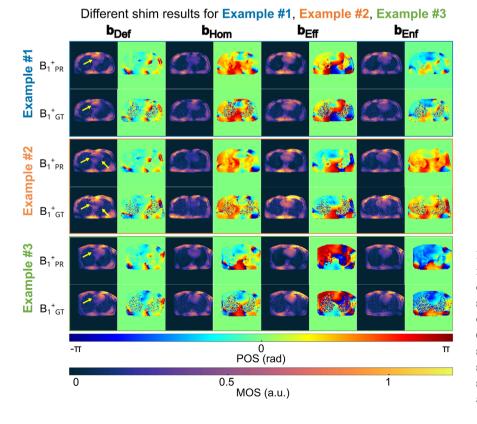


FIGURE 7 Different B_1^+ -shimming results for the three unseen test cases. The default shim setting $\mathbf{b}_{\mathrm{Def}}$, the homogeneous shim $\mathbf{b}_{\mathrm{Hom}}$, the efficiency shim $\mathbf{b}_{\mathrm{Eff}}$, and the enforced shim $\mathbf{b}_{\mathrm{Enf}}$ applied to the predicted $(B_{1}^+P_R)$ and the ground truth $(B_{1}^+G_T)$ data are shown for the magnitude of the complex sum (MOS), and the phase of the complex sum (POS). Dropouts using the default shim are highlighted with yellow arrows.

FIGURE 8 (A) Coefficient of variation (CV) in the heart region of interest (ROI) for the default shim setting $\mathbf{b}_{\mathrm{Def}}$ and the homogenous shim setting \mathbf{b}_{Hom} , optimized on the generated B_1^+ -maps and applied to the predicted $(B_1^+)_{\mathrm{PR}}$) and the ground truth $(B_1^+)_{\mathrm{GT}}$ data. (B) Boxplot summarizing the mean transmit efficiency in the ROI for the default shim setting $\mathbf{b}_{\mathrm{Def}}$ and the homogenous shim setting $\mathbf{b}_{\mathrm{Eff}}$, optimized on the generated B_1^+ -maps and applied to the predicted and the ground truth data.

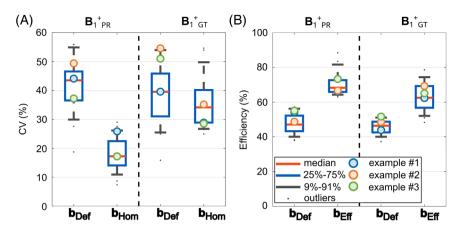
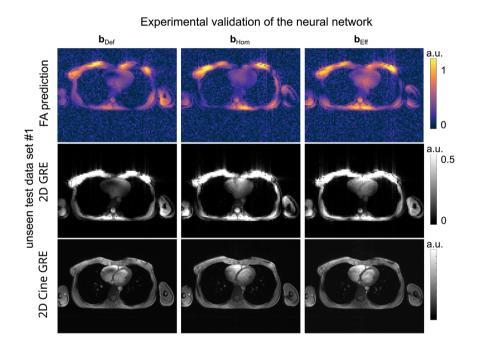


FIGURE 9 Flip angle (FA) prediction for the measured channel-combined B_1^+ -maps, the reconstructed 2D gradient-echo (GRE) image, and cine GRE for the unseen test case 1. The used parallel transmission pulses for the default, homogenous, and efficiency settings were calculated on the deep learning-based B_1^+ -maps. All test subjects were not part of the cross-validation process.



coil. In the Tx case, four elements (1 dipole and 3 loops) are combined with a fixed phase setting, whereas all elements acquire independently in the Rx case. This might be beneficial for this type of application because the 8 Tx B₁⁺-maps are derived from 32 Rx maps. Future investigations could also include other types of coils, for example, pure 8 Tx/8 Rx transceiver body coil arrays. Furthermore, the presented work investigates only 2D transversal slices, but oblique slices are typically required for practical cardiac applications. It is expected that changing the orientation will require new training data, and the performance of the method in oblique slices may be different. However, extending the method toward 3D coverage that allows FA optimization over a 3D volume can be considered. Covering multiple slices of 2D transversal B₁+-maps is expected to be feasible, for example, by relying on serial data processing. The extension of this technique by estimating 3D

B₁⁺-maps from 3D localizer data may be more challenging and requires further investigation.

As for other DL approaches, the performance of the present technique depends on the type of training data used. In the presented case, the GT B_1^+ -maps and, therefore, the PR were relative³⁰ and biased by the square root of the proton density.¹⁹ Despite this limitation, such maps were chosen for this work because they have proven to perform well for various B_1^+ -shimming applications targeting the human heart or body.^{7,31,32} Future work may include the investigation of other B_1^+ -maps obtained by different mapping techniques.

Although further investigations are needed to investigate the full potential of this method, this proof-of-principle study demonstrates the feasibility of overcoming long calibration scans in a subject-specific calibration pipeline using DL. Together with other DL

methods, this technique could serve as a plug-and-play like calibration solution for UHF imaging.

5 | CONCLUSION

This study successfully demonstrates that DL approaches are highly suitable to predict 2D relative B_1^+ -maps from initial localizer scans in the human heart at 7T. The proposed approach reduces the calibration time for subject-specific pTx to less than a second. This work is expected to impact the progress of UHF body applications, which are hindered by stronger B_1^+ -variations compared to the brain and by longer calibration times. Based on this approach, a push-button in situ optimization embedded in the scanner's calibration routine may be feasible, potentially promoting the clinical applicability of body imaging at UHF in the future.

ACKNOWLEDGMENT

We gratefully acknowledge funding from the German Research Foundation SCHM 2677/2-1, SCHM 2677/4-1, and GRK2260-BIOQIC. Open Access funding enabled and organized by Projekt DEAL. Open Access funding enabled and organized by Projekt DEAL.

FUNDING INFORMATION

The German Research Foundation, Grant/Award Numbers: SCHM 2677/2-1, SCHM 2677/4-1 and GRK2260-BIOQIC.

ORCID

Felix Krueger https://orcid.org/0000-0001-9453-8992 Christoph Stefan Aigner https://orcid.org/0000-0003-3618-9610

Kerstin Hammernik https://orcid.org/0000-0002-2734-1409

Sebastian Dietrich https://orcid.org/0000-0002-1610-909X

Jeanette Schulz-Menger https://orcid.org/0000-0003-3100-1092

Tobias Schaeffter https://orcid.org/0000-0003-1310-2631

Sebastian Schmitter https://orcid.org/0000-0003-4410-6790

REFERENCES

- Ladd ME, Bachert P, Meyerspeer M, et al. Pros and cons of ultra-high-field MRI/MRS for human application. Prog Nucl Magn Reson Spectrosc. 2018;109:1-50.
- 2. Padormo F, Beqiri A, Hajnal J, Malik S. Parallel transmission for ultrahigh-field imaging. *NMR Biomed*. 2016;29:1145-1161.
- 3. Marques JP, Khabipova D, Gruetter R. Studying cyto and myeloarchitecture of the human cortex at ultra-high field with

- quantitative imaging: R1, R2* and magnetic susceptibility. *NeuroImage*. 2017;147:152-163.
- Wu X, Schmitter S, Auerbach EJ, Uğurbil K, Van de Moortele PF. Mitigating transmit B 1 inhomogeneity in the liver at 7T using multi-spoke parallel transmit RF pulse design. *Quant Imaging Med Surg.* 2014;4:4-10.
- 5. Padormo F, Hess AT, Aljabar P, et al. Large dynamic range relative B1+ mapping. *Magn Reson Med*. 2016;76:490-499.
- Aigner CS, Dietrich S, Schmitter S. Three-dimensional static and dynamic parallel transmission of the human heart at 7T. NMR Biomed. 2021;34:1-15.
- 7. Metzger GJ, Snyder C, Akgun C, Vaughan T, Ugurbil K, Van De Moortele PF. Local B1+ shimming for prostate imaging with transceiver arrays at 7T based on subject-dependent transmit phase measurements. *Magn Reson Med.* 2008;59:396-409.
- 8. Winter L, Kellman P, Renz W, et al. Comparison of three multichannel transmit/receive radiofrequency coil configurations for anatomic and functional cardiac MRI at 7.0T: implications for clinical imaging. *Eur Radiol*. 2012;22:2211-2220.
- Orzada S, Solbach K, Gratz M, et al. A 32-channel parallel transmit system add-on for 7T MRI. PLoS One. 2019;14:1-20.
- 10. Brunheim S, Gratz M, Johst S, et al. Fast and accurate multi-channel B1+ mapping based on the TIAMO technique for 7T UHF body MRI. *Magn Reson Med.* 2018;79:2652-2664.
- 11. Gras V, Vignaud A, Amadon A, Le Bihan D, Boulant N. Universal pulses: a new concept for calibration-free parallel transmission. *Magn Reson Med.* 2017;77:635-643.
- 12. Aigner CS, Dietrich S, Schaeffter T, Schmitter S. Calibration-free pTx of the human heart at 7T via 3D universal pulses. *Magn Reson Med.* 2022;87:70-84.
- 13. Herrler J, Liebig P, Gumbrecht R, et al. Fast online-customized (FOCUS) parallel transmission pulses: a combination of universal pulses and individual optimization. *Magn Reson Med*. 2021;85:3140-3153.
- 14. Niendorf T, Graessl A, Thalhammer C, et al. Progress and promises of human cardiac magnetic resonance at ultrahigh fields: a physics perspective. *J Magn Reson*. 2013;229:208-222.
- 15. Nehrke K, Börnert P. DREAM-a novel approach for robust, ultrafast, multislice B1 mapping. *Magn Reson Med*. 2012;68:1517-1526.
- Sacolick LI, Wiesinger F, Hancu I, Vogel MW. B1 mapping by Bloch-Siegert shift. Magn Reson Med. 2010;63:1315-1322.
- 17. Schmitter S, Moeller S, Wu X, Auerbach EJ, Metzger GJ. Simultaneous multislice imaging in dynamic cardiac MRI at 7T using parallel transmission. *Magn Reson Imaging*. 2017;77:1010-1020.
- Schmitter S, Adriany G, Waks M, et al. Bilateral multiband 4D flow MRI of the carotid arteries at 7T. Magn Reson Med. 2020;84:1947-1960.
- 19. Dietrich S, Aigner CS, Kolbitsch C, et al. 3D free-breathing multichannel absolute B1+ mapping in the human body at 7T. *Magn Reson Med.* 2021;85:2552-2567.
- Kent JL, Ladislav V, Iulius D, Hess AT. Rapid B1+ Mapping using a Sandwiched Train Satureated TurboFLASH Sequence at 7T. In: Proceedings of the 31th Annual Meeting of ISMRM, London, United Kingdom: Abstract 0456; 2022.
- Kent JL, Ladislav V, Dragonu I, Hess AT. Acceleration Methods to Acquire Fast 3D Multi-Channel Absoloute B1+ Maps at 7T for 8-Channel pTx. In: Proceedings of the 31th Annual Meeting of ISMRM, London, United Kingdom: Abstract 2864; 2022.

- Hess AT, Iulius D, Chiew M. Accelerated calibrationless parallel transmit mapping using joint transmit and receive low-rank tensor completion. *Magn Reson Med.* 2021;86: 2454-2467.
- 23. Vinding MS, Aigner CS, Schmitter S, Lund TE. DeepControl: 2DRF pulses facilitating B1+ inhomogeneity and B0 off-resonance compensation in vivo at 7T. *Magn Reson Med*. 2021;85:3308-3317.
- Ianni JD, Cao Z, Grissom WA. Machine learning RF shimming: prediction by iteratively projected ridge regression. *Magn Reson Med.* 2018;80:1871-1881.
- 25. Eberhardt B, Poser BA, Shah NJ, Felder J. B1 field map synthesis with generative deep learning used in the design of parallel-transmit RF pulses for ultra-high field MRI. *Z Med Phys*. 2022;1–12:334-345.
- 26. Wu Y, Ma Y, Du J, Xing L. Accelerating quantitative MR imaging with the incorporation of B1 compensation using deep learning. *Magn Reson Imaging*. 2020;72:78-86.
- 27. Bring W, Staring M, Remis R, Webb A. Fast Subject-Specific Local SAR and B1+ Prediction for PTx at 7T using only an Initial Localizer Scan. In: Proceedings of the 31th Annual Meet. ISMRM, London, United Kingdom: Abstract 0387; 2022.
- 28. Abbasi-Rad S, O'Brien K, Kelly S, et al. Improving FLAIR SAR efficiency at 7T by adaptive tailoring of adiabatic pulse power through deep learning B1+ estimation. *Magn Reson Med*. 2021;85:2462-2476.
- Plumley A, Watkins L, Treder M, Liebig P, Murphy K, Kopanoglu E. Rigid motion-resolved B1+ prediction using deep learning for real-time parallel-transmission pulse design. *Magn Reson Med*. 2022;87:2254-2270.
- Van de Moortele PF, Ugurbil K. Very fast multi channel B1 calibration at high field in the small flip angle regime. In: Proceedings of the 17th Annual Meeting of ISMRM, Honolulu, USA: Abstract 0367; 2009.
- Hess AT, Bissell MM, Ntusi NAB, et al. Aortic 4D flow: quantification o.f signal-to-noise ratio as a function of field strength and contrast enhancement for 1.5T, 3T, and 7T. Magn Reson Med. 2015;73:1864-1871.
- 32. Schmitter S, Delabarre L, Wu X, et al. Cardiac imaging at 7 tesla: single- and two-spoke radiofrequency pulse design with 16-channel parallel excitation. *Magn Reson Med*. 2013;70:1210-1219.
- 33. Van de Moortele PF, Snyder C, DelaBarre L, Adriany G, Vaughan T, Ugurbil K. Calibration tools for RF shim at very high field with multiple element RF coils: from ultra fast local relative phase to absolute magnitude B1+ mapping. In: Proceedings of the 15th Annual Meeting of ISMRM, Berlin, Germany: Abstract 1676; 2007.
- Ronneberger O, Fischer P, Brox T. U-net: convolutional networks for biomedical image segmentation. In: International Conference on Medical Image Computing and Computer-Assisted Intervention. Vol. 9351. Springer; 2015. pp. 234–241.
- Ye JC, Sung WK. Understanding geometry of encoder-decoder CNNs. In 36th International Conference on Machine Learning ICML 2019 2019;2019-June:12245–12254.
- Küstner T, Fuin N, Hammernik K, et al. CINENet: deep learning-based 3D cardiac CINE MRI reconstruction with multi-coil complex-valued 4D spatio-temporal convolutions. Sci Rep. 2020;10:1-13.

- 37. Kofler A, Dewey M, Schaeffter T, Wald C, Kolbitsch C. Spatio-temporal deep learning-based Undersampling artefact reduction for 2D radial cine MRI with limited training data. *IEEE Trans Med Imaging*. 2020;39:703-717.
- 38. Shih SF, Kafali SG, Calkins KL, Wu HH. Uncertainty-Aware Physics-Driven Deep Learning Network for Fat and R2* Quantification in Self-Gated Free-Breathing Stack-of-Radial MRI. In: Proceedings of the 31th Annual Meeting of ISMRM, London, United Kingdom, Abstract 0433; 2022.
- 39. Cho J, Zhang J, Spincemaille P, et al. QQ-NET—using deep learning to solve quantitative susceptibility mapping and quantitative blood oxygen level dependent magnitude (QSM+qBOLD or QQ) based oxygen extraction fraction (OEF) mapping. *Magn Reson Med.* 2022;87:1583-1594.
- Hu Z, Wang Y, Zhang Z, et al. Distortion correction of single-shot EPI enabled by deep-learning. *NeuroImage*. 2020; 221:117170.
- 41. He K, Zhang X, Ren S, Sun J. Deep residual learning for image recognition. *Proc IEEE Conf Comput Vis Pattern Recognit*. 2016;770-778.
- 42. Abadi M, Agarwal A, Barham P, et al. TensorFlow: large-scale machine learning on heterogeneous distributed systems. *arXiv*. 2016;1603.04467.
- 43. Terpstra ML, Maspero M, Sbrizzi A, van den Berg CAT. ⊥-loss: a symmetric loss function for magnetic resonance imaging reconstruction and image registration with deep learning. *Med Image Anal.* 2022;80:102509.
- 44. You K, Long M, Wang J, Jordan MI. How does learning rate decay help modern neural networks? *arXiv.* 2019; 1908.01878.
- 45. Isola P, Zhu JY, Zhou T, Efros A. Image-to-voxel model translation with conditional adversarial networks. *Proc IEEE Conf Comput Vis Pattern Recognit*. 2017;1125-1134.
- 46. Fischer B, Sedlmeier AM, Hartwig S, et al. Anthropometric measures in the German National Cohort—more than weight and height. *Bundesgesundheitsblatt Gesundheitsforschung Gesundheitsschutz*. 2020;63:290-300.

SUPPORTING INFORMATION

Additional supporting information may be found in the online version of the article at the publisher's website.

FIGURE S1 B_1^+ -magnitude and phase maps for the prediction of the neural network $B_1^+_{PR}$ compared to the ground truth (GT) $B_1^+_{GT}$ for the unseen example 2 with a medium SSIM value. The absolute error ΔB_1^+ between the prediction and GT shows a higher residual error for the first transmission (Tx) channel and the eighth channel as compared to channels 2–7. Overall, the prediction qualitatively matches the GT for both the magnitude and the phase.

FIGURE S2 B_1^+ -magnitude and phase maps for the prediction of the neural network $B_1^+_{PR}$ compared to the ground truth (GT) $B_1^+_{GT}$ for the unseen example 3 with a low SSIM value. The absolute error ΔB_1^+ between the prediction and GT shows a higher residual error for the first transmission (Tx) channel and the eighth channel as

-Magnetic Resonance in Medicine

compared to channels 2–7. Overall, the prediction qualitatively matches the GT for both the magnitude and the phase.

FIGURE S3 Combined B_1^+ maps for the unseen test case example 2 and example 3. The predicted $(B_1^+_{PR})$ and the ground truth $(B_1^+_{GT})$ data are shown for the sum of magnitudes (SOM) over the 8 Tx channels, the magnitude of sum (MOS), and the phase of the summed-up data (POS). When evaluating the ROI over the heart the average error is 4.65% regarding the SOM for example 2 and 1.07% for 3, 4.81% for the MOS for example 2 and 1.92% for 3, as well as a mean difference for the POS of 0.010 rad for example 2 and 0.0124 rad for 3. The absolute error ΔB_1^+ between the prediction and the ground truth, as well as local signal dropouts marked by the yellow arrows, are presented.

FIGURE S4 FA prediction for the measured channel-combined B_1^+ -maps, the reconstructed 2D GRE image, and cine GRE for the unseen test cases 2 and 3.

The used B_1^+ -shims for the default, homogenous, and efficiency settings were calculated on the PR B_1^+ -maps. The 2D cine GRE images for unseen test case 2 have been acquired with a higher FA.

Table S1 RMSE, SSIM and \perp loss values obtained from the evaluation of the five networks applied to the corresponding five test subsets when performing a 5-fold cross-validation using all thorax geometries. Network #5 evaluated on Subset #5 is used for the in vivo application because it results in the highest mean SSIM.

How to cite this article: Krueger F, Aigner CS, Hammernik K, et al. Rapid estimation of 2D relative B₁⁺-maps from localizers in the human heart at 7T using deep learning. *Magn Reson Med.* 2022;1-14. doi: 10.1002/mrm.29510