

Photoacoustic digital brain and deep-learning-assisted image reconstruction

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ABSTRACT

Photoacoustic tomography (PAT) is a newly developed medical imaging modality, which combines the advantages of pure optical imaging and ultrasound imaging, owning both high optical contrast and deep penetration depth. Very recently, PAT is studied in human brain imaging. Nevertheless, while ultrasound waves are passing through the human skull tissues, the strong acoustic attenuation and aberration will happen, which causes photoacoustic signals' distortion. In this work, we use 180 T1 weighted magnetic resonance imaging (MRI) human brain volumes along with the corresponding magnetic resonance angiography (MRA) brain volumes, and segment them to generate the 2D human brain numerical phantoms for PAT. The numerical phantoms contain six kinds of tissues, which are scalp, skull, white matter, gray matter, blood vessel and cerebrospinal fluid. For every numerical phantom, Monte-Carlo based optical simulation is deployed to obtain the photoacoustic initial pressure based on optical properties of human brain. Then, two different k-wave models are used for the skull-involved acoustic simulation, which are fluid media model and viscoelastic media model. The former one only considers longitudinal wave propagation, and the latter model takes shear wave into consideration. Then, the PA sinograms with skull-induced aberration is taken as the input of U-net, and the skull-stripped ones are regarded as the supervision of U-net to train the network. Experimental result shows that the skull's acoustic aberration can be effectively alleviated after U-net correction, achieving conspicuous improvement in quality of PAT human brain images reconstructed from the corrected PA signals, which can clearly show the cerebral artery distribution inside the human skull.

1. Introduction

Cerebral vascular diseases have become one of the most dangerous killers in the world. Especially, stroke has ranked among the top ten pathologies of death [1]. Effective and timely detection of the cerebral vascular stenosis/broken position or oxygen supply can greatly reduce the mortality of patients. Clinically, there are three major diagnosis methods to detect cerebral vascular disease. They are magnetic resonance angiography (MRA) [2], computed tomography angiography (CTA) [3] and digital subtraction angiography (DSA) [4]. Though the

above three methods can reveal the clear vascular structure, they have their own drawbacks and limitations. The biggest drawback of MRA is its low imaging speed, which makes it less useful in diagnosis of acute patients. Although CTA saves much more scanning time, frequent exposing to radiation environment is still a big concern for many people. DSA is the golden standard of cerebral vascular diagnosis in real life. However, it needs high-dose radiography operation to show the perfusion process. The cost of DSA is also much more expensive than the other two, requiring hospitalization for 1–2 days. These limitations intrigue us to find new non-invasive and real-time imaging technologies.

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As a hybrid imaging modality, photoacoustic (PA) imaging combines the advantages of both pure optical imaging (high optical contrast) and ultrasound imaging (deep penetration depth) [5]. Benefited from its vascular detection sensitivity and blood oxygen saturation quantification, PA imaging has become a promising non-invasive medical imaging modality in both preclinical and clinical application in recent decades. From now on, PA imaging has shown great potential in breast tumor classification, PA guided needle tip tracking, skin cancer detection and so on.

Considering that PA imaging plays an outstanding role in revealing vasculature, it seems that PA imaging can be competent in the cerebral vascular diagnosis task. Liming Nie et al. proved the feasibility that using a photon recycler can enable light to pass through the whole adult human skulls for the first time [6]. It starts the new era in human brain PA imaging. However, the aberration from wave reflection and refraction caused by skull tissue makes PA image reconstruction much difficult. In transcranial PA imaging, the targeted objects are cerebral vessels. The acoustic wavelength of the PA signals generated by micro vessels are at the same order as the size of skull pore, so the scattering mode is Mie scattering rather than Rayleigh scattering. In this scattering mode, some side-lobe signals are generated. The superimposition of main lobe and side lobes makes the reconstruction process more complicated [7]. In this regard, more works appear to correct aberration and make a big step towards the clinical PA human brain imaging. Chao Huang et al. proposed to combine PAT with the skull composition and morphology from adjunct X-ray CT modality to correct aberration [8]. This work performs well in both phantoms and monkey head skull-induced distortions. Leila Mohammadi et al. proposed a deterministic ray-tracing simulation framework [9]. Their model contains the attenuation and dispersion effects from wave reflection and refraction. They implement their work on the large 3D phantoms and get pretty good results with high computation speed. Shuai Na et al. proposed to take skull heterogeneity into account to build a three-layer human brain model [10]. With wave reflection and refraction coefficients modified, the proposed layered universal back-projection can correct the acoustic aberration. Then they prove the effectiveness on real ex vivo adult human skull with 64-element PAT system. Another way to solve the problem is model-based image reconstruction methods [11]. The wave propagation in heterogeneous and lossy medium is modeled and discretized to form a forward operator. Mitsuhashi et al. proposed a discrete forward operator and the corresponding adjoint operator for transcranial PACT image reconstruction based on elastic wave equations [12]. Poudel et al. proposed an optimization-based image reconstruction method [13]. The forward-adjoint operator pair is used to compute the regularizing penalty term, which is the least square estimates of the initial pressure distribution. However, this study didn't consider the acoustic heterogeneities within the skull. Poudel et al. further developed the joint reconstruction method that recovers both the initial pressure distribution and the spatial distribution of skull acoustic parameters [14]. With a more accurate estimate of the acoustic properties, the initial pressure distribution with higher quality could be achieved.

All abovementioned works focus on building a mathematical model as accurate as possible to correct the influence caused by skull-distortion. However, to the best of our knowledge, applying data-driven approach to correct the acoustic aberration of human brain is still an untapped area to be explored. In this paper, we propose to build high-quality PA digital brain phantom, and use deep learning network (e.g. U-net [15]) to correct distorted PA signals. We list the contributions of this paper as follows:

- We generate a realistic 2D human brain PA numerical phantom dataset for imaging and vascular disease diagnosis research based on MRA and T1 weighted images in IXI-dataset (<https://brain-development.org/ixi-dataset/>).
- We propose to use simulated PA sinograms obtained from realistic numerical human brain phantoms to train deep learning model to

correct the skull-induced acoustic aberration. With recovered PA signals, the distortion is effectively alleviated in realistic reconstructed PA human brain images.

- We use the generated numerical phantoms to prove the effectiveness of proposed method with good imaging quality.

The remainder of this paper is organized as follow: First, we will introduce some related work in Section 2. Next, details of our work are presented in Section 3. In Section 4, we show the experiment results. Finally, we draw conclusion in Section 5.

2. Related work

2.1. Photoacoustic numerical phantoms

The well-designed PA numerical phantoms are important for performing a simulation study, when lacking of enough experimental data, especially for PA imaging of human. Here we will summarize some PA numerical phantoms generation work and list parts of their database website links.

- (1) Human breast numerical phantoms: Lou et al. proposed to extract four parts (vessel, skin, fat and fibro glandular tissue) from contrast-enhanced magnetic resonance images (MRI) modality with threshold segmentation method [16]. But they only generate a few 2D numerical phantoms from 50 patients. Based on this work, Dantuma et al. proposed to add breast tumor tissue from mice additionally [17]. They also extend the 2D breast numerical phantoms to 3D, which makes simulation more realistic. Then Ma et al. proposed to extract four parts of breast (skin, fat, tumor and fibro glandular tissue) from mammography [18]. The first three tissues are manually segmented and the fibro glandular is segmented by deep neural network. This work can quickly generate large amounts of 2D breast numerical phantoms. But, lacking breast vessels greatly limits the usefulness of the dataset. Later, Han et al. proposed to extract four parts (skin, adipose, tumor and fibro glandular tissue) from a series of 2D US B-scan slices with manual segmentation [19]. The reconstructed 3D breast numerical phantoms are generated, but lacking vessel tissue either. After analyzing all the work above, Bao et al. proposed to generate a more realistic breast numerical phantoms including various kinds of breast types [20]. They use a software (VICTRE breast phantom) to generate a comprehensive breast model including skin, nipple, lactiferous duct, terminal duct lobular unit, interlobular gland tissue, fat, suspensory ligament, muscle, artery, and vein.
- (2) Human skin numerical phantoms: Lu et al. proposed to build two-layer skin model volumes (epidermis and underlying dermis) with random generation [21]. The structure is relatively simple but useful to optical properties analysis. Based on this, Lyu et al. proposed to extract vessel tissue from 3D lung CT scans with Frangi filter segmentation [22]. Then by embedding the vessel tissue to a three-layer cube (epidermis, dermis and hypodermis), they generate 3890 3D skin tissue numerical phantoms, which is available on the website: <https://iee-dataport.org/documents/3d-skin-tissue-vessel-models-medical-image-analysis>.
- (3) Human brain numerical phantoms: Firouzi et al. proposed to generate a numerical brain phantom from MRI and MRA [23]. They segmented six kinds of biological tissue (skull, white matter, gray matter, cerebrospinal fluid, edema, tumor) with 3DSlicer software package from MRI and segmented vascular tissue with level-set algorithm from MRA. It is suitable to investigate the human brain tumor diseases instead of cerebral vascular diseases. Later, Na et al. proposed to generate 2D human brain numerical phantoms by embedding a small piece of vessel model to oval skull models, which are both manually designed [10]. These

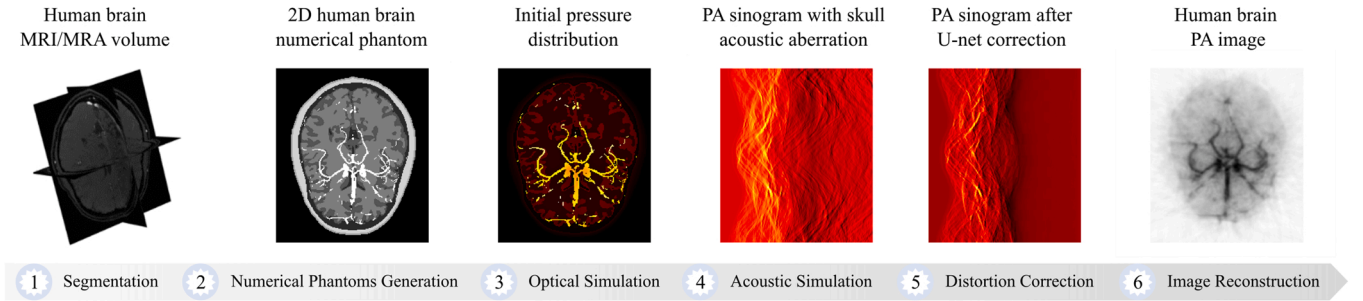


Fig. 1. The whole process of proposed method.

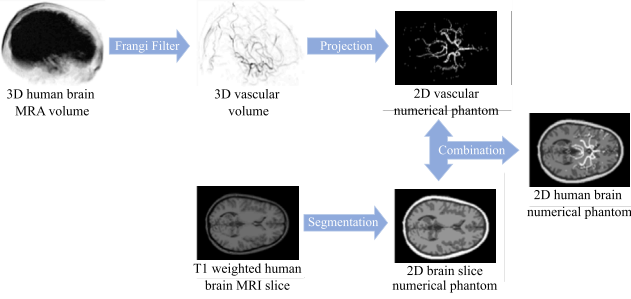


Fig. 2. 2D human brain numerical phantom generation process.

phantoms are relatively simple and discard lots of other brain tissue, like white matter.

2.2. Deep learning-based signal processing

Deep learning has achieved great success in signal processing (e.g. enhancement, denoising and recovery). Gutta et al. proposed to use a standard five layer fully connection neural network to enhance the bandwidth of PA signals [24]. The proposed deep neural network (DNN) can compensate the bandwidth loss caused by ultrasound transducers. With the bandwidth enhanced PA signals, they get the improved reconstructed PA images without significant computation burden. Based on this work, Awasthi et al. proposed to use convolutional neural network (CNN) to enhance the PA signals [25]. Combined with U-net (a well-known CNN structure) and proposed Elu activation function, the method can compensate both PA signal bandwidth and limited-view information loss. The results show that the method works well in PAT system with less root mean square error (RMSE) and higher signal-to-noise ratio (SNR) value. Ben Luijten et al. proposed to use DNN to learn an optimal set of apodization weights, which can adaptively enhance the ultrasound signals [26]. With back projection methods, the proposed deep learning-based adaptive signal processing framework achieves data-efficiency and robustness in ultrasound signal enhancement and image reconstruction. Allman et al. proposed to use CNN to distinguish PA source from reflection artifacts with the sampled PA signals [27]. This work greatly decreases the artifact influence in PA images with high accuracy.

3. Method

In this section, we will make detailed description about the proposed method. It includes five parts: human brain numerical phantoms generation, optical simulation, acoustic simulation, acoustic aberration correction using U-net, and PA image reconstruction. As illustrated in Fig. 1, detailed description of the proposed method will be introduced in following sections.

3.1. PA human brain numerical phantom generation

We propose to generate the human brain numerical phantoms from other medical image modalities. Considering MRA can provide high contrast with rich vessel information, we choose 3D MRA images with T1 weighted MRI volume in IXI dataset to generate the 2D human brain numerical phantoms.

To better simulate realistic human brain, our PA numerical phantom consists of six parts, which are scalp, skull, vessel, gray matter, white matter and cerebrospinal fluid. However, segmenting human brain volumes without labels is a challenging task. In additional, segmentation via deep learning is barely impossible since the amount of data is limited. Consequently, we choose to combine traditional image processing methods with manual segmentation. In MRA, vessels show higher intensity compared with other tissues caused by contrast agent, which makes vessels can be easily segmented with thresholding or filtering methods. After comparing the segmentation results, we choose 3D Frangi filter to segment vessel part [28]. Because the segmentation process of the 3D volume ($150 \times 256 \times 256$) is time consuming, we only choose an upper slice (256×256) with the largest brain area to generate 2D numerical phantoms. Thus, we take a proper axial slice of each T1 weighted MRI volumes and use multiplicative intrinsic component optimization (MICO) algorithm [29] for rough segmentation. Simultaneously, some manual adjustments are implemented on the result obtained from MICO to acquire a more precise segmentation of T1 weighted human brain MRI image. Assuming that the PA signal propagation is also affected by scalp according to the tissues' optical and acoustic properties, we manually add an additional layer of scalp outside the skull. Then we combine the 2D projection of 3D vessel information with other tissue parts to generate the final human brain numerical phantoms. The whole human brain numerical phantom generation process is illustrated in Fig. 2.

Considering light will pass through several layers of different types of tissues from top of the brain, we build pseudo-3D human brain numerical phantoms to make the optical simulation more realistic. Pseudo-3D means the numerical phantom is not generated from the real 3D MRA source. It is spliced with several 2D numerical phantom slices. As shown in Fig. 3, the light will go through the scalp layer (Layer 1: only with scalp), the skull layer (Layer 2: with scalp and skull), the medium layer (Layer 3: with scalp, skull and cerebrospinal fluid) to reach the final layer (Layer 4: complete 2D numerical human brain phantom that involves all 6 kinds of tissues). Layer 4 is the generated human brain numerical phantom illustrated in Fig. 2. Because different tissue has different size, we build the pseudo-3D human brain numerical phantom sizes $256 \times 256 \times 12$ (Two Layer 1 on the top, five Layer 2 under Layer 1, four Layer 3 under Layer 2 and a single Layer 4 at the bottom). Vessel is within the human brain phantom in the layer 4 at the bottom which has 11 layers above it.

3.2. Optical simulation

The optical fluence simulation is deployed using the 3D Monte Carlo

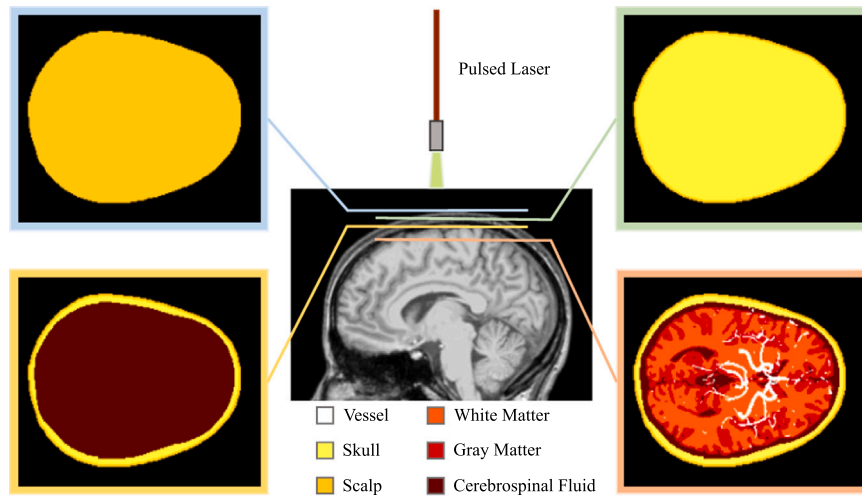


Fig. 3. Four kinds of layers and their positions from the pseudo-3D human brain numerical phantoms.

Table 1

Optical properties of the tissues in human brain numerical phantom [31].

	μ_a (mm ⁻¹)	μ_s (mm ⁻¹)	g	n
White matter	0.014 ± 0.002	91.00 ± 13.65	0.90	1.37
Gray matter	0.036 ± 0.005	22.00 ± 3.30	0.90	1.37
Cerebrospinal fluid	0.004 ± 0.001	2.40 ± 0.36	0.90	1.33
Scalp	0.018 ± 0.003	19.00 ± 2.85	0.90	1.37
Skull	0.016 ± 0.002	16.00 ± 2.40	0.90	1.43
Vessel	0.238 ± 0.036	52.20 ± 7.80	0.99	1.40

method [30]. The optical properties of the six kinds of tissues are shown in Table 1 including optical absorption coefficients (μ_a), reduced scattering coefficients (μ_s), anisotropy (g) and refractive index (n). To more realistically simulate the variation of tissue's optical properties among different human individuals, the value of optical absorption coefficients (μ_a) and reduced scattering coefficients (μ_s) assigned to tissues of different phantoms varies within a certain range. We use the MATLAB package MCXLAB to simulate the photons' propagation inside the human brain numerical phantoms illuminated by the pulsed laser.

Without loss of generality, our optical simulation setup based on 3D Monte Carlo method is as follows: 1) the shape of the light source is set to

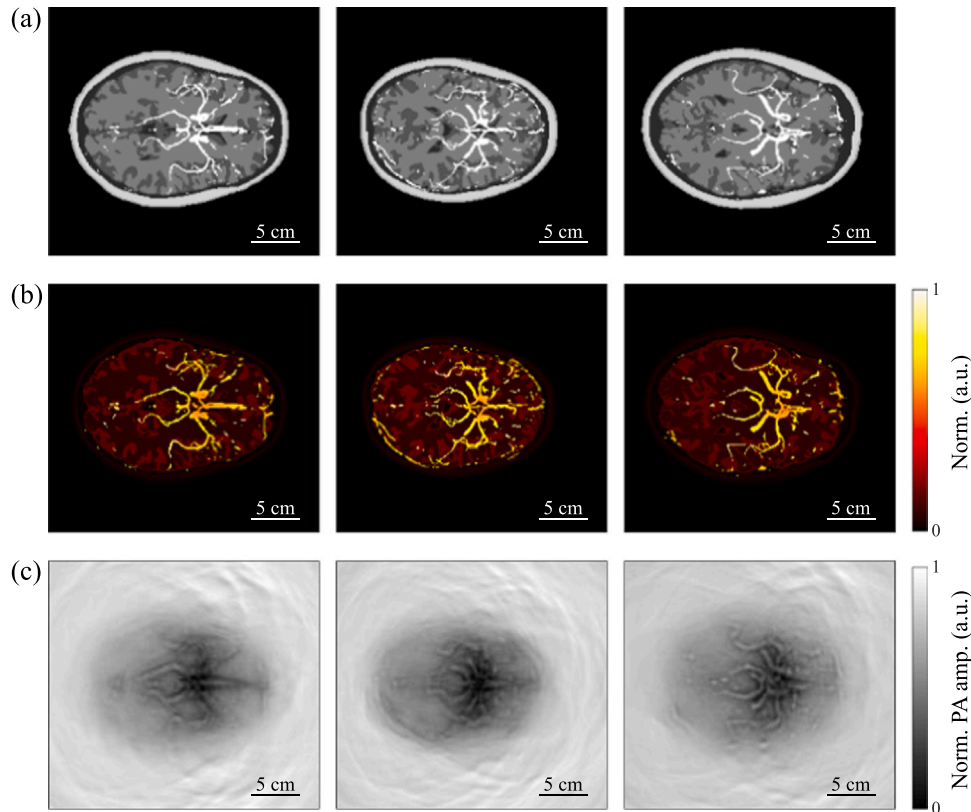


Fig. 4. Three examples of 2D human brain numerical phantoms along with their optical and acoustic simulation results. (a) 2D human brain numerical phantoms generated from different samples. (b) Normalized initial pressure distribution from 3D Monte Carlo optical simulation on phantoms in (a). (c) Normalized DAS Image reconstructions of skull-distorted PA signals obtained from fluid acoustic model using k-wave simulation.

Table 2

Part of acoustic properties of the tissues involved in human brain numerical phantom [34].

	C_{\min} (m/s)	C_{\max} (m/s)	d_{\min} (kg/m ³)	d_{\max} (kg/m ³)
White matter	1532	1573	1040	1043
Gray matter	1500	1500	1039	1050
Cerebrospinal fluid	1502	1507	1007	1007
Scalp	1537	1720	1100	1125
Skull	2190	3360	1800	2100
Vessel	1559	1575	1056	1147

be planar; 2) the total number of photons to be simulated is 1 billion; 3) the position and incident vector of the source is set properly to make sure that light will cover all the regions of interest; 4) the starting time and the ending time of the simulation is set to 0 and 1 nanoseconds, respectively, and the time-gate width of the simulation is set to 0.01 nanoseconds. According to Table 1, the optical fluence through the six kinds of human brain tissues is computed and obtained given the pulsed laser excitation. In the Monte Carlo optical simulation, the vertical thickness of each layer in the generated pseudo-3D human brain numerical phantoms is 1 mm, which means the total vertical thickness of a single pseudo-3D numerical phantom is 12 mm.

This optical simulation is run on a Linux server with 4 GTX 1080Ti GPUs. For each human brain numerical phantom, the photon

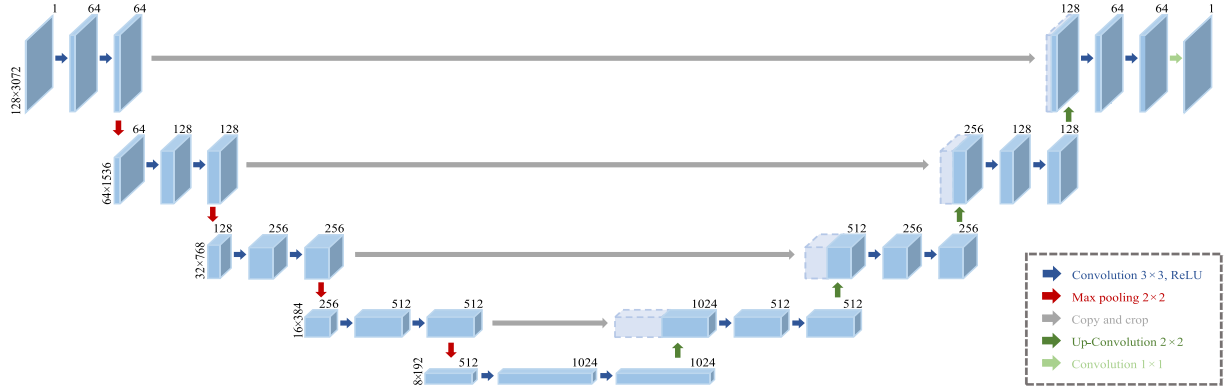
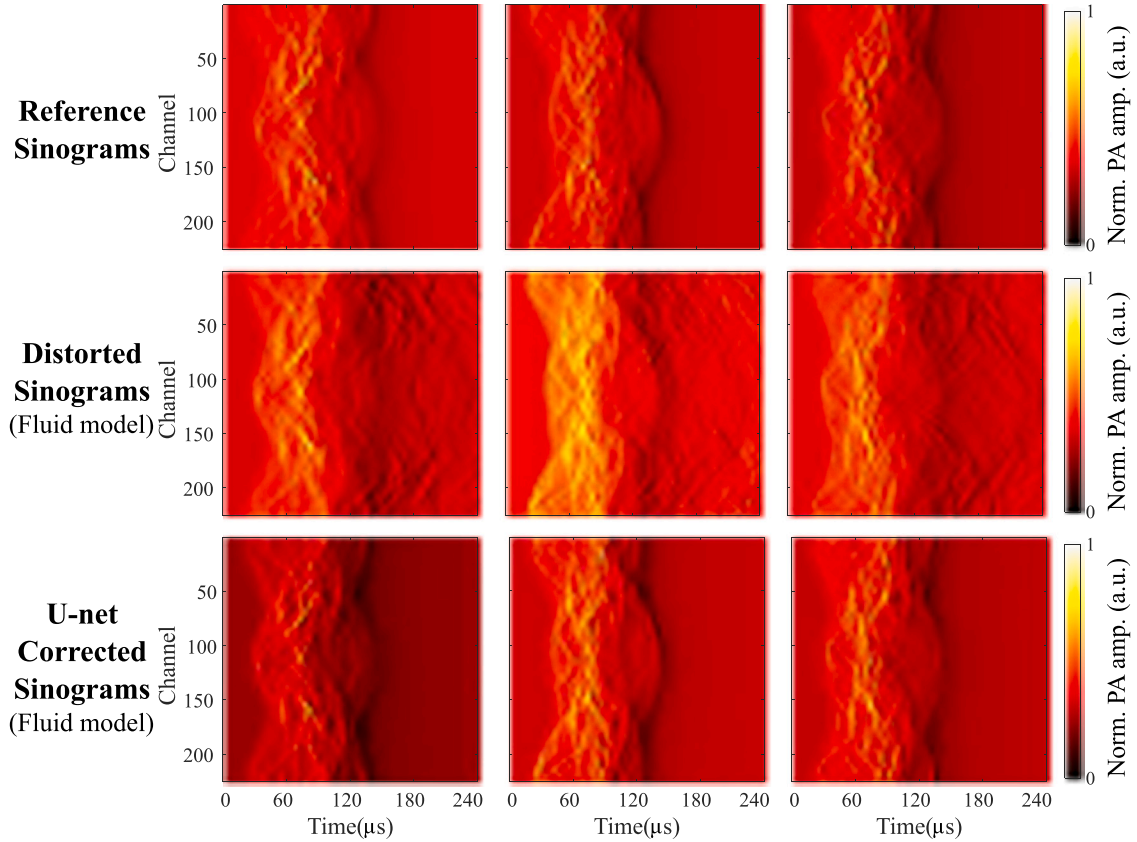
**Fig. 5.** The structure of U-net used in this paper.

Fig. 6. Three examples of normalized PA sinogram in acoustic aberration correction process. The top row shows the PA sinograms which are used as supervision during U-net training. The second row are the inputs of U-net that are PA sinograms with acoustic aberration from skull. The U-net corrected PA sonograms are at the bottom row.

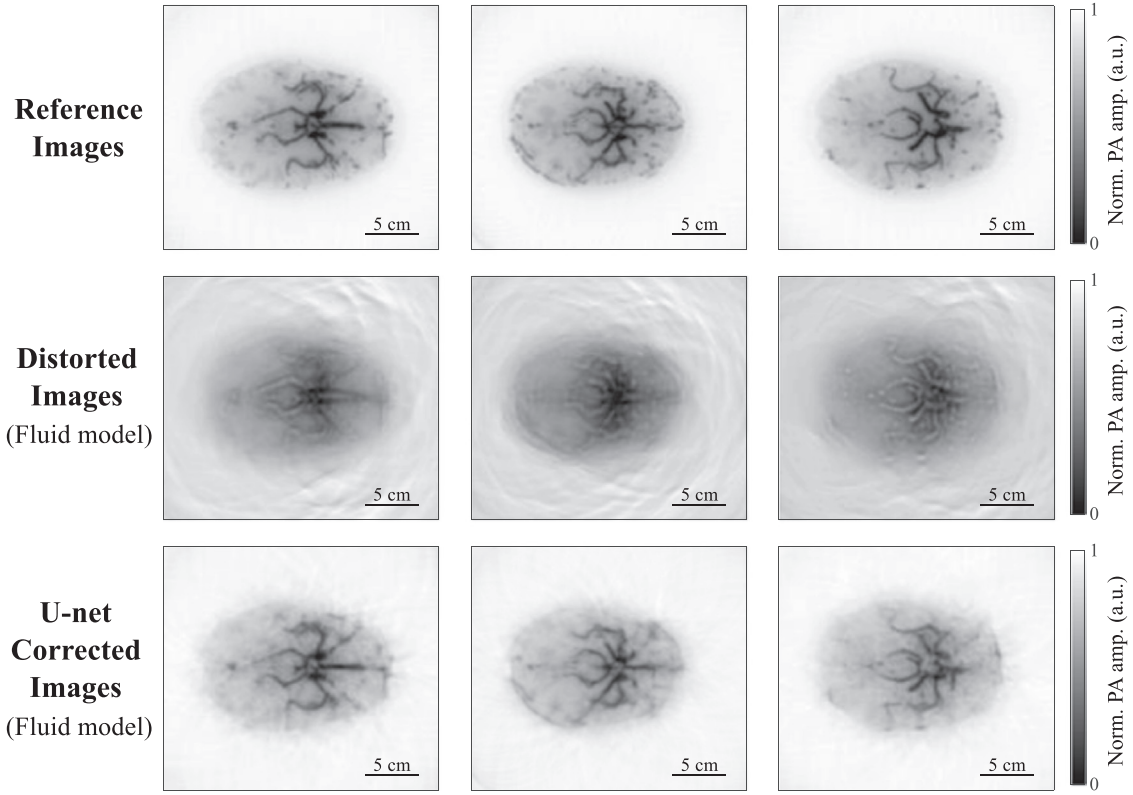


Fig. 7. Corresponding normalized human brain images reconstructed from PA sinograms with delay-and-sum (DAS) algorithms.

Table 3

Quantitative comparison of results from fluid media acoustic model with three different evaluation indices.

	SSIM	PSNR	MAE
Original (Normalized PA sinogram)	0.526	19.871	0.083
U-net corrected (Normalized PA sinogram)	0.832	25.638	0.044
Original (Normalized PA image)	0.718	24.099	0.052
U-net corrected (Normalized PA image)	0.939	33.424	0.012

Table 4

Quantitative comparison of results from viscoelastic media acoustic model with three different evaluation indices.

	SSIM	PSNR	MAE
Original (Normalized PA sinogram)	0.499	21.058	0.066
U-net corrected (Normalized PA sinogram)	0.805	25.015	0.045
Original (Normalized PA image)	0.732	25.315	0.042
U-net corrected (Normalized PA image)	0.921	31.950	0.014

propagation simulation process takes about 5 min to obtain an optical fluence volume. Only the optical fluence map of obtained optical fluence volumes' 12th slice is needed for the acoustic simulation, since our human brain numerical phantoms are built to be pseudo 3D. Through multiplying the obtained optical fluence maps by the corresponding optical absorption coefficients (μ_a), we can get the initial acoustic pressure distribution which is used for the following acoustic simulation. Fig. 4(b) shows three images of normalized initial pressure distribution

of the corresponding 2D numerical phantoms in Fig. 4(a).

3.3. Acoustic simulation

The acoustic simulation is conducted through the k-Wave toolbox developed by Treeby et al. [32]. Both fluid media acoustic model and viscoelastic media acoustic model are considered in this simulation. It deserves noting that shear waves is involved in viscoelastic acoustic simulation with Kelvin-Voigt model of viscoelasticity [33]. The parameters of k-Wave computational grid is set as follows: 1) the PML size is 17; 2) the number of grid points in the x direction (N_x) and the y direction (N_y) is 256 and 256 respectively; 3) the total grid size in the x direction (x) and y direction (y) is both 256 mm; 4) the grid point spacing in the x direction (dx) and y direction (dy) are both 1 mm.

The maximum and minimum speed of longitudinal acoustic waves (c), as well as the maximum and minimum medium density (d) of the different tissues in human brain numerical phantom, are presented in Table 2. Similar to the optical simulation, we let the speed of sound and density of tissues in different human brain numerical phantoms be the random values, which are between the maximum and minimum values for simulating the inter-individual variation of tissues in acoustic properties. In viscoelastic media acoustic model, speed of shear waves in skull is set to be between 1360 m/s and 1640 m/s [35], compressional and shear acoustic attenuation coefficients of skull are set to be 7.75 dB/(MHz² cm) and 16.70 dB/(MHz² cm) [36]. We use point-like ultrasound transducers with infinite bandwidth that minimize the measurement errors in the simulation, which help us to focus on the de-aberration of skull's acoustic heterogeneity. The 256-elements ring array has a radius of 125 mm, and the brain phantom is placed in the center of the ring array.

With 4 GTX 1080Ti GPUs, it takes about 0.3 s for fluid media acoustic simulation of each PA human brain numerical phantom, while viscoelastic media acoustic simulation will take about 1 min for each phantom. After obtaining the sensor data, we use the delay-and-sum

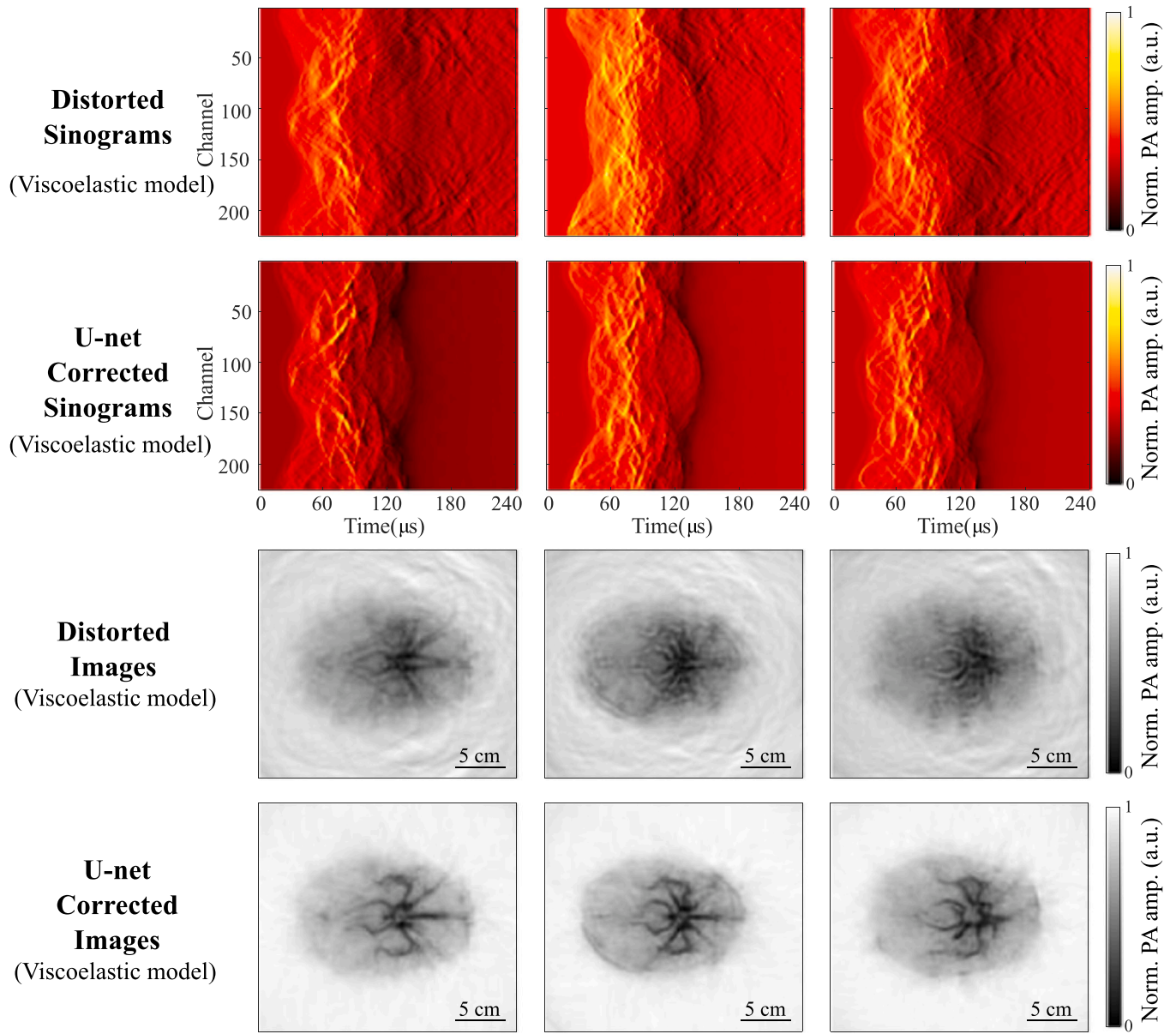


Fig. 8. Normalized PA sinograms and normalized DAS reconstructed human brain PAT images from viscoelastic media acoustic model.

(DAS) algorithm to reconstruct the original image. Fig. 4(c) shows the normalized images reconstructed from acoustic simulation results of fluid media model, in which we could clearly see that the reconstructed image is distorted due to the acoustic aberration of the skull tissue. To acquire the ideal distortion free PA sinograms as supervision term for deep learning-based algorithm, we manually remove the skull and scalp of the numerical human brain phantoms through setting the acoustic properties of these two tissues to be the same as the background, and then repeat the acoustic simulation process with fluid media model. Then, we can obtain the pseudo-ideal sensor data as the reference PA sinograms in which PA signals are free of acoustic aberration caused by skull and scalp. The skull-aberrated signals of two different acoustic simulation models are regarded as the raw data, which are used along with the reference PA sinograms for the following acoustic aberration correction study.

3.4. Acoustic aberration correction using U-Net

With the obtained PA sinogram from the acoustic simulation, we propose to use deep neural networks to correct the skull acoustic

aberration, which is illustrated in Fig. 1 (step 5).

First, we have to pre-process the data and build training and testing datasets. With the acoustic simulation, we can get the paired data (PA sinogram with acoustic aberration and PA sinogram without acoustic aberration). To keep the consistency of dimension after each pooling operation, we manually extend the PA sinogram in training set to 256×3072 by adding zeros at the end of each channel signal.

In this work, we choose U-net for this task for three reasons. First, the input sinogram is similar in the first half of the data with outputs, and the end of the input sinogram is unwanted reflection aberration signals. So, residual structure is suitable to this work. Interestingly, U-net can be regarded as a big residual structure in some way. Second, the special skip-connection can extract multi-scale features to improve correction outputs. Third, compared with other CNN structures, such as AlexNet, U-net needs less training samples and can process images with less training time and computation cost. U-net structure used in this work is illustrated in Fig. 5.

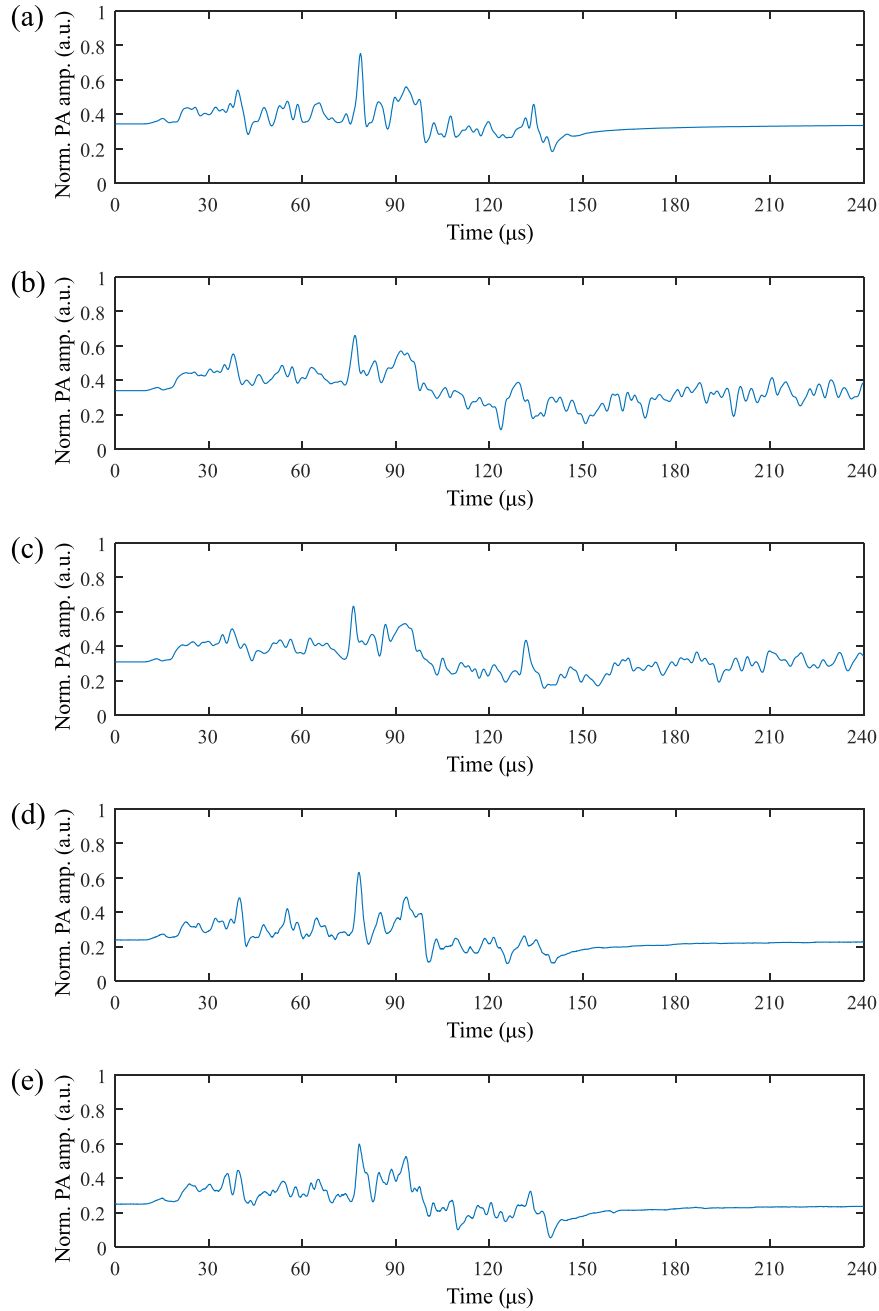


Fig. 9. Normalized PA signals taken from the first channel of the right-column sample shown in Fig. 4(a). (a) Reference PA signal. (b) PA signal with skull aberration obtained from fluid media acoustic model. (c) PA signal with skull aberration obtained from viscoelastic media acoustic model. (d) PA signal in (b) after U-net correction. (e) PA signal in (c) after U-net correction.

3.5. PA image reconstruction

The PA sonograms (with acoustic aberration correction) are used to reconstruct PA images. Because the U-net has eliminated the skull impact, we can regard the sensor data is generated from homogeneous medium. Then we can apply a lot of classical reconstruction algorithms, such as delay-and-sum, time reversal, model-based reconstruction and so on. In order to guarantee the reconstruction speed, we choose DAS as the PA image reconstruction algorithm in this work.

4. Experiment and results

We choose a subset of IXI dataset for experiment that is from Hamersmith Hospital (HH), which are used to generate 180 sets of human

brain PA sonograms with size of 256×3000 . The obtained sonograms are randomly divided into 120 training sets and 60 testing sets. Since the sensors in acoustic simulation is set to be circular and data in each channel of the obtained PA sonograms is from the corresponding sensor, we uniformly separate each extended PA sinogram in the training set into 8 segments with 32 channels each based on sensor positions. Then we sequentially choose 16 channels from each segment and concatenate the chosen data together to get a 128×3072 sub-sinogram. By overlapping channels in each segment, we can have 16 combinations of sub-sinogram from an original extended PA sinogram. The data augmentation eventually gives 1920 PA sinograms (128×3072) as training set. PA signals obtained from fluid media acoustic model and viscoelastic media acoustic model are trained separately but with the same U-net structure and settings. The iteration is set to 400 epochs and batch size is

set to 16. We choose ADAM optimizer and set the learning rate to 0.001. Mean square error is used as the loss in U-net training process. The training process takes 5 min per epoch and is conducted on 4 NVIDIA TITAN RTX GPU with 24 GB of RAM each. Testing process is conducted on a single NVIDIA TITAN RTX GPU and takes about 0.3 s for correcting one PA sinogram.

The normalized fluid media acoustic model's testing results of three samples shown in Fig. 4(a) are presented in Fig. 6. We could see that the corrected sensor data is much more similar to the reference PA signals. Since we want to make the corrections that are based on the physical features of the PA signals rather than the texture feature of images, the U-net used in this paper is trained with PA sinograms instead of the reconstructed PA images. In order to figure out whether the imaging quality is also improved with U-net correction, we use DAS to reconstruct the image from the PA sinograms presented in Fig. 6. Fig. 7 shows the reconstructed gray-scale human brain PA images with normalization. Compared with the reconstructed images without correction, we can clearly see the main cerebral vessels' shape and the cerebrovascular distribution from the reconstructed images of the U-net corrected results.

To better demonstrate the improvement on signal and image quality, we use structural similarity (SSIM), peak signal-to-noise ratio (PSNR) and mean absolute error (MAE) of the normalized PA sinograms and normalized grayscale DAS reconstructed PA images for quantitative evaluation. As shown in Table 3, three indices are greatly improved in comparison of both PA sinogram and reconstructed image. Furthermore, the largest improvement is on the structural similarity, which means U-net can effectively help identify the anatomical structure of main cerebral vessel without removing the skull.

For results of viscoelastic media model, the quantitative evaluation is presented in Table 4. Similar to the results of fluid media model, indices of both PA sinograms and PAT reconstruction images are conspicuously improved after the correction with U-net. The corresponding normalized PA sinograms and normalized PAT images reconstructed with DAS are illustrated in Fig. 8. Five normalized PA signals taken from the first channel of PA sinograms acquired from the rightmost object in Fig. 4(a) in different status are shown in Fig. 9, in which we can see that U-net filters out the redundant part of the PA signals and corrects the signal waveform, while the remained PA signals are still slightly distorted compared with the reference PA signal.

5. Discussion and conclusion

Previously reported model-based inversion can iteratively recover the optical absorption distribution by overcoming the acoustic aberration, which however is time-consuming and sometimes suffers convergence issue. Deep learning based method can be quite fast when it is well trained, which is more suitable for real-time imaging applications.

In this paper, we built PA digital brain dataset, and proposed to correct the human skull acoustic aberration through deep learning method for human brain PAT imaging. We designed and made 180 human brain numerical phantoms generated from MRA and T1 weighted MRI modality. Optical and acoustic simulations were performed on our human brain numerical phantoms to simulate the PA signal generation process. Based on pseudo-3D human brain numerical phantoms manually generated from 2D numerical phantoms, 3D Monte Carlo optical simulation was used to acquire the optical fluence maps for calculating the initial pressure distribution used in the acoustic simulation. In the acoustic simulation part, we designed two experiments to simulate the PA signals propagation process in human brain with and without distortion of skull. During the simulation with skull, we use heterogeneous acoustic properties of tissues for obtaining the sensor data with skull aberration. Moreover, both fluid media model and viscoelastic media model are taken into consideration. Then, the augmented PA sinograms with skull aberration were used as input for training the U-net model that is supervised with reference PA data

acquired in the skull-free acoustic simulation. The result shows that the deep learning assisted method can effectively improve the quality of reconstructed PAT images and correct the acoustic aberration caused by human skull with and without taking the propagation of shear waves into consideration.

Compared with methods based on physical model, deep learning assisted methods are with high-efficiency. However, there still exist some limitations when facing the clinical situation such as lack of high quality data, poor interpretability and so on. In the future work, we will continue exploring deep learning assisted methods that are used to improve the PAT image quality and optimize the experiment designs to make it better mimic realistic circumstances. In addition, our further works may be also validated by ex vivo experiments using a monkey (*Macaca fascicularis*)'s skull, and in vivo experiment in monkey and human brain in the long term.

Declaration of Competing Interest

The authors declare no conflicts of interest.

Data Availability

Data will be made available on request.

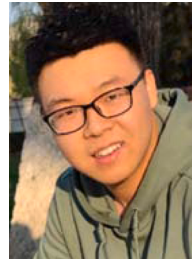
Acknowledgments

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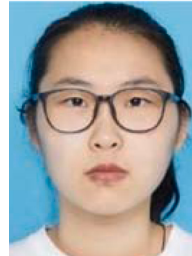
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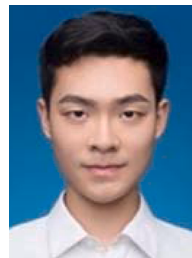
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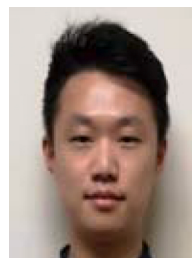
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