Brain Injury Localization in Electromagnetic Imaging using Symmetric Crossing Lines Method

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Abstract—To avoid death or disability, patients with brain injury should undertake a diagnosis at the earliest time and accept frequent monitoring after starting any medical intervention. This paper presents a novel approach to localize brain injury using the intersection of pairs of signals from symmetrical antennas based on the hypothesis that healthy brains are approximately symmetric that the bleeding targets will lead to significantly different amplitude and phase changes if a pair of symmetrical transmit signals cross targets with the same distances. The scattered signals (S-parameters) across the band 0.5-2 GHz are acquired using 100 realistic brain models and 150 experimental data measurements. Each signal is divided into two frequency bands and then they are converted into two graphs. The neighbour pair signals from symmetric hemisphere are evaluated to detect and localize the potential abnormalities in the brain. The higher weighted pairs across the axes of symmetry are detected and used to localize any abnormal targets. The results indicate that crossing pairs of antenna signals from the hemisphere with a blood mass exhibit significantly different signal amplitude in the graph features compared to those without the target (p<0.003). The experiments show that our novel localization algorithm can achieve an accuracy of 0.85 ± 0.08 Dice similarity coefficient based on 150 experimental measurements using an elliptical container, which is suitable for brain injury localization.

Index Terms—Brain injury, electromagnetic imaging, complex networks, symmetrical crossing lines, differential strengths.

I. INTRODUCTION

ELECTROMAGNETIC (EM) imaging technologies have been attracting widespread interest from researchers around the world due to their potential as a portable, low-cost and non-ionizing medical diagnostic system [1]–[5]. This technique has recently been implemented for different medical applications, such as the detection of stroke [1], [3], breast cancer [5], [6] and skin cancer [7]. Two important applications are the detection of breast cancer [8] or traumatic brain injury (TBI) at an early stage. TBI is a major cause of mortality and disability. Intracranial haemorrhage (ICH) is the most important complication of TBI. A mass lesion refers to an area of localized brain injury that must be removed at the earliest time possible for reducing the mortality rate. In most scenarios, people suffering from TBI are not administered timely treatment because the symptoms, such as decreased level of consciousness, cannot always be observed in the early stages. Clinical experiences indicate that the size and location of ICH are critical for clinicians in an operation, therefore

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a fast and accurate method for ICH localization and size estimation is of prominent importance.

Currently, most studies based on EM imaging aim to identify the types of tumors, stroke or intracranial bleedings (IBs) [1], [2], [8]–[11]. The state-of-the-art methods for brain injury localization can be categorized into three types: radar-based, tomography, and machine learning based methods. The radar-based methods and relevant results were reported in [4], [12], [13]. Tomography-based methods are capable of mapping the dielectric properties of brain tissues, including the abnormal ones, hence providing the localization of brain bleedings [3], [14]. In general, these methods need to solve complex inverse problem to identify the target from the solved dielectric distribution [15]. Furthermore, it was indicated in [16] that when certain a-priori information was used, such as the patient's magnetic resonance imaging (MRI) scan, results from tomography approaches could obtain 0.24 relative error on 35 subjects. Machine learning-based methods [2], [17]–[19] can avoid the computationally expensive solving of the inverse problem. However, these methods are more suitable for stroke sub-type classification [2], [18]. Apart from stroke classification, the studies in [17], [19] also attempted to localize the stroke position using support vector machines (SVMs) or the artificial fish swarm algorithm. Unfortunately, both methods require a huge amount of data for training even with a single phantom or simulated brain model. These requirements are usually difficult to satisfy in clinical scenarios [2]. In fact, while these three approaches have their own merits, they also have drawbacks. For example, the tomography-based methods are ill-posed and may not converge to the correct solution, in addition to taking long computational time. The radar-based methods often fail to correctly localize and shape the target, particularly in the case of heterogeneous tissues [20]. EM imaging using machine learning methods is still not accurate in localizing brain injury without large amounts of training data. These limitations restrict practical utilization of portable electromagnetic systems, especially in emergency applications. In addition, all these methods require precise and permanent positioning of the antennas [21]. This requirement could degrade the performance of these methods during practical applications.

This paper presents a novel symmetrical crossing line algorithm (SCLA) to localize the position and predict the size of a TBI. This is based on the hypothesis that a healthy brain is nearly symmetric with respect to the left-right axes [22], [23]. The assumption gives rise to the potential that anomalies can be detected based on the significant difference in symmetric
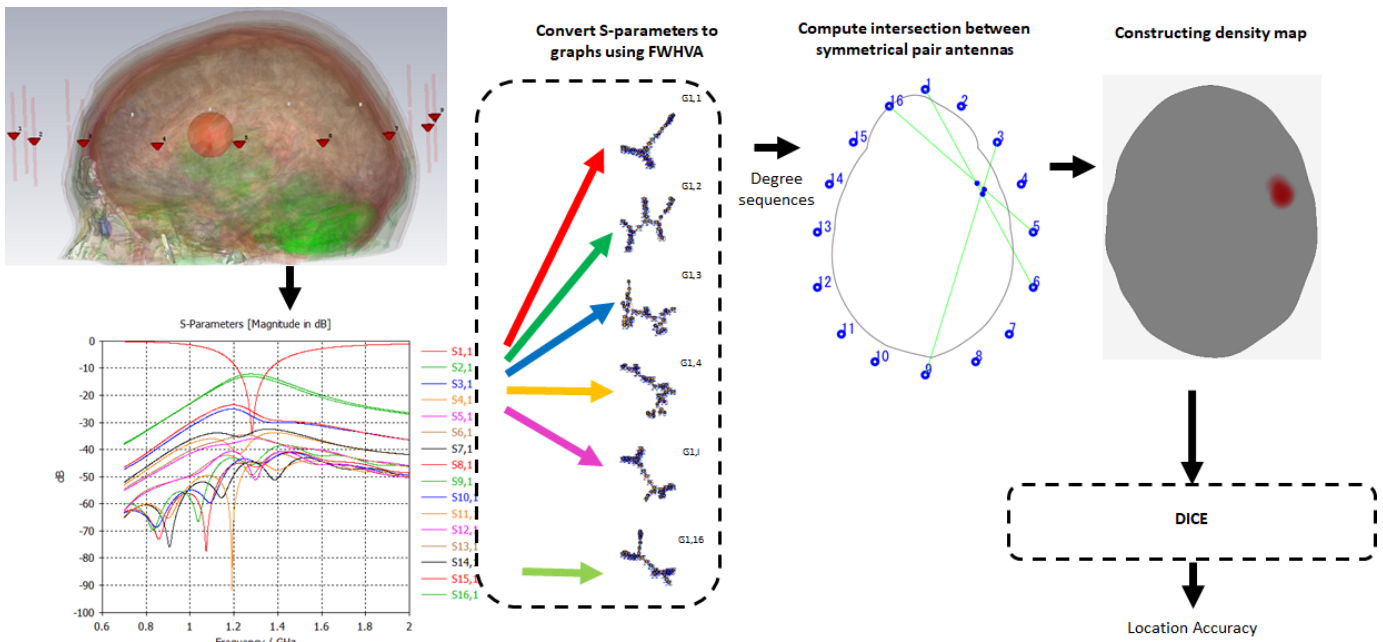
Fig. 1: Flowchart of the proposed localization procedure.

pairs of signals across this axis. The inputs to the algorithm are the S-parameters collected from an antenna array surrounding the head as well as the approximate positions of the antenna ports. The array used to test the technique is composed of 16 antennas which are approximately equally spaced around the head. only transmitted signal are utilized as the transmitters, whereas all reflected signals are ignored. To avoid the influence from amplitude variations, the signal measured from each antenna is converted into a weighted graph. Electromagnetic waves transmitted between two antennas whose signal path pass through bleeding area will have a significantly different weight of the graph features compared with signals in the mirrored antenna pair in the other hemisphere. Based on this idea, the IB can be localized by detecting the intersection of two transmitting lines with the highest weighting coefficient. A size constructing algorithm (SCA) helps to estimate the target centre and the size from three crossing lines. In total, 100 simulated data and 150 experimental data are used to evaluate the algorithm.

II. EXPERIMENTAL DATA AND METHODOLOGY

Fig 1 shows the work process adopted in this study, which can be summarized as follows:

- A patient is monitored by an EM imaging device with n antennas after a suspected traumatic brain injury
- The scattered data ($S_{i,j}$, where $i,j \in [1 \ldots n]$) is collected by an antenna array
- An inverse FFT is applied to convert these signals to $n \times n$ time series signals, and each time series is transformed into a graph using the Fast Weighted Horizontal Visibility Algorithm (FWHVA) [24].
- Each symmetric pair of signals is compared to determine mismatches, which are then used to localize the bleeding based on the intersection of the direct signal paths:

  - The centre of the bleed is obtained by a symmetric crossing line algorithm (SCLA).
  - The size of the target is estimated using a size constructing algorithm (SCA) based on the SCLA output.
- The Dice similarity coefficient (DICE) is computed to calculate the localization accuracy.

A. Brain models and simulation set-up

This study investigates the proposed SCLA algorithm by evaluating the simulation results of eight realistic brain models. The brain models used in this study include detailed Zubal [25] and Duke [26] phantoms that were derived from multiple imaging modalities. In addition, 6 other brain models (named P103111 to P111716), which were built from segmented MRI images [27], are utilized. Each model includes eight tissues (blood, skin, skull, CSF, grey matter, white matter, cerebellum, and ventricles), The details of the phantoms are listed in Table I. The number of voxels indicate the dimensions of the phantoms e.g. $280 \times 330 \times 280$ shows that the X, Y slices are 280×330 pixels, and the 280 indicates the number of slices in the Z-direction.

The proposed SCLA algorithm is evaluated using the electromagnetic simulator, CST Microwave Studio. To build phantoms with intracranial bleeding from the healthy models of Table I, each healthy model was injected with an elliptical shape target (blood), which emulates a TBI, for which each axis was randomly assigned a length of 1 mm to 10 mm and the position was randomly distributed in the X and Y plane limited from 76 to 152 for Zubal and 76 to 218 for others respectively. Z-direction slices ranging from 100 to 230 were used.

TABLE I: Details of three type of models

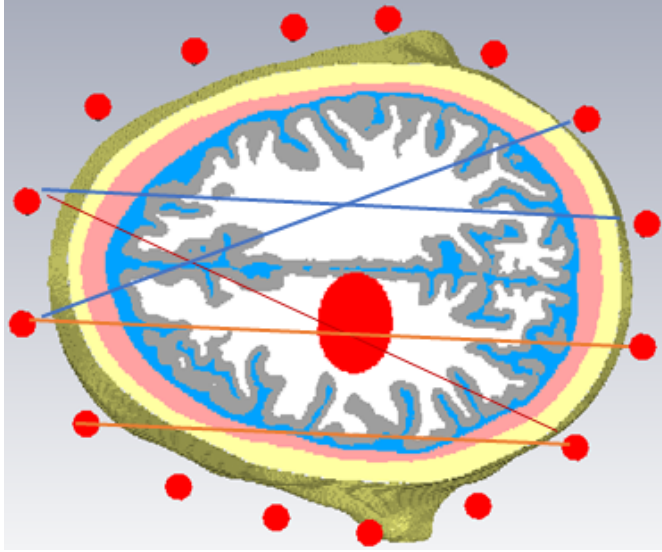| Phantom IDs	Number of tissues	Resolution (mm)	Number of voxels
Zubal [25]	58	1.1×1.1×1.4	256×256×128
Duke [26]	52	1.0×1.0×1.0	610×310×161
P103111-P111716	8	0.7×0.7×0.7	280×330×280

Fig. 2: A brain surrounded by a 16-element electromagnetic-imaging antenna array showing the direct signal paths between antenna pairs. The blood mass is indicated as a red ellipsoid inside the brain.

Fig. 3: The phase of pair antennas $1 - 8$ and $1 - 10$ of (and their mirror $16 - 9$ and $16 - 7$). The difference of pair signals crossing the bleeding are significantly larger than those that do not cross the bleeding both in low and high frequency bands.

B. Hypothesis

The hypothesis of this paper is that a brain containing an intracranial bleed will have an imbalance between symmetric pairs of signals, based on the assumption that a healthy brain is approximately symmetric [22], [23]. As electromagnetic waves pass through a domain, the signals are altered in amplitude and phase owing to different dielectric properties of the media that the signals pass through. Thus, when signals arrive at two antennas with the same distances, the amplitudes or phases will be significantly different if only one of them passes through a high contrast bleeding. In additional, when size of IB is large, both high frequency and lower frequency have the same trends compared with left or right hemisphere. Conversely, only high frequency has the significant changes if the size of IB is small.

Fig. 2 shows a 16-antenna array with antennas positioned symmetrically around the left-right axis of the head. The locations of these antennas are provided in supplementary Table 1. We denote $o_{10,1}$ as the signal travelling from antenna 1 to antenna 10 where the former antenna is used as the transmitter and the latter antenna is used as the receiver. Due to reciprocity, $o_{10,1} = o_{1,10}$, Based on the symmetry hypothesis, it can be anticipated that $o_{10,1}$ will be significantly different from $o_{16,7}$ due to the presence of the IB. Similarly, the similarity of $o_{16,9}$ and $o_{15,10}$ should have a lower difference
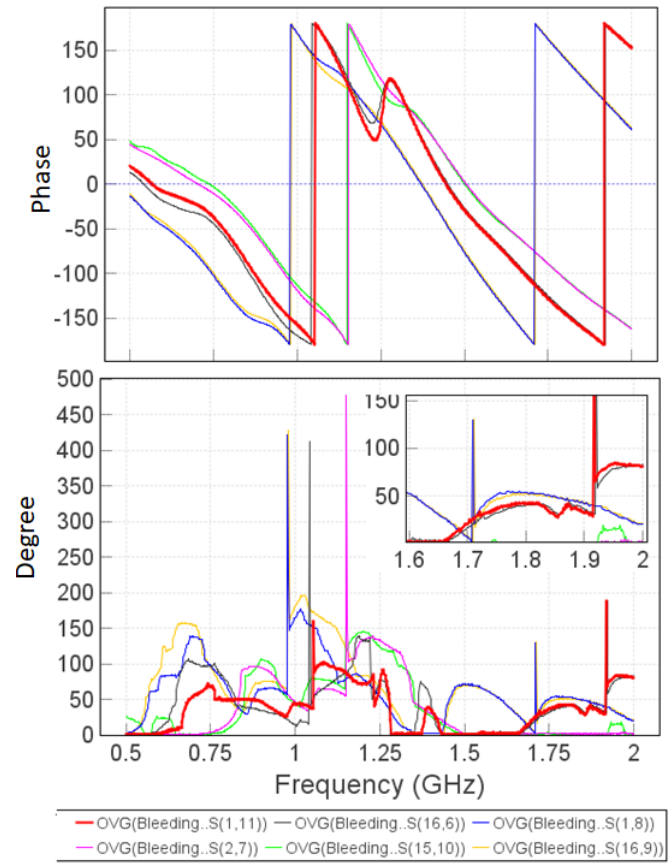
than opposite of the neighbouring antenna pairs ($o_{8,1}$ and $o_{7,2}$).

To verify this hypothesis, the Phase and the corresponding graph features from the pairs antenna $1 - 8$, $2 - 7$, and $1 - 10$ and their mirror pairs antenna $16 - 9$, $15 - 10$, and $16 - 7$ are plotted in Fig. 3. It is easy to see that the phase of opposite pair antennas (such as $1, 8$ and $16, 9$) are similar. On the other had, the graph feature of opposite pairs (such as $2, 7$ and $15, 10$) shows difference in both low frequency (0.5G-1.2G) and high frequency (about 1.92G). In addition, the opposite signals $1, 11$ and $16, 6$ shows a significant difference about $1G$. Thus, bleeding localization can be achieved by identifying the intersection of crossing pair lines (direct signal paths), such as $1, 10$ and $16, 9$ which are both affected by the blood mass in the example shown.

C. Normalizing electromagnetic signals using graph features

Regarding the example brain in Fig. 2, the signals from different antennas have a different amplitude and phase. More importantly, different head sizes also affect these parameters. To normalize the amplitude and align the different phases, a fast weighted horizontal visibility algorithm (FWHVA) [24] is employed to convert each time series into a weighted horizontal visibility graph (WHVGs). WHVGs have been widely used in biomedical signal processing, such as EEG
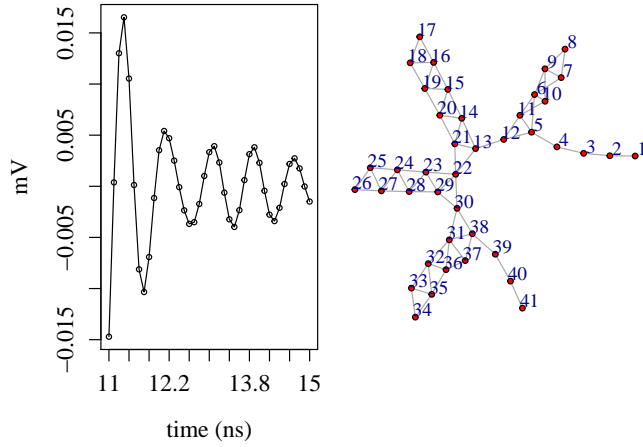
Fig. 4: (a) 41 points of transmission signals from the $O_{1,2}$ from 1.1ns to 15ns, and (b) A HVG mapped from the signal in (a).

[28], [29], EMG [30] and Pulse wave [31]. This paper applies it to electromagnetic signal processing.

From a mathematical point of view, an electromagnetic signal time series, $X_j(j = 1, \ldots, t)$, is mapped to a graph $G(V, E)$ by the following steps: a data point x_j is mapped into a node v_j in G; an edge e_{jk} between (v_j, v_k) from two points $(x_j; x_k)$ exists if and only if:

$$e_{jk} = \begin{cases} 1, & x_k > max(x_{(j+1)\ldots(k-1)}) \\ 1, & j+1 = k \\ 0, & otherwise. \end{cases} \quad (1)$$

where $e_{jk} = 0$ implies that the edge does not exist, otherwise it does.

Fig. 4 shows how signals from a real system between 11 ns to 15 ns are converted into a horizontal visibility graph (HVG). In Fig. 4 (a), a portion of the signal received by the first antenna and sent by the second antenna is shown. In this plot, the 5th point can horizontally see the 12th point but cannot horizontally see the 13th point because the 12th point blocks the visibility between 5th point and the latter. It can be seen that the signals decay rapidly in the time domain as shown in Fig. 4 (a). However, HVG can effectively detect the sine wave and abnormal signals.

In a complex network such as a horizontal visibility graph (HVG), node degree and degree sequence are two basic characteristics which can be used to describe the graph. The degree $d(v_i)$ of node v_i is the number of connected edges from v_i. As shown in Fig 4(b), most of the nodes have degree two, such as nodes 2, 4, 6 and so on, which represents the minimal values of Fig 4(a), although those minimal values are different.

D. Network topology

A time series can be characterized by the network topology, using measures such as the mean degree, clustering coefficient,

average path etc. [32], [33]. In addition to the node degree, the average degree $\bar{d}$ of a graph G with n nodes is defined as:

$$\bar{d} = \frac{1}{n} \sum_{i=1}^{n} d_i \quad (2)$$

A degree sequence (DS) is a sequence of the graph degrees concatenated in order of node number. This concepts has been used for stroke classification [34]. The similarity of two graphs G_x and G_y can be measured from two degree sequences DS_x amd DS_y

$$\bar{S}_j = \frac{1}{n} \sum_{k=1}^{n} (DS_x(i, j) - DS_x(i, j))/(DS_x(i, j) + DS_x(i, j)) \quad (3)$$

It is obviously that the range of equation 3 is in $[-1, 1]$ which could guarantee that the output can be comparable even through the amplitudes of two signals are significantly different.

III. SCLA ALGORITHM

Based on the hypothesis that a brain containing intracranial bleeding will have a significant difference in the measured signals of symmetric pairs of antennas, we can locate the bleeding based on which pairs exhibit large differences. Fig. 2 demonstrates that two pairs of signals cross a blood mass. However, directly using time series will be affected by noise and also the amplitude will be different in different head sizes. To avoid the challenges of time series analysis, we use graph degree and similarity to identify the blood location. The SCLA algorithm is divided into three steps: mapping the time series into a graph, identifying the correct quadrant of the bleeding location as in the left or right hemisphere, front or back, and picking candidate intersection points of antennas pairs from the graph. The first step is based on FWHVA. The next two steps are described in the following sections.

Firstly, we define a differential opposite pairs (DOP) and differential neighbour pairs (DNP) associated with two antennas as equation 4 and equation 5, respectively.

$$DOP_{ij} = |(S_{ij} - cal_{ij}) - Opp(S_{ij} - cal_{ij})| \quad (4)$$

$$DNP_{ij} = |(S_{ij} - cal_{ij}) - (S_{i+1,j-1}, -cal_{i+1,j-1})| \quad (5)$$

where S_{ij} is the graph similarity between pairs antennas i and j and Opp() indicates the symmetrically opposite antenna pair (mirror pair), originating from the same antenna i. where cal_{ij} is the features from the scattered signals measured with an elliptical homogeneous isotropic media in both simulation and experiment, without a head present. Certainly, regarding the value from low frequency (0.5-1.25 GHz), the symbol is labeled as $LDOP$ and the $HDOP$ is from high frequency bound (1.25-2 GHz). For example, $LDOP_{1,8}$ when transmitting from antenna pair 1 and 8 and opposite antennas transmitting from antenna 16 and 9 in lower frequency band. On the other hand, the $HDNP_{18,27}$ is the differential between $1 - 8$ and $2 - 7$ from high frequency band.

A. Localizing the hemisphere of the bleeding targets

In clinical cases, it is also important to detect which hemisphere (left/right) the bleeding lesion occurs. Although experts might detect it based on clinical symptoms. A naïve approach to detect whether the blood mass is in the left or right hemisphere is by comparing the features (such as amplitude, mean degree or strength) from the left signal to those of the right. However, this method can rarely provide an accurate answer due to the slight asymmetry of the head. This section presents a simple method to detect whether the bleeding is in the left or right hemisphere.

The algorithm first detect whether the bleeding target is close to the antenna $1 - 9$ or the opposite $16 - 8$. It calculates two parameters $LDOP$, $LDNP$ and $HDNP$ using equation 6.

$$DS_{1,16} = f(s_{1,9}) - f(cal_{16,8})) \quad (6)$$

where f is the function of graph transfer, and the phase of $S_{i,j}$.

$$f = \begin{cases} Left, & DS(1,9) > 0, LDNP(1,8) > LDNP(16,9) \\ Right, & DS(1,9) < 0, LDNP(1,8) < LDNP(16,9) \\ middle, & otherwise. \end{cases} \quad (7)$$

B. Picking the intersection of an antenna pair and its mirror

The antenna pairs that cross a bleeding from the ith antenna can be detected using (8), where % represents the modulo operation. Note that we don't consider neighbouring antennas as the information gained from them is less than the information from antennas spaced further apart.

$$max \sum_{j=(i+3)\%n}^{(n/2+3)\%n} DW_{ij} \quad (8)$$

The output is a set which contains the pairs of antennas and the assigned weights. The weights imply the probability that one of those pairs of signals are likely to cross a bleeding. Larger values imply that the signals pass a bleeding with high probability, and the position of the bleeding can be estimated based on the intersection with other signals that pass the bleeding. Thus, we can determine the antenna pairs whose signals cross the bleeding (or any abnormality causing an asymmetricity in the signals) and their mirrored pairs as well, but we need to determine which of these pairs cross the blood mass. For example, in Fig 2, pair $(1-7)$, $(9-4)$ on the left brain and their mirror pairs $(1-11)$, $(9-14)$ will be selected because their differential weighted nodes are significantly different from the other pairs. Then, the intersection co-ordinates (p_x, p_y) can be calculated based on (9) and (10) when four antenna positions (x_1,y_1), (x_2,y_2), (x_3,y_3) and (x_4,y_4) are provided.

$$p_x = \frac{\begin{vmatrix} \begin{vmatrix} x_1 & y_1 \\ x_2 & y_2 \\ x_3 & y_3 \\ x_4 & y_4 \end{vmatrix} & \begin{vmatrix} x_1 & 1 \\ x_2 & 1 \\ x_3 & 1 \\ x_4 & 1 \end{vmatrix} \end{vmatrix}}{\begin{vmatrix} \begin{vmatrix} x_1 & 1 \\ x_2 & 1 \\ x_3 & 1 \\ x_4 & 1 \end{vmatrix} & \begin{vmatrix} y_1 & 1 \\ y_2 & 1 \\ y_3 & 1 \\ y_4 & 1 \end{vmatrix} \end{vmatrix}} \quad (9)$$

$$p_y = \frac{\begin{vmatrix} \begin{vmatrix} x_1 & y_1 \\ x_2 & y_2 \\ x_3 & y_3 \\ x_4 & y_4 \end{vmatrix} & \begin{vmatrix} y_1 & 1 \\ y_2 & 1 \\ y_3 & 1 \\ y_4 & 1 \end{vmatrix} \end{vmatrix}}{\begin{vmatrix} \begin{vmatrix} x_1 & 1 \\ x_2 & 1 \\ x_3 & 1 \\ x_4 & 1 \end{vmatrix} & \begin{vmatrix} y_1 & 1 \\ y_2 & 1 \\ y_3 & 1 \\ y_4 & 1 \end{vmatrix} \end{vmatrix}} \quad (10)$$

C. Heat map constructing algorithm

This section describes a constructing heat map algorithm to obtain the size and the shape of the bleeding from the intersection of lines pairs from section III-B. The algorithm uses intersected heat mapped shape as a lesion, which requires the output of subsection III-B as input values, such as I={R_0, R_1, R_2}. The pseudo code is listed below.

(a) Calculate the maximum distance between the intersections I from the SCLA algorithm. For example, there are three intersections $R_0 = (x_0, y_0)$, $R_1 = (x_1, y_1)$, and $R_2 = (x_2, y_2)$, the maximum distance $md = |R0 - R1|$.

(b) if the md is larger than $10mm$ then the heat radius is assigned as $20mm$ otherwise $40mm$.

(c) Draw each intersection point with the assigned heat radius.

D. DICE

To evaluate the performance of localization, the Dice similarity coefficient (DICE) is used to calculate the overlap of the estimated size of the IB (S_t) with the real IB area (S_d). The DICE, also called the overlap index, is the most used metric in MRI segmentation. It is defined by

$$DICE = \frac{2|S_t \cap S_d|}{|S_t| + |S_d|} \quad (11)$$

The DICE can be viewed as a similarity measure over two bleeding sets. If two image contains the same area coordinate points, the DICE is equal to 1.0, while if two bleeding sets have no overlap, it is equal to 0.0.

E. Statistical analysis of head rotation

To investigate the difference between objects with targets and those without targets, a quantitative statistical test was performed on the two sets of graph features. A p-value< 0.05 was obtained, demonstrating that there is a significant difference in the graph features of objects with vs without targets.

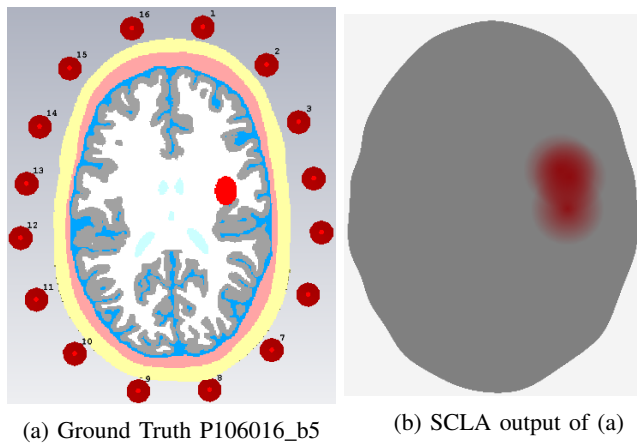(a) Ground Truth P106016_b5   (b) SCLA output of (a)

Fig. 5: A test from model P106016_b5, where the ground truth positions with tissues are shown on the left and the SCLA algorithm result is shown without tissues on the right.

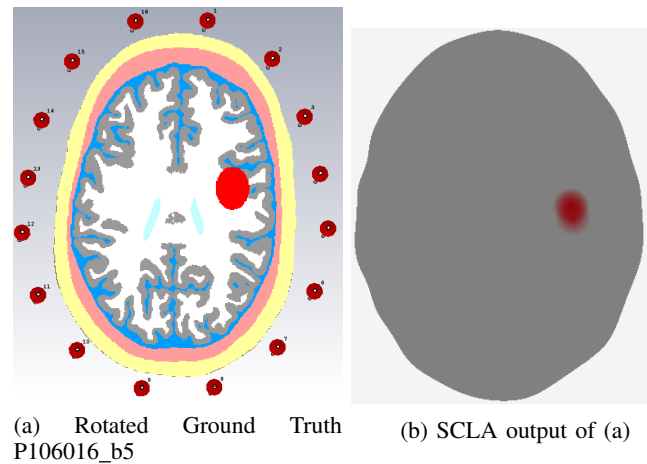(a) Rotated Ground Truth P106016_b5   (b) SCLA output of (a)

Fig. 6: A test from model P106016_b5 is the same as Fig 5 but all antenna positions are rotated 3 degrees, where the ground truth positions with tissues are shown on the left and the SCLA algorithm result is shown without tissue on the right.

IV. RESULTS ON SIMULATED DATA

The algorithms are implemented in C and R languages. Eight phantoms were employed for testing. Each phantom is constructed with five different bleeding observations by injecting an elliptical shaped blood mass with the dielectric properties of blood. In total, 28 brain model observations were used for evaluating the algorithm performance in simulations. In addition, a simple phantom with a small glass cylinder filled with distilled water to emulate a TBI is used to verify the algorithm with a 14 antenna array in experiments. The glass cylinder is moved clockwise to two different distances per antenna, and the S-parameter data is collected twice to measure the anti-noise performance of SCLA. Thus, a total of 56 simulated data are collected and tested. The mean DICE value of 94 testing samples is 0.78 and the maximum DICE value is 0.92.

A. Test the algorithm when rotating antenna

To test the tolerance of the algorithm to the antenna positions, this subsection evaluates the antenna Z-rotation tolerance in the simulation experiment. Rotating the antenna array along the Z-axis will significantly affect the algorithm performance because the other two dimensions (X and Y axis) can be easily aligned by ears and eyebrows. In experiments with healthy volunteers, we found that subjects were biased to left or right on the Z-axis but few of them rotated their heads on the X or Y-axis.

Figures 5 shows an observation from a brain model from the seven tissue brain models. The red target of Figures 5(a) indicates a bleed target on the ground truth, where the Figures 5(b) shows the SCLA algorithms' output.

Fig 6 shows the same brain model as Fig 5 but the antenna array has three degrees of rotation in Figure 6. The DICE between the SCLA output and the ground truth is approximately 0.08 in Fig 6, while the DICE is about 0.84 in the symmetrical case (Fig 5). Thus, this algorithm can tolerate a two-degree antenna rotating in all directions.

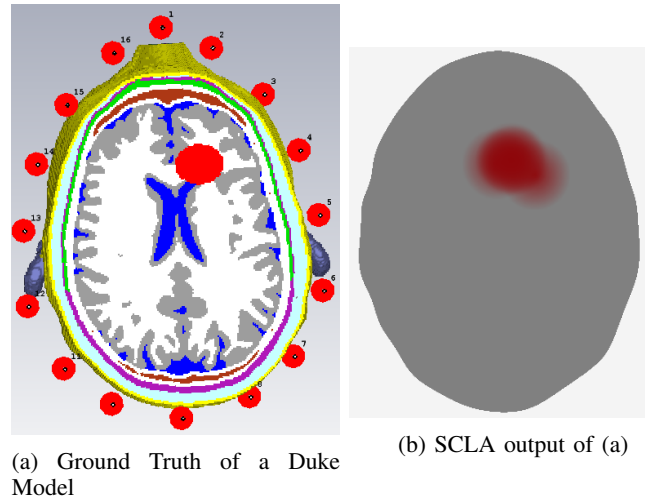(a) Ground Truth of a Duke Model   (b) SCLA output of (a)

Fig. 7: A test result on the Duke brain model. The ground truth positions with tissues are shown on the left and the SCLA algorithm result is shown without tissues on the right.

B. Testing different antenna positions

Figures 7 and 8 show two observations from two brain models with intracranial bleeding (Duke and Zubal brain models, respectively) with different distances between the head and antennas. The green line indicates the direct signal path from antennas 1, 5, and 9. The crossing points are plotted in blue, whereas the estimated targets are drawn in red.

It can be seen that the Duke and Zubal brain models are significantly different. In addition, the gap between head boundary and antennas in Fig 7 is smaller than those in Fig 8. Our SCLA was able to obtain the approximate location of the targets even though the two antennas have different placement and the distances between the antennas and the head boundary are also different.

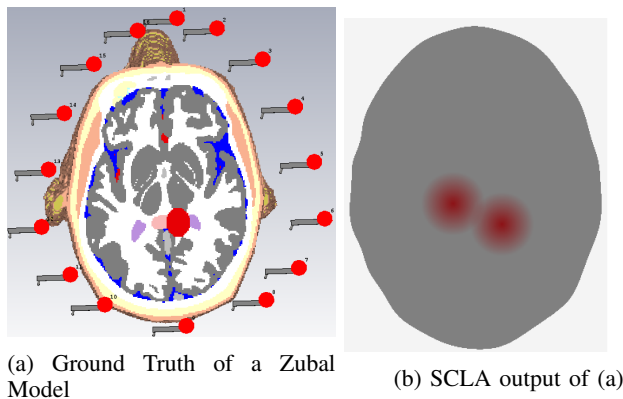(a) Ground Truth of a Zubal Model

(b) SCLA output of (a)

Fig. 8: A test result of the Zubal brain model. The ground truth positions with tissues are shown on the left and the SCLA algorithm result is shown without tissues on the right.
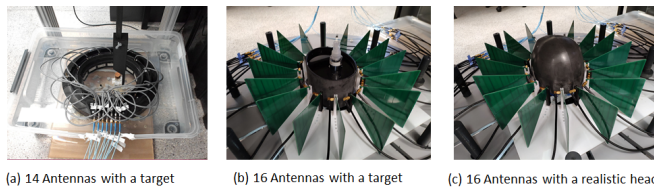

(a) 14 Antennas with a target   (b) 16 Antennas with a target   (c) 16 Antennas with a realistic head

Fig. 9: Three experimental platforms with 14 and 16 antennas respectively.

V. RESULTS ON EXPERIMENTAL DATA

A. Experiments data

The SCLA was applied to locate simple targets in two experiments. One is a dielectric-loaded 14-waveguide antenna array system. Each waveguide antenna is filled with a coupling medium with a dielectric permittivity of 33 and a conductivity of 0.1. An elliptical container filled with 60:40 glycerol water mixture is used to mimic a head, and a water-filled glass cylinder with an outer diameter of 22 mm and wall thickness of 1 mm is used as a target. The second is a 16 Log-Periodic array system. The experimental device is shown in Fig 9(a and b).

The target in Fig 9(a) is moved clockwise in front of each antenna with 10 mm and 14 mm distances, respectively. The S-parameter data was collected twice. Thus, a total of 56 targets were measured. A Keysight M9037A VNA was used to measure the signals, and a frequency scan from 0.5 GHz to 2 GHz with 2 MHz step size was used to provide a total of 751 frequency points.

The second experiment evaluated the algorithm performance using a CNC machine with a EM head scanner. There are 16 antenna array put on a Skin-Skull Bucket (SSB) as shown in Fig 10(a). The SSB was placed within the array imaging domain upside-down and filled with liquid which has average brain emulating dielectric property. It was placed on a centring mount within the array, to hold the SSB in the centre of the antennas. The pressure of the array membrane was then increased to 1 PSI. Following this, a solid haemorrhage emulating target (Fig 10(c)) was measured inside the SSB. These targets were moved along a 10mm spaced grid pattern

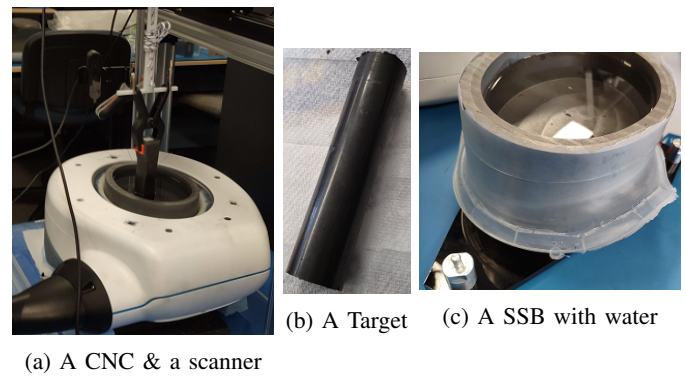(b) A Target   (c) A SSB with water

(a) A CNC & a scanner

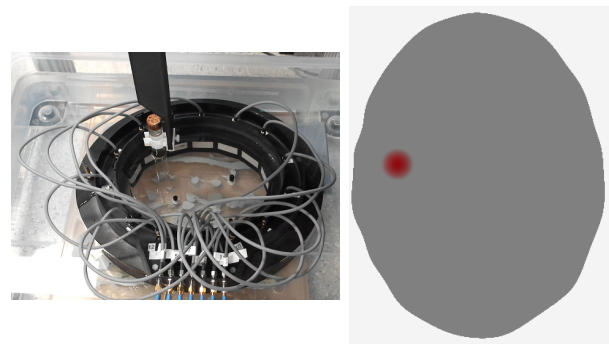Fig. 10: The SCLA output of two realistic phantoms.

Fig. 11: The experimental setup. The glass cylinder filled with distilled water was moved into a stable position near antenna 11 in a water filled container on the left and the SCLA algorithm result is shown in red on the right.

in X and Y, with a fixed Z as low as possible within the bucket. In total, there were 94 points measured inside the SSB.

For all these realistic systems, the measurement time is approximately 1.5 seconds per measurement on 16 antennas. The system uses a transmit power of approximately +10 dBm (10 mW), which is less than that used in mobile phones. Compared with an existing study which performed experiments at only a single frequency point [35]. Our system is slower due to a large number of frequency points (751 steps).

B. Locating the target in a realistic container

As introduced in Section V-A, an elliptic container filled with distilled water was used to test a target which is a glass cylinder with an outer diameter of 22 mm and wall thickness of 1 mm, filled with distilled water. During each measurement, a CNC machine moves the target to one of 28 positions, and the distance between the between the target and the antenna aperture is 10 mm and 14 mm respectively. A movie was made in the Supporting files to illustrate the target movements. Fig 11 shows a SCLA output of a measurement, where the container with target was used for calibration in equation 4. The SCLA can correctly localize the target according to the collected S-parameters. The average DICE score for these 56 measurements was 0.85 ± 0.08.
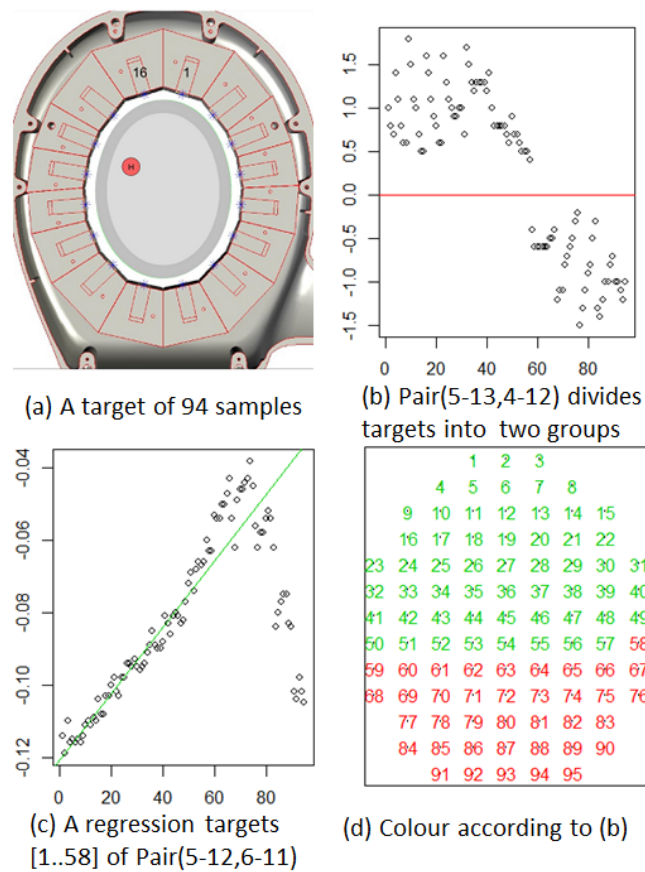
(a) A target of 94 samples

(b) Pair(5-13,4-12) divides targets into two groups

(c) A regression targets [1..58] of Pair(5-12,6-11)

(d) Colour according to (b)

Fig. 12: Two pairs antennas analysis of 94 points in a SSB container

C. Locating targets in asymmetric head phantoms

This section test the algorithms using 94 ICH targets which is placed in 94 positions inside a SSB container. One of the original position is shown in Fig 12 (a). It is clearly that the symmetric pair $4-12$ and $5-13$ can divide those targets into two groups as shown in Fig 12(b). For example, the 87th target is 100% at the back and the 23st target is 100% on the front. Then, the other pairs such as pairs $1-9$ and $16-8$ could divide the target into left or right hemisphere.

It is easy to find that the measurement results support that the real head is not completely symmetrical. Fig 12 (c) shows that the differential degree features from neighbour pair $5-12$ and $6-11$. The value increase when target is near the transmit line between pair $5-12$ and $6-11$, such as targets 60 or 75. On the other hand, the position of target $1-58$ has a linear regression relationship with those of $DNP_{5,6}$. In that case, it is easy to obtain the target $23^r d$ position by using the regression in Fig 12 (c) as we could detect the target is really in front area by detecting the $DOP_{4,5}$ as shown in Fig 12 (d). Certainly, any target in back area could have the similar linear relationship with the pair of $4-13$ and $3-14$.

VI. DISCUSSION

In this study, we presented a novel symmetrical crossing line algorithm (SCLA) which only requires the antenna positions

and measured signals to localize intracranial bleeding in the brain. The physical principle is that, compared to signals passing through two similar areas (non-target areas), a pair of symmetrical transmitted signals passing through the targets (bleeding) will generate significantly different amplitude and phase changes. Unlike conventional imaging algorithms, the output of the SCLA is the coordinates of the target instead of an image. Compared to other methods which need human or machine learning technology to provide the real coordinates from the medical image, our algorithm directly provides the real coordinates.

In fact, the SCLA might open a new direction for electromagnetic imaging which relies on the intersection of the direct signal paths between antenna pairs. Compared with radar-based detection methods, such as confocal imaging [36], the SCLA algorithm can obtain the location without knowing the boundary of the head and without estimating the average permittivity of the imaging subject. In fact, the SCLA algorithm is only based on the intersection of the two signal paths, which can obtain high localization accuracy even when the head boundary is unknown.

Compared with using multi-core CPUs [8] or using GPUs tomography-based methods, the SCLA method has much lower computational cost (less than 1 minute on an Intel Core i7-2600 CPU @ 3.40 GHz with 16G RAM), making it more suitable for use in emergency scenarios. In addition, the tomography-based methods are usually influenced by the selection of the antennas. For example, in [3], [37], antennas with TM_z polarization needed to be used to satisfy the integral equations, and in [38], directive waveguide antennas were used to restrict the radiation pattern so that the scattering problem could be approximated as a 2-D problem.

Compared with the machine learning-based methods which require a huge training data set [17], SCLA is based only on the on-site measurement and it is not a learning-based method thereby the substantial training data are not required.

Also, human heads have different sizes and shapes. Many existing algorithms fail to give an accurate image when the brain sizes and shapes are different. However, our SCLA algorithm is not affected by the boundary, size or shape. The only requirement is the positions of the antennas which is easy to obtain in a clinical situation. More importantly, the container and the target were filled with the same distilled water in Fig 11. Despite this, our proposed algorithm is still able to efficiently detect the changes caused by the 1 mm thick glass cylinder only.

The hypothesis of this method is assuming that the healthy human head is approximately symmetric across the major and minor axes (i.e., across the sagittal plane, separating the left-right of the brain, and across the central coronal plane, separating the front and back of the brain). Symmetric pairs of antennas across these lines of symmetry should have similar signals within a healthy head. If symmetric pairs of antennas measure different signals, this indicates an abnormality such as bleeding, and this observation can be used to facilitate localisation. In our realistic system, each antenna is just 10 mW and the measurement is about 1.5 ns. There are 8 transmitting antennas in total. This is very safe because it is

only the equivalent to 0.01s of a mobile phone transmission time.

Sections IV and V demonstrate that SLCA can obtain cerebral hemorrhage images from experimental phantoms and simulation data. However, two reasons could lead to difficulty in measuring the DICE in experimental data compared with those of simulated cases. Firstly, it is that the CNC can introduce error during movements, while the simulation model has none of this type of error. Secondly, shifts or rotations of the phantom or the human head might introduce errors during the measurement as shown in section IV-A. Currently, there is no effective tool to detect whether the phantom or head is placed in the center of the antenna array. It can only rely on manual and visual inspection to observe whether the head or phantom is rotated or shifted. Therefore, it is possible to obtain low or zero DICE in real experiments despite high DICE being achieved in simulation in the same situation. To overcome this limitation and to acquire a higher DICE score, we can use multiple measurements to determine whether the crossing line of the algorithm are from the same pair of antennas. In the future, a method to automatically detect rotation or shifts of a head or phantom will be developed.

Currently, the algorithm does not calculate the electrical properties of the bleeds. However, it can identify target localization using differential degree values between two antennas. Thus, it is possible to measure a target in 3-dimensional imaging by moving the system up or down on the z-axis using a 3-dimensional calibration. Each scan is a slice, and then all slices could be merged into a 3-dimensional case. In addition, it is possible to detect other abnormal tissues, such as tumors or ischemic in the future.

VII. CONCLUSION

Using a novel symmetrical crossing line algorithm, this paper is the first to localize an intracranial bleeding mass without using radar, tomography or machine learning methods. Based on the hypothesis that healthy brains are approximately symmetric, the algorithm is able to accurately localize and predict the shape of bleeding masses that cause a non-symmetric response in the electromagnetic signals. The algorithm was evaluated on 150 measurements with head phantoms of varying size, shape and resolution, and antenna types, and was able to accurately locate bleeding masses with an average Dice score of 0.85. Compared with existing electromagnetic imaging and localization algorithms, our approach is less sensitive to changes in head size, coupling mediums and antenna positions, directly outputs the coordinates of the bleeding mass, and does not require any training data. It is also significantly faster, localising the bleeding mass position within one minute. The speed and accuracy of the method shows great potential for use in emergency clinical applications.

REFERENCES

[1] A. Abbosh, "Microwave systems for head imaging: Challenges and recent developments," in *Microwave Workshop Series on RF and Wireless Technologies for Biomedical and Healthcare Applications (IMWS-BIO), 2013 IEEE MTT-S International*. IEEE, 2013, pp. 1–3.

[2] M. Persson, A. Fhager, H. D. Trefná, Y. Yu, T. McKelvey, G. Pegenius, J.-E. Karlsson, and M. Elam, "Microwave-based stroke diagnosis making global prehospital thrombolytic treatment possible," *IEEE Transactions on Biomedical Engineering*, vol. 61, no. 11, pp. 2806–2817, 2014.

[3] L. Guo and A. Abbosh, "Stroke localization and classification using microwave tomography with k-means clustering and support vector machine," *Bioelectromagnetics*, vol. 39, no. 4, pp. 312–324, 2018.

[4] E. Ricci, S. Di Domenico, E. Cianca, T. Rossi, and M. Diomedi, "PCA-based artifact removal algorithm for stroke detection using UWB radar imaging," *Medical & biological engineering & computing*, vol. 55, no. 6, pp. 909–921, 2017.

[5] B. Oliveira, D. Godinho, M. OHalloran, M. Glavin, E. Jones, and R. Conceição, "Diagnosing breast cancer with microwave technology: Remaining challenges and potential solutions with machine learning," *Diagnostics*, vol. 8, no. 2, p. 36, 2018.

[6] P. T. Nguyen, A. M. Abbosh, and S. Crozier, "3-d focused microwave hyperthermia for breast cancer treatment with experimental validation," *IEEE Transactions on Antennas and Propagation*, vol. 65, no. 7, pp. 3489–3500, 2017.

[7] N. Ghavami, G. Tiberi, M. Ghavami, S. Dudley, and M. Lane, "Huygens principle based uwb microwave imaging method for skin cancer detection," in *2016 10th International Symposium on Communication Systems, Networks and Digital Signal Processing (CSNDSP)*. IEEE, 2016, pp. 1–4.

[8] N. Simonov, B.-R. Kim, K.-J. Lee, S.-I. Jeon, and S.-H. Son, "Advanced fast 3-d electromagnetic solver for microwave tomography imaging," *IEEE transactions on medical imaging*, vol. 36, no. 10, pp. 2160–2170, 2017.

[9] M. Oziel, R. Korenstein, and B. Rubinsky, "Radar based technology for non-contact monitoring of accumulation of blood in the head: A numerical study," *PloS one*, vol. 12, no. 10, p. e0186381, 2017.

[10] A. Zamani, A. M. Abbosh, and A. T. Mobashsher, "Fast frequency-based multistatic microwave imaging algorithm with application to brain injury detection," *IEEE Transactions on Microwave Theory and Techniques*, vol. 64, no. 2, pp. 653–662, 2016.

[11] S. Candefjord, J. Winges, A. A. Malik, Y. Yu, T. Rylander, T. McKelvey, A. Fhager, M. Elam, and M. Persson, "Microwave technology for detecting traumatic intracranial bleedings: tests on phantom of subdural hematoma and numerical simulations," *Medical & Biological Engineering & Computing*, vol. 55, no. 8, pp. 1177–1188, Aug 2017. [Online]. Available: https://doi.org/10.1007/s11517-016-1578-6

[12] W. Shao, T. R. McCollough, and W. J. McCollough, "A phase shift and sum method for uwb radar imaging in dispersive media," *IEEE Transactions on Microwave Theory and Techniques*, 2019.

[13] B. Mohammed, K. Bialkowski, and A. Abbosh, "Radar-based time-domain head imaging using database of effective dielectric constant," *Electronics Letters*, vol. 51, no. 20, pp. 1574–1576, 2015.

[14] Q. Mahmood and A. Basit, "Prediction of ischemic stroke lesion and clinical outcome in multi-modal mri images using random forests," in *International Workshop on Brainlesion: Glioma, Multiple Sclerosis, Stroke and Traumatic Brain Injuries*. Springer, 2016, pp. 244–255.

[15] R. Benny, T. A. Anjit, and P. Mythili, "An overview of microwave imaging for breast tumor detection," *Progress In Electromagnetics Research*, vol. 87, pp. 61–91, 2020.

[16] Q. Mahmood, S. Li, A. Fhager, S. Candefjord, A. Chodorowski, A. Mehnert, and M. Persson, "A comparative study of automated segmentation methods for use in a microwave tomography system for imaging intracerebral hemorrhage in stroke patients," *Journal of Electromagnetic Analysis and Applications*, vol. 7, no. 05, p. 152, 2015.

[17] M. Salucci, J. Vrba, I. Merunka, and A. Massa, "Real-time brain stroke detection through a learning-by-examples techniquea n experimental assessment," *Microwave and Optical Technology Letters*, vol. 59, no. 11, pp. 2796–2799, 2017.

[18] P. Govindarajan, R. K. Soundarapandian, A. H. Gandomi, R. Patan, P. Jayaraman, and R. Manikandan, "Classification of stroke disease using machine learning algorithms," *Neural Computing and Applications*, Jan 2019. [Online]. Available: https://doi.org/10.1007/s00521-019-04041-y

[19] J.-B. Li, M.-D. Zhu, Y.-Z. Wu, and S. Ye, "Stroke detection based on an improved artificial fish swarm algorithm," in *2017 IEEE International Symposium on Antennas and Propagation & USNC/URSI National Radio Science Meeting*. IEEE, 2017, pp. 789–790.

[20] M. Elahi, D. OLoughlin, B. Lavoie, M. Glavin, E. Jones, E. Fear, and M. OHalloran, "Evaluation of image reconstruction algorithms for confocal microwave imaging: Application to patient data," *Sensors*, vol. 18, no. 6, p. 1678, 2018.

[21] J. Yang, H. Zhao, H. Li, G. Liu, Q. Ran, J. Chen, Z. Bai, G. Jin, J. Sun, J. Xu, M. Qin *et al.*, "An experimental study on the early diagnosis of traumatic

brain injury in rabbits based on a noncontact and portable system," *PeerJ*, vol. 7, p. e6717, 2019.

[22] K. Hugdahl, "Symmetry and asymmetry in the human brain," *European Review*, vol. 13, no. S2, pp. 119–133, 2005.

[23] S. X. Liu, "Symmetry and asymmetry analysis and its implications to computer-aided diagnosis: A review of the literature," *Journal of biomedical informatics*, vol. 42, no. 6, pp. 1056–1064, 2009.

[24] G. Zhu, Y. Li, and P. P. Wen, "Epileptic seizure detection in EEGs signals using a fast weighted horizontal visibility algorithm," *Computer methods and programs in biomedicine*, vol. 115, no. 2, pp. 64–75, 2014.

[25] I. G. Zubal, C. R. Harrell, E. O. Smith, Z. Rattner, G. Gindi, and P. B. Hoffer, "Computerized three-dimensional segmented human anatomy," *Medical physics*, vol. 21, no. 2, pp. 299–302, 1994.

[26] A. Christ, W. Kainz, E. G. Hahn, K. Honegger, M. Zefferer, E. Neufeld, W. Rascher, R. Janka, W. Bautz, J. Chen *et al.*, "The virtual familyde-velopment of surface-based anatomical models of two adults and two children for dosimetric simulations," *Physics in Medicine & Biology*, vol. 55, no. 2, p. N23, 2009.

[27] E. Lee, W. Duffy, R. Hadimani, M. Waris, W. Siddiqui, F. Islam, M. Rajamani, R. Nathan, and D. Jiles, "Investigational effect of brain-scalp distance on the efficacy of transcranial magnetic stimulation treatment in depression," *IEEE Transactions on Magnetics*, vol. 52, no. 7, pp. 1–4, 2016.

[28] G. Zhu, Y. Li, and P. P. Wen, "Analysis and classification of sleep stages based on difference visibility graphs from a single-channel EEG signal," *IEEE Journal of Biomedical and Health Informatics*, vol. 18, no. 6, pp. 1813–1821, 2014.

[29] A. Supratak, H. Dong, C. Wu, and Y. Guo, "Deepsleepnet: A model for automatic sleep stage scoring based on raw single-channel eeg," *IEEE Transactions on Neural Systems and Rehabilitation Engineering*, vol. 25, no. 11, pp. 1998–2008, Nov 2017.

[30] P. Artameeyanant, S. Sultornsanee, and K. Chamnongthai, "An emg-based feature extraction method using a normalized weight vertical vis-ibility algorithm for myopathy and neuropathy detection," *SpringerPlus*, vol. 5, no. 1, p. 2101, 2016.

[31] Z. Hai, D. Sheng-chang, D. Wei, and C. Xing-chi, "Complex network analysis of pulse wave," *Journal Of Northeastern University Nature Science*, vol. 37, no. 12, pp. 1696–1699, 2016.

[32] G. Zhu, Y. Li, P. P. Wen, and S. Wang, "Analysis of alcoholic EEG signals based on horizontal visibility graph entropy," *Brain informatics*, vol. 1, no. 1-4, pp. 19–25, 2014.

[33] S. Supriya, S. Siuly, H. Wang, and Y. Zhang, "Eeg sleep stages analysis and classification based on weighed complex network features," *IEEE Transactions on Emerging Topics in Computational Intelligence*, 2018.

[34] G. Zhu, A. Bialkowski, L. Guo, B. Mohammed, and A. Abbosh, "Stroke classification in simulated electromagnetic imaging using graph approaches," *IEEE Journal of Electromagnetics, RF and Microwaves in Medicine and Biology*, vol. 5, no. 1, pp. 46–53, 2021.

[35] A. Hamidipour, T. Henriksson, M. Hopfer, R. Planas, and S. Semenov, "Electromagnetic tomography for brain imaging and stroke diagnostics: Progress towards clinical application," in *Emerging Electromagnetic Technologies for Brain Diseases Diagnostics, Monitoring and Therapy*. Springer, 2018, pp. 59–86.

[36] L. Guo and A. M. Abbosh, "Optimization-based confocal microwave imaging in medical applications," *IEEE Transactions on Antennas and Propagation*, vol. 63, no. 8, pp. 3531–3539, 2015.

[37] C.-C. Chiu and R.-H. Yang, "Electromagnetic imaging for complex cylindrical objects," *IEEE transactions on medical imaging*, vol. 14, no. 4, pp. 752–756, 1995.

[38] M. Hopfer, R. Planas, A. Hamidipour, T. Henriksson, and S. Semenov, "Electromagnetic tomography for detection, differentiation, and moni-toring of brain stroke: A virtual data and human head phantom study." *IEEE Antennas and Propagation Magazine*, vol. 59, no. 5, pp. 86–97, 2017.