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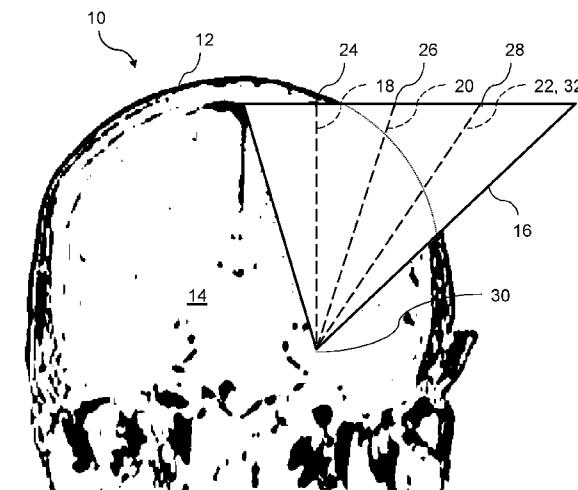


Fig.1

(57) Abstract: The invention relates to a method for determining at least one applicable path (32) for the movement of an object, especially of a surgical and/or diagnostical device, in human tissue (14) or animal tissue by means of a data set of intensity data obtained by a 3D imaging technique, the applicable path (32) of movement connecting a starting position (28) of the device with a defined target location (30). According to the invention the method comprises the steps: -defining the target location (30) of a reference point of the device and choosing at least one possible starting position (24, 26, 28) of the reference point of the device; -determining a candidate path of movement (18, 20, 22) between the corresponding possible starting position (24, 26, 28) and the defined target location (30); and -evaluating the candidate path of movement (18, 20, 22) as being an applicable path (32) depending on information about local intensity extrema and/or intensity variation resulting from the intensity data along the candidate path of movement (18, 20, 22). The invention further relates to a corresponding computer-readable medium, a corresponding computer program product, and a corresponding computerized system.

## METHOD FOR DETERMINING AT LEAST ONE APPLICABLE PATH OF MOVEMENT FOR AN OBJECT IN TISSUE

### FIELD OF THE INVENTION

The invention relates to a method for determining at least one applicable path for the movement of an object, especially of a surgical and/or diagnostical device, in human tissue or animal tissue by means of a data set of intensity data obtained by a 3D imaging

5 technique, the path of movement connecting a starting position of the device with a defined target location. The invention also relates to a computer-readable medium such as a storage device a floppy disk, CD, DVD, Blue Ray disk, or a random access memory (RAM), as well as a computer program product comprising a computer usable medium including computer usable program code and a computerized system.

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### BACKGROUND OF THE INVENTION

Healthcare diagnosis decision support systems or computer-aided diagnosis or computer-assisted diagnosis (CAD) systems are used in medicine to assist users like medical experts or physicians in the interpretation of medical images. Imaging techniques in X-ray, 15 magnetic resonance imaging (MRI), and Ultrasound diagnostics yield a great deal of information, which the user has to analyze and evaluate comprehensively in a short time.

CAD systems help scan digital images, e.g. from magnetic resonance imaging, for typical appearances and to highlight conspicuous structures, such as vessels, nerve pathways, ventricles, functionally eloquent regions and/or tumor regions. Usually, machine-learning 20 technologies, such as a decision tree and neural network, are utilized to build classifiers based on a large number of known cases with ground truth, i.e., cases for which the diagnosis has been confirmed by pathology. The classifier bases its diagnosis on a computational structure built from known cases and inputted features for the unknown structure case. The classifier output indicates the estimated nature of the unknown structure and optionally a confidence

25 value. As the precision of medical imaging facilities improves to detect very small structures, and as the number of digital images to be processed increases this type of CAD becomes increasingly important as a tool to assist users like physicians. The computer-produced classification is considered a second opinion to a user like a physician in order to raise the accuracy and confidence associated with diagnosis.

Computer assisted surgery (CAS) represents a surgical concept and set of methods, that use computer technology for presurgical planning, and for guiding or performing surgical interventions. CAS is also known as computer aided surgery, computer assisted intervention, image guided surgery and surgical navigation, but these terms that are 5 more or less synonyms with CAS.

The traditional approach of determining an applicable path of movement for a surgical and/or diagnostical device (a safe surgical trajectory) in an image guided therapy like MRI always comprises the two basic steps of: segmenting each critical structure in a spatial region defining the possible path(s) of movement around these regions and afterwards 10 determining a corresponding safe or applicable path. One important area of application is brain surgery. In detail, the traditional surgery planning mainly follows the following steps:

In a first step, the target location is defined manually, or automatically, or in a semi-automated manner. This either involves registering the magnetic resonance (MR) 15 volume to a template, often in the stereotactic coordinate system, and detection of anatomical structures from this transformation, or identifying some point and plane landmarks, such as Mid-sagittal plane, and AC/PC points to determine the location of the target. Once the target is determined, the planning reduces to the detection of an entry point. In many cases, the path between the entry and target points has to be straight and should not hit the critical structures.

The second step is the identification of critical structures, such as vessels, gyri 20 and sulci, ventricles, and some functionally eloquent regions in the brain. This involves segmenting these structures for example from contrast-enhanced T<sub>1</sub> weighted MR images by applying a set of image segmentation algorithms. Functionally eloquent regions are determined by analyzing fMRI (functional Magnetic Resonance images) and/or DTI (diffusion tensor images) data.

25

## SUMMARY OF THE INVENTION

It is an object of the invention to provide an enhanced method and an enhanced system for determining at least one applicable path of movement for a surgical and/or diagnostical device in tissue by means of a data set obtained by a 3D imaging 30 technique.

This object is achieved by the present invention as defined in the independent claims.

The method according to the invention comprises the steps: (i) defining the target location of a reference point of the device and choosing at least one possible starting

position of the reference point of the device; (ii) determining a candidate path of movement between the corresponding possible starting position and the defined target location; and (iii) evaluating the candidate path of movement as being an applicable path depending on information about local intensity extrema and/or intensity variation resulting from the 5 intensity data along the candidate path of movement.

Instead of the earlier mentioned two-step surgery planning approach of first determining the critical structures to avoid by segmenting them in multiple steps, and then determining the applicable path of movement (the safe trajectory) as a function of these segmentation results, a direct search for the applicable path of movement by skipping the 10 segmentation step is performed in the method according to the invention. In other words, the method according to the invention does not consider the position and extension of critical structures as a whole, but simply looks for information about the intensity maxima and/or intensity minima and/or intensity variation (intensity information) resulting from the intensity data along the candidate path of movement indicating the presence of a critical structure in a 15 domain of the candidate path. The critical structures are structures within the tissue, such as vessels, nerve pathways, ventricles, and/or functionally eloquent regions. The information about the intensity extrema and/or intensity variation of the data set is therefore directly used to determine at least one applicable path of movement. The method according to the invention has *inter alia* the following advantages:

20 direct evaluation of path candidates by intensity characteristics without need for segmentation process;

automatic adaptation to local intensity characteristics thereby avoiding the intensity inhomogeneity problems affecting e.g. MR images;

25 flexibility to add other types of constraints, such as integration of functional information, in the searching step of determining the possible path(s) of movement, and

support for non-straight paths by extending the method to a search for multiple line segments;

option for providing an explanatory display of the path information as intensity plots and image-based thumbnails.

30 To determine the safety level of a candidate path, an intensity profile along the possible path of movement based on the information about the intensity extrema and/or intensity variation is analyzed. A safe or applicable path must not have large maxima or large minima in the path region. These maxima and/or minima are detected by a so called sliding window approach, where for each point, a window is centered and the maximum and the

minimum intensity within the window are computed. If the center pixel is either the maximum or the minimum, it can be a critical structure. To verify this, the difference of the maximum and the minimum in the window is compared with the intensity at the target location to determine the saliency of the peak and valley. Afterwards, considering some 5 spatial constraints, the trajectory is determined to be safe or not. A candidate path determined as safe is an applicable path.

According to a preferred embodiment of the invention, the candidate path of movement is a linear path of movement connecting the starting position and the target location along the shortest route. The linear path of movement is the most widely used type 10 of path. A typical surgical and/or diagnostical device using a linear path is a needle shaped or rod shaped device.

According to another preferred embodiment of the invention, the evaluation considers an expert assessment of the intensity data in an area around the candidate path of movement. The expert assessment preferably is a case-by-case decision of an expert whether 15 an intensity variation or a local maximum (peak) or a local minimum (valley) results from a critical structure. With respect to the present invention, the term "expert" therefore relates to a person having the knowledge to interpret intensity data with reference to the displayed structures.

According to yet another preferred embodiment of the invention, the 20 evaluation of the candidate path of movement is an automatic evaluation performed by a computerized system. This computerized system preferably is a CAD (computer-assisted diagnosis) or CAS (computer assisted surgery) system.

Preferably, the expert assessment together with the corresponding information about the intensity and/or intensity variation is used for a training (teaching) of the 25 computerized system. The computerized system uses an adaptive algorithm to the intensity variations by learning the dynamic range of the intensity of the input data. The corresponding computerized system is based on a machine-learning technology, such as a decision tree, a support vector machine, adaboost, or a neural network. The classifier of this system bases its diagnosis on a computational structure built from known cases and inputted intensity extrema 30 and/or intensity variation along the candidate path of movement.

The 3D imaging technique preferably is a medical imaging technique to generate a three-dimensional image of the inside of an object from a large series of two-dimensional X-ray images taken around a single axis of rotation. The 3D imaging technique is for example an X-ray computed tomography (CT) technique or an imaging ultrasound

technique. According to a preferred embodiment of the invention the 3D imaging technique is a magnetic resonance imaging (MRI) technique, a nuclear magnetic resonance imaging (NMRI) technique, or a magnetic resonance tomography (MRT) technique. The intensity data are derived from  $T_1$ -weighted MR scans and/or  $T_2$ -weighted MR scans and/or  $T^*_2$ -weighted MR scans and/or by magnetization transfer MRI and/or by fluid attenuated inversion recovery (FLAIR) and/or by magnetic resonance angiography and/or by magnetic resonance gated intracranial CSF dynamics (MR-GILD).

According to a preferred embodiment of the invention, the human or animal tissue is brain tissue. An important application of the path determining method is brain surgery planning with the corresponding tissue being brain tissue. The critical structures within this tissue are vessels, gyri and sulci, ventricles, and/or functionally eloquent regions.

In a first alternative, allowable yaw and pitch angles of the allowable path are determined. The selected angles satisfy the two constraints: 1) the starting position should be on the same hemisphere as the target location, and 2) the search should be towards the top of the brain rather than to the sides or below the target location. Having determined the yaw and pitch angles, within that preset angle range, rays from the target location towards the surface of the brain are casted where potential starting positions are located. Finally, the intensity profile of each candidate path (candidate trajectory) is analyzed and labeled as safe or not, and/or given a continuous safety value. A candidate path labeled as safe or having a corresponding safety value is an applicable path.

In a second approach, a slice in the axial direction that corresponds to the top of the head (immediately satisfying the constraint that the search should be towards the top of the brain rather than to the sides or below the target location) is determined. For a target location on the left side of the brain, we choose the left half of the slice, and for a target location on the right, only the region of the slice corresponding to the right brain hemisphere is chosen (satisfying the constraint that the starting position should be on the same hemisphere as the target location). After that, the lines between each eligible point on the slice and the predetermined target location as described above form the candidate paths.

According to another embodiment of the invention, the method comprises the further step of displaying the at least one candidate path of movement or at least the starting position of the at least one candidate path of movement. In a preferred mode of graphic representation of the possible starting positions, a region of interest of possible starting position(s) is/are displayed, wherein the starting position(s) of applicable paths is/are displayed in a first color and the starting position(s) of the other paths is/are displayed in a

second color. Especially, this can be in the form of a colored cortical surface map, where, for example, the first color is green, indicating applicable paths that satisfy the constraints of avoiding critical structures and the second color is red, indicating the other (unsafe) starting positions. If the tissue is brain tissue, the colored cortical surface map displays candidate 5 entry positions on the skull.

Preferably, the candidate path of movement is displayed as an intensity profile. More preferably, the intensity profile is displayed in association with at least one image of an area around the local extrema, e.g. as a thumbnail image.

According to another preferred embodiment of the invention, an acceptance 10 threshold of the information about the local intensity extrema and/or intensity variation for a given candidate path is determined or at least in part determined by the adjacent paths of said given candidate path.

According to another aspect of the present invention, the intensity profile is calculated by integrating the intensity data over the maximum cross section of the device 15 perpendicular to a direction of motion along the candidate path of movement. This integration may be implemented in various ways. In one embodiment, the intensity value that results in the largest intensity change along the path is used to represent the cross-section. In another embodiment, multiple paths are considered separately to decide the safety of each individual path. Afterwards, the safety of the cross-section is computed as the aggregate of all 20 single paths.

According to another aspect of the invention, a cross section can be defined to refer to a safety margin (defined as the distance to the nearest critical structure) around a candidate path. Within the pre-defined cross sectional area, no critical structure indicated by certain intensity characteristics exists.

In another aspect of the invention, all candidate paths are evaluated as safe or 25 not. Afterwards, for each path, a safety margin is computed by detecting the closest path that are labeled as unsafe. Then, for the safe paths, the entry points on the surface of the brain can be color coded as a function of the computed safety margin. For the unsafe paths, a distinct color coding can be used that can be a function of the total number of intensity minima and 30 maxima along the path, or a color coding scheme that discriminate whether only intensity minima, or intensity maxima or both are observable along the path.

The invention further relates to a computer-readable medium such as a storage device, a floppy disk, CD, DVD, Blue Ray disk, or a random access memory (RAM), containing a set of instruction that causes a computer to perform an aforementioned method

and a computer program product comprising a computer usable medium including computer usable program code, wherein the computer usable program code is adapted to execute the aforementioned method.

The invention finally relates to a computerized system comprising a

5 processing unit, a memory, a data interface, a display and an input device, wherein the system is adapted for performing the aforementioned method. Preferably the computerized system is a CAD (computer-assisted diagnosis) or CAS (computer assisted surgery) system.

## BRIEF DESCRIPTION OF THE DRAWINGS

10 These and other aspects of the invention will be apparent from and elucidated with reference to the embodiments described hereinafter.

In the drawings:

Fig. 1 shows a search region view from a coronal slice of a human head. A

15 region of possible paths of movement is constructed by many possible paths (three of them explicitly shown) connecting corresponding starting positions with a common target location;

Fig. 2 the intensity profile of a “safe” possible path of movement therefore being an applicable path of movement;

20 Fig. 3 the intensity profile of an “unsafe” possible path of movement; and

Fig. 4 a top view on the brain with a visualization of the region of possible paths subdivided into applicable paths and unsafe paths.

## DETAILED DESCRIPTION OF EMBODIMENTS

25 Fig. 1 shows a search region view from a coronal slice of a human head 10. The image shows the top of the skull 12 and the tissue 14 of the brain inside the skull 12. A region of candidate paths 16 for the movement of an object like a surgical or diagnostical device (not shown) in the brain tissue 14 is constructed by many candidate paths 18, 20, 22 (three of them explicitly shown) connecting corresponding possible starting positions 24, 26, 30 28 with a common target location 30.

An applicable path 32 for the movement of the device in the tissue 14 is determined by means of a data set of intensity data obtained by the 3D magnetic resonance imaging technique, the applicable path 32 of movement is connecting a starting position 28 of the device with a defined target location 30. The corresponding determination method

comprises the steps:

defining the target location 30 of a reference point of the device and choosing at least one possible starting position 24, 26, 28 of the reference point of the device;

determining the at least one candidate path of movement 18, 20, 22 between

5 the corresponding possible starting position 24, 26, 28 and the defined common target location 30; and

evaluating the candidate path of movement 18, 20, 22 as being the applicable path 32 depending on information about the local intensity minima/maxima (extrema) and/or intensity variation resulting from the intensity data along the candidate path of movement 18,

10 20, 22.

The applicable path 32 for the movement of the device in the tissue 14 is a safe path which does not interfere with a critical structure (depicted in the insets of Fig. 3).

To determine the safety level of a candidate path 18, 20, 22, an intensity profile 34, 36 (shown in Figs. 2 and 3) along the candidate path of movement 18, 20, 22 based on the information about the intensity extrema and/or intensity variation is analyzed. A safe applicable path 32 must not have large maxima or large minima in the (brain) tissue region. In general there are various options to define the feature “large”.

A preferred approach for detecting these kind of maxima and/or minima is a sliding window approach. In a sliding window approach for each point or voxel (volume element represented by the point), a window is centered and the maximum and the minimum intensity within the window are computed to make sure that the corresponding point/voxel needs to be locally maximum or minimum to be a global maximum or minimum. Afterwards, any combination of the following steps can be done:

- the absolute difference between the intensity of the current point or voxel and the intensity of the target location 30 should be within  $K^*$  intensity standard deviation of the whole tissue (e.g. the brain volume). Factor K can be specified individually (e.g.  $K = 1$ ) and defines an upper and a lower intensity threshold. The current point or voxel is a large minimum if the intensity of this point/voxel is lower than the lower intensity threshold and a large maximum if higher than the upper intensity threshold.

-  $X, y, z$  gradient of a volume surrounding the current point/voxel is computed by taking the  $x, y, z$  derivatives, respectively. From the derivatives, gradient magnitude is computed for each point/voxel. Then, a histogram is built from gradient magnitudes. Some percentile (e.g.  $P = 80\%$ ) of the histogram is computed and assigned as threshold value. The above absolute difference between the intensity of the current point/voxel and the intensity at

the target location 30 is compared with this threshold value to find the minimum or maximum.

5 The target location 30 is given initially (or found elsewhere) and the search for possible starting position 24, 26, 28 starts from this target location 30. As such, the intensity value at the target location 30 is already known and can be used for the detection of the minima and maxima.

10 If the pixel/voxel is either the maximum or the minimum, it indicates a critical structure. Afterwards, considering some spatial constraints, the candidate paths 18, 20, 22 are determined to be safe or not. The candidate path of movement 22 determined as safe is the applicable path 32. The critical structures within the shown brain tissue are vessels, gyri and sulci, ventricles, and/or functionally eloquent regions.

15 In a first alternative, allowable yaw and pitch angles of the applicable path 32 are determined. The selected angles satisfy the two constraints: 1) the starting position should be on the same hemisphere as the target location 30, and 2) the search should be towards the top of the brain rather than to the sides or below the target location 30. Having determined the yaw and pitch angles, within that preset angle range, rays from the target location 30 towards the surface of the brain are casted where the possible starting positions 24, 26, 28 are located. Finally, the intensity profile of each candidate path (candidate trajectory) 18, 20, 22 is analyzed and labeled as safe or not, and/or given a continuous safety value. A candidate 20 path 18, 20, 22 labeled as safe or having a corresponding safety value is an applicable path 32.

25 In a second approach, a slice in the axial direction that corresponds to the top of the head (immediately satisfying the constraint that the search should be towards the top of the brain rather than to the sides or below the target location) is determined. For a target location 30 on the left side of the brain, the left half of the slice is chosen, and for a target location 30 on the right, only the region of the slice corresponding to the right brain hemisphere is chosen (satisfying the constraint that the starting position should be on the same hemisphere as the target location). After that, the lines between each eligible point on the slice and the predetermined target location 30 as described above form the candidate 30 paths 18, 20, 22.

Fig. 2 depicts the intensity profile 34 of a safe candidate path of movement 22; Fig. 3 depicts the intensity profile 36 of an unsafe candidate path of movement 20. The corresponding path of both intensity profiles start at the target location 30. The intensity profile 34 in Fig. 2 does not show large fluctuations (large minima or large maxima outside

the intensity thresholds given by  $K^*$  intensity standard deviation around the intensity of the target location 30) in the intensity until the end of the path, where there is a salient maximum in the intensity profile 34. This is expected as it corresponds to the intensity of the skull 12 and not the brain tissue. As a result, this candidate path 22 is labeled as safe. The candidate 5 path of Fig. 3 has peaks (local maxima) within the brain region; therefore, it is considered as unsafe. Only the safe candidate path of movement 22 corresponding to the intensity profile 34 shown in Fig. 2 is an applicable path 32.

A further aspect of the method is an enhanced visualization strategy. In addition to the intensity profiles 34, 36, the user can also see an image 38 (a thumbnail 10 image) of an area around the local maxima or local minima by clicking on the graphs as shown in Figs 2 and 3. In both Figures, only maxima are associated with a thumbnail image; however, both maxima and minima can be visualized in the same way. Furthermore, one can enlarge the size of the thumbnail image to get more information on the spatial context.

In another embodiment, one can add non-intensity constraints to the searched 15 trajectories. This can be in the form of defining a search region as explained in the candidate entry point detection step. A search region can also be defined by using the result of another method in the form of a map that defines allowable and unacceptable points. This map can be generated from the analysis of a different modality, such as fMRI, DTI,  $T_2$  etc. It can also be generated from the same MR  $T_1$  contrast used for trajectory analysis. For example, one can 20 use the result of cortical segmentation and brain mask for constraining the search area for the safe trajectory detection.

Fig. 4 shows a top view on the brain with a visualization of a region of interest 40 consisting of possible starting positions 24, 26, 28 of the candidate paths 18, 20, 22 within the region of candidate paths 16, the region of interest 40 subdivided into areas 42 of possible 25 starting positions 28 of applicable paths 32 and a remaining area 44 of unsafe candidate paths 18, 20.

A user of a corresponding computer system like for example a surgeon can have a view of the brain on a display of the system depicting the region of interest (ROI) 40 of possible starting positions 24, 26, 28 comprising areas 42 colored green to indicate starting 30 positions 28 of save applicable paths 32 and another area 44 colored red to indicate starting positions 24, 26 of unsafe paths.

If the user places the cursor 46 on a possible starting position, the corresponding intensity plot (as shown in Figs. 2 and 3, is shown in one embodiment of the invention. In the example shown in Fig. 4, the cursor 46 points at one of the areas 42 of

possible starting positions 28 of applicable paths 32. The system would depict the corresponding intensity profile of a “safe” applicable path of movement 32 like shown in Fig. 2. In a computer system being a computer assisted surgery (CAS) system, the cursor 46 will follow the movement of the surgical and/or diagnostical device.

5 While the invention has been illustrated and described in detail in the drawings and foregoing description, such illustration and description are to be considered illustrative or exemplary and not restrictive; the invention is not limited to the disclosed embodiments. Other variations to the disclosed embodiments can be understood and effected by those skilled in the art in practicing the claimed invention, from a study of the drawings, the 10 disclosure, and the appended claims. In the claims, the word "comprising" does not exclude other elements or steps, and the indefinite article "a" or "an" does not exclude a plurality. The mere fact that certain measures are recited in mutually different dependent claims does not indicate that a combination of these measures cannot be used to advantage. Any reference signs in the claims should not be construed as limiting the scope.

## CLAIMS:

1. Method for determining at least one applicable path (32) for the movement of an object, especially of a surgical and/or diagnostical device, in human tissue (14) or animal tissue by means of a data set of intensity data obtained by a 3D imaging technique, the applicable path (32) of movement connecting a starting position (28) of the device with a defined target location (30), comprising the steps:

- defining the target location (30) of a reference point of the device and choosing at least one possible starting position (24, 26, 28) of the reference point of the device;
- determining a candidate path of movement (18, 20, 22) between the corresponding possible starting position (24, 26, 28) and the defined target location (30); and
- evaluating the candidate path of movement (18, 20, 22) as being an applicable path (32) depending on information about local intensity extrema and/or intensity variation resulting from the intensity data along the candidate path of movement (18, 20, 22).

15 2. The method according to claim 1, wherein the candidate path of movement (18, 20, 22) is a linear path of movement connecting the possible starting position (24, 26, 28) and the target location (30) along the shortest route.

20 3. The method according to claim 1, wherein the evaluation of each of the candidate paths of movement (18, 20, 22) further considers an expert assessment of the intensity data in an area around said candidate path of movement (18, 20, 22).

25 4. The method according to claim 1, wherein the evaluation of the candidate path of movement (18, 20, 22) is an automatic evaluation performed by a computerized system.

5. The method according to claim 4, wherein the expert assessment together with the corresponding information about the local intensity extrema and/or intensity variation is used for a training of the computerized system.

6. The method according to claim 1, wherein the 3D imaging technique is a magnetic resonance imaging technique, a nuclear magnetic resonance imaging technique, or a magnetic resonance tomography technique.

5

7. The method according to claim 1, comprising the further step of: displaying the candidate path (18, 20, 22) or at least the possible starting position (24, 26, 28) of the candidate path (18, 20, 22).

10 8. The method according to claim 7, wherein a region of interest (40) of the possible starting position (24, 26, 28) is displayed, wherein the starting position (28) of applicable path (32) is displayed in a first color and the starting position (24, 26) of the other path (18, 22) is displayed in a second color.

15 9. The method according to claim 7, wherein the candidate path (18, 20, 22) is displayed as an intensity profile (34, 36).

10. The method according to claim 7, wherein the intensity profile (34, 36) is displayed in association with at least one image (38) of an area around the local extrema of 20 the intensity profile (34, 36).

11. The method according to claim 1, wherein an acceptance threshold of the information about the local intensity extrema and/or intensity variation for a given candidate path of movement (18, 20, 22) is determined or at least in part determined by the adjacent 25 paths of said given candidate path of movement (18, 20, 22).

12. The method according to claim 1, wherein the information about the intensity and/or intensity variation is calculated by integrating the intensity data over the maximum cross section of the device perpendicular to a direction of motion along the candidate path of 30 movement (18, 20, 22).

13. A computer-readable medium such as a storage device, a floppy disk, CD, DVD, Blue Ray disk, or a random access memory, containing a set of instruction that causes a computer to perform a method according to one of the preceding method claims.

14. A computer program product comprising a computer usable medium including computer usable program code, wherein the computer usable program code is adapted to  
5 execute a method of any of the preceding method claims.

15. A computerized system comprising a processing unit, a memory, a data interface, a display and an input device, wherein the system is adapted for performing a method of any of the preceding method claims.

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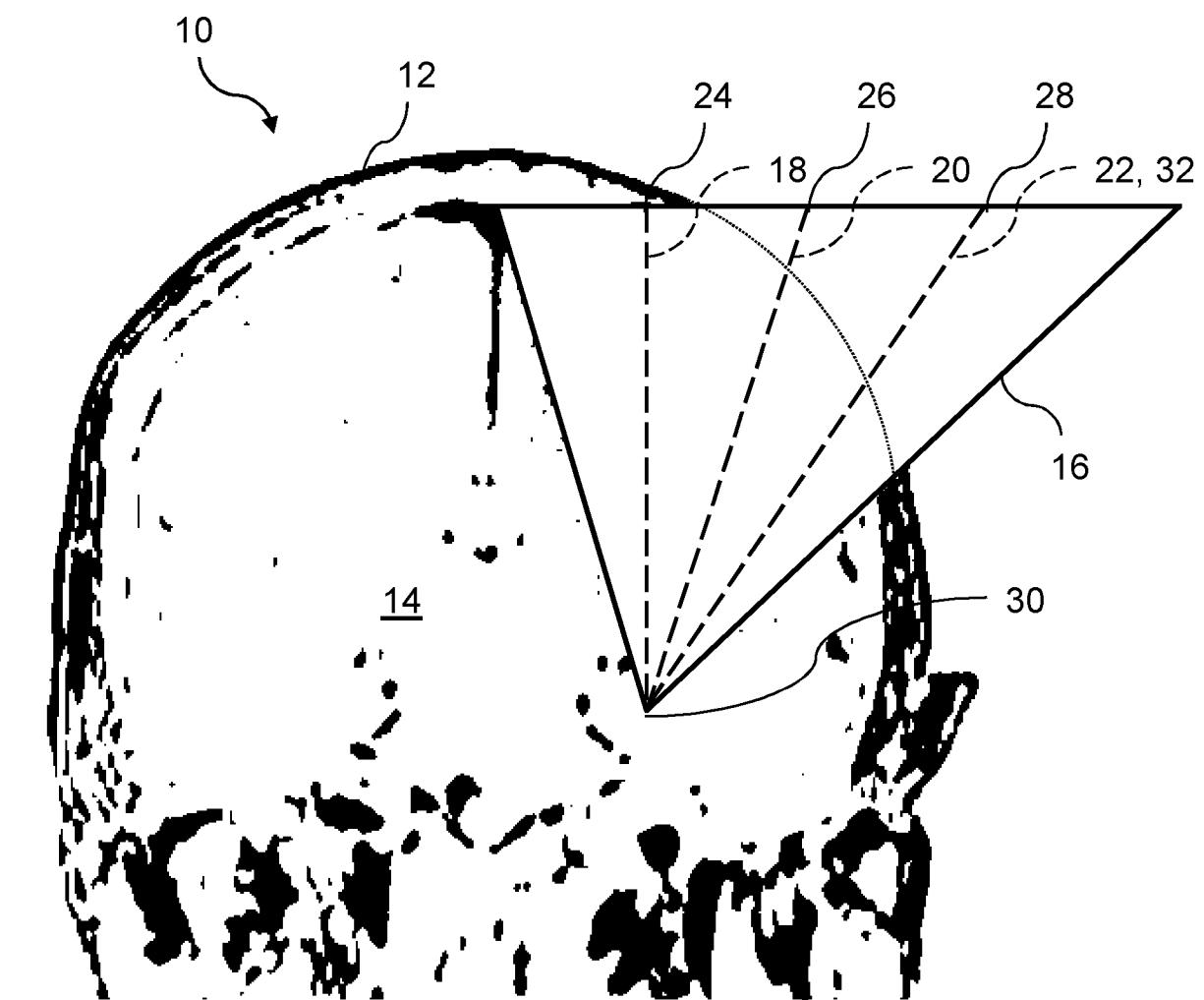


Fig. 1

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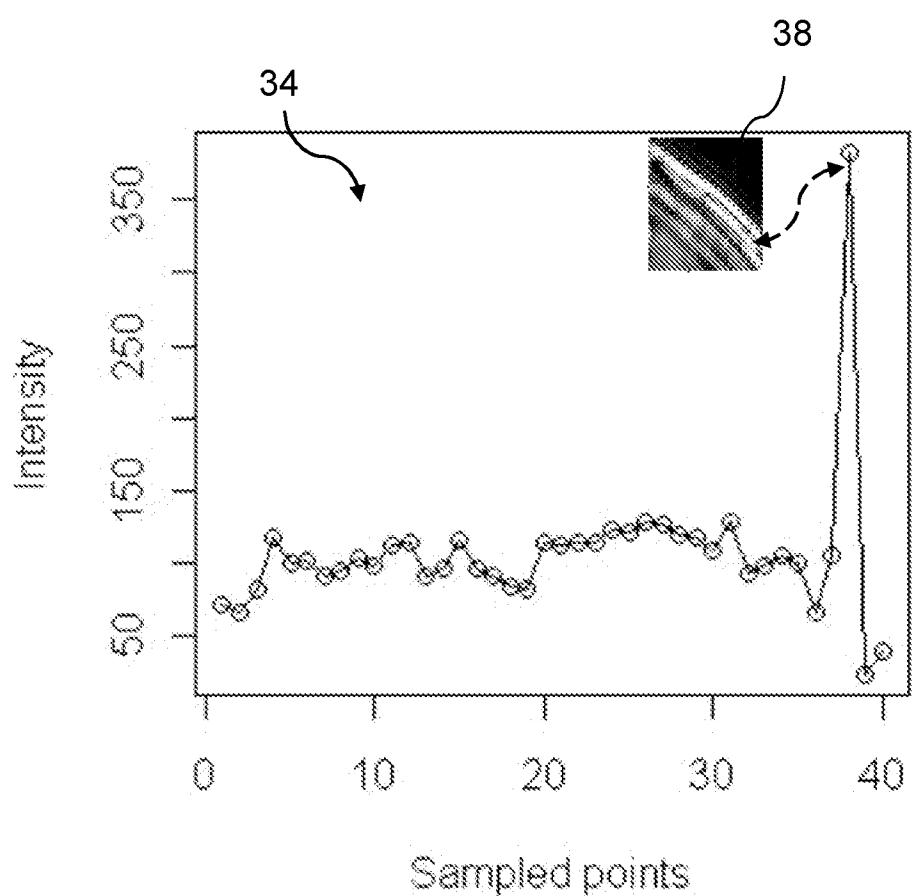


Fig. 2

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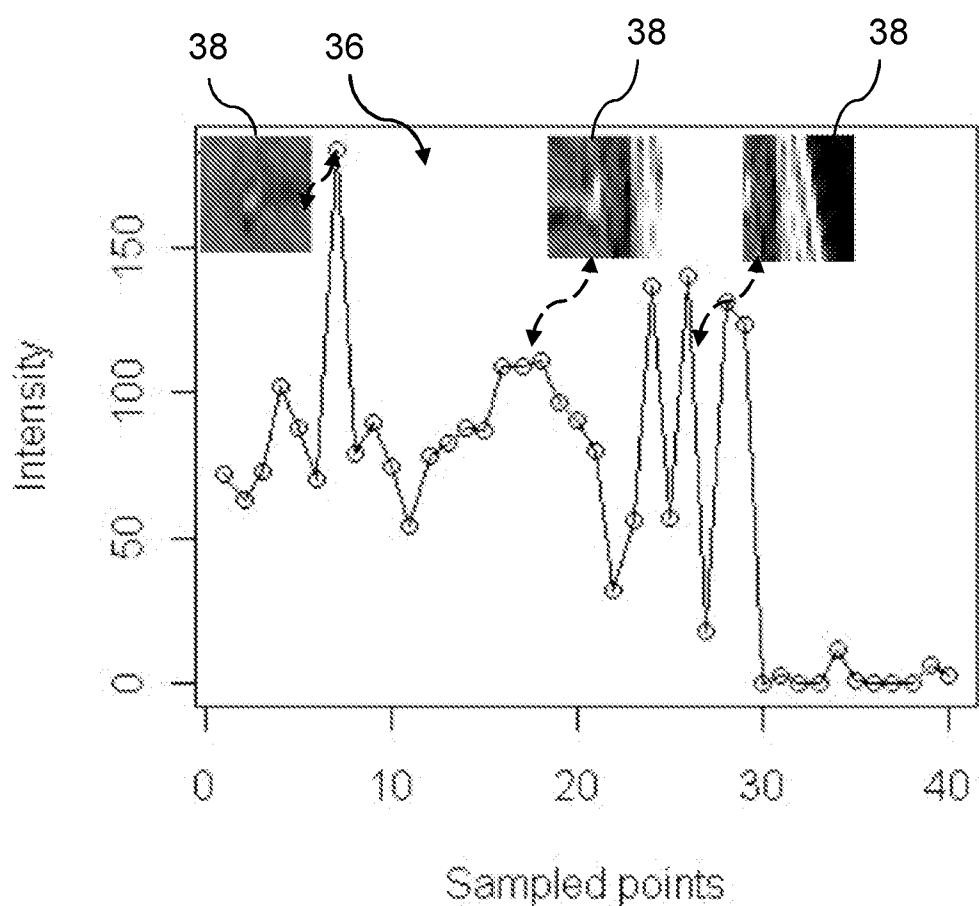


Fig. 3

4/4

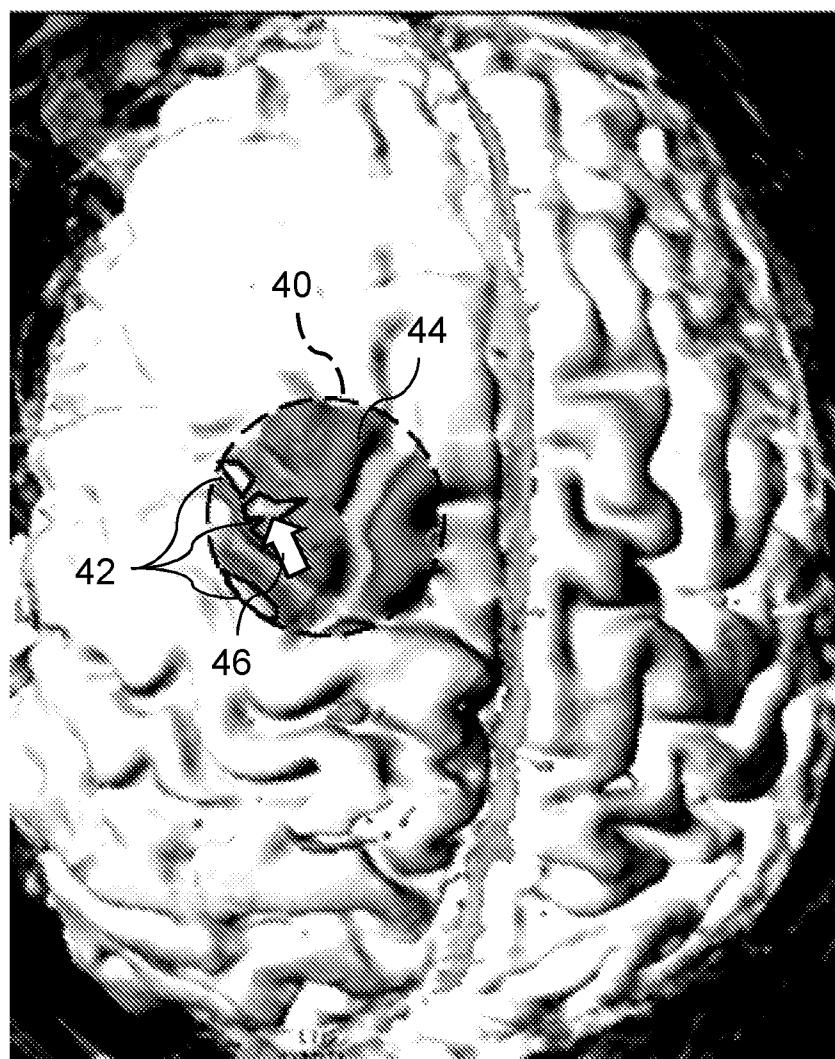


Fig. 4

# INTERNATIONAL SEARCH REPORT

International application No  
PCT/IB2012/050169

**A. CLASSIFICATION OF SUBJECT MATTER**  
INV. A61B19/00 A61B17/34 G06T7/00  
ADD.

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)

A61B G06T

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

EPO-Internal, WPI Data

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 2008/123922 A1 (GIELEN FRANS L H [NL] ET AL) 29 May 2008 (2008-05-29)	1,2,4, 6-8,11, 13-15
Y	paragraphs [0027], [0063] - [0073] -----	9,10,12
X	US 2011/007071 A1 (PFISTER MARCUS [DE]) 13 January 2011 (2011-01-13) paragraphs [0020], [0025]; claim 20 -----	1
Y	US 2003/068074 A1 (HAHN HORST [DE]) 10 April 2003 (2003-04-10) paragraph [0167] -----	9,10,12



Further documents are listed in the continuation of Box C.



See patent family annex.

\* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier application or patent but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
30 April 2012	08/05/2012

Name and mailing address of the ISA/  
European Patent Office, P.B. 5818 Patentlaan 2  
NL - 2280 HV Rijswijk  
Tel. (+31-70) 340-2040,  
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Authorized officer

Angeli, Markus

## INTERNATIONAL SEARCH REPORT

International application No.  
PCT/IB2012/050169

### Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1.  Claims Nos.: **3, 5**  
because they relate to subject matter not required to be searched by this Authority, namely:  
**Rule 39.1(iii) PCT - Scheme, rules and method for doing business**
2.  Claims Nos.:  
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3.  Claims Nos.:  
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

### Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1.  As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2.  As all searchable claims could be searched without effort justifying an additional fees, this Authority did not invite payment of additional fees.
3.  As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4.  No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

#### Remark on Protest

The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee.

The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.

No protest accompanied the payment of additional search fees.

**INTERNATIONAL SEARCH REPORT**

Information on patent family members

International application No  
PCT/IB2012/050169

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 2008123922	A1 29-05-2008	NONE	
US 2011007071	A1 13-01-2011	NONE	
US 2003068074	A1 10-04-2003	NONE	