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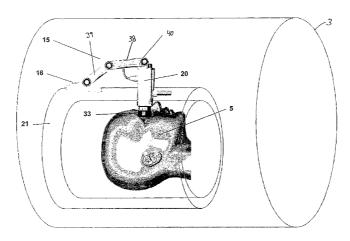
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(54) Title: APPARATUS FOR PROVIDING HIGH RESOLUTION IMAGES IN A MRI-DEVICE



(57) Abstract: The present invention relates to an apparatus for providing high resolution images to patients positioned in a magnetic resonance imaging (MRI) device (3), the MRI device (3) comprising a head coil (21), the head coil arranged to surround a patient's (5) head and to provide MRI images thereof, the apparatus comprising means (12, 13, 14) for receiving video or picture image signals from an external source. According to the present invention, the apparatus further comprises means (14, 28, 34, 29) for displaying a video or picture image, said display means being arranged in a housing (20), said housing (20) being suspended in an arm comprising at least two successive members (38, 39), wherein a joint (40) between the housing (20) and the adjacent member (38), a joint or joints (41) between the successive members (38, 39), and a joint (42) between an attachment element (16) for attaching the apparatus to the head coil (21), or other part of the MRI device (3), and the member (39) adjacent to the coil attachment element (16), each is hinged to allow rotation of the joints (40, 41, 42).



# APPARATUS FOR PROVIDING HIGH RESOLUTION IMAGES IN A MRI-DEVICE

#### FIELD OF THE INVENTION

The present invention relates generally to those products that are used to provide visual stimuli for testing and comforting patients undergoing diagnostic treatment. More specifically, the present invention relates to an apparatus for providing high resolution images to patients positioned in a magnetic resonance imaging (MRI) device.

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# BACKGROUND OF INVENTION

In the medical field, magnetic resonance imaging (MRI) is a commonly used non-invasive technique to diagnose the medical condition of a patient. MRI has the ability to distinguish healthy and diseased tissue, fat and muscle, and between adjacent structures within the body which other imaging modalities cannot demonstrate. MRI utilizes safe radio waves and a magnetic field to generate the images processed by a computer. Typically, the patient is placed within a large homogeneous magnetic field and is subjected to a set of gradient fields and radio frequency (RF) fields. The various fields are accurately controlled to cause nuclei within a selected slice of the patient to precess about an axis and to emit RF signals. These signals are then used to reconstruct an image of the slice. By varying the gradient fields, images of the patient at different slices may be captured. The separate slices can then be combined to form a complete scan of the patient.

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Generally, with respect to the use of MRI scanners, video systems are employed for both (a) patient comfort and (b) functional imaging applications. With respect to patient comfort, the concern is directed to anxious or claustrophobic patients who resist entering the tunnel of the MRI scanner. The capability to adequately display visual information for viewing is an important factor for relief for the anxious or claustrophobic patient. The second use of video systems in MRI scanners is directed to functional imaging applications. In some instances, the diagnostic procedure performed with the MRI is used to evaluate a patient's response to

specific visual stimuli. The operator sends a series of images to a screen which is seen by the patient during the MRI procedure and the patient's responses are included in the MRI report.

A problem with introducing conventional video signals into an MRI device is that very small magnetic fields generated by another device can destroy the images generated by the MRI device. Conversely, the strong fields generated by the MRI device may prevent the normal operation of certain devices, such as a cathode ray tube (CRT) or liquid display panel (LCD), within the vicinity of the MRI device.

Therefore, any type of system used to present video signals to the patient must not generate any stray magnetic fields in the vicinity of the MRI device and should be shielded from the magnetic fields generated by the MRI device.

Another problem is that the MRI device is based on the use of radio frequencies that may disrupt signal modulation. For these reasons, the video signal must be in a form that is not affected by the radio frequency and transmitted by a system that is not easily magnetized.

The most common method for presentation of visual stimuli inside the MR scanner is to generate an image outside the magnetic field of the MR machine and have a mirror or prism for reflecting the image to the patient. For instance, viewing systems as described in U.S. Pat. No. 5,076,275 to Bechor et al., U.S. Pat. No. 6,774,929 to Kopp and an MRI video system disclosed in a Nuclear Associates brochure all reflect images generated from a video source located away from the patient into the eyes of the patient. The projection is achieved within the magnetic environment by employing an MRI-compatible LCD screen, or by using a video projector and a translucent screen. The screen is positioned in the proximity of the MR scanner. The projector or LCD screen is positioned either inside or outside the MR room. The video information is viewed by the patient with the aid of adjustable light reflecting mirrors or through a prism. The utility of this method of visual activation is limited by the position of the patient within the scanner tunnel. Further, the level of ambient light in the MRI magnet room will affect the quality of the image that the patient sees on the screen. A high level of ambient light will cause the screen image to be washed out. Also, the time required to adjust the light

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reflecting mirrors with respect to the screen is determined by the position of the patient inside the scanner tunnel. For functional magnetic resonance imaging, it is ideal to cover the entire patient field-of-view with the MRI screen or display.

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The effectiveness of this method of visual activation is further reduced by an open field of view (e.g., the screen is outside of the tunnel) which enables the patient to be aware of her surroundings. Therefore, the patient may find it difficult to focus on the video images and may therefore find it difficult to completely relax. This may be especially true for systems which reflect the video images from behind the MRI device to the patient. With this type of system, the patient may be distracted by items which are adjacent to the display screen or by people working behind the patient. Thus, the possibility of being distracted by the external surroundings in addition to the interior of the tunnel further limits the usefulness of this technique for the reduction of anxiety and claustrophobia in patients. It would therefore be desirable to have the patient focus on the video images during the MRI procedure so that the patient is able to relax.

An attempt to address this problem is found in U.S. Pat. No. 4,901,141 which utilizes a fibre optic taper positioned within the bore of an MRI apparatus. In order to isolate the video system from the fields generated by the MRI device and to prevent any magnetic fields from affecting the MRI device, this system pipes in video images to the patient while the patient is within the MRI device. A CRT produced image is delivered to the fibre optic taper through a coherent image guide. The fibre optic taper expands the end of the image guide so as to provide a larger viewing surface for the patient. The problem with the fibre optic taper is that it is stationary and the patient must be positioned in a fixed location so as to be able to see the end of the optic taper. Further, to prevent distortion the patient must be located directly beneath the isocenter of the taper. Thus, the disclosure does not address different size patients, patient positioning, or near and far sighted patients. For instance, a tall person may lay with their head partially outside the bore during diagnostics of the lower body whereas a child may be well encapsulated by the bore, neither of which could properly see a fixed fibre optic taper. In addition; the use of a fixed taper will interfere with auxiliary coils, such as

head and c-spine coils, that require close proximate to the body. Current construction of head and c-spine coils is such that the visual field as needed for viewing a fixed positioned fibre taper is either obscured or completely blocked if the fibre taper is utilized.

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Another prior art device is disclosed in U.S. Pat. No. 5,414,459 directed to a pair of glasses worn by the patient. The glasses receive the video picture by fibre optic guide.

- In both theses devices the installation is permanent with a fibre optic connection between the shielded MRI room and a remote location housing the operating elements of the system. The connection requires the shielding which surrounds the MRI room to be breeched and that penetration must be adequately protected.
  - Current MRI fibre optic systems that position the LCD screen within the scanner room (but outside the bore of the MRI scanner) are extremely useful and provide a definite advance in the art. Notwithstanding, certain features of this design could be improved. In particular, the length of the fibre optic bundle employed to carry the video images from the LCD screen to the eyepiece for viewing by the patient is of concern. As with all transmission systems, a portion of the transmitted parameter is lost during transmission and the longer the transmission path, the greater the loss. For long fibre optic bundles, it is known that the loss of as much as forty percent (40%) of the transmitted video image can occur. This loss affects the resolution and brightness of the transmitted video image. Therefore, the resolution and brightness of the transmitted video image is limited by the length of the fibre optic bundle. Additionally, the longer the fibre optic bundle, the more cumbersome it is to carry the bundle and associated fibre optic equipment into and out of the MRI scanner tunnel.
- A fibre optic bundle is comprised of a plurality of optical fibres. When an optical fibre is interrupted, the pixels of light of the transmitted image carried by the interrupted fibre are blocked. This situation results in dead pixels, e.g., black spots that appear on the video display. As the length of the fibre optic bundle is

increased, the probability that individual fibres will be broken increases. Further, as the fibre optic bundle is bent and manipulated over a period of time, the number of broken fibres increases. An increasing number of broken fibres results in a greater number of black spots appearing on the video display. Eventually, the transmitted image becomes inadequate and distorted. Thus, long fibre optic bundles are not cost effective.

During an MRI examination, the patient is positioned upon an examination table which can be moved into and out of the MRI scanner tunnel. When lying upon the examination table within the scanner tunnel, the patient's head is positioned within a head coil. The head coil is arranged to surround the patient's head and to provide MRI images thereof. An advanced design of MRI scanner head coils minimizes the distance between the patient's eyes and the top of the head coil. The limited distance between the patient's head and the head coil would be inadequate to accommodate the goggles employed by known MRI fibre optic systems that (a) position the image from the LCD display within the scanner tunnel or (b) employ a reflecting mirror over the patient's eyes.

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The advance of the functional imaging field requires implementation of visual activation paradigms that are becoming more sophisticated. During functional imaging, the best results are achieved when the visual stimulus is controlled which is inconsistent with an open field of view. Further, this method of visual activation does not include the ability to generate three-dimensional (3D) images for patient viewing since the image is projected onto a single screen. The inability to create a condition is which the eye and brain perceive a 3D effect prevents virtual reality from being achieved.

Further, the development of new and smaller head coils limits the distance between the patient's head and the head coil, putting restraints on the size of the goggles to be used within the head coil. Together with the introduction of MR machines with higher field strength both in the clinical and research field, the shielding of the MR goggles to avoid generation of any stray magnetic fields or

disruption of signal modulation by radio frequency is becoming increasingly important.

The use of functional imaging in clinical work also requires devices that are fast and easy to set up and operate in a tight clinical schedule. Easy positioning of the device and effective eye correction features are crucial elements to achieve a satisfactory clinical workflow.

Thus, there is a need in the art for an improvement in video systems for use with MRI scanners which provide high resolution video images with a three-dimensional effect, shortens the transmission paths that the video image must travel, eliminates the problems associated with fibre optic bundles, is sized to fit within the limited space of modern head coil designs, is sufficiently shielded to avoid image artefacts and can be mounted and operated within the MRI magnetic field.

#### **SUMMARY**

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According to the present invention, the above mentioned improvements are solved by means of an apparatus according to the characterizing clauses of claims 1 and 6. Further preferred embodiments and improvements are obtained by the features given in the dependent claims.

### BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 is a view of a preferred embodiment of the present invention illustrating the entire visual system separated between a control room and a magnet room.

- FIG. 2 is a view of a preferred embodiment of the visual system showed used in a MRI system bore.
- FIG. 3 is a view of a preferred embodiment of the visual system showing the adjustable arm, the housing of the micro displays chip and optics, the pupil distance adjustment knob and the dioptic compensator knobs.

FIG. 4 is a view of the adjustment arm and pupil distance adjuster

FIG. 5 is an exploded view of the arrangement of the micro display housing.

# 5 DETAILED DESCRIPTION OF THE INVENTION

The following detailed description outlines an MRI compatible visual system having a head coil mounted micro display. In the following description, numerous details such as specific materials and configurations are set forth in order to provide a more complete understanding of the present invention. But it is understood by those skilled in the art that the present invention can be practiced without these specific details. In other instances, well known elements are not described in detail so as not to obscure the present invention. In any event, the scope of the invention is best determined by reference to the appended claims.

# 15 GENERAL ARRANGEMENT

In a preferred embodiment, the present invention provides an MRI compatible visual system. FIG. 1 gives a general overview of how the present invention system is set up in relation to the MRI system, which is disposed partly in a magnet room 1 and partly in a control room 2.

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One portion of the present invention system is located inside the MRI control room 2. That portion of the system includes a fibre optical transmitter 9. The control room 2 also contains a PC 8. Dashed lines in FIG. 1 circumscribe the borders of that room. Everything outside the dashed lines represents the examination or magnet room 1. The other portion of the system that includes a head coil mounted micro display and its circuitry are located within the magnet room 1. As the name implies, the magnet room 1 contains a main magnet 3 of the MRI device that generates a strong magnetic field.

Continuing with the general overview, FIG. 1 shows that the system contained in the magnet room 1 is again divided such that certain parts of the system are mounted inside the bore of the main magnet and other parts remain outside the bore. The parts inside the bore include adjustable head coil mounted micro

displays 4. Outside the bore, but still inside the magnet room 1 includes fibre optical receiver unit 6 and micro display control electronics 7.

#### CONTROL ROOM PART

In the control room 2 a fibre optical transmission unit 9 receives signals from a PC 8. The fibre optical transmitter converts electrical signals into light signals. A fibre optic cable 10 is used to bring the light signal into the magnet room 1. The fibre cable 10 is mounted via a wave guide 11. A typical MRI signal is very sensitive to the electrical noise around the procession frequency of a hydrogen proton, wherein this frequency varies from 12 MHz to 130 MHz depending on the field strength of the magnet. This relationship is generally expressed as:

f=42.5\*B

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wherein B is the field strength in Tesla and f is the frequency in Megahertz.

Mindful of the foregoing relationship, the dimensions of the wave guide is calculated so that only non-disturbing frequencies will enter the magnet room. The wave guide 11 is typically a tube mounted on the Faraday cage that surrounds the magnet room (not shown).

#### MAGNET ROOM PART

The fibre optic cable 10 is fed into the magnet room 1 where the other part of the visual system is located. In the magnet room 1 there is a fibre optic receiver unit 6. This converts the light signals into electrical signals. The signals enter a micro display driver unit 7. This unit controls the micro display chips 14 (Fig 5) and controls image rotation, colour adjustments, automatic shut down and other functions.

The micro display chips are mounted on an adjustable arm 15 (Fig 2). The mechanism is designed to make it easy for the patient 5 to adjust the visual system into the right angle for preferred view.

Between the micro display driver unit 7 and the micro display chips 14 there are a shielded cable 12. The shielded cable 12 brings the electrical signals into a small faradays cage 13 (Fig 5) that contains the micro display 14.

# 5 ADJUSTMENT OF THE MICRO DISPLAY HOUSING.

Main viewing angle adjustment

FIG. 2 illustrates the adjustment mechanism on the head coil mounted micro display 4. The mechanism consists of a coil attachment part 16 two distance arms 38, 39 with friction links 40, 41, 42 and friction adjustment knobs 19 (Fig 4).

The adjustable arm allows people lying in different positions in the head coil to have a quick adjustment to get the micro display housing 20 (Fig 3) in the right position for best angle of view. The distance arms 38, 39 in combination with the friction links 40, 41, 42 let the micro display housing 20 move both in horizontal and vertical direction. The angle of the micro display housing 20 can also be changed. The coil attachment element 16 is arranged to fit different head coils 21. The system is arranged to fit several different kinds of coil attachment elements 16, thereby facilitating fitment in different head coil systems.

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Pupil distance adjustment mechanism.

According to the present invention, the system has adjustments for pupil distance. The distance is adjusted by turning knob 22. The knob 22 will move the micro display housing 20 symmetrical away from and against each other.

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The knob 22 is fixed to a shaft 23. The shaft can rotate inside the mid section 24. The movement of the micro display housings 20 is fixed into one direction with two bolts 25. The shaft 23 is threaded with opposite threads on each side of mid section 24. Turning the shaft 23 will make a symmetrical movement on the micro display housings 20.

Micro display housing arrangement

The micro display chips 14 are mounted and fixed inside Faraday cages 13 mounted in the micro display housings 20. The Faraday cages 13 shield the sensitive electronics against disturbances generated by the RF pulses from the scanner 3. The Faraday cages 13 also shield against electromagnetic noise generated by the micro display 14. In the lover part of the Faraday cage 13, there is arranged a shielded window 28. The shielded window 28 comprises wires or a mesh of metal that keeps the shielding function intact, but also let the patient 5 see the image on the micro display 14.

Between the patient's eye and the micro display there are arranged optical elements 29 that enlarge the image from the micro display 14. The optical elements 29 are designed so the micro display 14 is mounted a couple of centimetres away from the patient's eye. The design of the optical elements decreases the problem of distortion on the MRI system caused by the electronics and display chip 14.

The knob 30 facilitates dioptic correction for patients that normally wear glasses. Ajusting the knob 30 will change the distance between the micro display 14 and the optical elements 29. The motion of the Faraday cage 13 is locked into one direction by two bolts 31. The Faraday cage 13 is threaded inside. The knob 30 is fixed to the end of a shaft 32. The shaft 32 is threaded, and by turning the knob 30, the Faraday cage will slide along the bars 31. The micro display 14 is fixed inside the Faraday cage 13, thereby ensuring that the movement of the micro display 14 will equal to the movement of the Faraday cage 13. The optical elements 29 are arranged to ensure that a linear movement of the micro display 14 will result a dioptic correction that follows the normal steps used in the dioptic scale. For example, turning the knob 30 half a turn may give an adjustment of one step on the dioptic scale.

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Around the housing of the optical elements 37 there are softer materials 33 that will cover the patient's eye completely. The cover avoids the patient to be disturbed by ambient light.

As a part of the optical system there is mounted a beam splitter. The beam splitter is designed to let the visible light pass, but the light in the infra red frequency area is led into another optical direction, the eye tracker channel 35. The beam splitter allows an infrared image of the patient's eye to enter the eye tracker channel 35.

The image of the patient eye is fed into the end of a coherent image guide 36. In the other end of the coherent image guide 36, it is possible to connect an infrared camera to capture the movement of the patient's gaze. To ensure a crisp and bright image in the infrared camera end of the coherent image guide, the patient's eye is lit up with infrared light. This infrared light is fed into the magnet bore by one ore more fibre optic cables (not shown).

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- 1. Apparatus for providing high resolution images to patients positioned in a magnetic resonance imaging (MRI) device (3), the MRI device (3) comprising a head coil (21), the head coil arranged to surround a patient's (5) head and to provide MRI images thereof, the apparatus comprising means (12, 13, 14) for receiving video or picture image signals from an external source, c h a r c t e r i z e d in that the apparatus further comprises means (14, 28, 34, 29) for displaying a video or picture image, said display means being arranged in a housing (20), said housing (20) being suspended in an arm (15) comprising at least two successive members (38, 39), wherein a joint (40) between the housing (20) and the adjacent member (38), a joint or joints (41) between the successive members (38, 39), and a joint (42) between an attachment element (16) for attaching the apparatus to the head coil (21), or other part of the MRI device (3), and the member (39) adjacent to the coil attachment element (16), each is hinged to allow rotation of the joints (40, 41, 42).
  - 2. Apparatus according to claim 1, c h a r c t e r i z e d i n that the housing (20) comprises two optical apertures (33), the distance between said optical apertures (33) being adjustable.
  - 3. Apparatus according to claim 1 or 2, c h a r c t e r i z e d i n that the hinged joints (40, 41, 42) comprise friction links.
- 4. Apparatus according to one of the previous claims, c h a r c t e r i z e d i n that the coil attachment element (16) is arranged to fit different head coils (21) or other part of the MRI device (3).
  - 5. Apparatus according to one of the previous claims, c h a r c t e r i z e d i n that the apparatus further comprises means (14, 28, 34, 29) for displaying a video or picture image, said display means (14, 28, 34, 29) being arranged in an upper part of a housing (20), said display means (14, 28, 34, 29) comprising a micro display chip (14) inside a Faraday cage (13), the micro display chip (14)

comprising means for converting a video or picture image signal to a visible image on a small screen, the Faraday cage at its lower end comprising a shielded window (28), the apparatus between the patient's eye(s) and the micro display(s) (14) comprising optical elements (29) in a lower part of the housing (20) that adjustably enlarges the image from the micro display (14), wherein the distance between the display means (14, 28, 34, 29) in the upper part of the housing (20), and the optical elements (29) in a lower part of the housing (20) is such that no sensitive or disturbing electronic parts are located inside the head coil (21) during the operation of the MRI device (3), while the part of the housing (20) that is inside the head coil (21) during operation of the MRI device (3) only comprises elements that do not significantly affect the MRI and are not significantly affected by the magnetic and RF fields inside the head coil (21).

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Apparatus for providing high resolution images to patients positioned in a 6. magnetic resonance imaging (MRI) device (3), the MRI device (3) comprising a head coil (21), the head coil arranged to surround a patient's (5) head and to provide MRI images thereof, the apparatus comprising means (12, 13, 14) for receiving video or picture image signals from an external source, c h a r c t e r i z e d in that the apparatus further comprises means (14, 28, 34, 29) for displaying a video or picture image, said display means (14, 28, 34, 29) being arranged in an upper part of a housing (20), said display means (14, 28, 34, 29) comprising a micro display chip (14) inside a Faraday cage (13), the micro display chip (14) comprising means for converting a video or picture image signal to a visible image on a small screen, the Faraday cage at its lower end comprising a shielded window (28), the apparatus between the patient's eye(s) and the micro display(s) (14) comprising optical elements (29) in a lower part of the housing (20) that adjustably enlarges the image from the micro display (14), wherein the distance between the display means (14, 28, 34, 29) in the upper part of the housing (20), and the optical elements (29) in a lower part of the housing (20) is such that no sensitive or disturbing electronic parts are located inside the head coil (21) during operation of the MRI device (3), while the part of the housing (20) that is inside the head coil (21) during the operation of the MRI device (3) only comprises elements

that do not significantly affect the MRI and are not significantly affected by the magnetic and RF fields inside the head coil (21).

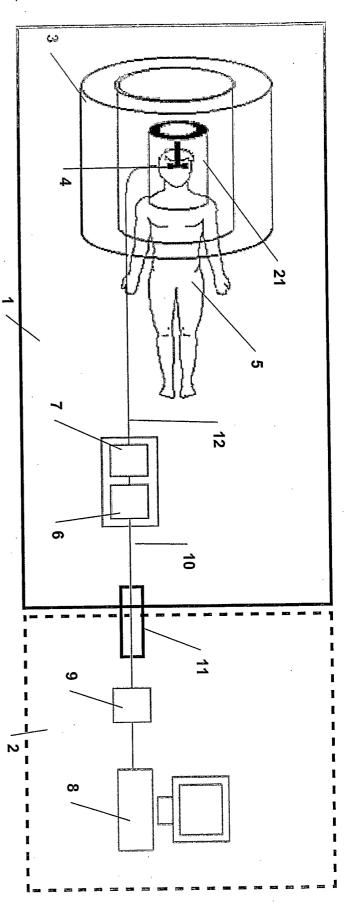
7. Apparatus according to claim 6, c h a r a c t e r i z e d i n that the distance between the upper and lower part of the housing (20) can be adjusted.

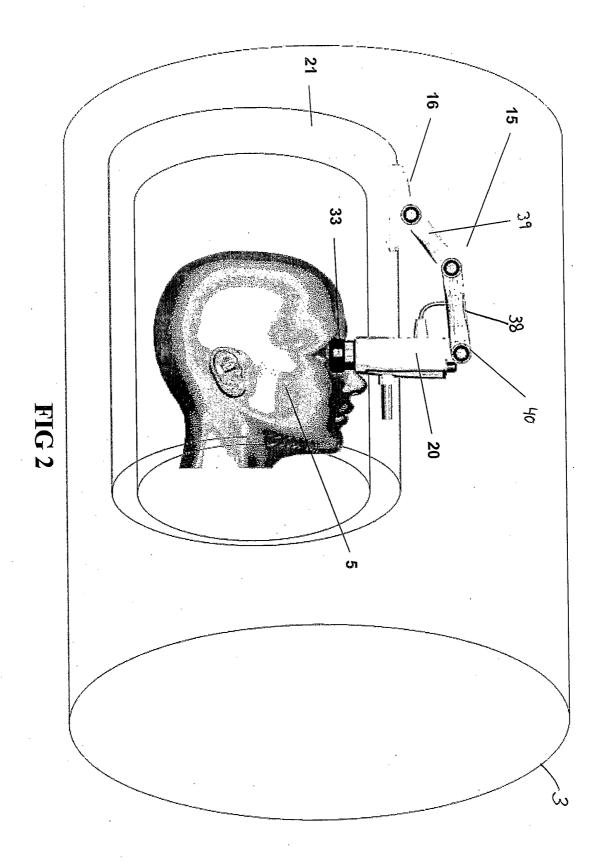
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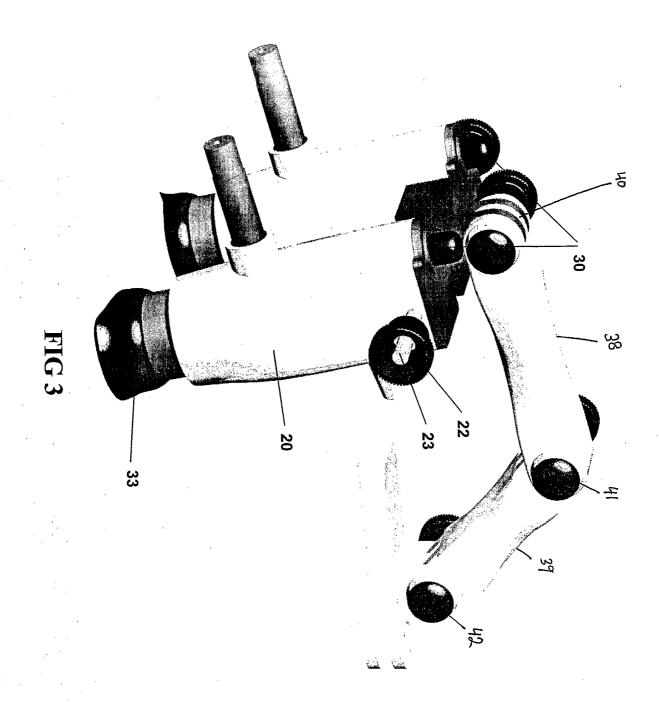
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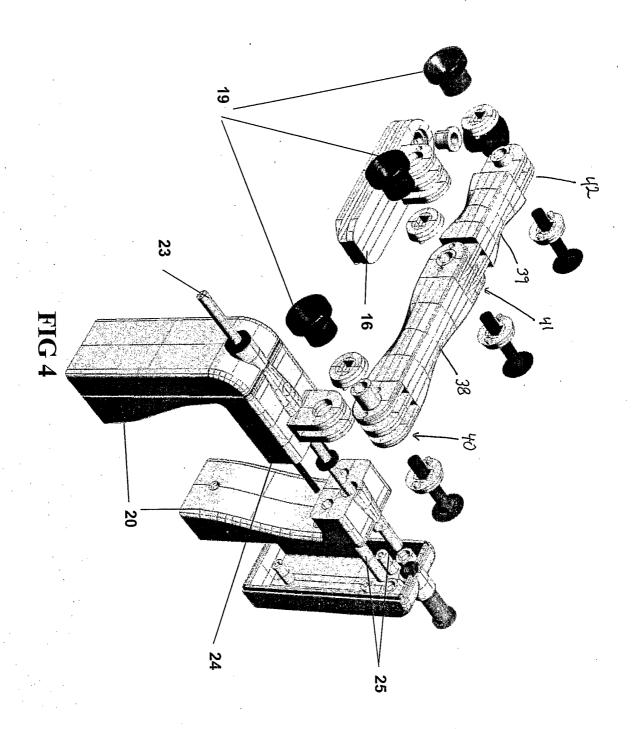
8. Apparatus according to claim 6 or 7, c h a r a c t e r i z e d i n that the apparatus further comprises means (14, 28, 34, 29) for displaying a video or picture image, said display means being arranged in a housing (20), said housing (20) being suspended in an arm (15) comprising at least two successive members (38, 39), wherein a joint (40) between the housing (20) and the adjacent member (38), a joint or joints (41) between the successive members (38, 39), and a joint (42) between an attachment element (16) for attaching the apparatus to the head coil (21), or other part of the MRI device (3), and the member (39) adjacent to the coil attachment element (16), each is hinged to allow rotation of the joints (40, 41, 42).

FIG 1











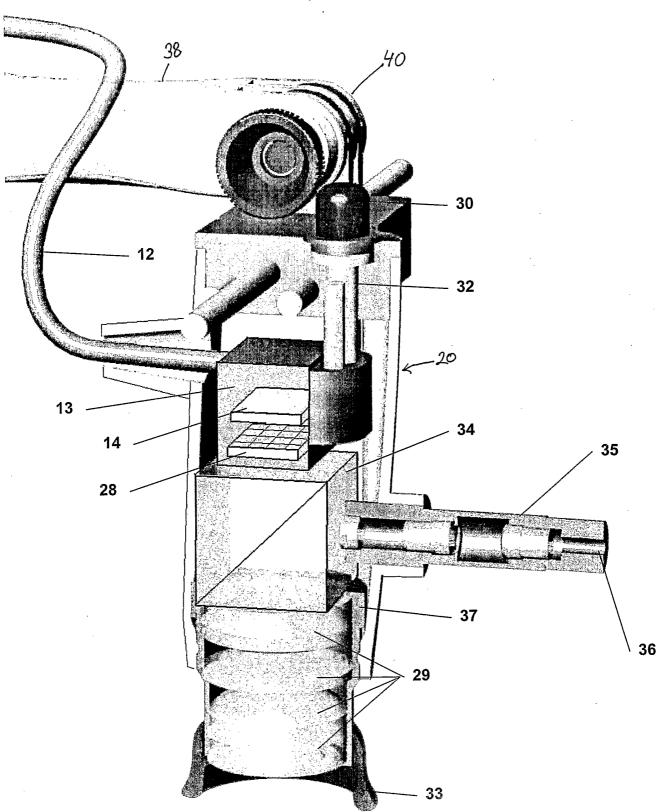


FIG 5

#### INTERNATIONAL SEARCH REPORT

International application No.

#### PCT/NO2006/000211

### A. CLASSIFICATION OF SUBJECT MATTER

IPC: see extra sheet
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)

#### IPC: A61B, G01R, G09G

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

#### SE,DK,FI,NO classes as above

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

#### EPO-INTERNAL, WPI DATA, PAJ

Facsimile No. +46 8 666 02 86

#### C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Х	US 5877732 A (MOKHTAR ZIARATI), 2 March 1999 (02.03.1999)	6-7
Y		1-5,8
Ρ,Χ	WO 2005119284 A1 (INVITO CORPORATION), 15 December 2005 (15.12.2005)	6-7
Ρ,Υ		1-5,8
	<b></b>	
Y	US 20020163499 A1 (FRANK SAUER), 7 November 2002 (07.11.2002)	1-5,8
A		6-7

Х	Further documents are listed in the continuation of Box	<b>C.</b>	See patent family annex.	
*	Special categories of cited documents:	"T"	later document published after the international filing date or priority	
"A"	document defining the general state of the art which is not considered to be of particular relevance		date and not in conflict with the application but cited to understand the principle or theory underlying the invention	
"E"	earlier application or patent but published on or after the international filing date		document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive	
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12	12 Sept 2006		<b>2</b> 6 -09- 2006	
Name and mailing address of the ISA/		Authorized officer		
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# INTERNATIONAL SEARCH REPORT

International application No. PCT/N02006/000211

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C (Continu	nation). DOCUMENTS CONSIDERED TO BE RELEVANT	
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US 4852500 A (JON B RYBURG ET AL), 1 August 1989 (01.08.1989)	1-5,8
A		6-7
	<del></del>	
A	US 5706070 A (STANLEY M. REICH ET AL), 6 January 1998 (06.01.1998), abstract	1-8
	<del></del>	
A	US 5339813 A (EDGAR A. DEYOE ET AL), 23 August 1994 (23.08.1994), abstract	1-8
	bang bart	
A	DATABASE WPI Week 200363 Derwent Publications Ltd., London, GB; Class P31, AN 2003-667305 & JP 203190112A (GE MEDICAL SYSTEMS GLOBAL TECHNOLOGY CO) 8 July 2003 (2003-07-08) abstract	1-8
	· <b>e</b> va sen	
A	US 5864331 A (PREM K. ANAND ET AL), 26 January 1999 (26.01.1999), abstract	1-8
A	DE 3844482 C1 (BRUKER MEDIZINTECHNIK GMBH), 31 December 1988 (31.12.1988), abstract	1-8
	•	
	·	

#### International patent classification (IPC)

A61B 5/055 (2006.01) G01R 33/28 (2006.01)

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Cited literature, if any, will be enclosed in paper form.

# INTERNATIONAL SEARCH REPORT Information on patent family members

04/03/2006

International application No. PCT/N02006/000211

US	5877732	Α	02/03/1999	NONE
WO	2005119284	A1	15/12/2005	US 20050273000 A 08/12/2005
US	20020163499	A1	07/11/2002	US 6919867 B 19/07/2005 US 20020082498 A 27/06/2002
US	4852500	A	01/08/1989	CA 1294025 A,C 07/01/1992 DE 3862133 D 00/00/0000 EP 0283016 A,B 21/09/1988 JP 2620801 B 18/06/1997 JP 63231607 A 27/09/1988
US	5706070	Α	06/01/1998	NONE
US	5339813	Α	23/08/1994	NONE -
US	5864331	Α	26/01/1999	US 5861865 A 19/01/1999
DE	3844482	C1	31/12/1988	WO 9007301 A 12/07/1990