

AZIENDA USL TOSCANA SUD EST

GROSSETO

"Diagnostic Imaging Area" Radiology Department

Director: Dr. LUCA FRANCI U.O.S.D. Neuro-Radiology, resp: Dr. MARCO CIRINEI Tel 0564/485238

MEDICAL HISTORY FORM AND INFORMED CONSENT FOR MRI SCAN TEST

(Name, Surname, Profes	ssion of the MRI team staff member)
Patient: Surname	
Date and Place of birth	Heiaht (ka) Weiaht
(m) Address	Tel
MRI Exam Type Ward/Doctor requirng MRI scan	

Informations about MRI scan test

Magnetic Resonance Imaging (MRI) is a diagnostic technique that does not use ionizing radiation or radioactive substances. MRI diagnostics exploit the use of intense static fields of magnetic induction and RF electromagnetic waves. In some types of investigation, some substances with paramagnetic properties can also be administered intravenously to the patient as a contrast medium. With the exception of these cases, MRI is configured as a non-invasive diagnostic test.

In case of examinations on pregnant women (pregnancy confirmed or presumed), particular attention, especially in cases of urgency, is paid to the justification and to the optimization of the MRI examination, with particular regard of the patient and the unborn child. During the MRI examination, the onset of adverse reactions is very rare. The most probable occurrence is represented by a crisis of claustrophobia of a transient nature.

The use of a contrast agent based on paramagnetic substances is generally well tolerated and does not cause any particular sensation. Rarely however, episodes of hypersensitivity as urticaria or other allergic phenomena may occur.

In very rare cases episodes of anaphylactic shock have been reported.

In cases of previous or suspected allergy to investigations with MDC o with bronchial asthma or urticaria, angioedema or a history of ideopathic anaphylaxis can be used premedication according to the outline of the prepared form.

The RM site always guarantees the presence of specialized medical personnel ready to intervene in case of medical emergencies of this kind.

MRI SCAN TEST

Patients can do the MRI scan only after excluding any possible contraindication to the exam. That needs to be checked in advance by the Doctor in charge of the diagnostic service (MRP), using the specific medical history questionnaire and informed consent form.

To carry out the MRI examination it is necessary that the patient, where necessary supported by the MRI team staff:

- remove any face make-up and hairspray;
- deposits in the locker room or in the special lockers any metal, ferromagnetic or magnetic holder
 (e.g. mobile phones, coins, watches, keys, earrings, brooches, jewelry, paper clips hair, magnetic cards, credit cards, pocket knives, money clips, automatic hooks, brooches, zipped clothes, metal tweezers, files, scissors, etc.);
- remove any dental prostheses and hearing aids;
- · remove contact lenses or glasses;
- undress, and then put on the special disposable gown provided by the service staff;
- use the headset or ear plugs provided by the MRI team staff member.

The average duration of the MRI exam is approximately 30 minutes, but it can vary according to clinical needs and the number of anatomical districts to check. During the data acquisition phase of the MRI exam, rhythmic noises of variable intensity are audible. That noises are caused by the normal operation of the MRI scanner.

Ventilation, lighting and temperature conditions are provided to ensure a maximum well-being and reduce possible claustrophobic effects. During the exam phase you need to remain calm and keep the maximum degree of immobility due not to compromise the diagnostic result of the images. Breath regularly and swallowing saliva do not disturb the examination.

However in some types of MRI scan test it may be required to collaborate with respiratory acts and short periods of apnea for improving the quality of the imaging diagnostics.

In the control room there is always service personnel ready to intervene in case of any need. The patient is always in vocal, acoustic and visual contact with the operators, who carry out constant checks throughout the examination phase. If you experience discomfort, such as a feeling of claustrophobia, heat, itching, breathlessness, palpitations or fainting, the patient should notify the staff as soon as possible present, using the appropriate signaling devices.

PRELIMINARY QUESTIONNAIRE

The anamnestic questionnaire aims to ascertain the absence of contraindications to the MRI examination or the non-pertinence of specific preventive insights. This questionnaire must be carefully filled out and signed by the Doctor in charge of the Diagnostic Service, who, in relation to the answers provided by the patient, can conclude that there are no contraindications to the MRI examination. The countersignature of the patient on foot of the same page, at the bottom of the consent formula, guarantees - among others, also its full consent awareness of the serious consequences that false or mendacious answers to the questions submitted to him may have.

□ Have you previously had MRI exams?	Yes	No
□ Have you had any allergic reactions after administration		
of the contrast medium?	Yes	No
□ Do you suffer from claustrophobia?	Yes	No
□ Have you ever worked (or do you work) as a welder,	103	140
turner, coachbuilder?	Yes	No
□ Has he ever had road accidents, hunting accidents?	Yes	No
□ Was he a victim of trauma from an explosion?		
Last monetrial period.	Yes	No
□ Last menstrual period:		

□ You had surgery on:		
head neck		
abdomen extremities		
chest other:		
□ Are you aware that you have one or more medical		
devices or metal bodies inside your body?	Yes	No
□ Do you have a cardiac pacemaker or other types		
of cardiac catheters?	Yes	No
□ Are you carrying splinters or metal fragments?	Yes	No
□ Do you have Clips on aneurysms (blood vessels).		
aorta, brain?	Yes	No
□ Heart valves?	Yes	No
□ Stents?	Yes	No
□ Implanted defibrillators?	Yes	No
□ Spine distractors?	Yes	No
□ Infusion pump for insulin or other medications?	Yes	No
□ Metal bodies in the ears or hearing implants?	Yes	No
□ Neurostimulators, electrodes implanted in the		
brain or subdurals?	Yes	No
□ Other types of stimulators?	Yes	No
□ Intrauterine bodies?	Yes	No
□ Spinal or ventricular shunt?	Yes	No
□ Fixed or mobile dentures?	Yes	No
□ Metal prostheses (for previous fractures,		
joint corrective operations, etc.), screws, nails, wire, etc.?	Yes	No
□ Other prostheses?	Yes	No
Localization:		
□ Do you believe you may have prostheses/devices or other		
metallic bodies inside your body that you're NOT aware of?	Yes	No
Additional information		
□ Do you have sickle cell anemia?	Yes	No
□ Are you a wearer of a lens prosthesis?	Yes	No
□ Are you a piercing wearer?	Yes	No
Localization:		
□ Do you have tattoos?	Yes	No
Localization:		
□ Are you using medical patches?	Yes	No

To carry out the MRI examination it is necessary to remove any: contact lenses; hearing aids; dentures; mobile temporary crowns; hernial girdle; clips for hair; clothespins; eyeglasses; jewels; clocks; credit cards or other magnetic cards; pocket knives; stop money; coins; keys; hooks; automatic; metal buttons; pins; clothes with zipper; nylon stockings; garments in acrylic; metal tweezers; metal points; limes; scissors; any other metal objects. Before undergo the examination please remove cosmetics from the face.

The Radiologist Responsible for carrying out the MRI examination (*) having acknowledged the answers provided by the patient and once the possible medical examination and/or further preliminary diagnostic investigations have been carried out,

AUTHORIZES the execution of the MR investigation.

Doctor's signature	Date
	······································
INFORM	MED CONSENT TO THE MRI EXAMINATION
The patient believes has sufficie to exposure to electromagnetic field	ently been informed about the associated risks and contraindications ds generated by the MRI equipment. Therefore, aware of the importance of the answers provided,
Α	AGREES to the execution of the exam.
Patient's signature (**)	Date
INFORMED CONSENT	TO THE ADMINISTRATION OF A CONTRAST MEDICATION
of the contrast medication. Made a	ntly been informed about the risks associated with the administration ware, by the Doctor in charge of the diagnostic performance, about the n of the diagnostic benefits and related risks,
	AGREES to their administration.
Patient's signature (**)	Date
that upon administration of the contrast medium, the is then recommended; if ap any further consents signed by	rge of the diagnostic performance and the informed consents linked both to the execution of the MRI investigation e signature of the patient must necessarily be affixed on a single sheet; possibly also in duplex mode. It oplicable; to predict consecutively (on the same sheet or on a separate model) the patient, made in the same way as those shown here by way of example only. If a minor patient, the signature of a parent or guardian is required.
	Further informed consents
INFORMED CONSEI (Add an expla	NT TOanatory note on the risks associated with the specific procedure)
associated to	onsiders himself sufficiently informed on the risks
	GIVES the consent.
Patient's signature (**)	Date

Appendix

ACCESS CARD (*) (**)

Reserved for visitors, companions and all those who access the CONTROLLED AREA

The medical history check has the purpose of ascertaining the absence of contraindications exposure to risks associated with intense electromagnetic fields present in the ZONES CHECK within the RM SITE. This questionnaire must be carefully completed and signed at the bottom by the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND THE DIAGNOSTIC EFFECTIVENESS OF THE EQUIPMENT RM or another delegated doctor, who, in relation to the answers provided, is required to assess whether there are any contraindications to access.

SURNAME NAME		
BORN (state/city) D	ATE OF BIRTH	
(Indicate if visitor, companion or other)		
□ Have you ever worked (or do you work) as a v		
turner, coachbuilder?	Yes	No
☐ Has he ever had road accidents, hunting accidents		No
□ Was he a victim of trauma from an explosion?□ You had surgery on:	Yes	No
head neck		
abdomen extremities		
chest o other:		
□ Are you aware that you have one or more med		
devices or metal bodies inside your body?	Yes	No
□ Do you have a cardiac pacemaker or other typ		
of cardiac catheters?	Yes	No
Are you carrying splinters or metal fragments?	Yes	No
☐ Do you have Clips on aneurysms (blood vesse	els),	
aorta, brain?	Yes	No
□ Heart valves?	Yes	No
□ Stents?	Yes	No
□ Implanted defibrillators?	Yes	No
□ Spine distractors?	Yes	No
□ Infusion pump for insulin or other medications'		No
□ Metal bodies in the ears or hearing implants?	Yes	No
Neurostimulators, electrodes implanted in the brain or subdurals?	V	
□ Other types of stimulators?	Yes Yes	No No
□ Intrauterine bodies?	Yes	No
□ Spinal or ventricular shunt?	Yes	No
□ Fixed or mobile dentures?	Yes	No
	103	110
□ Metal prostheses (for previous fractures,		
joint corrective operations, etc.), screws, nails,	wire, etc.? Yes	No
Other prostheses?	Yes	No
Localization:		
Do you believe you may have prostheses/devi		
metallic bodies inside your body that you're NO	T aware of? Yes	No

□ Are you a wearer of a lens prosthesis? □ Are you a piercing wearer? Localization: □ Are you using medical patches? □ Additional information			
□ Are you a wearer of a lens prostness? □ Are you a piercing wearer? Localization: □ Are you using medical patches? □ Additional information			
□ Are you a piercing wearer? Localization: □ Are you using medical patches? □ Additional information	- Are you a wearer of a lens prosthesis?		
Localization: Are you using medical patches? Additional information	Are you a piercing wearer?	Yes	No
□ Are you using medical patches? □ Additional information	Tocalization.		
□ Are you using medical patches? □ Additional information	Localization		
To access the CONTROLLED AREA you need to remove: any contact lenses - hearing aids - dentures - temporary mobile crowns - hernia belt - hair clips - barrettes - glasses - jewelery - watches - credit cards or other magnetic cards - pocket knives - money clips - coins - keys - hooks - press-studs - metal buttons - brooches - clothes with zips - nylon stockings – clothes in acrylic - metal tweezers - staples - files - scissors - any other objects metallic. Objects or devices cannot be brought into the CONTROLLED AREA electrical unless expressly authorized by the personnel present on the designated RM site accompanying the subject during his presence. The residence time within the risk zones it must be limited to what is strictly necessary to carry out the activities for which access has been allowed and in any case in the maximum conditions of optimization of its security. THE RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOSTIC EFFECTIVENESS OF THE MRI EQUIPMENT (*) or its delegate acknowledged the answers provided by the subject and carried out all the necessary checks authorize access to the RM site Signature of the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOST EFFECTIVENESS OF THE MRI EQUIPMENT or its delegate Doctor's signature Date Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated and the MIA engineers. Therefore, aware of the importance of the answers provided, he logs	Are you using medical patches?	Yes	No
To access the CONTROLLED AREA you need to remove: any contact lenses - hearing aids - dentures - temporary mobile crowns - hernia belt - hair clips - barrettes - glasses - jewelery - watches - credit cards or other magnetic cards - pocket knives - money clips - coins - keys - hooks - press-studs - metal buttons - brooches - clothes with zips - nylon stockings - clothes in acrylic - metal tweezers - staples - files - scissors - any other objects metallic. Objects or devices cannot be brought into the CONTROLLED AREA electrical unless expressly authorized by the personnel present on the designated RM site accompanying the subject during his presence. The residence time within the risk zones it must be limited to what is strictly necessary to carry out the activities for which access has been allowed and in any case in the maximum conditions of optimization of its security. THE RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOSTIC EFFECTIVENESS OF THE MRI EQUIPMENT (*) or its delegate acknowledged the answers provided by the subject and carried out all the necessary checks authorize access to the RM site Signature of the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOST EFFECTIVENESS OF THE MRI EQUIPMENT or its delegate Doctor's signature Date Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated The MEN accidence of the answers provided, he logs	Additional information		
any contact lenses - hearing aids - dentures - temporary mionic crowns' hernia belt - hair clips - barrettes - glasses - jewelery - watches - credit cards or other magnetic cards - pocket knives - money clips - coins - keys - hooks - press-studs - metal buttons - brooches - clothes with zips - nylon stockings - clothes in acrylic - metal tweezers - staples - files - scissors - any other objects metallic. Objects or devices cannot be brought into the CONTROLLED AREA electrical unless expressly authorized by the personnel present on the designated RM site accompanying the subject during his presence. The residence time within the risk zones it must be limited to what is strictly necessary to carry out the activities for which access has been allowed and in any case in the maximum conditions of optimization of its security. THE RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOSTIC EFFECTIVENESS OF THE MRI EQUIPMENT (*) or its delegate acknowledged the answers provided by the subject and carried out all the necessary checks authorize access to the RM site Signature of the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOST EFFECTIVENESS OF THE MRI EQUIPMENT or its delegate Doctor's signature Date Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated and the MRI Requirement Therefore, aware of the importance of the answers provided, he logs the MRI Requirement Therefore aware of the importance of the answers provided, he logs the MRI Requirement Therefore aware of the importance of the answers provided, he logs the MRI Requirement of the importance of the answers provided, he logs the MRI Requirement of the importance of the answers provided, he logs the MRI Requirement of the importance of the answers provided, he logs the manufacture of the answers provided, he logs the manufacture of the answers provided.			
any contact lenses - hearing aids - dentures - temporary mionic crowns' hernia belt - hair clips - barrettes - glasses - jewelery - watches - credit cards or other magnetic cards - pocket knives - money clips - coins - keys - hooks - press-studs - metal buttons - brooches - clothes with zips - nylon stockings - clothes in acrylic - metal tweezers - staples - files - scissors - any other objects metallic. Objects or devices cannot be brought into the CONTROLLED AREA electrical unless expressly authorized by the personnel present on the designated RM site accompanying the subject during his presence. The residence time within the risk zones it must be limited to what is strictly necessary to carry out the activities for which access has been allowed and in any case in the maximum conditions of optimization of its security. THE RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOSTIC EFFECTIVENESS OF THE MRI EQUIPMENT (*) or its delegate acknowledged the answers provided by the subject and carried out all the necessary checks authorize access to the RM site Signature of the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOST EFFECTIVENESS OF THE MRI EQUIPMENT or its delegate Doctor's signature Date Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated and the MRI Requirement Therefore, aware of the importance of the answers provided, he logs the MRI Requirement Therefore aware of the importance of the answers provided, he logs the MRI Requirement Therefore aware of the importance of the answers provided, he logs the MRI Requirement of the importance of the answers provided, he logs the MRI Requirement of the importance of the answers provided, he logs the MRI Requirement of the importance of the answers provided, he logs the manufacture of the answers provided, he logs the manufacture of the answers provided.	그리카를 가장 살아서 그는 가장 살아 있다면 하는 것이 없었다.		
any contact lenses - hearing aids - dentures - temporary mionic crowns' hernia belt - hair clips - barrettes - glasses - jewelery - watches - credit cards or other magnetic cards - pocket knives - money clips - coins - keys - hooks - press-studs - metal buttons - brooches - clothes with zips - nylon stockings - clothes in acrylic - metal tweezers - staples - files - scissors - any other objects metallic. Objects or devices cannot be brought into the CONTROLLED AREA electrical unless expressly authorized by the personnel present on the designated RM site accompanying the subject during his presence. The residence time within the risk zones it must be limited to what is strictly necessary to carry out the activities for which access has been allowed and in any case in the maximum conditions of optimization of its security. THE RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOSTIC EFFECTIVENESS OF THE MRI EQUIPMENT (*) or its delegate acknowledged the answers provided by the subject and carried out all the necessary checks authorize access to the RM site Signature of the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOST EFFECTIVENESS OF THE MRI EQUIPMENT or its delegate Doctor's signature Date Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated and the MRI Requirement Therefore, aware of the importance of the answers provided, he logs the MRI Requirement Therefore aware of the importance of the answers provided, he logs the MRI Requirement Therefore aware of the importance of the answers provided, he logs the MRI Requirement of the importance of the answers provided, he logs the MRI Requirement of the importance of the answers provided, he logs the MRI Requirement of the importance of the answers provided, he logs the manufacture of the answers provided, he logs the manufacture of the answers provided.	To access the CONTROLLED AREA you need to remove:		
hernia belt - hair clips - barrettes - giasses - jewelery other magnetic cards - pocket knives - money clips - coins - keys - hooks - press-studs - metal buttons - brooches - clothes with zips - nylon stockings – clothes in acrylic - metal tweezers - staples - files - scissors - any other objects metallic. Objects or devices cannot be brought into the CONTROLLED AREA electrical unless expressly authorized by the personnel present on the designated RM site accompanying the subject during his presence. The residence time within the risk zones it must be limited to what is strictly necessary to carry out the activities for which access has been allowed and in any case in the maximum conditions of optimization of its security. THE RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOSTIC EFFECTIVENESS OF THE MRI EQUIPMENT (*) or its delegate acknowledged the answers provided by the subject and carried out all the necessary checks authorize access to the RM site Signature of the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOST EFFECTIVENESS OF THE MRI EQUIPMENT or its delegate Doctor's signature Doctor's signature Date Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated The person in Charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated	dentures - temporary mou	olle crowns -	
other magnetic cards - pocket knives - money clipts - boths - keys - keys - clothes in press-studs - metal buttons - brooches - clothes with zips - nylon stockings - clothes in acrylic - metal tweezers - staples - files - scissors - any other objects metallic. Objects or devices cannot be brought into the CONTROLLED AREA electrical unless expressly authorized by the personnel present on the designated RM site accompanying the subject during his presence. The residence time within the risk zones it must be limited to what is strictly necessary to carry out the activities for which access has been allowed and in any case in the maximum conditions of optimization of its security. THE RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOSTIC EFFECTIVENESS OF THE MRI EQUIPMENT (*) or its delegate acknowledged the answers provided by the subject and carried out all the necessary checks authorize access to the RM site Signature of the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOST EFFECTIVENESS OF THE MRI EQUIPMENT or its delegate Doctor's signature Date Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated and the contraindications related to exposure to the electromagnetic fields generated for the MRI equipment. Therefore, aware of the importance of the answers provided, he logs			,
press-studs - metal buttons - brooches - clothes wild zpps - Hybrids - Middle - Clothes wild zpps - Hybrids - Middle - Clothes wild zpps - Hybrids - Scissors - any other objects metallic. Objects or devices cannot be brought into the CONTROLLED AREA electrical unless expressly authorized by the personnel present on the designated RM site accompanying the subject during his presence. The residence time within the risk zones it must be limited to what is strictly necessary to carry out the activities for which access has been allowed and in any case in the maximum conditions of optimization of its security. THE RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOSTIC EFFECTIVENESS OF THE MRI EQUIPMENT (*) or its delegate acknowledged the answers provided by the subject and carried out all the necessary checks authorize access to the RM site Signature of the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOST EFFECTIVENESS OF THE MRI EQUIPMENT or its delegate Doctor's signature Date Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated to the total contraindications related to exposure to the electromagnetic fields generated to the text of the importance of the answers provided, he logs			
acrylic - metal tweezers - staples - files - scissors - any other objects metallic. Objects or devices cannot be brought into the CONTROLLED AREA electrical unless expressly authorized by the personnel present on the designated RM site accompanying the subject during his presence. The residence time within the risk zones it must be limited to what is strictly necessary to carry out the activities for which access has been allowed and in any case in the maximum conditions of optimization of its security. THE RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOSTIC EFFECTIVENESS OF THE MRI EQUIPMENT (*) or its delegate acknowledged the answers provided by the subject and carried out all the necessary checks authorize access to the RM site Signature of the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOST EFFECTIVENESS OF THE MRI EQUIPMENT or its delegate Doctor's signature Date Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated to the total contraindications related to exposure to the electromagnetic fields generated to the stable of the answers provided, he logs	motal buttons - brooches - Cloudes Will 2105 - 11	ylori otooimige	
Metallic. Objects or devices cannot be brought into the CONTROLLED AREA electrical unless expressly authorized by the personnel present on the designated RM site accompanying the subject during his presence. The residence time within the risk zones it must be limited to what is strictly necessary to carry out the activities for which access has been allowed and in any case in the maximum conditions of optimization of its security. THE RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOSTIC EFFECTIVENESS OF THE MRI EQUIPMENT (*) or its delegate acknowledged the answers provided by the subject and carried out all the necessary checks authorize access to the RM site Signature of the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOST EFFECTIVENESS OF THE MRI EQUIPMENT or its delegate Doctor's signature Date Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated for the MRI equipment. Therefore aware of the importance of the answers provided, he logs	acrylic - metal tweezers - staples - files - scissors - any other	objects	
electrical unless expressly authorized by the personner presence. The residence time within the risk zones it must be limited to what is strictly necessary to carry out the activities for which access has been allowed and in any case in the maximum conditions of optimization of its security. THE RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOSTIC EFFECTIVENESS OF THE MRI EQUIPMENT (*) or its delegate acknowledged the answers provided by the subject and carried out all the necessary checks authorize access to the RM site Signature of the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOST EFFECTIVENESS OF THE MRI EQUIPMENT or its delegate Doctor's signature Date Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated the supportance of the answers provided, he logs	metallic.		
electrical unless expressly authorized by the personner presence. The residence time within the risk zones it must be limited to what is strictly necessary to carry out the activities for which access has been allowed and in any case in the maximum conditions of optimization of its security. THE RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOSTIC EFFECTIVENESS OF THE MRI EQUIPMENT (*) or its delegate acknowledged the answers provided by the subject and carried out all the necessary checks authorize access to the RM site Signature of the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOST EFFECTIVENESS OF THE MRI EQUIPMENT or its delegate Doctor's signature Date Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated the supportance of the answers provided, he logs	CONTROLLE	DARFA	
accompanying the subject during his presence. The testication within the risk zones it must be limited to what is strictly necessary to carry out the activities for which access has been allowed and in any case in the maximum conditions of optimization of its security. THE RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOSTIC EFFECTIVENESS OF THE MRI EQUIPMENT (*) or its delegate acknowledged the answers provided by the subject and carried out all the necessary checks authorize access to the RM site Signature of the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOST EFFECTIVENESS OF THE MRI EQUIPMENT or its delegate Doctor's signature Date Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated that the MRI equipment Therefore aware of the importance of the answers provided, he logs	Objects or devices cannot be brought into the CONTROLLER	ent on the design	ated RM site
within the risk zones it must be limited to what is strictly necessary in activities for which access has been allowed and in any case in the maximum conditions of optimization of its security. THE RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOSTIC EFFECTIVENESS OF THE MRI EQUIPMENT (*) or its delegate acknowledged the answers provided by the subject and carried out all the necessary checks authorize access to the RM site Signature of the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOST EFFECTIVENESS OF THE MRI EQUIPMENT or its delegate Doctor's signature Date Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated the MRI equipment. Therefore aware of the importance of the answers provided, he logs	electrical unless expressly authorized by the personner president	ce time	
activities for which access has been allowed and in any case in the measure optimization of its security. THE RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOSTIC EFFECTIVENESS OF THE MRI EQUIPMENT (*) or its delegate acknowledged the answers provided by the subject and carried out all the necessary checks authorize access to the RM site Signature of the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOST EFFECTIVENESS OF THE MRI EQUIPMENT or its delegate Doctor's signature Date Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated to the MDI contraindications related to exposure to the electromagnetic fields generated the MDI contraindications related to exposure to the electromagnetic fields generated the MDI contraindications related to exposure to the electromagnetic fields generated the MDI contraindications related to exposure to the electromagnetic fields generated the MDI contraindications related to exposure to the electromagnetic fields generated the MDI contraindications related to exposure to the electromagnetic fields generated the MDI contraindications related to exposure to the electromagnetic fields generated the most of the importance of the answers provided, he logs			it the
THE RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOSTIC EFFECTIVENESS OF THE MRI EQUIPMENT (*) or its delegate acknowledged the answers provided by the subject and carried out all the necessary checks authorize access to the RM site Signature of the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOST EFFECTIVENESS OF THE MRI EQUIPMENT or its delegate Doctor's signature Date Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated and exposure to the electromagnetic fields generated and exposure to the electromagnetic fields generated to exposure to the electromagnetic fields generated and exposure to the electromagnetic fields generated to exposure to the electromagnetic fields generated	within the risk zones it must be limited to what is stretty need	in the maximum	conditions of
THE RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOSTIC EFFECTIVENESS OF THE MRI EQUIPMENT (*) or its delegate acknowledged the answers provided by the subject and carried out all the necessary checks authorize access to the RM site Signature of the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOST EFFECTIVENESS OF THE MRI EQUIPMENT or its delegate Doctor's signature Date Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated and exposure and	activities for which access has been allowed and in any sass		
acknowledged the answers provided by the subject and carried out all the necessary checks authorize access to the RM site Signature of the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOST EFFECTIVENESS OF THE MRI EQUIPMENT or its delegate Doctor's signature Date Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated for the MRI agripment. Therefore, aware of the importance of the answers provided, he logs	optimization of its security.		
acknowledged the answers provided by the subject and carried out all the necessary checks authorize access to the RM site Signature of the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOST EFFECTIVENESS OF THE MRI EQUIPMENT or its delegate Doctor's signature Date Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated for the MRI agripment. Therefore, aware of the importance of the answers provided, he logs			-1100710
acknowledged the answers provided by the subject and carried out all the necessary checks authorize access to the RM site Signature of the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOST EFFECTIVENESS OF THE MRI EQUIPMENT or its delegate Doctor's signature Date Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated and on the contraindications related to exposure to the electromagnetic fields generated for the MRI agripment. Therefore, aware of the importance of the answers provided, he logs	THE RADIOLOGIST RESPONSIBLE FOR CLINICA	AL SAFETY AND	OF THE DIAGNOSTIC
acknowledged the answers provided by the subject and carried out all the necessary checks authorize access to the RM site Signature of the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOST EFFECTIVENESS OF THE MRI EQUIPMENT or its delegate Doctor's signature Date Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated to the person of the answers provided, he logs	FFFECTIVENESS OF THE MRI EQU	JIPMENT (*) or its	s delegate
Signature of the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOST EFFECTIVENESS OF THE MRI EQUIPMENT or its delegate Doctor's signature Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated to the answers provided, he logs			
Signature of the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOST EFFECTIVENESS OF THE MRI EQUIPMENT or its delegate Doctor's signature Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated to the answers provided, he logs	acknowledged the answers provided by the subject	and carried out	all the necessary checks
Signature of the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND OF THE DIAGNOST EFFECTIVENESS OF THE MRI EQUIPMENT or its delegate Doctor's signature Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated to the MRI agreement. Therefore, aware of the importance of the answers provided, he logs			
Doctor's signature Date Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated to exposure to the answers provided, he logs	authorize access to t	THE INVISITE	
Doctor's signature Date Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated to exposure to the answers provided, he logs			
Doctor's signature Date Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated to exposure to the answers provided, he logs	THE PARIOLOGIST RESPONSIBLE FOR CLIN	NICAL SAFETY A	ND OF THE DIAGNOSTIC
Doctor's signature Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated to exposure of the electromagnetic fields generated to exposure to exposure to the electromagnetic fields generated to exposure to exposure to exposure to exposure t	Signature of the RADIOLOGIST RESPONSIBLE FOR SELECTION OF THE MPI FOLLIPMENT or its delega	ite	
Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated to exposure of the enswers provided, he logs	EFFECTIVENESS OF THE WIRI EQUIL MELLE STATES		
Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated to exposure of the enswers provided, he logs			
Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated to exposure to the electromagnetic fields generated the MRI organization. Therefore, aware of the importance of the answers provided, he logs	Doctor's signature	Dat	e
Informed consent The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated and the MRI organization. Therefore, aware of the importance of the answers provided, he logs	Doctor a signature		
The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated			
The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated			
The person in charge of access believes he has been sufficiently informed about the risks and on the contraindications related to exposure to the electromagnetic fields generated	L.f. mad on	ncont	
and on the contraindications related to exposure to the electromagnetic management and the MRI agreement. Therefore, aware of the importance of the answers provided, he logs	나이 이 그렇게 잘 맛있었다. 얼마 그렇지 그 없는 그는 그렇게 되었다. 그 그 그 그 그 그 그 그 그 그 그 그 그 그 그 그 그 그 그		
and on the contraindications related to exposure to the electromagnetic management and the MRI agreement. Therefore, aware of the importance of the answers provided, he logs	- : I was of seeses holioves he has	heen sufficiently	informed about the risks
form the MDI equipment Therefore aware of the Importance of the difference	The person in charge of access believes the trace	e to the electroma	agnetic fields generated
to the RM site aware of the risks present.	and on the contraindications related to exposure	portance of the a	nswers provided, he logs o
	to the RM site aware of	the risks present.	
	to the rain one division		
게 크리겠다면서 2000년 1월 12 - 12 - 12 - 12 - 12 - 12 - 12 - 1			Date
Signature of the person in charge of access (**)	Signature of the person in charge of access (**)		Date

.......

(*) Verification of medical history signed by the RADIOLOGIST RESPONSIBLE FOR CLINICAL SAFETY AND EFFECTIVENESS

MRI EQUIPMENT DIAGNOSTICS and the informed consent signed by the person in charge of access must be

necessarily affixed to a single sheet, possibly also in front/back mode.

(**) In order to simplify the access procedures, once the form has been filled in for the first time, for subsequent accesses it is possible to foresee

also the possibility of confirming at each subsequent entry that nothing has changed in the subject for the purpose of verifying the

contraindications provided in the anamnestic questionnaire, his awareness of the risks present in the MRI site and of the

knowledge of the procedures to be followed, providing for the signature of the subject, of the RADIOLOGICAL DOCTOR RESPONSIBLE FOR

CLINICAL SAFETY AND DIAGNOSTIC EFFECTIVENESS OF THE MRI EQUIPMENT that authorized access and date.