

Cochlear™ Nucleus® Implants MRI Safety Checklist

This form guides you through the critical aspects of performing an MRI scan safely for patients with a Cochlear[™] Nucleus[®] implant. Before using this form, review the Cochlear Nucleus MRI Guidelines, available on the website:

www.cochlear.us/mri

Ensure the implant site has healed before an MRI scan is performed. The cochlear implant physician should be consulted if there are any concerns.

Follow these steps prior to inviting the patient into the MRI room and before performing the MRI scan.

Identify and record the Cochlear Nucleus Implant model using the Cochlear patient identification card, X-ray or surgical/clinical notes in the spaces provided.
Implant model number for left ear:
Implant model number for right ear:
2 Determine if implant magnet removal is required, or if a Cochlear Nucleus MRI Kit is necessary.

- Implant magnet removal may be mandatory at certain field strengths for specific implant types. Refer to **Table 2: MRI safety information and recommended SAR limits**.
- Implant magnet removal may be necessary to reduce the artefact. Refer to the MRI Guidelines for artefact sizes. Metal Artefact Reduction Sequences (MARS) are recommended for optimal results.
- Implant magnet removal may be avoided at certain field strengths with use of a Cochlear Nucleus MRI Kit.

The intent of an MRI Kit is to provide pressure over the implant magnet – not the implant body. Refer to **Table 2: MRI safety information and recommended SAR limits**.



Note

If there is no implant magnet present, then an MRI Kit is not required.

Circle the applicable option:

Implant magnet will be in place for scan / Implant magnet will not be in place for scan

Circle the applicable option:

MRI Kit is required for scan / MRI Kit is not required for scan

	3 Record the MR parameters and transmit coil to be used for this scan in the spaces provided.						
	MRI field strength:						
	Maximum spatial gradient the patient will encounter entering the MRI bore:						
	Circle the applicable transmit coil to be used: Whole body coil / Head coil / Other local cylindrical transmit coil						
	Identify and record the SAR limit and maximum allowable spatial gradient for the implant type and MRI RF transmit coil in the spaces provided. Refer to Table 1 and Table 2 .						
	Note If bilaterally implanted, a patient may have two different cochlear implant types. Comply with the lowest SAR limit and the lowest allowable spatial gradient of the two devices.						
	Lowest SAR limit for recipient's implant types and scan conditions:						
	Lowest allowable spatial gradient for recipient's implant types:						
	Counsel the patient on sensations and risks.						
	 For patients where an implant magnet is in place, explain that they might feel the implant magnet moving slightly and might sense resistance to movement as pressure on the skin. 						
	 For devices which require an MRI Kit, the MRI Kit will reduce the likelihood of the implant magnet moving. The sensation will be similar to pressing down firmly on the skin with the thumb. 						
	Refer to the MRI Guidelines for a complete list of Warnings and Cautions.						
	Remove the sound processor and any accessories before entering the MRI room. The sound processor is MR Unsafe.						
	Note The patient may no longer be able to hear instructions with the sound processor removed.						
	7 Prior to entering the MRI room, apply the MRI Kit if determined it is required in step 2.						
	Ensure you have the contents of the MRI Kit available and within easy reach.						
	 Full instructions are supplied with MRI Kits, or are listed in the MRI Guidelines. 						
	• Apply the MRI Kit contents to the implant site or sites in accordance with the information in Table 2 .						
8	Comply with patient positioning requirements.						
	 For safety, the patient should be in a supine position, lying flat on back with face upward, prior to entering the MRI bore. 						
	 Align the patient's head with the bore axis of the MRI machine (max 15° deviation allowed from the z-axis). 						
	 Advise the patient to lie as still as possible and to not move their head during the MRI scan. 						
	 Correctly position the patient prior to the MRI scan to minimise discomfort and reduce the risk of 						

implant magnet demagnetisation.

Scan conditions and SAR limits

- The MRI safety information in the tables below only applies to 1.5 T and 3 T MRI horizontal scanners (closed bore or wide bore) with a circularly polarised (CP) RF field.
- MRI scans at 3 T must be performed in quadrature mode or CP mode for the radio frequency (RF) transmit coil.
- Using a multichannel mode may result in localised heating above safe levels.
- Maximum active scan time of 60 minutes with the SAR limits in **Table 2** below.
- The SAR limit depends on the selection of MRI RF transmit coil.
 Refer to the Specific conditions for RF transmit coil in Table 1 below.

RF transmit coil	Specific conditions				
Main scanner body coil	 Local cylindrical RF receive only coils can be placed anywhere, with respect to the implant. Comply with the whole body average SAR limit for the relevant implant type, field strength and landmark location. See Table 2. 				
Transmit/receive head coil	Comply with the head SAR limit for the relevant implant model and field strength. See Table 2 .				
Other local transmit/ receive coils (e.g. knee)	 Ensure distance between coil and implant is greater than the coil radius. There are no added SAR restrictions due to the presence of the implant. Limit SAR as you would for a typical patient who does not have an implantable device. 				

Table 1: Specific conditions for RF transmit coil

Implant type	MRI field strength (T)	ength implant magnet		Maximum allowable spatial gradient	Head average SAR limit (W/kg) Using transmit /	Whole body average SAR limit (W/kg) Landmark location <40 cm from ≥40cm from					
	()		Yes/No	field (T/m)	receive head coil	top of head	top of head				
CI600 Series implants											
CI612, CI622, CI624, CI632	1.5	No	No	20	<2	<1	<2				
CI612	3	No	No	20	<1	<0.5	<1				
CI622, CI624, CI632	3	No	No	20	<1	<0.4	<1				
CI500 Series implants											
CI512, CI522, CI532, ABI541	1.5	No	Yes	20	<2	<1	<2				
CI512, ABI541	3	Yes	No	20	<1	<0.5	<1				
CI522, CI532	3	Yes	No	20	<1	<0.4	<1				
CI24RE Series implants											
CI422, CI24REH (Hybrid L24)	1.5	No	Yes	20	<2	<1	<2				
CI24RE (CA), CI24RE (ST)	3	Yes	No	20	<1	<0.5	<1				
CI24R & CI24M Series implants											
CI24R (CA), CI24R (CS),	1.5	No	Yes	20	<2	<1	<2				
CI24R (ST), CI24M, ABI24M	3	Yes	No	20	<1	<0.5	<1				
CI 11+11+2M	1.5	No	Yes	20	<1	<0.5	<1				
	3	MRI is contraindicated									
CI22M Series implants											
CI22M with	1.5	No	Yes	20	<2	<1	<2				
removable magnet	3	MRI is contraindicated									
CI22M without	1.5	MRI is contraindicated									
removable magnet	3										

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As the global leader in implantable hearing solutions, Cochlear is dedicated to helping people with moderate to profound hearing loss experience a life full of hearing. We have provided more than 600,000 implantable devices, helping people of all ages to hear and connect with life's opportunities.

We aim to give people the best lifelong hearing experience and access to innovative future technologies. We collaborate with the industry's best clinical, research and support networks.

That's why more people choose Cochlear than any other hearing implant company.

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ACE, Advance Off-Stylet, AOS, Ardium, AutoNRT, Autosensitivity, Baha, Baha SoftWear, BCDrive, Beam, Bring Back the Beat, Button, Carina, Cochlear, 科利耳, コクレア, 코클리어, Cochlear SoftWear, Contour, コントゥア, Contour Advance, Custom Sound, DermaLock, Freedom, Hear now. And always, Hugfit, Human Design, Hybrid, Invisible Hearing, Kanso, LowPro, MET, MP3000, myCochlear, mySmartSound, NRT, Nucleus, Osia, Outcome Focused Fitting, Off-Stylet, Piezo Power, Profile, Slimline, SmartSound, Softip, SoundArc, True Wireless, the elliptical logo, Vistafix, Whisper, WindShield and Xidium are either trademarks or registered trademarks of the Cochlear group of companies.

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