

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:

<u>www.cochlear.com/mri</u>

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 Implant Card, X-ray or surgical/clinical notes in the spaces provided.

- Implant model number for left ear:
- Implant model number for right ear:

Determine if implant magnet removal is required or if a Cochlear MRI Kit (bandage and splint) 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 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**.



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



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 guadrature mode or CP mode for the radio free
- 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 RE transmit coil

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	 Keep local planar (flat linearly polarised) receive only RF coils more than 10 cm away from cochlear implant. 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	Remove implant magnet Yes/No	MRI Kit required Yes/No	Maximum allowable spatial gradient field (T/m)	Head average SAR limit (W/kg) Using transmit / receive head coil	Whole body average SAR limit (W/kg) Landmark location			
	(T)					<40 cm from top of head	≥40cm from 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		
		C	124RE Ser	ies implants					
CI422, CI24REH (Hybrid L24) CI24RE (CA), CI24RE (ST)	1.5	No	Yes	20	<2	<1	<2		
	3	Yes	No	20	<1	<0.5	<1		
		CI24R	& CI24N	l Series implan	ts				
CI24R (CA), CI24R (CS), CI24R (ST), CI24M, ABI24M	1.5	No	Yes	20	<2	<1	<2		
	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							
		C	I22M Ser	ies implants					
CI22M with removable magnet	1.5	No	Yes	20	<2	<1	<2		
	3	MRI is contraindicated							
CI22M without	1.5	MRI is contraindicated							
removable magnet	3								

Table 2: MRI safety information and recommended SAR limits*

* Not all products are available in all countries. Please contact your local Cochlear representative for product information.

Hear now. And always

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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This material is intended for health professionals. If you are a consumer, please seek advice from your health professional about treatments for hearing loss. Outcomes may vary, and your health professional will advise you about the factors which could affect your outcome. Always read the instructions for use. Not all products are available in all countries. Please contact your local Cochlear representative for product information.

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