### MRI Checklist for MED-EL CI models

Mi1200 SYNCHRONY | Mi1200 SYNCHRONY PIN | Mi1210 SYNCHRONY ST Mi1250 SYNCHRONY 2 | Mi1250 SYNCHRONY 2 PIN

If the conditions or instructions herein are NOT followed, INJURY to the patient and/or DAMAGE to the implant may result!

- ightarrow VALID for all intracochlear electrode variants
- ightarrow VALID for all body regions

In case of additional implants, e.g. a hearing implant in the other ear: MRI safety guidelines for this implant need to be considered in addition.

The external components of the MED-EL Implant System (audio processor and accessories) are MR Unsafe and need to be removed prior to scanning.

The implant components of the MED-EL Implant System are MR Conditional.



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#### GENERAL CONDITIONS

GENERAL CONDI					
→ PERMITTED STATIC MAGNETIC FIELD STRENGTH			0.2 T, 1.0 T, 1.5 T, 3.0 T		
ightarrow maximum per	RMITTED SAR (at 0.2T, 1.0T, 1.5T)		NORMAL OPERATING MODE 3.2 W/kg (Head), 2.0 W/kg (Whole-Body)		
ightarrow maximum per	RMITTED SAR (at 3.0T)	Hea Whole Body <35 cm from the top of the hea Whole Body ≥35 cm from the top of the hea			
		that are less than 35 cm from the top of the on rate (SAR) or to display the estimated ma			
PREPARATION					
1. PATIENT ID CAR Request patient ID	RD card in order to identify the implant type	2	(	🔿 ок	
2. IMAGE ARTEFACT       ○ YES → continue (with step 3.)         Is an accurate diagnosis possible even with the expected image artefact?       ○ NO → decide if the magnet should be rem					
MAGNET REMOVAL $\bigcirc$ YES $\rightarrow$ continueHas the implant magnet been surgically removed? $\bigcirc$ NO $\rightarrow$ STOP			→ continue (with step 3.) > STOP		
<ol> <li>AUDITORY/NON-AUDITORY SENSATIONS         Inform the patient about possible auditory and non-auditory sensations during the examination.     </li> <li>NOTE: The likelihood and intensity of auditory and non-auditory sensations can be reduced by selecting sequences with a lower Specific Absorption Rate (SAR) and slower gradient slew rates.     </li> </ol>					
<ol> <li>HEAD ORIENTATION (only applicable at 1.0T, 1.5T, 3.0T) Inform the patient not to tilt their head to either side.</li> </ol>				⊖ ок	
5. EXTERNAL COMPONENTS Remove audio processor and accessories before entering the scanner room.				🔿 ок	
<ul> <li>OPTIONAL HEAD BANDAGE</li> <li>A supportive head bandage over the implant using an elastic bandage wrapped tightly around the head for at least three times can optionally be used.</li> </ul>					
NOTE: In rare cases the patient might perceive a clicking sound upon entry in the MRI scanner tube.					
EXECUTION					
NOTE: To reduce the likelihood and degree of patient discomfort, the patient should keep their head away from the scanner wall near the entrance of the scanner.					
<ol> <li>PATIENT POSITIONING (only applicable at 1.0T, 1.5T, 3.0T) The patient should be lying in the scanner in a supine, prone or side position with the head kept straight.</li> <li>NOTE: When lower extremities are to be examined, it is recommended that patient's legs are positioned in the scanner first.</li> </ol>				⊖ ок	
2. OPERATING MODE (only applicable at 0.2T, 1.0T, 1.5T) Run sequences in "Normal Operating Mode" only. NOTE: max. 3.2W/kg for Head scans, 2.0W/kg for Whole-Body scans				⊖ ок	
3. OPERATING MODE (only applicable at 3.0T)         Apply maximum permitted SAR according to following table only:         SAR (Head)       SAR (Whole-Body)				⊖ ок	
	<35 cm from the top of the head	≥35 cm from the top of the head			
1.6 W/kg	1.0 W/kg	2.0 W/kg			
4. ACCESSORIES (only applicable at 3.0T) Do not utilise head transmit coils or multi-channel transmit coils.				⊖ ок	

NOTE: In rare cases temporary pain or discomfort may occur in the implant area during MRI even if all protocols and bandaging instructions are followed.

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## MRI Checklist for MED-EL ABI models

Mi1200 SYNCHRONY ABI | Mi1200 SYNCHRONY PIN ABI

The external components of the MED-EL Implant If the conditions or instructions herein are NOT followed, INIURY to the MR System (audio processor and accessories) are MR patient and/or DAMAGE to the implant may result! Unsafe and need to be removed prior to scanning. → VALID for ABI electrode variant  $\rightarrow$  VALID for all body regions In case of additional implants, e.g. a hearing implant in the other ear: The implant components of the MED-EL Implant MRI safety guidelines for this implant need to be considered in addition. System are MR Conditional. GENERAL CONDITIONS → **PERMITTED** STATIC MAGNETIC FIELD STRENGTH 0.2 T. 1.0 T. 1.5 T

- → MAXIMUM PERMITTED SAR NORMAL OPERATING MODE 3.2 W/kg (Head), 2.0 W/kg (Whole-Body) PREPARATION 1. PATIENT ID CARD Request patient ID card in order to identify the implant type 2. IMAGE ARTEFACT  $\bigcirc$  YES  $\rightarrow$  continue (with step 3.) Is an accurate diagnosis possible even with the expected image artefact?  $\bigcirc$  NO  $\rightarrow$  decide if the magnet should be removed MAGNET REMOVAL  $\bigcirc$  YES  $\rightarrow$  continue (with step 3.) Has the implant magnet been surgically removed?  $\bigcirc$  NO  $\rightarrow$  STOP 3. AUDITORY/NON-AUDITORY SENSATIONS Inform the patient about possible auditory and non-auditory sensations during the examination. NOTE: The likelihood and intensity of auditory and non-auditory sensations can be reduced by selecting sequences with a lower Specific Absorption Rate (SAR) and slower gradient slew rates. 4. HEAD ORIENTATION (only applicable at 1.0T, 1.5T) Inform the patient not to tilt their head to either side. 5. EXTERNAL COMPONENTS Remove audio processor and accessories before entering the scanner room. - OPTIONAL HEAD BANDAGE A supportive head bandage over the implant using an elastic bandage wrapped tightly around the head for at least three times can be used. NOTE: In rare cases the patient might perceive a clicking sound upon entry in the MRI scanner tube. NOTE: To reduce the likelihood and degree of patient discomfort, the patient should keep their head away from the scanner wall
- near the entrance of the scanner. 1. PATIENT POSITIONING (only applicable at 1.0T, 1.5T) O OK The patient should be lying in the scanner in a supine, prone or side position with the head kept straight. NOTE: When lower extremities are to be examined, it is recommended that patient's legs are positioned in the scanner first. О ОК 2. OPERATING MODE Apply "Normal Operating Mode" only. NOTE: max. 3.2 W/kg for Head scans, 2.0 W/kg for Whole-Body scans

NOTE: In rare cases temporary pain or discomfort may occur in the implant area during MRI even if all protocols and bandaging instructions are followed.

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### MRI Checklist for MED-EL CI and ABI models

Mi1000 CONCERTO | Mi1000 CONCERTO PIN | SONATA Mi1000 CONCERTO ABI | Mi1000 CONCERTO PIN ABI

If the conditions or instructions herein are NOT followed, INJURY to the patient and/or DAMAGE to the implant may result!

- $\rightarrow$  VALID for all electrode variants
- ightarrow VALID for all body regions

GENERAL CONDITIONS

In case of additional implants, e.g. a hearing implant in the other ear: MRI safety guidelines for this implant need to be considered in addition.

The external components of the MED-EL Implant System (audio processor and accessories) are MR Unsafe and need to be removed prior to scanning.

The implant components of the MED-EL Implant System are MR Conditional.



#### → PERMITTED STATIC MAGNETIC FIELD STRENGTH 0.2 T, 1.0 T, 1.5 T → MAXIMUM PERMITTED SAR NORMAL OPERATING MODE, i.e. 3.2 W/kg (Head), 2.0 W/kg (Whole-Body) PREPARATION 1. PATIENT ID CARD O OK Request patient ID card in order to identify the implant type 2. IMAGE ARTEFACT $\bigcirc$ YES $\rightarrow$ continue Is an accurate diagnosis possible even with the expected image artefact? $\bigcirc$ NO $\rightarrow$ Stop 3. AUDITORY/NON-AUDITORY SENSATIONS О ОК Inform the patient about possible auditory and non-auditory sensations during the examination. NOTE: The likelihood and intensity of auditory and non-auditory sensations can be reduced by selecting sequences with a lower Specific Absorption Rate (SAR) and slower gradient slew rates. 4. HEAD ORIENTATION (only applicable at 1.0T, 1.5T) O OK Inform the patient not to tilt their head to either side. 5. EXTERNAL COMPONENTS ○ OK Remove audio processor and accessories before entering the scanner room. 6. HEAD BANDAGE (only applicable at 1.0T, 1.5T) O OK Place a supportive headband over the implant. NOTE: The bandage may be an elastic bandage wrapped tightly around the head at least three times. EXECUTION NOTE: To reduce the likelihood and degree of patient discomfort, the patient should keep their head away from the scanner wall near the entrance of the scanner. 1. PATIENT POSITIONING (only applicable at 1.0T, 1.5T) О ОК The patient should be lying in the scanner in a supine, prone or side position with the head kept straight. NOTE: When lower extremities are to be examined, it is recommended that patient's legs are positioned in the scanner first to minimise any risk of weakening the implant magnet. 2. OPERATING MODE О ОК Run sequences in "Normal Operating Mode" only.

Run sequences in "Normal Operating Mode" only. NOTE: max. 3.2W/kg for Head scans, 2.0W/kg for Whole-Body scans

NOTE: In rare cases temporary pain or discomfort may occur in the implant area during MRI even if all protocols and bandaging instructions are followed.

wi1000 concerto abi i mi1000 concerto pin abi

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### MRI Checklist for MED-EL CI and ABI models

PULSAR | C40+ | C40 PULSAR ABI | C40+ ABI

If the conditions or instructions herein are NOT followed, INJURY to the patient and/or DAMAGE to the implant may result!  $\rightarrow$  VALID for all electrode variants

- $\rightarrow$  VALID for all body regions
- $\rightarrow$  VALID for unilateral as well as bilateral implant provision



The external components of the MED-EL Implant System (audio processor and accessories) are MR Unsafe and need to be removed prior to scanning.

The implant components of the MED-EL Implant System are MR Conditional.



GENERAL CONDITIONS					
→ PERMITTED STATIC MAGNETIC FIELD STRENGTH	0.2 Т, 1.0 Т, 1.5 Т				
→ MAXIMUM PERMITTED SAR	NORMAL OPERATING MODE, i.e. 3.2W/kg (Head), 2.0W/kg (Whole-Body)				
PREPARATION					
1. PATIENT ID CARD Request patient ID card in order to identify the implant type	○ ОК				
2. IMPLANT CONDITION Is the implant housing mechanically intact? (not fractured or shattered)	$\bigcirc YES \rightarrow continue \\ \bigcirc NO \rightarrow STOP$				
3. IMPLANTATION STATUS Has the implant been implanted for at least six months?	$\bigcirc YES \rightarrow continue \\ \bigcirc NO \rightarrow STOP$				
4. BONE THICKNESS Is the bone underneath the implant at least 0.4 mm thick?	$\bigcirc YES \rightarrow continue \\ \bigcirc NO \rightarrow STOP$				
5. IMAGE ARTEFACT Is an accurate diagnosis possible even with the expected image artefact?	$\bigcirc YES \rightarrow continue \\ \bigcirc NO \rightarrow STOP$				
6. AUDITORY/NON-AUDITORY SENSATIONS Inform the patient about possible auditory and non-auditory sensations during the examination. NOTE: The likelihood and intensity of auditory and non-auditory sensations can be reduced by selecting sequences with a lower Specific Absorption Rate (SAR) and slower gradient slew rates.					
7. HEAD ORIENTATION (only applicable at 1.0T, 1.5T) Inform the patient not to tilt their head to either side.	⊖ ок				
8. EXTERNAL COMPONENTS Remove audio processor and accessories before entering the scanner room.	○ ОК				
<ul> <li>9. HEAD BANDAGE (only applicable at 1.0T, 1.5T)</li> <li>Place a supportive headband over the implant.</li> <li>NOTE: The bandage may be an elastic bandage wrapped tightly around the head at least three times.</li> </ul>	○ ок				
EXECUTION					
NOTE: To reduce the likelihood and degree of patient discomfort, the patient should keep their head away from the scanner wall near the entrance of the scanner.					
<ol> <li>PATIENT POSITIONING (only applicable at 1.0T, 1.5T) The patient should be lying in the scanner in a supine, prone or side position with the head kept straight.</li> <li>NOTE: When lower extremities are to be examined, it is recommended that patient's legs are positioned in the scanner first to minimise any risk of weakening the implant magnet.</li> </ol>					
2. OPERATING MODE Run sequences in "Normal Operating Mode" only. NOTE: max. 3.2W/kg for Head scans, 2.0W/kg for Whole-Body scans					

NOTE: In rare cases temporary pain or discomfort may occur in the implant area during MRI even if all protocols and bandaging instructions are followed.

OULSAR ABI | C40+ ABI

Checklist available for download in printout format on www.medel.com/isi/