





MRI SAFETY INFORMATION

OPTIMIZER® Smart Mini Implantable Pulse Generator (Model: CCM X11)

MRI Safety Information

For Whole-Body MR Examinations: A person implanted with the OPTIMIZER Smart Mini System may be safely scanned anywhere in the body at 1.5T or 3.0T under the following conditions. Failure to follow these conditions may result in injury.

Parameter	Condition			
Device	The OPTIMIZER Smart Mini IPG must be implanted with two			
	ventricular leads and an optional atrial MR-conditional lead from the			
	list below. When combined, the OPTIMIZER Smart Mini IPG and			
	such leads constitute an MR conditional device system.			
	Lead [Manufacturer, Model] Length Lead's whole-			
		[cm]	body SAR limit	
			(See OEM lead	
			MRI labeling for	
			additional	
			information)	
	Medtronic, CapSureFix Novus	52	1.5T: 2.0 W/kg	
	MRI™ SureScan 5076	58	3T: B _{1+RMS} ≤ 2.8 µT	
	Medtronic, SelectSecure MRI™	59	1.5T: 2.0 W/kg	
	SureScan 3830	69	3T: B _{1+RMS} ≤ 2.8 µT	
	Abbott, 2088TC Tendril STS	52	2.0 W/kg	
		58		
	Boston Scientific, Ingevity+ 7841	52	2.0 W/kg	
	Boston Scientific, Ingevity+ 7842	59		
	Biotronik, Solia S-53	53	2.0 W/kg	
	Biotronik, Solia S-60	60	2.0 W/kg	
Device Configuration	OOO mode (CCM Stimulation OFF, Sensing OFF)			
Static Magnetic Field Strength	1.5T and 3T			
(B ₀)				
Type of Nuclei	Hydrogen			
MR Scanner Type	Cylindrical closed-bore magnet			
B ₀ Field Orientation	Horizontal			
Maximum Spatial Field	40 T/m			
Gradient				
Maximum Gradient Slew Rate	200 T/m/s per axis			
RF Excitation	Circularly Polarized (CP)			
RF Transmit Coil Type	Integrated Whole Body Transmit Coil. Local transmit coils may be			
	used but should not be placed directly over the OPTIMIZER Smart			
	Mini System.			

Parameter	Condition	
RF Receive Coil Type	Any	
Operating Mode	Normal Operating Mode	
RF Conditions	Whole-body SAR ≤ 2.0 W/kg or ≤ 3.2 W/kg head absorption rate for local head transmit coil.	
Scan Duration	Scan for up to 30 minutes. Wait 30 minutes before the next imaging session	
Scan Regions	Any landmark is acceptable	
Image Artifact	The presence of the OPTIMIZER Smart Mini System may produce an image artifact. In non-clinical testing, the maximum image artifact size was seen on the gradient echo pulse sequence at 1.5T and extends by approximately 6.7 cm from the boundary of the implant. At 3T the maximum image artifact extends by approximately 4.6 cm from the boundary of the implant. Some manipulation of scan parameters may be needed to compensate for the artifact.	

Restrictions for the Patient and the Implanted System

• There are no other active or abandoned cardiac implants (e.g., lead extensions, lead adapters, or abandoned leads) in the patient's body.

WARNING: Do not bring any system components that are not marked MR-safe or MR-conditional into the MRI suite.

- Other active or passive implants are permitted if they are identified as MR conditional by the manufacturer.
- At least six (6) weeks have elapsed since the OPTIMIZER Smart Mini IPG and/or lead implantation and/or any electrode revision or surgical modification.
- The device system is implanted pectorally.
- Patient does not have elevated body temperature or compromised thermoregulation at time of scan.
- **WARNING:** Do not scan a patient with an elevated body temperature.

Restrictions During the MRI scan

- Emergency equipment for resuscitation must be kept at hand and properly certified staff must be available.
- Patient must be continuously monitored by pulse oximetry and electrocardiography (ECG).

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