

Precision Montage™ MRI ImageReady MRI Full Body Patient Eligibility

This form provides information about the patient's implanted Precision Montage MRI Spinal Cord Stimulator System and MRI scan eligibility. Determination of whether an MRI is appropriate must be determined by the patient's physician.

- Prior to performing an MRI Scan, confirm that the patient's stimulation is OFF
- Refer to www.bostonscientific.com/imageready for labeling and safety conditions

Patient Name: _____

Date: _____

Physician Name: _____

Office Address: _____

Phone: _____

A.	MR CONDITIONAL PRECISION MONTAGE MRI SYSTEM INFORMATION	MODEL #	MRI FULL BODY ELIGIBLE	NOT MRI ELIGIBLE
1.	Implantable Pulse Generator (IPG)			
	• Precision Montage MRI IPG, 16-contact IPG	SC-1200	<input type="checkbox"/>	
	NOTE: If you have another model number IPG, please refer to the labeling specific to your IPG model number.			
2.	Percutaneous and/or surgical paddle leads (check all that apply)			
	• Avista™ MRI Percutaneous Lead, 8-contact lead, 56 cm	SC-2408-56	<input type="checkbox"/>	
	• Avista MRI Percutaneous Lead, 8-contact lead, 74 cm	SC-2408-74	<input type="checkbox"/>	
	• Linear™ Percutaneous Leads, 50 cm	SC-2158-50 SC-2138-50	<input type="checkbox"/> <input type="checkbox"/>	
	• Linear Percutaneous Leads, 70 cm	SC-2158-70 SC-2138-70	<input type="checkbox"/> <input type="checkbox"/>	
	• Linear ST Percutaneous Leads, 50 cm	SC-2218-50 SC-2208-50	<input type="checkbox"/> <input type="checkbox"/>	
	• Linear ST Percutaneous Leads, 70 cm	SC-2218-70 SC-2208-70	<input type="checkbox"/> <input type="checkbox"/>	
	• Linear 3-4 Percutaneous Leads, 50 cm	SC-2352-50	<input type="checkbox"/>	
	• Linear 3-4 Percutaneous Leads, 70 cm	SC-2352-70	<input type="checkbox"/>	
	• Linear 3-6 Percutaneous Leads, 50 cm	SC-2366-50	<input type="checkbox"/>	
	• Linear 3-6 Percutaneous Leads, 70 cm	SC-2366-70	<input type="checkbox"/>	
	• Infinion™ CX Percutaneous Leads, 50 cm	SC-2317-50	<input type="checkbox"/>	
	• Infinion CX Percutaneous Leads, 70 cm	SC-2317-70	<input type="checkbox"/>	
	• Artisan™ MRI Surgical Leads, 50 cm	SC-8416-50	<input type="checkbox"/>	
	• Artisan MRI Surgical Leads, 70 cm	SC-8416-70	<input type="checkbox"/>	
	• Artisan Surgical Leads, 50 cm	SC-8216-50 SC-8120-50 SC-8116-50	<input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>	
	• Artisan Surgical Leads, 70 cm	SC-8216-70 SC-8120-70 SC-8116-70	<input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>	
	• Other Lead(s): _____		<input type="checkbox"/>	
	• Adapters, Extensions, or Splitters:		<input type="checkbox"/>	
3.	Surgical Accessories (check all that apply)			
	• Clik™ X MRI Anchor	SC-4319	<input type="checkbox"/>	
	• Clik X Anchor	SC-4318	<input type="checkbox"/>	
	• Clik Anchor	SC-4316	<input type="checkbox"/>	
	• Med-A	SC-4320	<input type="checkbox"/>	
	• Silicone Suture Sleeves		<input type="checkbox"/>	
	• Other: _____			

Note: Leads should be connected directly into the IPG, Patient should not be implanted with lead extensions, splitters, or adapters.

B. PATIENT IMPLANT CONFIGURATION INFORMATION (ALL QUESTIONS MUST BE ANSWERED)		MRI FULL BODY ELIGIBLE	NOT MRI ELIGIBLE
1.	The lead implant location is epidural	Yes	No
2.	The IPG is implanted in the upper buttock or lower flank	Yes	No
3.	Patient does not have abandoned leads or IPGs (i.e. leads or IPGs that are not connected to the functioning Precision Montage MRI System)	Yes	No
4.	No evidence can be found of fractured leads or compromised IPG-lead system integrity	Yes	No

C. INSTRUCTIONS FOR THE PATIENT PRIOR TO THE MRI EXAM		MRI FULL BODY ELIGIBLE	NOT MRI ELIGIBLE
1.	Instruct the patient to fully charge their IPG (IPG charge shown as 3 bars on the Remote Control) and bring the Charger to the MRI Center (in case charging is necessary)	Yes	No
2.	Instruct the patient to bring their Remote Control to the MRI exam and turn stimulation off before the MRI Scan	Yes	No

Note: The Charger and Remote Control are MR Unsafe and must not be brought into the MRI Scanner Room/Scanner Room.

US Indications for Use Statement

Indications for Use. The Boston Scientific Neuromodulation Spinal Cord Stimulator (SCS) Systems are indicated as an aid in the management of chronic intractable pain of the trunk and/or limbs, including unilateral or bilateral pain associated with the following: failed back surgery syndrome, intractable low back pain, and leg pain.

Contraindications, warnings, precautions, side effects. The SCS Systems are contraindicated for patients who: are unable to operate the SCS System, have failed trial stimulation by failing to receive effective pain relief, are poor surgical risks, or are pregnant. Refer to the Instructions for Use provided with the SCS System or ControlYourPain.com for potential adverse effects, warnings, and precautions prior to using this product.

Caution: Federal (U.S.) law restricts this device to sale by or on the order of a physician.

The Precision Montage MRI SCS System with ImageReady MRI Technology is "MR-Conditional" only when exposed to the MRI environment under the specific conditions defined in the ImageReady™ MRI Full Body Guidelines for Precision Montage™ MRI Spinal Cord Stimulator System.

**Boston
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