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WAVEWRITER ALPHA™ IMAGEREADY MRI FULL BODY PATIENT ELIGIBILITY

This form provides information about the patient's implanted WaveWriter Alpha or WaveWriter Alpha Prime Spinal Cord Stimulator System and MRI scan eligibility. It may be provided to the radiologist to support the confirmation of the patient's scan.

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

Patient's Name: _____ Date: _____

Physician's Name: _____

Office Address: _____ Phone: _____

A. MR CONDITIONAL WAVEWRITER ALPHA MRI SYSTEM INFORMATION	MODEL #	MRI FULL BODY ELIGIBLE	NOT MRI ELIGIBLE
1. Implantable Pulse Generator (IPG)			
WaveWriter Alpha IPG, 32 contact IPG	SC-1232	☐	
WaveWriter Alpha 16 IPG, 16 contact IPG	SC-1216	☐	
WaveWriter Alpha Prime IPG, 32 contact IPG	SC-1432	☐	
WaveWriter Alpha Prime 16 IPG, 16 contact IPG	SC-1416	☐	

Note: if you have another model number IPG, please refer to the labeling specific to your IPG model number.

A. MR CONDITIONAL WAVEWRITER ALPHA MRI SYSTEM INFORMATION	MODEL #	MRI FULL BODY ELIGIBLE	NOT MRI ELIGIBLE
2. Percutaneous and/or surgical paddle leads (check all that apply)			
Avista™ MRI Percutaneous Lead, 8-contact lead, 56 cm	SC-2408-56	☐	
Avista MRI Percutanious Lead, 8-contact lead, 74 cm	SC-2408-74	☐	
Linear™ Percutaneous Leads, 50 cm	SC-2158-50	☐	
Linear Percutaneous Leads, 70 cm	SC-2158-70	☐	
Linear ST Percutaneous Leads, 50 cm	SC-2218-50	☐	
Linear ST Percutaneous Leads, 70 cm	SC-2218-70	☐	
Linear 3-4 Percutaneous Leads, 50 cm	SC-2352-50	☐	
Linear 3-4 Percutaneous Leads, 70 cm	SC-2352-70	☐	
Linear 3-6 Percutaneous Leads, 50 cm	SC-2366-50	☐	
Linear 3-6 Percutaneous Leads, 70 cm	SC-2366-70	☐	
Infinion™ CX Percutaneous Leads, 50 cm	SC-2317-50	☐	
Infinion CX Percutaneous Leads, 70 cm	SC-2317-70	☐	
Artisan™ MRI Surgical Leads, 50 cm	SC-8416-50	☐	
Artisan MRI Surgical Leads, 70 cm	SC-8416-70	☐	
Artisan Surgical Leads, 50 cm	SC-8216-50	☐	
Artisan Surgical Leads, 70 cm	SC-8216-70	☐	

A. MR CONDITIONAL WAVEWRITER ALPHA MRI SYSTEM INFORMATION	MODEL #	MRI FULL BODY ELIGIBLE	NOT MRI ELIGIBLE
2. Percutaneous and/or surgical paddle leads continued (check all that apply)			
CoverEdge™ 32 Surgical Leads, 50 cm	SC-8336-50	☐	
CoverEdge™ 32 Surgical Leads, 70 cm	SC-8336-70	☐	
CoverEdge X 32 Surgical Lead, 50 cm	SC-8352-50	☐	
CoverEdge X 32 Surgical Lead, 70 cm	SC-8352-70	☐	
CoverEdge 32 MRI Surgical Lead, 50 cm	SC-8436-50	☐	
CoverEdge 32 MRI Surgical Lead, 70 cm	SC-8436-70	☐	
CoverEdge X 32 MRI Surgical Lead, 50 cm	SC-8452-50	☐	
CoverEdge X 32 MRI Surgical Lead, 70 cm	SC-8452-70	☐	
Other Lead(s):			☐
Adapters, Extensions, or Splitters:			☐
3. Surgical Accessories (check all that apply)			
Clik X MRI Anchor	SC-4319	☐	
Clik X Anchor	SC-4318	☐	
Clik Anchor	SC-4316	☐	
Med-A	SC-4320	☐	
Silicone Suture Sleeves		☐	
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 has no abandoned leads or IPGs (i.e. leads or IPGS that are not connected to the functioning WaveWriter Alpha or WaveWriter Alpha Prime 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
1. For patients with a rechargeable IPG (SC-1232 or SC-1216), instruct the patient to fully charge their IPG and bring the Charger to the MRI Center (in case charging is necessary)
2. Instruct the patient to bring their Remote Control to the MRI exam and to enable MRI mode before the MRI Scan

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

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Indications for Use: Boston Scientific Neuromodulation Spinal Cord Stimulator Systems are indicated as an aid in the management of chronic intractable pain. **CAUTION:** The law restricts these devices to sale by or on the order of a physician. Indications, contraindications, warnings and instructions for use can be found in the product labelling supplied with each device. Informational use only in countries with applicable health authority registrations. Material not intended for use in France. WaveWriter Alpha and WaveWriter Alpha Prime 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 WaveWriter Alpha and WaveWriter Alpha Prime Spinal Cord Stimulator Systems.

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