bioventus®
Innovations For Active Healing

StimRouter® Neuromodulation System and Supplies, Physician Statement of Medical Necessity/Prescription

This section may be completed by Bioventus

Patient Name: First	MI.	Last	
Street Address:	City:	State:	Zip:
Patient DOB: MM/DD/YYYY	Phone:		

This section MUST be completed by practitioner/staff

Primary Diagnosis (mark all that apply):

Pain _____ ICD-10 code Pain, not classified _____ ICD-10 code Other _____ ICD-10 code

Location: ☐ Shoulder ☐ Arm ☐ Elbow ☐ Wrist ☐ Hand/Fingers ☐ Unspecified
☐ Hip ☐ Knee ☐ Ankle/Foot ☐ Leg/Thigh

Functional Limitations:

Patient's Need (mark all that apply):

☐ Pain Reduction, localized ☐ Improve Functional Capabilities
☐ Reduce Dependency on Oral Medications ☐ Other (explain)

Length of Need:	Prognosis:

Physician's Signature:	Date:

State law requires renewal on said item every 12 months. Length of need is dictated based on state standard of 1 year unless indicated above. I certify that the above-prescribed equipment is medically indicated and in my opinion is reasonable and necessary for this patient's treatment.

This section may be completed by Bioventus

PHYSICIAN INFORMATION		
Physician:	License #:	NPI #:
Address:	Phone:	Fax:
City, State, Zip:	Office Contact:	

Upon completion, fax this form to Bioventus Customer Service Department
Fax: 661.310.9623 | Email: PNSorders@Bioventus.com | Phone: 888.453.2136

The StimRouter Neuromodulation System is indicated for pain management in adults who have severe intractable chronic pain of peripheral nerve origin, as an adjunct to other modes of therapy (e.g., medications). The StimRouter is not intended to treat pain in the craniofacial region.

Do not use the StimRouter Neuromodulation System in users who have an implanted demand cardiac pacemaker, implanted cardioverter defibrillator (ICD), or other implanted active device, or who have bleeding disorders that cannot be stopped in advance of the StimRouter implantation procedure. Do not use the system where a metallic implant or a cancerous lesion is present in the immediate implant area. Effects of stimulation during pregnancy are not known. StimRouter is capable of producing skin irritation and muscle ache in the area of stimulation.