



Authorization Request Form - **Part 2**

**Spinal Cord Stimulators**

Complete and fax the clinical worksheet immediately following the Part 1 authorization request fax form, including any substantiating clinical documentation. Your responses enable faster processing of authorization requests and reduces the likelihood we may require you to submit additional clinical documentation to complete our review.

Please fill in each question option completely    ☐ → ☒

Question 1	<p>Which type of stimulator device is being considered? (Required, fill in one option)</p> <div><input type="radio"/> Spinal Cord Stimulator</div> <div><input type="radio"/> Dorsal Root Ganglion stimulator (DRG)</div> <div><input type="radio"/> Bladder Stimulator</div> <div><input type="radio"/> Gastric Stimulator</div> <div><input type="radio"/> Optic Nerve Stimulator</div> <div><input type="radio"/> Phrenic Nerve Stimulator</div> <div><input type="radio"/> Peripheral Nerve Stimulator</div> <div><input type="radio"/> Other Neurostimulator</div>
Question 2	<p>Which of the following is being requested? (Required, fill in one option)</p> <div><input type="radio"/> Trial placement of stimulator</div> <div><input type="radio"/> Permanent implantation of stimulator</div>
Question 3	<p>Does the patient have any of the following? (Required, fill in all that apply)</p> <div><input type="radio"/> Complex regional pain syndrome (CRPS)/reflex sympathetic dystrophy</div> <div><input type="radio"/> Failed back surgery syndrome (FBSS) with primarily radicular pain</div> <div><input type="radio"/> Inoperable chronic limb ischemia</div> <div><input type="radio"/> Chronic pain syndrome</div> <div><input type="radio"/> Chronic radicular pain</div> <div><input type="radio"/> Cervical level spinal cord stimulation</div> <div><input type="radio"/> Dorsal root ganglion stimulation</div> <div><input type="radio"/> Simultaneous use of SCS and intrathecal opioid therapy/implantable infusion pump</div> <div><input type="radio"/> Individuals with cardiac pacemakers and/or defibrillators</div> <div><input type="radio"/> None of the above</div>
Question 4	<p>In the past 12 months, how long has the patient been receiving conservative care? (Required, fill in one option)</p> <div><input type="radio"/> No conservative care occurred</div> <div><input type="radio"/> Greater than 6 weeks</div> <div><input type="radio"/> Greater than 12 weeks</div> <div><input type="radio"/> Greater than 6 months</div> <div><input type="radio"/> None of the above options apply</div>
Question 5	<p>What type of other treatment modalities have been attempted? (Required, fill in all that apply)</p> <div><input type="radio"/> Pharmacological</div> <div><input type="radio"/> Failed Surgery or not a candidate for surgical intervention</div> <div><input type="radio"/> Physical Therapy</div> <div><input type="radio"/> Psychological (cognitive behavioral therapy)</div> <div><input type="radio"/> No modalities used since no conservative care attempted</div>
Question 6	<p>Has pain persisted despite other treatment modalities (pharmacological, surgical, physical and psychological therapies) that have been tried for six consecutive months? (Required, fill in one option)</p> <div><input type="radio"/> Yes, pain has persisted despite conservative care</div> <div><input type="radio"/> No, pain has persisted BUT they have not attempted conservative care</div> <div><input type="radio"/> No, pain has not persisted</div>
Question 7	<p>Has a psychological evaluation been obtained and determined that the individual is an acceptable candidate? (Required, fill in one option)</p> <div><input type="radio"/> Yes</div> <div><input type="radio"/> No</div>

Questions continued on following page

 Please fill in each question option completely    ☐ → ☒

Question 8	Has the individual undergone careful screening, physical evaluation and diagnosis by a multidisciplinary team prior to implantation? (Required, fill in one option)  <input type="radio"/> Yes <input type="radio"/> No
Question 9	If this is a request for a permanent electrode placement, has a temporary trial of at least two days duration been completed? (Required, fill in one option)  <input type="radio"/> Yes, a temporary trial of two days has been completed <input type="radio"/> No, a temporary trial of two days has not been completed <input type="radio"/> No, this is a request for a Temporary Percutaneous Electrode Placement
Question 10	Demonstration of at least 50% reduction in symptoms and improved function with the temporary implanted electrode prior to the permanent implantation? (Required, fill in one option)  <input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Not applicable, this is the first request
Question 11	Which manufacturer’s device is being used? (Required, fill in one option)  <input type="radio"/> Boston Scientific <input type="radio"/> Medtronic <input type="radio"/> Nevro <input type="radio"/> Abbot <input type="radio"/> Other
Question 12	Does the surgeon have a preference for where the patient is discharged for post-acute care (if still appropriate at the time of discharge)? (Required, fill in one option)  <input type="radio"/> Discharge home, no post-acute services required <input type="radio"/> Discharge home, outpatient Physical Therapy services required <input type="radio"/> Discharge home, Home Health Agency (HHA) services required <input type="radio"/> Discharge to Skilled Nursing Facility <input type="radio"/> No discharge preference indicated
Question 13	Did the patient’s primary care physician provide preoperative medical clearance for this patient? (Optional, fill in one option)  <input type="radio"/> Yes <input type="radio"/> No