



Provider Tip Sheet: Overcoming Patient Objections to Spinal Cord Stimulation for Painful Peripheral Neuropathy

Peripheral neuropathy can cause debilitating pain in the hands and feet, significantly impacting a patient's quality of life. Spinal cord stimulation (SCS), a minimally invasive treatment, offers relief by interrupting pain signals; however, some patients have misperceptions or concerns. This sheet can help you address some common concerns.

Key Talking Points

- **Effective Alternative When Medications Fail:** SCS offers an alternative when conservative treatments fail, providing hope for patients struggling with chronic pain. SCS has been FDA-approved since 1989 to treat chronic pain and has evolved to become smaller, safer, and more convenient. The FDA expanded approval in 2023 to treat **painful diabetic neuropathy**. SCS was found to have positive long-term benefits for many patients with painful diabetic neuropathy, reducing pain by up to 75% as well as decreasing hospitalizations.
- **Minimally Invasive Procedure:** SCS is a minimally invasive procedure. SCS consists of a small paddle or leads that are placed on the spinal cord and controlled by a pacemaker-like generator implanted under the skin. The device sends tiny electrical impulses, which reduce or block pain signals from reaching the brain. Patients can adjust stimulation levels, using a wireless remote, even turning it off when not needed.
- **Trial Period Assurance:** All qualified patients receive a temporary external trial, which ensures they are happy with the results before implantation.
- **Patient Support:** Patients receive support from a team, including the expertise of interventional pain management physicians and neurosurgeons, and assistance from device representatives for programming and adapting to the stimulation.
- **Pain Relief Expectations:** Manage patient expectations by explaining that the **goal is to reduce pain by 50-75%**, recognizing that each individual's response may vary.

Addressing Common Concerns

Concern: Fear of constant electrical stimulation.

- **Response:** Patients can control when and how much stimulation they receive, making it adaptable to their pain levels and activities.

Concern: Uncertainty about the effectiveness of spinal cord stimulation.

- **Response:** Many patients experience a significant reduction in pain, with up to 75% relief. A trial period is conducted before permanent implantation to ensure efficacy.
 - A 2021 randomized controlled trial for painful diabetic neuropathy reported patient's mean pain score went from 7.6 at baseline to 1.7 at 6 months with high-frequency spinal cord stimulation.

Concern: Apprehension about surgery and recovery.

- **Response:** The surgical procedure is outpatient, taking 1-2 hours, and patients often return home the same day. Recovery is relatively quick, with most patients resuming light activities within two weeks.