

Case Report

Neuromodulation of the Cervical Spinal Cord in the Treatment of Chronic Intractable Neck and Upper Extremity Pain: A Case Series and Review of the Literature

Ricardo Vallejo^{1,2}, MD, PhD, Jeffery Kramer^{1,2,3}, PhD, and Ramsin Benyamin^{1,2,3}, MD

From: ¹ Millennium Pain Center, ² Illinois State University, Bloomington, IL, ³ University of Illinois, Urbana-Champaign, IL. Dr. Vallejo¹ is Director of Research, Staff Pain Medicine, Millennium Pain Center, and Adjunct Professor of Biology, Illinois State University, Bloomington, IL. Dr. Kramer² is with the Department of Biology, Illinois State University and the College of Medicine at Peoria, Department of Cancer Biology and Pharmacology, Peoria, IL. Dr. Benyamin³ is President, Millennium Pain Center, Bloomington, IL, and Clinical Instructor, Department of Surgery, College of Medicine, University of Illinois, Urbana-Champaign, IL.
Address Correspondence:
Ricardo Vallejo, MD, PhD
Millennium Pain Center
1015 S. Mercer Ave.
Bloomington, IL 61701
E-mail: vallejo@millenniumpaincenter.com

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Electrical spinal neuromodulation in the form of spinal cord stimulation is currently used for treating chronic painful conditions such as complex regional pain syndrome, diabetic neuropathy, postherpetic neuralgia, peripheral ischemia, low back pain, and other conditions refractory to more conservative treatments. To date, there are very few published reports documenting the use of spinal cord stimulation in the treatment of head/neck and upper limb pain. This paper reports a case series of 5 consecutive patients outlining the use of spinal cord stimulation to treat upper extremity pain. All subjects had previously undergone cervical fusion surgery to treat chronic neck and upper limb pain. Patients were referred following failure of the surgery to manage their painful conditions. Spinal cord stimulators were placed in the cervical epidural space through a thoracic needle placement. Stimulation parameters were adjusted to capture as much of the painful area(s) as possible. In total, 4 out of 5 patients moved to implantation. In all cases, patients reported significant (70–90%) reductions in pain, including axial neck pain and upper extremity pain. Interestingly, 2 patients with associated headache and lower extremity pain obtained relief after paresthesia-steering reportedly covered those areas. Moreover, 2 patients reported that cervical spinal cord stimulation significantly improved axial low back pain. Patients continue to report excellent pain relief up to 9 months following implantation. This case series documents the successful treatment of neck and upper extremity pain following unsuccessful cervical spine fusion surgery. Given this initial success, prospective, controlled studies are warranted to more adequately assess the long term utility and cost effectiveness of electrical neuromodulation treatment of chronic neck and upper extremity pain.

Key words: spinal cord stimulator, cervical, neck pain, radicular pain, axial pain, headache, leg pain

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Spinal neuromodulation for the management of chronic pain conditions exists in 2 basic forms: chemical neuromodulation or electrical neuromodulation. Spinal chemical neuromodulation is exemplified by the use of epidural injections or the intrathecal delivery of drugs (1-3), whereas electrical spinal neuromodulation is produced by the use of spinal cord stimulation (SCS) devices (4,5). An increasing number of reviews on the use of this technology in the treatment of a variety of pain conditions have been written summarizing the short- and long-term effectiveness, purported mechanism(s) of action, and cost benefits (6-11). Despite the growing literature base documenting the use of SCS in the treatment of chronic pain conditions, to our knowledge, there have been very few reports detailing the use of spinal cord stimulation therapy to treat chronic pain following failed cervical spinal surgery (12).

One predominant indication for SCS is the relatively nebulous diagnosis of failed back surgery syndrome (FBSS) (8,13-17). In general, back surgery that does not adequately relieve pain in an individual is labeled as "failed." Reasons for the lack of success can be multifold including incorrect patient selection, poor technique, recurrent pathology, and a variety of other causes (18). Relatively recent randomized trials suggest that spinal cord stimulators provide an effective therapeutic option in the treatment of FBSS (14,15,19). In a prospective, controlled study by North et al (19), patients were randomized to either a spinal cord stimulation group or a re-operation group. Either group could cross over to the other if the initial randomized treatment failed to adequately control pain. A greater percentage of the re-operation patients crossed over into the spinal cord stimulation group as opposed to SCS subjects crossing over into the re-operation group. Moreover, spinal cord stimulation offered a greater chance of successfully controlling pain as opposed to re-operation. This study suggests that spinal cord stimulation may more effectively control the pain of FBSS.

In contrast to the amount of study devoted to the efficacy and cost-effectiveness of SCSs in the treatment of low back pain, relatively little is published about the use of spinal cord stimulators in the treatment of neck and upper limb pain. In upper extremity pain most studies have focused on a variety of chronic pain conditions such as diabetic neuropathy (20-22), complex regional pain syndrome (8,23,24), Raynaud's phenomenon (25-27), painful peripheral vascular disorders (28-31), and pain resulting from brachial plexus avulsion (32), just

to name a few. In particular, there is little published about the effectiveness of spinal cord stimulators in the treatment of neck (axial) and upper limb pain after neck surgery. In this paper we present a series of 5 consecutive cases where neck and/or upper limb pain were treated with cervical spinal electrical neuromodulation. In particular, we focused on patients that had a prior neck surgery without sufficient pain relief. Outcomes, complications, and recommendations are discussed.

CASE SERIES

Five consecutive patients treated with spinal cord stimulators epidurally placed in the cervical spine region for the treatment of neck and/or upper limb pain were identified through chart review and included in the study. All patients had uncontrolled neck and/or upper extremity pain despite aggressive interventional procedures prior to being considered candidates for cervical spinal electrical neuromodulation. In all cases, radiological reports (including x-rays, MRIs, and CTs) were mostly unremarkable. No significant structural instability, stenosis, or cord impingement was noted. In most cases, slight cervical disc degeneration was noted at 1 or more levels with or without a slight disc bulge or very slight disc herniation. In no cases were pseudoarthrosis or myeloradiculopathy noted. Physical exams including a neuromuscular exam were normal with normal deep tendon reflexes, muscle strength, and gait stability.

Three out of 5 patients underwent diagnostic cervical facet joint injections prior to spinal cord stimulator trials in order to rule out sources of pain amenable to treatment with radiofrequency lesioning. Although 3 of the patients reported significant relief following diagnostic cervical facet medial branch blocks, subsequent radiofrequency denervation failed to achieve sustained reductions in pain. In all cases, patients had previously undergone anterior cervical fusion surgeries without successful reductions or adequate control of neck and/or upper limb pain. Based on history, physical examination as well as imaging, patients were not considered surgical candidates. Because we were unable to further rule out other sources of pain and, similar to FBSS, accounted the pain to an idiopathic, chronically intractable neck pain syndrome in which the pain following the so-called failed neck surgeries persisted for at least 3 months.

In both the trial phase and the implant phase (if warranted) patients were brought to the fluoroscopy/surgery suite and placed in the prone position. One

gram of intravenous cefazolin was given prophylactically and 1% lidocaine was administered for skin and subcutaneous anesthesia. Using a sterile technique and under direct fluoroscopic visualization, a #14 epidural needle via a paramedian approach was used to identify the T3-4 epidural space by using the loss of resistance technique. Leads were typically advanced in the dorsal epidural space to the C2-4 level. Sensory stimulation and programming was completed to capture as much of the painful regions as possible. Trials lasted for between 5 and 7 days at which point patients rated pain relief on a 100 point numerical rating scale (percentage). If trials produced at least 50% pain relief then the option to move forward with the implantation phase was discussed.

RESULTS

A total of 5 patients were trialed with circumferential style leads manufactured by Advanced Bionics (2 patients), Advanced Neuromodulation Systems (2 patients), or Medtronic (1 patient). Patient details are outlined in Table 1. In most cases (4/5; 80%) patients indicated significant pain relief (>50%; mean 82.5% ± 6%, range 50%–90%). In 1 case the patient indicated that he did not obtain at least 50% pain relief, predominantly because of persistent complaints of pain over the spinous process at C7. This pain was not alleviated with local infiltration of local anesthetic and methylprednisolone (to rule out neuroma). Interestingly, this patient had a diagnosed C7 radiculopathy with a potential deafferentation syndrome. As a result

Table 1. Summary of patient information including pain locations, prior surgeries, pharmacotherapy, interventional procedures attempted, lead positions and outcomes. TENS = transcutaneous electrical nerve stimulation, CESI = cervical epidural steroid injection, CFDB = cervical facet diagnostic blocks, CFRFL = cervical facet radiofrequency lesion.

Patient	Age	Gender	Prior Surgeries	Pain Location(s)	Medications	Attempted Treatments	Trial Relief	Trial / Implant Lead Position	Implant Relief
A	39	M	C7-T1 Foramenotomy C7-T1 Fusion	Neck, Left Arm (C3-C7 Distribution)	Opiates NSAIDs	Physical Therapy TENS CESI CFDB CFRFL	< 50% Neck & Arm	C3 Dual 8-contact Trial Leads	N/A
B	56	M	C6-7 Discectomy C6-7 Fusion	Neck, Back, Feet Headache (C2-7 & L4-S1 Distribution)	Topromide	Physical Therapy TENS CFDB CESI CFRFL	70% Total Body	C2-3 / C3 Single 8-Contact Lead	>90% in neck, back, feet (9 months); some headache relief
C	57	M	C5-7 Fusion	Neck, Bilateral Shoulder, Feet (C3-7 & L2-S1 Distribution)	Opiates	Physical Therapy TENS Acupuncture CESI	80% Neck & Shoulder	C3 / C3 Single 8-Contact Lead	70% (1 month)
D	72	M	C5-7 Fusion	Neck and Bilateral Shoulder and Arm (C4-C8 & L2-L5 Distribution)	Opiates NSAIDs	Physical Therapy CESI Trigger Point BoTox Suprascapular Block CFDB CFRFL	80% > Shoulder & Arm < Neck	C3 / C4 Single 8-Contact Lead	>50% (8 months)
E	44	F	C5-6 Fusion	Neck, Shoulder, Right Arm, Low Back (C2-7 Distribution)	Opiates Pregabalin Muscle Relaxants	TENS Physical Therapy	100% Total Body	C2/C3 Dual 8-Contact Leads	>90% (2 months)

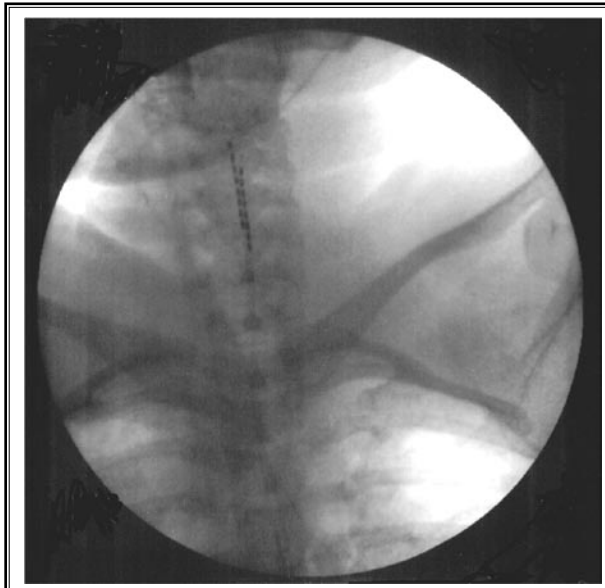


Fig. 1. Cervical placement of dual spinal cord stimulator leads. This patient obtained very good relief of elbow and hand pains. Interestingly, this patient could also feel paresthesia in the mandible.

of the poor relief obtained during the trial, the patient did not want to move on to the implant phase. Including the patient that had an unsuccessful trial, the mean trial pain relief was $75\% \pm 8.9\%$. In another case, the device failed following implantation and a revision had to be completed with a different lead type. The longest follow-up time recorded is 8 months with the patient still reporting $>50\%$ relief. The range of follow-up is 1 month to 9 months with 4/4 patients reporting an average pain relief at approximately 70%. It is unclear if there is any "creep" in pain levels, also termed "tolerance," over time as reported in other papers (33). However, 2 patients have reported high ($>90\%$) and moderately high ($>70\%$) levels of pain control at 6 months and longer. It is important to recognize that in 4 out of 5 patients, spinal cord stimulation was able to significantly alleviate not only upper extremity pain in the shoulders and arms but also in an axial distribution. Primarily this axial pain reduction was in the neck, however 2 patients reported axial relief (both $>50\%$) in the low back with a spinal cord stimulator placed in the cervical region. Pain relief was, for the most part, similar between the axial and distal regions with the exception of 1 patient that found somewhat better relief in the upper extremities compared to the neck.

Most patients report paresthesia mapping along a C2-C3 dermatome pattern. In all cases, the neck and upper limbs (from the shoulder to the fingers) could be captured. In some cases, paresthesia was reported in the occipital region of the head and in another case in the mandible region, which is again consistent with a C2 dermatome sensory innervation (Fig.1). In 3 cases patients reported head to toe paresthesias which were able to cover other more distal pain regions including low back, legs, and feet. Two patients reported the ability to drive the paresthesia supraoccipitally to the parietal region of the head. This coverage was able to help alleviate cervicogenic headache pain in tandem with upper and lower extremity pain. In no cases did we observe lead migration. In cases where optimal paresthesia coverage was lost, we could always recapture the painful regions by reprogramming.

DISCUSSION

The use of spinal electrical neuromodulation as an advanced treatment to alleviate chronically intractable painful conditions is steadily increasing in both the number of patients being treated as well as an ever-expanding list of indications for which spinal cord stimulation is effective. One predominant use of spinal cord stimulators is in the treatment of FBSS (8,16,19,34). In the current case series, we present 5 patients that underwent spinal cord stimulation following a so-called failed neck surgery. In all cases, anterior cervical spinal fusions failed to adequately control or alleviate pain in the neck and/or upper limbs. This condition is analogous to FBSS in the fact that a surgical procedure failed to control pain and there exists, basically, an idiopathic painful condition. Moreover, despite that 60% (3 out of 5) of the cases obtained relief after dual facet medial branch blocks, none of them responded to radiofrequency ablation. The reason for the lack of response to radiofrequency ablation following dual medial branch blocks is unclear. Previous papers have documented approximately 75–85% dual block specificity (35, 36). The discrepancy could be partly explained by multiple false positive results. Recently, a preliminary report of a randomized, controlled trial by Manchikanti et al (37) demonstrated significant long-term pain relief following medial branch blocks without steroids (13.4 ± 3.5 weeks). This duration of pain relief is significantly longer than the local anesthetic duration of action and suggests that facet joint pain may have a neuropathic component (38). This latter point would help explain the lack of

efficacy following radiofrequency ablation and yet the success of spinal cord stimulation therapy.

Since the epidural space in the cervical region is relatively narrow compared to the thoracic or lumbar regions, lead placement can be met with some resistance, especially when attempting to place 2 leads. Also, the second cervical vertebral level is currently about the highest the leads can be placed. Some benefit might be gained by capturing higher dermatomal levels although the placement of leads in this upper cervical region would be technically difficult given the anatomy of the epidural space and connectivity of the dura in the C2 spinal region. In all cases, generators were implanted in the hip/buttock region. Despite this relatively long distance from the lead tips to the implanted pulse generator, we observed no lead migration. It should be noted that all of the patients represented in this case series had undergone anterior spinal fusion(s). It is possible that posterior fusions might lead to difficulties in lead placements; however, our data cannot directly address this potential limitation.

Our observation of the ability to capture the occipital regions of the head is not surprising. The greater and lesser occipital nerves emanate from the C2 root, and to a lesser extent the C3 root, and connect in the spinal cord within the spinal nucleus of the trigeminal nerve (which becomes the nucleus caudalis in the cervical region) (39, 40). Both human and animal studies suggest that this anatomical substrate may be responsible for the sensory convergence observed (41-44). This anatomical and functional overlap may also explain why 1 patient reported paresthesia in the mandible, a sensory region subserved by the trigeminal nerve. Deeper stimulation with increasing pulse widths may eventually stimulate neural structures associated with projections from the greater occipital nerve, thereby producing sensations seeming to be associated with parietal regions. Two patients in this case series noted significant improvement in headaches. This result is similar to those reported by Dario et al who reported treatment of cervicogenic headache via spinal cord stimulation with lead tips placed at the C3 level (12). Also, these results are similar to those observed when utilizing occipital nerve stimulation (45).

Stimulation of more distal structures such as the lower back, legs, and feet can be explained by the anatomical arrangement of sensory fibers within the dorsal columns. As sensory fibers project cephalad from more caudal regions of the body, they assume a

more medial position while other sensory fibers from upper extremities course more laterally. Feirabend et al published an elegant anatomical study on the morphometry of the spinal dorsal columns and concluded that lateral fibers demonstrate both an increasing fiber density as well as fiber frequency with a subsequent increase in collateralization (46). This anatomic arrangement would provide a layer of stimulation where weaker electrical fields would stimulate more lateral nerve fibers and other neural structures while stronger electrical fields would be able to recruit more medial fibers and, thus, more caudally located dermatomes. These findings also help explain our observations of a cephalad to caudal paresthesia with increasing current amplitudes or pulse widths. Moreover, stimulation of the cervical spinal cord could effectively manage pain in the lower extremities in conjunction with pain located in the neck and upper extremities.

These observations are in direct contrast to those of Eisenberg and Brecker who reported effective pain management of lower extremity pain (leg and foot pain of cervical origin) with a low thoracic placed spinal cord stimulator (47). These authors reported that, despite the clear cervical origin of the pain, placement of the lead to produce paresthesia in the affected limb could still relieve pain. In contrast, our observations indicate that pain presumably of thoracic or lumbar origin can be effectively treated with cervical lead placement. Thus, it seems that lead placement is less important than the location of the paresthesias that are generated.

In this case series we did not observe any adverse events such as infection or lead migration. Although we did not collect radiographic evidence of anatomical migration, upon reprogramming we could always functionally capture painful regions. Thus, our observations of a lack of lead migration are based upon dermatomal paresthesia mapping and not based upon radiographic imaging. One might anticipate an increased incidence of lead migration in the cervical versus thoracolumbar region due to the relatively greater flexibility and movement in the neck; however, it is not currently known if this is the case. Despite our and others observations of cervical spinal cord stimulation induced headache relief, Ward and Levin reported a case of lead migration causing cervicogenic headache (48). It is hypothesized that stimulation of the trigemino-vascular system following migration caused the headaches since repositioning the leads alleviated the headaches. Epidural cervical spinal stimulation also has

other measurable effects besides analgesia including increasing cerebral blood flow, as well as tumor oxygenation and metabolism (49-53). While interesting, it is unclear what clinical significance this may play.

Despite the often referenced classic publication by Wall and Melzack in 1965 (54) there still exists a very limited amount of basic research data on spinal neuromodulation in the treatment of chronically painful conditions (11, 55). Significantly more research needs to be conducted in order to determine the precise mechanisms of action underlying the analgesic effects of spinal cord stimulation. In particular, it would be helpful to know which spinal and super-spinal regions

are activated when leads are placed in the neck region versus thoracic regions. Moreover, it will be interesting to see if mechanisms of pain reduction differ between cervical and thoracic lead placement.

In conclusion, our case series suggests that spinal cord stimulation can be an effective treatment for patients with persistent axial neck pain with or without upper extremity pain following failed cervical fusion surgery. A more rigorous prospective study is warranted to directly determine the long-term efficacy as well as document the rate of complications associated with cervical spinal cord stimulation.

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