Although not commonly used, tractotomy-nucleotomy is a viable treatment option for patients with malignancy-related facial pain, neuropathic facial pain, postherpetic neuralgia, glossopharyngeal neuralgia, geniculate neuralgia, and refractory trigeminal neuralgia. Postherpetic neuralgia is an especially severe and difficult pain syndrome to treat and is occasionally associated with facial allodynia and hyperalgesia. Tractotomy has a durable success rate of greater than 80%. The procedure is typically performed with CT guidance in an awake and prone patient. Primary complications include temporary ataxia and motor dysfunction.

Some patients, however, have facial pain so profound that they are unable to tolerate awake prone positioning; for these patients, CT-guided tractotomy-nucleotomy might not be feasible. The authors describe 2 such patients, for whom percutaneous intraoperative CT-guided tractotomy-nucleotomy under general anesthesia was successful. One patient was a 79-year-old man with profound left facial postherpetic neuralgia, who was unable to tolerate awake CT-guided tractotomy-nucleotomy, and the other was a 45-year-old woman with intractable hemicranial pain that developed after a right frontal lesionectomy for epilepsy. Each patient underwent a percutaneous intraoperative CT-guided tractotomy-nucleotomy under general anesthesia. No complications occurred, and each patient reported excellent pain relief for up to 6 and 3 months after surgery, respectively. Percutaneous intraoperative CT-guided tractotomy-nucleotomy performed on anesthetized patients is effective for facial postherpetic neuralgia and postoperative hemicranial neuralgia.

For confirming the correct location of the radiofrequency electrode before creation of a lesion, percutaneous CT-guided trigeminal tractotomy–nucleotomy is most commonly performed with the patient prone and awake. However, for patients whose facial pain and hypersensitivity are so severe that the patients are unable to rest their face on a support (as required with prone positioning), awake CT-guided tractotomy-nucleotomy might not be feasible. The authors describe 2 such patients, for whom percutaneous intraoperative CT-guided tractotomy-nucleotomy under general anesthesia was successful. One patient was a 79-year-old man with profound left facial postherpetic neuralgia, who was unable to tolerate awake CT-guided tractotomy-nucleotomy, and the other was a 45-year-old woman with intractable hemicranial pain that developed after a right frontal lesionectomy for epilepsy. Each patient underwent a percutaneous intraoperative CT-guided tractotomy-nucleotomy under general anesthesia. No complications occurred, and each patient reported excellent pain relief for up to 6 and 3 months after surgery, respectively. Percutaneous intraoperative CT-guided tractotomy-nucleotomy performed on anesthetized patients is effective for facial postherpetic neuralgia and postoperative hemicranial neuralgia.

**Case 1**

A 79-year-old man sought care after a 7-year history of profound left-sided facial pain, particularly at the ophthalmic division of the trigeminal nerve, resulting from facial herpes zoster (shingles). He reported dull, aching background pain and episodic sharp, burning pain that was exacerbated by light touch. Previous treatments consisted of oral pregabalin and left supraorbital nerve stimulation, both of which minimally relieved the pain. Neurological examination results were nonfocal.

For this patient with a diagnosis of medically refractory postherpetic neuralgia, a decision was made to perform a left-sided CT-guided tractotomy-nucleotomy with the patient awake. However, during the procedure, the patient was unable to tolerate the facial pain caused by the CT scanner headrest; despite aggressive premedication and numerous attempts to reposition and pad the patient’s face, the procedure was aborted. A decision was then made to proceed with an intraoperative CT-guided tractotomy-nucleotomy with the patient under general anesthesia and use of somatosensory and motor evoked potentials of the extremities and of the lower cranial nerves.

**Key Words**

- trigeminal
- tractotomy-nucleotomy
- CT guidance
- general anesthesia
- postherpetic neuralgia
- hemicranial neuralgia
Impedance was measured by using a Radionics RFG-3C generator (formerly Radionics Inc.) (Fig. 1D) and was 300–400 Ω, which resulted in increased heart rate, suggestive of significant pain. Neuromonitoring was used during the procedure; however, the CT scanner and the Radionics generator produced enough electrical artifact to render neuromonitoring results unusable. Two lesions were then created, one at 90°C for 90 seconds and one at 95°C for 90 seconds. The patient was awakened from general anesthesia and extubated without problem. The patient immediately noted a slight decrease in touch-induced pain on the left side of his face, and the next day, he reported complete relief of his facial pain. Notably, he did experience mild ataxia and numbness of the left arm, which significantly improved 2 days after discharge. Six months after the procedure, the patient rated his facial pain as 2/10; before the procedure, he had rated it at 9/10.

Case 2

A 45-year-old woman sought care for right hemicranial pain in both trigeminal and occipital nerve distributions, which had developed after surgical resection of a right frontal seizure focus 13 years earlier. The patient had previously undergone occipital nerve stimulation and right C-2 gangliectomy 9 and 7 years earlier, respectively. The patient reported considerable pain that was resistant to a drug regimen consisting of carbamazepine, gabapentin, escitalopram oxalate, acetaminophen/hydrocodone, and diclofenac. Neurological examination findings were nonfocal. Neuromonitoring was not used for the significant electrical artifact problems encountered in Case 1. Initially, the patient was placed prone on the DORO radiolucent skull clamp; however, because of her physique, we were unable to visualize to C-1. Therefore, the patient was placed prone on a NeuroLogica Scan Board (NeuroLogica Corp.) (Fig. 2) with her shoulders gently retracted inferiorly with tape, which enabled visualization to C-2. As was done in Case 1, a CereTom CT scanner was used to plan the skin insertion site of the percutaneous needle approximately 2 cm left of the midline between the occiput and C-1. A thin, 1.25-mm-wide FOV scan through the foramen magnum to the bottom of C-1 was acquired (Fig. 1B). The skin-to-dura distance and the diameters of the upper cervical spinal cord were determined. The skin-to-dura distance was marked on the introducer cannula for the Levin-Cosman cordotomy electrode system (Cosman Medical). The electrode depth was adjusted to project 3 mm from the introducer metal cannula. The laser on the CT scanner was used to confirm correct electrode positioning. The tip of the Levin-Cosman cordotomy electrode was stimulated at approximately 0.1 V, which resulted in increased heart rate, suggestive of significant pain. Neuroradiological examination was performed before and during the procedure to ensure proper positioning of the electrode within the spinal cord, and 2 lesions were made.
The patient is placed on chest rolls to enable head flexion on the NeurolLogica Scan Board. The head is secured with tape, and the shoulders are gently held inferiorly to facilitate entry into the CereTom CT scanner and visualization down to C-2.

Immediately after the procedure, the patient was awakened from general anesthesia and extubated without problem. Immediately after the procedure, the patient reported significant pain relief. The patient in Case 1 (a long and thin neck) was such that use of the CereTom CT scanner in 3-point fixation was favorable. In Case 2 (a long and thin neck) was such that use of the CereTom CT scanner was not. In the patients reported on here, a CereTom CT scanner, which is specifically designed for cranial imaging and not spinal imaging, was used. The physique of the patient in Case 1 (a long and thin neck) was such that use of the CereTom CT scanner was favorable. The physique of the patient in Case 2 was less favorable, necessitating the use of a scan board designed to fit in the CereTom CT scanner. If these scanners are available, we recommend use of a whole-body or spine scanner such as the O-arm (Medtronic) or BodyTom (NeuroLogica Corp.) for CT-guided tractotomy-nucleotomy procedures.

After surgery, the patient in Case 1 experienced temporary ataxia of the left arm and leg and numbness of the left arm, which significantly improved after 2 days, and he reported complete relief of his postherpetic neuralgia pain. At the most recent visit, 6 months after surgery, the patient reported significant pain relief. The patient in Case 2 did not experience any adverse effects from the lesioning and was completely pain free 3 months after surgery.

The largest series of CT-guided tractotomy-nucleotomy procedures has been reported by Kanpolat et al. and Raslan. Use of general anesthesia for this procedure has not been reported. To our knowledge, this is the first report of percutaneous intraoperative CT-guided tractotomy-nucleotomy in fully anesthetized patients. Given the preliminary safety and success for this procedure in our 2 patients, we advocate the use of general anesthesia for CT-guided tractotomy-nucleotomy procedures.

Conclusions

Use of intraoperative CT imaging to gain adequate radiologic verification of electrode position enables percutaneous CT-guided tractotomy-nucleotomy to be performed on anesthetized patients. Percutaneous intraoperative CT-guided tractotomy-nucleotomy is an effective technique for relief of postherpetic neuralgia and postoperative hemicranial neuralgia.
Disclosure

The authors report no conflict of interest concerning the materials or methods used in this study or the findings specified in this paper.

Author contributions to the study and manuscript preparation include the following: Conception and design: Raslan. Acquisition of data: Raslan, Thompson. Analysis and interpretation of data: all authors. Drafting the article: all authors. Critically revising the article: all authors. Reviewed submitted version of manuscript: all authors. Approved the final version of the manuscript on behalf of all authors: Raslan.

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