

Background

- Occipital neuralgia is a paroxysmal jabbing pain in the distribution of the greater occipital nerve (GON) or lesser occipital nerve (LON) or of the third occipital nerve.¹
- When occipital neuralgia is medically intractable, favorable clinical evidence exists for the use of occipital nerve stimulation (ONS).²
- Communicating branches exist between the GON, the LON, and the auriculotemporal nerve (ATN).³
- Patients have described numbness of posterior auricular and temporal [and retro-orbital] regions after CT-guided C2 and C3 nerve blocks.⁴
- Neurostimulation techniques have been combined with ONS to manage headache pain.⁵
- ONS with infra- or supra- orbital stimulation.
- No neurostimulation techniques have been used in the treatment of complicated migraines.
- Complicated migraine is a non-standardized term describing rare migrainous syndromes which include episodic, transient, and self-limited paroxysmal neurologic dysfunction (prolonged auras), such as hemiplegic migraine, basilar-type migraine, retinal migraine and ophthalmoplegic "migraine."^{1,6}

Case Report

A 44-year old Caucasian woman described headaches with separate and distinct origins—sharp jabbing pain across the occipital base and a throbbing component that encompassed the temporal region bilaterally. Some headaches were accompanied by photophobia, loss of vision, slurred speech, a dropping right eye, and left leg weakness.

Past medical history: absence seizures with familial history.
Surgical history: Le Fort I osteotomy.
Social history: moderate cigarette smoking.

Headache profile: assigned by the referring neurologist:

- Occipital neuralgia, 4-5 episodes / week, 28 year history.
- Complicated migraine, 1 attack / month, 3 year history.

(No recognized triggering factors.)

Previous treatments:

- Occipital nerve blocks, botulinum toxin injections, and rescue medications.
- (Note: ergots and triptans are contraindicated due to the nature of complicated migraines.)

Trialing procedure



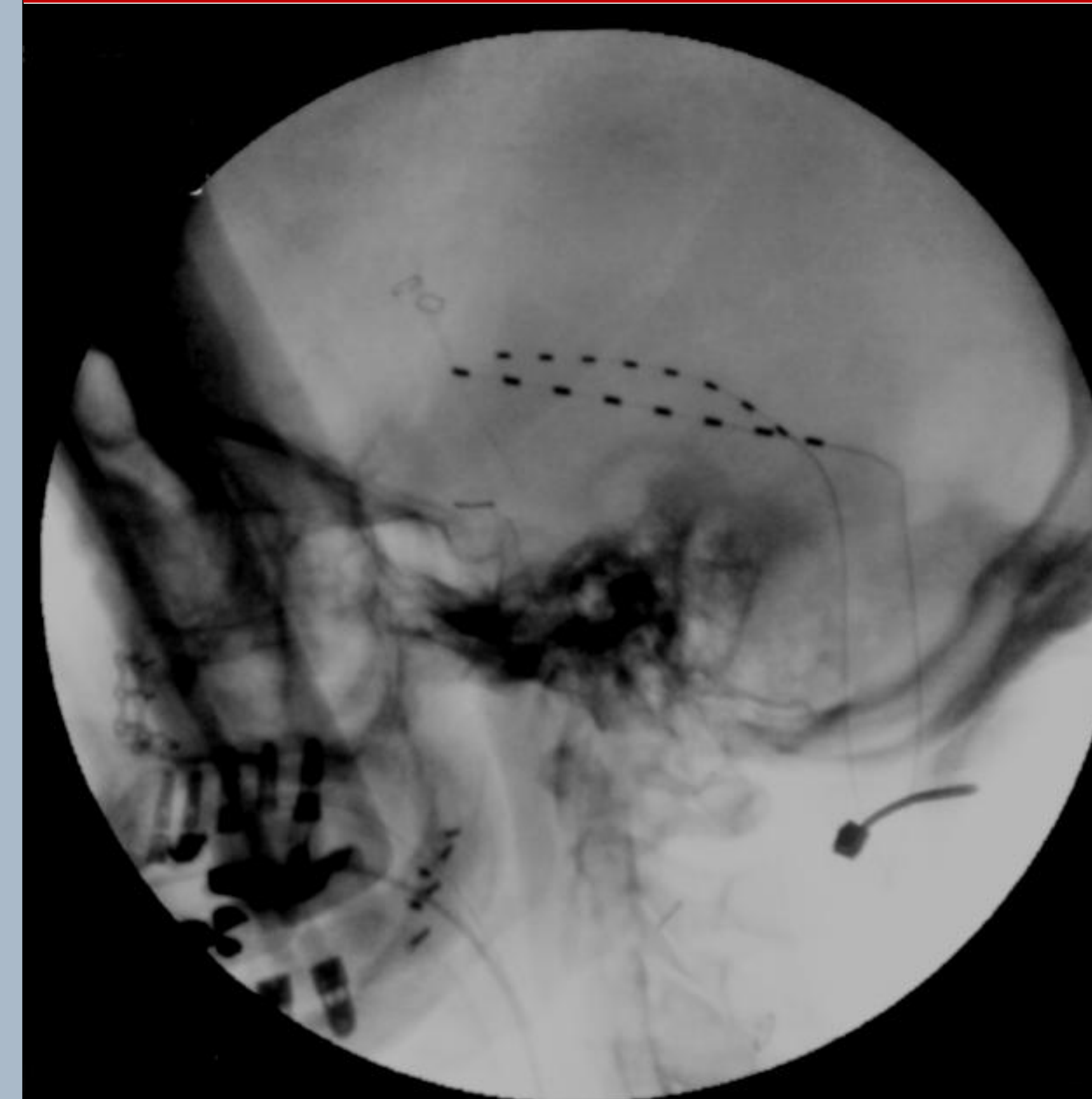
1x8 Standard™ lead over the occiput and 1x4 Quad™ leads bitemporally, Medtronic Neuromodulation, Minneapolis, MN, USA

Case Report – continued

Following a multidisciplinary approach, we focused on suppressing the dual location/complaints of headache pain (occipital and temporal regions).

- 7-day neurostimulation trial:
 - Technique: ONS in combination with bilateral subcutaneous electrical stimulation over the temporalis muscle.
 - Results: headache free during the trial period.
- Device implant: the above technique was implanted, differentiated by use of 1x8 leads over the temporalis muscles. Two Medtronic Restore Ultra™ neuropulse generators were also implanted, each into gluteal pockets created bilaterally.
- 18-months postimplant:
 - Only 2 "major" headaches, reducing the need for rescue medications.
 - No neurologic deficits.
 - 24-hour stimulator use.
 - Patient controls amplitude and rate: prefers slower "drumming" frequencies; amplitudes are consistent with levels reported elsewhere for ONS.⁷
 - No complications or adverse side-effects.

Implant procedure



1x8 Standard™ lead over the occiput and 1x8 Standard™ leads bitemporally, Medtronic Neuromodulation, Minneapolis, MN, USA

Discussion

Complicated migraine: theory of pathogenesis:

- Excessive activation of the sympathetic nervous system (SNS) in response to headache pain.^{8,9}
- Producing sympathetic-driven vasoconstriction and blood platelet activation.^{10,11}

Neurostimulation technique:

- ONS modulates the trigeminocervical complex (TCC).
- Subcutaneous stimulation of the temporal region:
 - Provides alternate routes for convergence on the TCC, due to interconnections between the GON, LON, and ATN.
 - Offers direct local anti-inflammatory and membrane-depolarizing effects, reducing the sensitivity of circulating catecholamines.¹²

With reduced activation of the nociceptive sensitized third and second order neurons throughout the trigeminovascular system, headache pain was reduced—decreasing the likelihood of increased SNS activation.

By "turning-off" the potential for a sympathetic activation-response, the chance for pronounced sympathetic-driven vasoconstriction and blood platelet activation was "turned-off" as well.

Conclusions

The neurostimulation technique suppressed occipital neuralgia, as well as the frequency of migraines.

- Notably, the neurologic deficits that accompanied the migraine attacks were suppressed.

Case presentation provides initial assessment of treatment safety, not conclusive evidence of treatment effectiveness.

- It is reasonable to suggest inclusion of this technique (ONS/subcutaneous temporal-region electrical stimulation) in future studies to better quantify outcomes.

Literature cited

1. International Headache Society. The International Classification of Headache Disorders (ICHD-2) website. <http://ihs-classification.org/en>.
2. Weiner RL, Reed KL. Peripheral neurostimulation for control of intractable occipital neuralgia. *Neuromodulation*. 1999;2:217-221.
3. Becser N, Bovim G, Sjaastad O. Extracranial nerves in the posterior part of the head. *Spine*. 1998;23:1435-1441.
4. Kapoor V, Rothfus WE, Grahovac SZ, Kassam SZA, Horowitz MB. Refractory occipital neuralgia: preoperative assessment with CT-guided nerve block prior to dorsal cervical rhizotomy. *AJNR Am J Neuroradiol*. 2003;24:2105-2110.
5. Slavin KV, Colpan ME, Munawar N, Wess C, Nersesyan H. Trigeminal and occipital peripheral nerve stimulation for craniofacial pain: a single-institutional experience and review of the literature. *Neurosurg Focus*. 2006;21:E6.
6. Rothner AD. Complicated migraine and migraine variants. *Curr Pain Headache Rep*. 2002;6:233-239.
7. Trentman TL, Zimmerman RS, Seth N, Hentz JG, Dodick DW. Stimulation ranges, usage ranges, and paresthesia mapping during occipital nerve stimulation. *Neuromodulation*. 2008;11:56-61.
8. Spierings ELH. New data on the relation between ischemic stroke and migraine. *Journal Watch Neurology*. March 8, 2002.
9. Milhaud D, Bogousslavsky J, van Melle G, Liot P. Ischemic stroke and active migraine. *Neurology*. 2001;57:1805-1811.
10. Röcker L, Drygas WK, Heyduck B. Blood platelet activation and increase in thrombin activity following a marathon race. *Eur J Appl Physiol*. 1986;55:374-380.
11. Visocchi M. Sympathetic activity does influence cerebral blood flow. Comments on "Point: Counterpoint: Sympathetic activity does/does not influence cerebral blood flow." *J Appl Physiol*. 2008;105:1369-1373.
12. Reverberi C, Bonezzi C, Demartini L. Peripheral subcutaneous neurostimulation in the management of neuropathic pain: five case reports. *Neuromodulation*. 2009;12:146-155.