



ISSN Online: 2329-3276 ISSN Print: 2329-3268

# OnabotulinumtoxinA in the Treatment of Occipital Neuralgia Following Gunshot Injury

# Andrew Ea1\*, Terence Gray2

<sup>1</sup>University of New England College of Osteopathic Medicine, Biddeford, USA

<sup>2</sup>Mercy Hospital, Interventional Pain Center, Portland, USA

Email: \*aea@une.edu, grayt@emhs.org

How to cite this paper: Ea, A. and Gray, T. (2016) OnabotulinumtoxinA in the Treatment of Occipital Neuralgia Following Gunshot Injury. *Pain Studies and Treatment*, **4**, 43-47.

http://dx.doi.org/10.4236/pst.2016.44007

Received: August 9, 2016 Accepted: September 4, 2016 Published: September 7, 2016

Copyright © 2016 by authors and Scientific Research Publishing Inc. This work is licensed under the Creative Commons Attribution International License (CC BY 4.0).

http://creativecommons.org/licenses/by/4.0/





# **Abstract**

Occipital neuralgia, while typically idiopathic in presentation, is a common form of posttraumatic headache. It is associated with severe pain in the greater, lesser, and/or third occipital nerves, and often accompanied by tenderness or trigger points in the surrounding musculature. OnabotulinumtoxinA (ONA) has been recently utilized in nerve blocks to treat occipital neuralgia, but current literature supporting such use is scarce. We describe a case of occipital neuralgia in a patient following C1 fracture and vertebral artery dissection due to gunshot injury. Successful treatment with bilateral ONA nerve blocks led to an 80% - 90% improvement in pain, with decreased Visual Analog Scale (VAS) pain scores immediately following treatment and upon follow-up 1 month later.

# **Keywords**

Occipital Neuralgia, OnabotulinumtoxinA, Botox, Bupivacaine, Lidocaine, Gunshot

#### 1. Introduction

Occipital neuralgia is defined by the International Committee for Headache Disorders (ICHD-III) as unilateral or bilateral pain in the distribution of the greater, lesser, and/or third occipital nerves. Diagnostic criteria include the presence of severe or paroxysmal pain, described as shooting, stabbing, and/or sharp in quality. Pain is typically associated with tenderness and trigger points over the affected nerve branches [1]. Occipital neuralgia is usually idiopathic, but is also considered a common form of post-traumatic headache [2]; it has also been associated with the elderly population [3], postoperative complications [4], vascular compression [5], and infectious diseases [6]. Nerve blocks with local anesthetic are commonly used in diagnosis and treatment, but procedures with the potential for more pronounced long-term relief have been explored

DOI: <u>10.4236/pst.2016.44007</u> September 7, 2016

recently, including nerve stimulation [7], pulsed radiofrequency [8], cryoablation [9], and onabotulinumtoxinA.

OnabotulinumtoxinA (ONA), brand name Botox, has been approved to treat a variety of neuropathic pain syndromes, including chronic migraine [10]. Its effect on pain reduction is understood to be primarily peripheral, via the decreased release of neurotransmitters [11], though recent evidence supports an additional central mechanism [12]. There is a dearth of current literature that explores the treatment of occipital neuralgia with ONA. A recent literature review discovered insufficient evidence for such treatment, due to the existence of only three case reports, one retrospective study, and one prospective study [13]. Of these case reports, two described successful ONA nerve blocks of the right greater occipital nerve [14] [15]. A third report detailed ONA treatment of right lesser occipital neuralgia, with limited relief [16].

This report is the first to present ONA treatment of the greater and lesser occipital nerves concurrently, and first to describe bilateral ONA injections for occipital neuralgia. We hope to add to the limited number of case reports detailing successful treatment of occipital neuralgia with onabotulinumtoxinA.

# 2. Case Description

A 50-year-old Caucasian female presented for initial evaluation with a 14-month history of pain in the right occiput radiating to the top of the right scalp. Pain was described as constant and throbbing, and rated as a 5 using the Visual Analog Scale (VAS) for pain. Secondary complaints included several tender points over the patient's right scapula, right arm weakness, and intermittent numbness and tingling in all digits bilaterally. The patient sustained a gunshot wound through the mouth 14 months earlier, fracturing the 1<sup>st</sup> cervical vertebra (C1) and leading to right vertebral artery dissection and aneurysm following stent placement. Cervical spine flexion and extension x-rays taken 3 months after the injury revealed evidence of comminuted C1 fracture and mild anterolisthesis of C2 on C3 upon flexion. Previous unsuccessful treatments included physical therapy, acupuncture, application of heat, and lidocaine injections in the right shoulder and neck. Pain medications prescribed prior to evaluation included aspirin 81 mg, gabapentin 300 + 600 mg, and oxycodone HCl 5 mg.

Diagnostic nerve blocks with local anesthetic were performed 3 weeks later. A total of 3 ml of 0.25 bupivacaine and 3 ml of 1% lidocaine plus 40 mg triamcinolone was injected into the right greater and lesser occipital nerves and the right auriculotemporal nerve. The patient reported a VAS score of 7 prior to injection, which decreased to 2 immediately following treatment. Upon follow-up 1 month later, she reported a VAS score of 5, reporting a 25% overall improvement in pain and functionality. Slightly greater than 50% relief was experienced for several weeks; the patient stated this to be the first period of pain relief since her injury. A diagnosis of occipital neuralgia was made based on ICHD-III criteria, including severe paroxysmal pain in the distribution of the greater and lesser occipital nerves, tenderness over the affected nerves, and temporary pain relief by local anesthetic block [1].

Right occipital and auriculotemporal nerve blocks were repeated after another 3 weeks, leading to a VAS score decrease from 7 to 0 following treatment. The patient reported 100% relief for 3 days during follow-up, with an overall improvement of 70% in right occiput and lateral neck, and 30% in the right parietal area. Stated VAS scores in the right occipital and right parietal areas were 1 and 4, respectively. The decision was made to progress to ONA treatment.

Occipital nerve blocks with onabotulinumtoxinA were performed 4 months after initial evaluation. 120 units of ONA were equally divided among 24 injection points, along the trapezius, occipitalis, temporalis, and cervical paraspinal muscles bilaterally. The patient reported immediate pain relief, from a pre-procedure VAS score of 5 to 0 post-procedure. Upon follow-up 1 month later, the patient stated that the ONA injections "were different" than previous treatment with local anesthetic, leading to complete resolution in the pain on the top and right side of her skull, in addition to her right shoulder, neck and arm. The only remaining pain was experienced in the right occiput in a 2 - 3 cm diameter, with a pain score of 3. The patient reported an overall improvement in symptoms of 80% - 90%, stating, "this is the closest to normal I have ever been". An additional follow-up after 4 - 6 weeks was recommended.

#### 3. Discussion

We describe a case of occipital neuralgia following C1 fracture and right vertebral artery dissection due to gunshot injury. After two successful occipital nerve blocks with local anesthetic, treatment with ONA resulted in greater overall improvement of symptoms, at 80% - 90%, and a decreased VAS score of 3 upon follow-up. These findings support the efficacy of onabotulinumtoxinA in treating the pain associated with occipital neuralgia, while providing a unique case of successful treatment following severe trauma. Sustained ONA treatment may provide long-term relief from occipital neuralgia and other pain conditions, while avoiding the potential side effects and increased morbidity of pharmacology or surgical intervention [17]. Further treatments are necessary to examine the results of repeated injections, which have been recently suggested to increase the duration of ONA's effects [18].

#### Disclosure

The authors have no conflicts of interest to report.

### References

- [1] Headache Classification Committee of the International Headache Society (IHS) (2013) The International Classification of Headache Disorders, 3rd edition. *Cephalalgia*, **33**, 629-808.
- [2] Evans, R.W. (2014) Posttraumatic Headaches in Civilians, Soldiers, and Athletes. *Neurologic Clinics*, **32**, 283-303. <a href="http://dx.doi.org/10.1016/j.ncl.2013.11.010">http://dx.doi.org/10.1016/j.ncl.2013.11.010</a>
- [3] Ruiz, M., Pedraza, M.I., de la Cruz, C., Barón, J., Muñoz, I., Rodríguez, C., Celorrio, M., Mulero, P., Herrero, S. and Guerrero, A.L. (2014) Headache in the Elderly: Characteristics in a Series of 262 Patients. *Neurologia*, **29**, 321-326.

#### http://dx.doi.org/10.1016/j.nrl.2013.07.007

- [4] Gautschi, O.P., Payer. M., Corniola, M.V., Smoll, N.R., Schaller, K. and Tessitore, E. (2014) Clinically Relevant Complications Related to Posterior Atlanto-Axial Fixation in Atlanto-Axial Instability and Their Management. *Clinical Neurology & Neurosurgery*, **123**, 131-135. <a href="http://dx.doi.org/10.1016/j.clineuro.2014.05.020">http://dx.doi.org/10.1016/j.clineuro.2014.05.020</a>
- [5] Shimizu, S., Oka, H., Osawa, S., et al. (2007) Can Proximity of the Occipital Artery to the Greater Occipital Nerve act as a Cause of Idiopathic Greater Occipital Neuralgia? An Anatomical and Histological Evaluation of the Artery-Nerve Relationship. Plastic and Reconstructive Surgery, 119, 2029-2034, discussion 2035-2036. <a href="http://dx.doi.org/10.1097/01.prs.0000260588.33902.23">http://dx.doi.org/10.1097/01.prs.0000260588.33902.23</a>
- [6] Kihara, T. and Shimohama, S. (2016) Occipital Neuralgia Evoked by Facial Herpes Zoster Infection. *Headache*, 46, 1590-1591. <a href="http://dx.doi.org/10.1111/j.1526-4610.2006.00616">http://dx.doi.org/10.1111/j.1526-4610.2006.00616</a> 2.x
- [7] Sweet, J.A., Mitchell, L.S., Narouze, S., Sharan, A.D., Falowski, S.M., Schwalb, J.M., Machado, A., Rosenow, J.M., Petersen, E.A., Hayek, S.M., Arle, J.E. and Pilitsis, J.G. (2015) Occipital Nerve Stimulation for the Treatment of Patients With Medically Refractory Occipital Neuralgia: Congress of Neurological Surgeons Systematic Review and Evidence-Based Guideline. *Neurosurgery*, 77, 332-341.
- [8] Manolitsis, N. and Elahi, F. (2014)Pulsed Radiofrequency for Occipital Neuralgia. *Pain Physician*, **17**, E709-E717.
- [9] Kim, C.H., Hu, W., Gao, J., Dragan, K., Whealton, T. and Julian, C. (2015) Cryoablation for the Treatment of Occipital-Neuralgia. *Pain Physician*, **18**, E363-E368.
- [10] Dodick, D.W., Turkel, C.C., DeGryse, R.E., Aurora, S.K., Silberstein, S.D., Lipton, R.B., Diener, H.C. and Brin, M.F., PREEMPT Chronic Migraine Study Group (2010) Onabotulinumtoxin A for Treatment of Chronic Migraine: Pooled Results from the Double-Blind, Randomized, Placebo-Controlled Phases of the PREEMPT Clinical Program. *Headache*, 50, 921-936. http://dx.doi.org/10.1111/j.1526-4610.2010.01678.x
- [11] Francisco, G.E., Tan, H. and Green, M. (2012) Do Botulinum Toxins Have a Role in the Management of Neuropathic pain?: A Focused Review. *American Journal of Physical Medicine & Rehabilitation*, **91**, 899-909. <a href="http://dx.doi.org/10.1097/PHM.0b013e31825a134b">http://dx.doi.org/10.1097/PHM.0b013e31825a134b</a>
- [12] Matak, I., Bach-Rojecky, L., Filipović, B. and Lacković, Z. (2011) Behavioral and Immuno-histochemical Evidence for Central Antinociceptive Activity of OnabotulinumtoxinA. *Neuroscience*, **186**, 201-207. http://dx.doi.org/10.1016/j.neuroscience.2011.04.026
- [13] Brown, E.A., Schütz, S.G. and Simpson, D.M. (2014)Botulinum Toxin for Neuropathic Pain and Spasticity: An Overview. *Pain Management*, 4, 129-151. <a href="http://dx.doi.org/10.2217/pmt.13.75">http://dx.doi.org/10.2217/pmt.13.75</a>
- [14] Narouze, S. and Souzdalnitski, D. (2013) Occipital Nerve Entrapment within the Semispinalis Capitis Muscle Diagnosed with Ultrasound. *Cephalalgia*, 33, 1358-1359. http://dx.doi.org/10.1177/0333102413492912
- [15] Volcy, M., Tepper, S.J., Rapoport, A.M., Sheftell, F.D. and Bigal, M.E. (2006) Botulinum Toxin A for the Treatment of Greater Occipital Neuralgia and Trigeminal Neuralgia: A Case Report with Pathophysiological Considerations. *Cephalalgia*, 26, 336-340. <a href="http://dx.doi.org/10.1111/j.1468-2982.2005.00959.x">http://dx.doi.org/10.1111/j.1468-2982.2005.00959.x</a>
- [16] Siefferman, J. and Khelemsky, Y. (2015) Occipital Neuralgia after Hair Transplantation and Its Treatment. *Case Reports in Neurological Medicine*, 2015, Article ID: 428413. <a href="http://dx.doi.org/10.1155/2015/428413">http://dx.doi.org/10.1155/2015/428413</a>
- [17] Alshadwi, A., Nadershah, M. and Osborn, T. (2015) Therapeutic Applications of Botulinum Neurotoxins in Head and Neck Disorders. *The Saudi Dental Journal*, **27**, 3-11. http://dx.doi.org/10.1016/j.sdentj.2014.10.001

[18] Lecouflet, M., Leux, C., Fenot, M., Célerier, P. and Maillard, H. (2014) Duration of Efficacy Increases with the Repetition of Botulinum Toxin A Injections in Primary Palmar Hyperhidrosis: A Study of 28 Patients. *Journal of the American Academy of Dematology*, 70, 1083-1087. http://dx.doi.org/10.1016/j.jaad.2013.12.035



# Submit or recommend next manuscript to SCIRP and we will provide best service for you:

Accepting pre-submission inquiries through Email, Facebook, LinkedIn, Twitter, etc.

A wide selection of journals (inclusive of 9 subjects, more than 200 journals)

Providing 24-hour high-quality service

User-friendly online submission system

Fair and swift peer-review system

Efficient typesetting and proofreading procedure

Display of the result of downloads and visits, as well as the number of cited articles

Maximum dissemination of your research work

Submit your manuscript at: <a href="http://papersubmission.scirp.org/">http://papersubmission.scirp.org/</a>