Diagnosis and treatment of cervicogenic headache are both clinically challenging; as a result, this disorder remains one of the most controversial in headache medicine. Between 50% and 75% of patients with whiplash injury may have headache related to a disorder of the neck, whereas in the general population, the prevalence of cervicogenic headache has been estimated to be 4.1%. Given that the pathology in cervicogenic headache is thought to lie in the cervical spine area, the mechanism of the actual head pain, which often involves the frontal and temporal areas, likely relates to convergence between cervical nociceptive and trigeminal afferents in the trigeminocervical complex at in the upper cervical spine.

Diagnostic criteria for cervicogenic headache were published by Sjaastad et al in 1998, and these were the diagnostic criteria used by Boudreau et al when selecting patients for their study, which is published in the Canadian Journal of Neurological Sciences. Boudreau et al required that all patients have strictly unilateral headache without side shift. Their patients also had to be severely affected in that at least 4 hours of continuous pain of moderate to severe intensity per day were required, and patients had to have headache on at least 15 days per month. Finally, patients had to have an abnormal neck examination with what they termed signs of neuropathic involvement in the territory of the upper cervical nerve roots. The presence of alldynia—in other words, some degree of central sensitization—appears to have been sufficient to meet this criterion.

The Third Edition of the International Classification of Headache Disorders (ICHD-3 beta) divides chronic daily headache related to the neck into two disorders. The first is persistent headache attributed to whiplash. It is defined as a new headache of any phenotype that begins within 7 days of a whiplash injury, lasts at least 3 months, and is not better accounted for by another diagnosis. The second headache type related to the neck is cervicogenic headache. This is defined as a headache that occurs in a patient with clinical, laboratory, or imaging evidence of a neck disorder known to cause headache. Several criteria are given that are considered to provide evidence of causation, and the headache should not be better accounted for by another diagnosis. Of note, in the study by Boudreau et al, at least 85% of patients were reported to have neck trauma as a triggering factor; therefore, the majority of their patients may have fallen into the category of persistent headache attributed to whiplash headache as defined by the ICHD-3. Both the Sjaastad criteria and the ICHD-3 criteria for cervicogenic headache stress the importance of pain abolition by anesthetic blockade of cervical structures, although in the ICHD-3 criteria such blockade is no longer mandatory for the diagnosis. Diagnostic nerve blocks were not used by Boudreau et al in this study, and in fact patients booked for somatic nerve blocks were excluded. Patients appear to have been diagnosed purely on clinical features.

The neck has many potential pain generators. Clinical studies have indicated that the C2-3 zygapophysial joints are one of the most common sources of pain in cervicogenic headache, particularly in those with whiplash injury or motor vehicle accidents, accounting for 50% of cases. Unfortunately, manual examination is not reliably accurate in detecting zygapophysial joint pain, nor is medical imaging beneficial in localizing the pain generator. Studies with magnetic resonance imaging in patients with cervicogenic headache reported no demonstrable differences in the appearance of cervical spine structures on magnetic resonance imaging scans in comparison with control subjects and subjects with other headache disorders. Imaging, therefore, plays a role primarily to assess other causes of headache that may require surgical treatment. To achieve pain control of cervicogenic headache, it would appear best to employ a multimodal approach with pharmacological, manipulative, and invasive interventions.

Medications alone are often ineffective or provide only modest benefit for cervicogenic headache. Medications commonly used include antidepressants, antiepileptic drugs, analgesics, and muscle relaxants, but none of these medications has been studied in controlled clinical trials. Medical management of cervicogenic headache therefore relies on the anecdotal experience of clinicians and/or is based on treatment of other headache disorders. A patient with headache following whiplash injury who demonstrated a substantial response to treatment with botulinum toxin has been reported, but a randomized controlled trial of botulinum toxin in cervicogenic headache was negative. It may be, however, that optimal injection protocols for cervicogenic headache have not yet been developed.

Boudreau et al report a single center, double-blind, randomized controlled study on the use of pregabalin in patients with cervicogenic headache in a relatively severely affected patient cohort. To be included, patients had to have proven refractory to commonly used medications including neuromodulators, muscle relaxants, tricyclics, and nonsteroidal anti-inflammatory drugs. Patients were excluded if they used rescue medications.

This manuscript served as a companion piece for the original article “Pregabalin for the Management of Cervicogenic Headache: A Double Blind Study” by Guy P. Boudreau and Luc Marchand, which was published in the September 2014 issue of The Canadian Journal of Neurological Sciences.

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Administration approval for use in neuropathic pain syndromes patients with neuropathic pain and has US Food and Drug therefore cannot be assumed. Pregabalin has been widely used in izability of their findings to all patients with cervicogenic headache hypoesthesia, dysesthesia, allodynia, or hyperalgesia. The general-pathic involvement in the C2 or C3 distribution with hyperesthesia, to previous trauma. It should be noted, however, that all patients in this trial were required to have evidence on examination of neuropathic involvement in the C2 or C3 distribution with hyperesthesia, hypoesthesia, dysesthesia, allodynia, or hyperalgesia. The general-izability of their findings to all patients with cervicogenic headache therefore cannot be assumed. Pregabalin has been widely used in patients with neuropathic pain and has US Food and Drug Administration approval for use in neuropathic pain syndromes including diabetic neuropathy, postherpetic neuralgia, and neuropathic pain in spinal injury. It is also known to have anxiolytic properties. These factors may explain the positive results of pregabalin on pain and mood in this study. It is unclear at this point whether pregabalin is beneficial in patients with cervicogenic headache without clinical neuropathic involvement. As the authors point out, further research and larger studies are needed to define the role of pregabalin in the treatment of cervicogenic headache, but the randomized controlled trial by Boudreau et al is certainly an important step in the right direction. The authors are to be congratulated on a study, which finally casts some scientific light on the medical treatment of this controversial and often refractory headache disorder.

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