



UNRAVELLING THE MYSTERY OF TRIGEMINAL NEURALGIA: INSIGHTS INTO DIAGNOSIS AND MANAGEMENT - AN OVERVIEW

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ABSTRACT

A rare and painful condition known as trigeminal neuralgia, this condition is typified by abrupt, unbearable, paroxysmal, piercing, stabbing pain that is frequently unilateral throughout the trigeminal dermatomes. While the illness generally is not lethal, it is debilitating and can occasionally end in psychological disorders. This disorder diminishes the quality of everyday life. To ease the pain and suffering of a patient's agony, a precise diagnosis and an effective therapeutic approach are essential. A comprehensive examination of the diagnosis, management, and rehabilitation of trigeminal neuropathy is given in this article. This includes pharmaceutical administration, surgical management, alternative recently discovered methods of treatment and a screening approach. Latest innovations and newer guidelines of treatment are also discussed to enhance clinical outcomes and improve patients' prognosis.

KEYWORDS : nerve compression, anticonvulsant, radiosurgery, facial pain, and surgical intervention

INTRODUCTION:

An uncommon dentofacial distress recognized as trigeminal neuralgia (TN) is frequently brought by compaction caused by arteries and veins. A global headache association explains it as repeated, unambiguous, brief, electric shock-like discomfort that stops on its own, is abruptly initiated, restricted to the branching of several nerve divisions, or is induced by harmless stimuli. Fothergill identified the ailment as *tic douloureux* after diagnosing the first case of it in 1756 initially he called it a painful affection of the face but later on, John Hunter clarified that he meant it to be a type of nerve illness. This condition was medically managed with powdered mercury, the drug, opium and arsenic in the 19th and 18th centuries.⁽¹⁾ In 1994, phenytoin was found to deliver useful outcomes for trigeminal neuralgia. The precise physiology or the development of this condition are still in debate. However thorough research has been conducted mostly on classical trigeminal neuralgia. However thorough research has been conducted mostly on classical trigeminal neuralgia. The condition starts when the trigeminal fibres in the pons or dorsal root entrance zone sustain damage to their core myelin; the "ephapsys" caused by these myelinated nerve fibre losses permit all tactile proprioceptive and temperature stimuli to be converted into pain. An overwhelming amount of pain stimuli sensitizes the trigeminal cells, causing them to fire continuously. Nerves become hyper-excited in the peripheral because of more pain stimuli, and sensitivity in the center advances. Non-painful stimuli can activate these hyperexcitable nerve cells, which causes them to react with semi-epileptic painful episodes. Paroxysmal attacks become less frequent as the damage gets worse, and persistent neuropathic pain starts.⁽²⁾ Neurovascular constriction induced by an aberrant artery or blood vessel, arteriovenous abnormalities, vestibular tumour, meningioma, epidermoid cyst and other cysts and tumours, and vessel aggregation in the area anterior to the pons at the nerve root entrance zone. While other arteries or veins may also contribute to 5th cranial nerve compaction.⁽³¹⁾

Along with identifiable sources of facial discomfort, this kind of disorder is typified by the existence of trigger zones devoid of any observable neurological deficiency. An innocent touch to the cheek, while cleaning teeth, or when opening the mouth may cause this pain. It is possible to characterize the discomfort as electric shock-like, acute, stabbing, sporadic, abrupt, severe, piercing, and paroxysmal. Before determining the exact cause, a physician must rule out other possible reasons for facial pain. These may include:

Crack Tooth: An incomplete breakage and crack in a crucial dental region affecting the connective tissue and nerves is known as crack tooth syndrome. When the patient expresses their suffering, it closely resembles trigeminal neuropathy. The pain is commonly characterized as burning or jabbing, and it is identified as TMJ disorders or similar oral-facial difficulties. Among widely accepted methods that rule out crack teeth syndrome were selective biting testing and transillumination.

Post-traumatic trigeminal nerve impairment is the term used to refer to discomfort that develops within locations related to the trigeminal

nerve post-injury to the dentofacial region and exhibits all the characteristics of facial nerve damage. This could be the consequence of regular surgeries like endodontic procedures, tooth extractions, or various iatrogenic and physical traumas. People who are affected frequently describe the discomfort as shooting, pyrogenic, or recurring. A precise diagnosis can be reached with the aid of a careful or thorough background recording.

Glossopharyngeal Neuralgia is a unique kind of mouth awkwardness that does not look like trigeminal nerve pain. It produces abrupt bursts of tenderness throughout the neck or throat region; individuals usually describe it as a stabbing knife-like feeling in these areas and sporadically extending discomfort to the ears. The position of the pain—which must occur in the rear segment of the lingual apparatus, tonsillar cavity, angle of the mandible, or the ears—is the main difference between trigeminal neuralgia (TN) and glossopharyngeal neuropathy.

Nervus Intermedius Neuralgia: A rare condition called nerve intermedius neuropathy can cause brief bursts of ear discomfort which may radiate to the parieto-occipital region. Once it leaves the brainstem, this neuron enters the internal auditory meatus. To differentiate between nervus intermedius neuralgia and trigeminal neuralgia (TN), it is critical to consider the primary complaint of the patient.

Short-lasting unilateral neuralgiform headache with conjunctival injections and tearing (SUNCT): An atypical entity⁽³⁸⁾ that mostly impacts individuals who are in the middle decades of life range. The pain with this condition produces is usually scorching, shooting, or electric, which can range in severity from moderately to extreme. It affects both the eye and periocular regions. The presence of conjunctival injections and tears are characteristics that set this illness apart from trigeminal neuralgia.

Postherpetic Neuralgia: In senior citizens, the reappearance of herpes zoster virus is prevalent, causing postherpetic neuralgia. Vesicular eruptions over those areas involved in trigeminal nerve terminals indicate its occurrence. It is linked to post-herpetic neuralgia, encephalitis and eyesight loss. It can occur from ageing or a bacteria-related infection. Antiviral drugs are frequently employed to treat this illness.

Post-operative complications, the Trigeminal nerve being almost entirely sensory, when undergoing surgeries often show hypoesthesia, dysesthesia, or loss of sensation. Invasive, non-ablative procedures have definitive deficits of cranial nerve VIII, VII, and IV in less than 5% of cases. Invasive ablative procedures can offer a degree of damage to the nerve as well, Corneal reflex loss, keratitis, and anesthesia dolorosa, which is quite rare, causing numbness associated with nerve damage and masticatory muscle weakness. In non-invasive ablative procedures diplopia, hearing loss, mastication muscle weakness can be seen.⁽³⁾

Therefore, if there are four or more unilateral bouts of face discomfort such as; it involves either all or some trigeminal nerve parts; both the beginning and end of pain are abrupt, whereas there is no pain radiation outside this nerve's dispersion. Pain that exhibits at least three of the following four features: it is intense, recurrent in paroxysmal episodes that can last up to 120 seconds, simulating electrical shockwave, firing, or knifing, and it is brought on by harmless inputs on a portion of the skin on the face that is impacted. Attacks are individual to all affected persons and neurologic deficit is absent in these individuals.⁽⁴⁾

Hence, additional neurological assessments are discussed below that can be used for further examination of certain clinical criteria as novel analytical techniques for trigeminal nerve pain (TN) as follows.

Computed Tomography (CT): With the use of digital geometric processing, a sequence of 2D x-ray images recorded around the axis of rotation can be converted into 3D images for use in medical imaging scans like computed tomography (CT). It is used to identify nerve damage brought on by multiple sclerosis, an inflammatory condition that causes nerve inflammation and the destruction of the myelin sheath. Trigeminal neuralgia and multiple sclerosis can occasionally coexist, and facial nerve tumours, sinusitis, and neurovascular conflicts can also result in nerve injury.

Magnetic Resonance Imaging (MRI): Utilizing fields of magnetic attraction and radiofrequency, the technique of magnetic resonance imaging (MRI) produces finely precise pictures that depict the body's inner organs. It is regarded as the gold standard for assessing diseases like trigeminal neuralgia (TN) because it can identify potential trigeminal nerve compression sites in the vasculature. Moreover, it can assist in ruling out any further reasons for TN, such as cysts or tumours that might be pinching the trigeminal nerve root. Individuals who have heart rate monitors or defibrillators are unable to have an MRI because magnetic waves can cause problems with such appliances, even though there is no biological risk involved.

Magnetic resonance angiography (MRA): An approach based on MRI principles seeks to identify any neurovascular problems using the foramen ovale. A substance that contrasts: gadolinium therapy or iodine is injected into Meckel's cave using this technique while few examples of gadolinium accumulation in the brain have been reported; these investigations have not shown any neurological harm.

Treating trigeminal neuralgia (TN) can be handled with two sorts of therapies: pharmaceuticals and operations, i.e. in first-line approaches, medical treatments should be the initial line of care; only if drugs do not work can surgery be considered. Roughly 33-50% of patients can need surgical intervention during some stage. In this article, we will examine the level as well as the duration of pain reduction associated with each course of therapy. As a result, several medications are utilized to relieve pain produced by neural illnesses. For decades, anticonvulsant medicines like Carbamazepine and oxcarbazepine have been effectively practiced as the earliest curative approach for trigeminal neuralgia, and they can reduce distress by up to 90% in most patients. It interacts strongly with other medicines that patients may be on, and oxcarbazepine causes substantial neural system depression.

The medication tolerability determines the progression of the therapy. Carbamazepine dosage spans 200-1200 mg per day, whereas oxcarbazepine dosage spans 600-1800 mg per day. Some of the most often utilized second-line approaches include gabapentin, lamotrigine, and baclofen. These medications have fewer undesirable effects and greater potency. These are used when the patient's primary-line medications are not well tolerated. Due to the presence of acute rashes and other harmful reactions, a slow-dose titration protocol is used. These medications work through three major mechanisms i.e. augmentation of GABA action, inhibition of sodium channel functions, and inhibition of calcium channel functions. Various first-line and second-line drug approaches are mentioned in table 1.⁽⁵⁾

A surgical procedure is frequently necessary to aid individuals who develop drug resistance or who have intolerable side effects. The present treatments are not lasting, but they can provide temporary pain relief without achieving full remission. The extent and length vary depending on the therapy's option. The several sorts of surgical operations are discussed below.

Microvascular Decompression: In 1925, Walter Dandy described microvascular decompression as a key factor in treating trigeminal

neuralgia (TN). It is strongly recommended as the treatment of choice due to its long-lasting pain relief, safety, and effectiveness. This procedure aims to target the area where the trigeminal nerve root intersects the nerve pons, requiring surgery on the posterior fossa involving suboccipital craniectomy and nerve combing. It is recommended only for cases of classical trigeminal neuralgia following a high-resolution contrast-enhanced MRI indicating a neurovascular conflict. CT can also be utilized to pinpoint the compressed region of the trigeminal nerve.⁽⁶⁾ Sometimes, a division of the basilar artery or the vena petrosa sparks compaction.⁽³²⁾⁽⁴⁰⁾ Over 70% of patients were symptom-free devoid of pharmacotherapy for 10 years term post-surgery, and 75-80% remained pain-free after 15 years.⁽⁶⁾ In a previous study, a fatality ratio between 0.2-1% was considered an operative deviation. However, recent findings from two large series involving 444⁽⁷⁾ and 1995⁽⁸⁾ patients showed no deaths. Even though the MVD's reputation has certainly grown over the last two decades, there is currently a discussion concerning its benefit from the neurodestructive method.

Percutaneous balloon compression (PBC) provides an unexplained solution, but it is unknown whether it is the most useful method. Each method has a set number of challenges and occurrences. Its earliest appearance was in 1978 and was issued by Mullan and Lichtor in 1983. The first trial included 50 patients who received gasserian ganglion compaction employing a Fogarty balloon. The operation is carried out upon spinal anaesthesia, with a fluoroscopically guided cannula inserted within a foramen ovale and never stretched further. The Fogarty catheter is placed into Meckel's cave until it fills the entire hollow. The catheter tips contain 0.5-1.0 mL of contrast-enhancing substance. The compaction is sufficient, lasting from one to six minutes in total. While this technique causes some little sensory impairment, nearly all patients get immediate pain alleviation.⁽⁹⁾ The benefits of the percutaneous approach are its ease and speed from a scientific point of view; practical identification does not require involvement; and its ability to be carried out on a patient throughout only a short time of widespread sedation with minimal pain. It is believed that all precautions should be taken to prevent hypotension and bradycardia during surgery.⁽¹⁰⁾

Radiofrequency ablation was discovered to be an efficient and non-intrusive method towards managing TN, especially contrasted with balloon compression and microvascular decompression. It is widely acknowledged for its security, potency, and successful percentage of happiness among affected individuals.⁽¹¹⁾ According to a 2016 comprehensive review research, radiofrequency ablation was found to contain an average of 5-10% incidences of reappearance and a between 85 and 90 per cent rate of survival for treating TN.⁽¹²⁾ A study done in 2019 including 100 participants that recently appeared in the Journal of Neurosurgery indicated that 98% of procedures were successful following an average monitoring of one year.⁽¹³⁾ In 2020 research reported that radiofrequency ablation had a more economically viable alternative for TN than microvascular decompression. Furthermore, individuals with haemophilia or specific types of tumors, should not undergo radiofrequency ablation.⁽¹³⁾ This could result in infections, hemorrhagic tendencies, infection, and injuries to the nerves, among possible side effects. Although these dangers are usually small, they are to be considered when assessing radiofrequency ablation as a substitute for remedy.⁽¹⁴⁾ Thus, radiofrequency ablation is a potent alternative for medication overall for persistent TN, but it has drawbacks. While undergoing therapy, it can also be costly, and healthcare may not cover treatment and therefore might be a barrier for certain individuals.

Glycerol Rhizotomy (GR): In 1904, Schloesser gave some of the earliest described administrations of alcohol through the trigeminal neurons. It has been found by coincidence; that glycerol has been utilized as an agent for injecting tantalum particles through the trigeminal canal.⁽¹⁵⁾ Precise process underlying glycerol injection-induced injury to big myelinated fibers is believed to be caused by a rapid shift in intracellular osmolarity, and axonal degeneration with shattering. In a study, it was found that the treatment provided mild-to-moderate pain relief, which usually lasted between 3 and 6 months with an average span of 11 months.⁽¹⁶⁾ Over one year, around 70% of the patients stated that they were able to manage their pain.⁽¹⁷⁾ In some patients, post-procedure symptoms such as reduced sensation and temporary facial paralysis were observed, usually from hours to days after the procedure.⁽¹⁷⁾⁽¹⁸⁾ While severe reductions in sensitivity and pain relief are rare, individuals who undergo ongoing treatments may be more vulnerable.⁽¹⁹⁾ However, the percentages of oral haemorrhage and

epithelial penetration are reported following the carotid stab rate.^(18,20)

Gamma Knife Radiosurgery: This protocol, invented by Lars Leksell⁽³³⁾ over half a century ago, is a radiosurgical tool that replaces open intracranial procedures in the treatment of cognitive neurological disorders. Nonetheless, the proximal trigeminal root close to the pons has been the goal in recent years, with substantially better results.⁽²¹⁾ In this method, the individual receiving treatment lies down and their forehead is protected by an electromagnetic collimator mask. Securing the frame is done under local anaesthesia, and irradiation is often done under intravenous or moderate mouth sedation.⁽²²⁾ Usually, relief from discomfort takes time to manifest. There was evidence of localized axonal degeneration for six months afterwards following 80 Gy to 100 Gy of recommended radiation. This seems to be associated with MRI enhancement of contrast observed several months following radiosurgery with no visible ganglion alterations.⁽²³⁾ As a result, it is now recognized as one of the main therapeutic methods for TN for individuals whose alternative therapies have failed. Among the other neurosurgical treatment techniques, it has the best level of security and is the least invasive. It has drawbacks, a delay until pain alleviation and an elevated probability of recurrence at prolonged examination. It is anticipated that current advances in neuroscience will enhance these methods and clinical results for TN individuals.⁽²⁴⁾

Peripheral Nerve Resection: Straightforward, relatively secure treatment for any terminating branch of all three trigeminal divisions. This operation can be performed without general anaesthesia in an outpatient environment. This type of surgery is known as postganglionic surgery and involves severing the nerve after it exits the skull. Peripheral nerve regeneration is the secondary cause of pain recurrence. Ali et al carried out a study in rural India where 14 individuals affected with trigeminal neuralgia, who were medically unfit for major surgeries and who could not afford the other treatments or patients who were unresponsive to conservative therapy were selected. Throughout 12 to 24 months beyond the operation, many patients experience pain recurrence. A second neurectomy may be used to address recurrent pain, however these operations frequently result in less long-lasting pain relief.⁽²⁵⁾

Cryotherapy: Lloyd in 1976 published the first description of cryotherapy for inhibiting peripheral nerves. Pradel et al invented a cryoprobe with a radius of 1.3mm which harnesses liquid nitrogen to cool the tip of the probe. It can reach up to a maximum of 158°C. The temperature reached at the nerve determines the duration of numbness and the speed of pain relief. The guidance structures for nerve fibre regeneration are critical to the healing phase with the resulting repair of perception. The probe's tip is placed transmucosal over the nerve with extreme accuracy. The ablated nerve stands anaesthetized and begins to regain its sensory output in the first 3 months. Few physicians use this to be a primary management protocol for TN-affected individuals since the results are generally less favourable in terms of pain-free longevity when compared to other percutaneous methods.⁽²⁶⁾

Alcohol Block: When treating trigeminal neuralgia, alcohol nerve block was often administered; however, these days, percutaneous techniques or microvascular decompression are more frequently utilized. While frequent administration of peripheral phenol or alcohol-based injections may induce adjacent membranous damage to the site of administration such as tenderness along with fibrotic scarring, there is a continuous discussion on the injection's suitability for managing trigeminal neuralgia, even though it offers moderate pain relief.⁽³⁴⁾⁽³⁶⁾ The principal adverse impact caused by alcohol doses is the transient weakening of the masticatory apparatus, as the motor root is near the inferior alveolar nerve.⁽³⁵⁾

Transcutaneous Electric Nerve Stimulation [TENS]: It is a treatment that essentially includes running an electronic impulse beneath the upper dermis area to stimulate receptors more thoroughly, decreasing discomfort. Patients have the potential to alleviate both severe and persistent pain with TENS therapy.⁽²⁷⁾ Several research studies compared the effects of two different TENS modes, constant mode and burst mode, over three weeks. A constant mode of TENS was shown to considerably reduce the Visual Analog Score from a mean value of 9 to 20 in 26 out of 31 patients, indicating considerable improvement. Both burst and steady modes of TENS were helpful; however, some studies of 30 to 40 patients using only the burst mode showed a noticeable rise in the degree of alleviating discomfort. It therefore serves as an appropriate care option for the condition

attributable to its non-invasiveness, affordability, security, and low adverse effects.⁽³⁷⁾ It can also be utilized for individuals who have intolerable side effects, are unable to tolerate medicine, or are not responding effectively to past therapy approaches.

Botulinum Toxins (BNT): In 2010, a whole new application for BT arose with the approval of BT medications for the therapy of severe vertigo. This condition can be safely and effectively controlled with an injectable botulinum toxin A; once the procedure terminated, the highest potency was observed between three and six weeks; most side effects disappeared in a week, and headaches in the symmetry of the face were relieved immediately after the dose. More research is needed to determine the longest time a patient can be pain-free, the most effective injection method, the optimal dosage of BT-A, any drug interactions with potential consequences, and to design the best course of care.⁽²⁸⁾ The studies reported significantly less discomfort, although additional double-masked trials will be essentially needed to verify the positive effects of the approach despite its drawbacks.⁽²⁹⁾

Transcranial Magnetic Stimulation (TMS): A developing innovative management for psychological distress and valuable in managing neurologic situations.⁽³⁹⁾ Based on this single-case investigation, persons with refractory TN may benefit from 10 Hz TMS as adjuvant care. Long-term recurrent exposure to 10 Hz is not linked to major problems. These findings offer promising new perspectives for creating neurological treatments that are customized for every individual to produce durable pain alleviation; however, these initial findings highlight the necessity for additional patient trials to look at the prospective medical advantages of long-term TMS use in TN.

Low-level Laser Therapy (LLLT): Many physicians employ lower-level wavelength laser therapy (LLLT), a type of electromagnetic radiation treatment, to treat an array of illnesses. It is applied by professionals to treat acute as well as persistent pain, by dentists to treat ulcerations and inflamed oral tissues, by dermatologists to treat burns, dermatitis, and oedema, and by rheumatologists to manage autoimmune medical conditions with prolonged inflammation. According to multiple authors, ectopic action potentials in this nerve and a lack of segmental inhibition of the craniofacial nucleus have been attributed to persistent discomfort of the trigeminal nerve. Certain study findings aligned with those of earlier studies. Eckerdal conducted a randomized, placebo-controlled, double-masked trial to examine the beneficial effects of LLLT in treating TN. For five weeks, 16 TN-affected individuals (830 nm, 30 mW) received laser radiation, and they were compared to fourteen patients in the control group. Post-assessment for a year, they concluded that LLLT is a great way to complement traditional TN therapy techniques while still being an effective treatment. Additionally, lasers can eliminate accumulated waste materials while simultaneously enhancing local microcirculation and the oxygen delivered to hypoxic cells in trigger point (TP) regions.⁽³⁰⁾ As for side effects, no occurrences were identified. The outcome derived from both current studies and earlier investigations concluded overall LLLT had been equivalent to a placebo in addition to laser application for activating sites that seemed preferable in brain pathway administration.

Table 1: Treatment of trigeminal neuralgia with medicine⁽⁵⁾

	Drugs	Dose	Side effects
First line drugs	• Carbamazepine	600-1200 mg/day	Headache decreased motor coordination in the muscles, dizziness, kidney pain, and skin reactions.
	• Oxcarbazepine	600-1800 mg/day	Like carbamazepine, however, without fewer serious adverse reactions reduced adverse effects than using carbamazepine
Second line drugs	• Gabapentin	12 mg/day	tiredness, diplopia, vertigo, decreased neuromuscular control, drowsiness, and hand tremor
	• Lamotrigine	400 mg/day	skin responses, nausea, dizziness, constipation, dysgeusia, and psychological distress
	• Baclofen	40-80 mg/day	fatigue, nausea, or muscle hypotonia. Errors or convulsions can occur with abrupt pharmaceutical cessation

CONCLUSION:

We would like to conclude that no single treatment is best for all TN patients. Every technique has benefits, drawbacks, uses, and contraindications. Thus, the course of treatment is solely determined by what the patient responds to. An early diagnosis makes a favourable prognosis for the possible treatment plan. Dentists are essential in diagnosing trigeminal neuralgia and referring patients to neurosurgeons since the condition frequently resembles dental pain. The effects of chronic pain on one's physical and mental well-being can be negative. Professional guidance, comfort, and assistance groups, particularly for individuals with trigeminal neuralgia, may offer insights, advice, and coping mechanisms to alleviate isolation. Additional investigations into the molecular mechanisms beneath trigeminal neuralgia will open pathways for innovative, profitable therapies with fewer invasive therapeutics.

REFERENCES:

- Xu R, Xie ME, Jackson CM. Trigeminal Neuralgia: Current Approaches and Emerging Interventions. *J Pain Res.* 2021 Nov 3; 14:3437-3463.
- Mol Pain. 2020; 16: 1744806920901890. Published online 2020 Jan 26.
- Zeme S (2016) The Surgical Treatment of Trigeminal Neuralgia. *J Pain Relief* 54: 006.
- Srivastava, Rahul & Jyoti, Bhuvan & Shukla, Ashutosh & Priyadarshi, Pankaj. (2015). Diagnostic criteria and management of trigeminal neuralgia: A review. *Asian pacific journal of health sciences*.
- Al-Quliti KW. Update on neuropathic pain treatment for trigeminal neuralgia. The pharmacological and surgical options. *Neurosciences (Riyadh)*. 2015 Apr;20(2):107-14.
- Barker FG 2nd, Jannetta PJ, Bissonette DJ, Larkins MV, Jho HD. The long-term outcome of microvascular decompression for trigeminal neuralgia. *N Engl J Med.* 1996 Apr 25;334(17):1077-83.
- Kondo A. Follow-up results of microvascular decompression in trigeminal neuralgia and hemifacial spasm. *Neurosurgery.* 1997 Jan;40(1):46-51; discussion 51-2.
- McLaughlin MR, Jannetta PJ, Clyde BL, Subach BR, Comey CH, Resnick DK. Microvascular decompression of cranial nerves: lessons learned after 4400 operations. *J Neurosurg.* 1999.
- Mullan S, Lichtor T. Percutaneous micro compression of the trigeminal ganglion for trigeminal neuralgia. *J Neurosurg* 1983; 59: 100712.
- Percutaneous Balloon Compression of Trigeminal Gasserian Ganglion for Idiopathic Trigeminal Neuralgia Kyu Sang Ahn, M.D., Myung Ki Lee, M.D., Sung Hyuck Hwang, M.D., Jae Eun Lee, M.D., Chang Weon Cho, M.D., Dae Jo Kim, M.D. Department of Neurosurgery, Maryknoll General Hospital, Busan, Korea
- Obermann M: Recent advances in understanding/managing trigeminal neuralgia. *F1000Res.* 2019, 8:10.12688/f1000research.16092.1].
- Araya EI, Claudino RF, Piovesan EJ, Chichorro JG: Trigeminal neuralgia: basic and clinical aspects. *Curr Neuropharmacol.* 2020, 18:109-19. 10.2174/1570159X17666191010094350].
- Orhurhu V, Khan F, Quispe RC, et al.: Use of radiofrequency ablation for the management of facial pain: a systematic review. *Pain Physician.* 2020, 23: E559-80.
- Eshraghi Y, Vitug SJ, Guirguis M: Interventional treatment options for trigeminal neuralgia. *Peripheral Nerve Disorders and Treatment.* Turker H, Benavides LG, Gallardo GR, Del Villar MM (ed): IntechOpen, London; 2019 27.
- Håkanson S: Trigeminal neuralgia treated by the injection of glycerol into the trigeminal cistern. *Neurosurgery* 9: 638-646, 1981
- Degn J, Brennum J: Surgical treatment of trigeminal neuralgia. Results from the use of glycerol injection, microvascular decompression, and rhizotomy. *Acta Neurochir (Wien)* 152: 2125-2132, 201020
- Burchiel KJ: Percutaneous retrogasserian glycerol rhizolysis in the management of trigeminal neuralgia. *J Neurosurg* 69: 361-366, 1988
- Blomstedt PC, Bergenheim AT: Technical difficulties and perioperative complications of retrogasserian glycerol rhizotomy for trigeminal neuralgia. *Stereotact Funct Neurosurg* 79: 168-181, 2002
- Rappaport ZH, Gomori JM: Recurrent trigeminal cistern glycerol injections for tic douloureux. *Acta Neurochir (Wien)* 90: 31-34, 1988
- Dieckmann G, Bockermann V, Heyer C, Henning J, Roosen M: Five-and a-half years' experience with percutaneous retrogasserian glycerol rhizotomy in treatment of trigeminal neuralgia. *Appl Neurophysiol* 50: 401-413, 1987
- Kondziolka D. Functional radiosurgery. *Neurosurgery.* 1999 Jan;44(1):12-20; discussion 20-2.
- Kondziolka D, Perez P, Flickinger JC, Habeck M, Lunsford LD Gamma knife radiosurgery for trigeminal neuralgia. *Arch Neurol* 1998; 55: 1524-9
- Young RF, Vermeules SS, Grimm P, Blasko J, Posewitz A. Gamma TGN+pathophysiology, diagnosis, and current treatment 131 knife radiosurgery for treatment of trigeminal neuralgia. Idiopathic and tumor-related. *Neurology* 1997; 48: 608±14
- Lee S, Lee JI. Gamma Knife Radiosurgery for Trigeminal Neuralgia : Review and Update. *J Korean Neurosurg Soc.* 2022 Sep;65(5):633-639
- Ali FM, Prasant M, Pai D, Aher VA, Kar S, Safiya T. Peripheral neurectomies: A treatment option for trigeminal neuralgia in rural practice. *J Neurosci Rural Pract.* 2012 May;3(2):152-7.
- Pradel W, Hlawitschka M, Eckelt U, Herzog R, Koch K. Cryosurgical treatment of genuine trigeminal neuralgia. *Br J Oral Maxillofac Surg.* 2002 Jun;40(3):244-7.
- Thorsen SW, Lumsden SG. Trigeminal neuralgia: sudden and long-term remission with transcutaneous electric nerve stimulation. *J Manipulative Physiol Ther* 1997; 20: 415-19.
- Rubis A, Juodzbals G. The Use of Botulinum Toxin A in the Management of Trigeminal Neuralgia: a Systematic Literature Review. *J Oral Maxillofac Res.* 2020 Jun 30;11(2):e2.
- Pearl C, Moxley B, Perry A, Demian N, Dallaire-Giroux C. Management of Trigeminal Neuralgia with Botulinum Toxin Type A: Report of Two Cases. *Dent J (Basel).* 2022 Nov 3;10(11):207
- Simunovic ZI. Low-level laser therapy with trigger points technique: a clinical study on 243 patients. *J Clin Laser Med Surg.* 1996 Aug;14(4):163-7.
- Chen GQ, Wang XS, Wang L, Zheng JP. Arterial compression of nerve is the primary cause of trigeminal neuralgia. *Neurol Sci.* 2014 Jan;35(1):61-6
- Attassi, M., Seegenschmied, M.H., Körner, M. (2008). Trigeminal Neuralgia. In: Seegenschmied, M.H., Makoski, H.B., Trott, K.R., Brady, L.W. (eds) Radiotherapy for Non-Malignant Disorders. Medical Radiology. Springer, Berlin, Heidelberg.
- Cheuk, A. V., Chin, L. S., Petit, J. H., Herman, J. M., Fang, H. B., & Regine, W. F. (2004). Gamma knife surgery for trigeminal neuralgia: Outcome, imaging, and brainstem correlates. *International Journal of Radiation Oncology Biology Physics*, 60(2), 537-540
- Malik N. Text Book of Oral and Maxillofacial Surgery. New Delhi, India: JP Brother Medical Publishers; 2008. p. 685-97.
- Wilkins RH. Historical overview of surgical techniques for trigeminal neuralgia. *Tech Neurosurg* 1999;5:202-17.
- Breivik, H., Nicholas, M., Campbell, W., & Newton-John, T. (2008). *Clinical Pain Management: Practice and Procedures* (2nd ed.). CRC Press
- Carayannopoulos, A. (2017). *Comprehensive Pain Management in the Rehabilitation Patient. Comprehensive Pain Management in the Rehabilitation Patient.*
- Russell Lane, Paul Davies. "Migraine", CRC Press, 2019
- Mark J. Stillman. "Abstracts and Citations", Headache The Journal of Head and Face Pain, 4/2008
- Mared Attassi. "Trigeminal Neuralgia", Medical Radiology, 2008