Peripheral nerve blocks for headache disorders

Linford Fernandes,1 Marc Randall,1,2 Luis Idrovo1,2

ABSTRACT

Headache is a common neurological referral and a frequent cause for acute hospital admissions. Despite peripheral nerve blocks being widely used in headache and pain services to treat patients with headache disorders, there is no readily accessible resource with instructions for the delivery of peripheral nerve blocks. Here we provide a practical approach for administering peripheral nerve blocks and cover the current evidence base for such procedures in different headache disorders. We provide instructions and an audiovisual guide for administering greater and lesser occipital, supratrochlear, supraorbital and auriculotemporal nerves blocks, and give information on their adverse effects and potential complications. This information will provide a reference for headache practitioners when giving peripheral nerve blocks safely to people with headache.

INTRODUCTION

Headache is one of the most prevalent, disabling and undertreated conditions in neurological clinical practice.1 Headache practitioners commonly administer peripheral nerve blocks to treat various headache disorders both in the acute and outpatient setting, often with rewarding results.2 There is no current national consensus on the technical aspects of delivering peripheral nerve blocks. However, a recent survey among UK headache practitioners showed that blocks have become relatively popular transitional treatments for cluster headache and chronic migraine.3 Here we aim to provide practical instructions for effective and safe delivery of the most common peripheral nerve blocks used in headache medicine. We briefly outline the evidence base for common indications and describe the peripheral nerve block method, including injection location, technique, drug constituents and potential pitfalls. This, together with the supplementary illustrative videos, should provide a comprehensive guide on nerve block delivery. This guide should be used to support, but not to replace, the experience obtained from clinical supervision by practised healthcare professionals.

EVIDENCE BASE FOR PERIPHERAL NERVE BLOCKS

Headache practitioners frequently target the greater occipital nerve (GON), but the treatment of both primary and secondary headaches might target other cervical and cranial nerves. Despite headache specialists seeing positive results in clinical practice, there is little high-quality information to support their widespread use.4 Several recent randomised controlled trials and cohort studies have studied the efficacy of peripheral nerve blocks, particularly GON blocks. The level of evidence for the effectiveness of peripheral nerve blocks for managing different headache disorders varies depending on the pericranial nerve targeted and the outcome measure used (table 1). GON blocks for the acute and preventative treatment of migraine and cluster headaches reduce headache days and give high levels of patient-reported efficacy.5,6 Furthermore, peripheral nerve blocks have been effective in managing acute or prolonged migrainous episodes that commonly present to the emergency department.7,8 The rapid onset of pain relief provided by anaesthetic nerve blocks makes them ideal for acute headache presentations, where timely management is essential, reducing the need for opiate-based therapies.

Although it is difficult to predict which patients will benefit significantly from peripheral nerve blocks, headache practitioners have developed a wealth of clinical experience in maximising their efficacy. In the trigeminal autonomic cephalalgias—predominantly unilateral headaches including cluster headache and hemicrania continua—ipsilateral greater and lesser occipital nerve (LON) blocks as first-line treatments may avoid the need for corticosteroids or indomethacin. Patients with headache who have reproducible pain with palpation over

- Supplemental material is published online only. To view please visit the journal online (http://dx.doi.org/10.1136/practneurol-2020-002612).
- Accepted 26 August 2020
- Published Online First 23 October 2020


© Author(s) (or their employer(s)) 2021. No commercial re-use. See rights and permissions. Published by BMJ.

Check for updates
Peripheral nerve blocks are also demonstrably effective in the older population with headache disorders, whose comorbidities might preclude the use of first-line preventative medications.

There are conflicting results about adding corticosteroid to nerve blocks in people with migraine, but evidence to support its efficacy in cluster headache.

Greater occipital neuralgia with or without another coexisting headache disorder is not uncommon. Giving a GON block to a patient with suspected GON neuralgia can be both diagnostic and therapeutic, usually conferring prolonged relief. The evidence base for using peripheral nerve blocks in other cranial neuralgias, such as auriculotemporal and supraorbital neuralgias, is predominantly anecdotal, from published case series.

Pregnant women with troublesome headaches can often be managed throughout pregnancy and the postpartum period with anaesthetic nerve blocks without corticosteroid, reducing the need for medications. The frequency of the nerve blocks can be tailored to the individual’s response duration but is usually 3 months or more. If the benefit lasts less than 2 months, then clinicians might consider other headache medications or interventions.

Some headache practitioners inject only the occipital nerves during the first session, which reduces the number of injections and allows for an assessment of initial response. If the patient reports some benefit but has residual facial pain, then at a subsequent session, the trigeminal nerves can be blocked as well as the occipital nerves. Consensus recommendations by the American Headache Society and the Spanish Headache Study Group have used this evidence base to provide guidance on the administration of peripheral nerve blocks for different headache disorders.

**GENERAL CONSIDERATIONS**

Knowledge of the anatomical landmarks of the occipital and superficial branches of the trigeminal nerve is important for effective nerve blockade, and to avoid possible complications such as nerve trauma, bleeding or inadvertent arterial injection of anaesthetic drug. People with headache disorders often describe pain over the forehead, behind their eyes, temples, occipital and upper cervical areas. The forehead and upper periorbital areas are innervated by peripheral branches of the first division of the trigeminal nerve (V1), mainly the supraorbital and the supratrochlear nerves. The temples are largely innervated by the auriculotemporal nerve branch from the mandibular division of the trigeminal nerve (V3). The upper cervical and occipital region is innervated by C2/C3 posterior cervical branches, mainly the greater, lesser and third occipital nerves.

Having identified someone as suitable for a peripheral nerve block, we find it helps to show them an illustration of the peripheral cranial nerve to be injected (figure 1). The written consent should include the known complications of any invasive procedure, such as bleeding or infection at the injection site, and some may find the procedure painful. Peripheral nerve blocks are contraindicated at any previous surgical site, for example, previous burr hole or previous haemorrhage.

---

**Table 1** Evidence base for the efficacy of peripheral nerve block in treating different headache disorders

<table>
<thead>
<tr>
<th>Headache disorder</th>
<th>Type of nerve block studied</th>
<th>Evidence level*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Acute migraine</td>
<td>GON</td>
<td>2B</td>
</tr>
<tr>
<td>Chronic migraine</td>
<td>GON</td>
<td>2A</td>
</tr>
<tr>
<td>Cluster headache</td>
<td>GON, suboccipital</td>
<td>1B</td>
</tr>
<tr>
<td>Occipital neuralgia</td>
<td>GON</td>
<td>2B</td>
</tr>
<tr>
<td>Chronic daily headache</td>
<td>GON</td>
<td>2B</td>
</tr>
<tr>
<td>Other trigeminal autonomic cephalalgias</td>
<td></td>
<td></td>
</tr>
<tr>
<td>SUNCT/SUNA</td>
<td>Supraorbital, supratrochlear</td>
<td>4</td>
</tr>
<tr>
<td>Paroxysmal hemicrania/ hemicrania continua</td>
<td>Supraorbital, supratrochlear</td>
<td>4</td>
</tr>
<tr>
<td>Other painful cranial neuralgias</td>
<td>Supraorbital, auriculotemporal</td>
<td>4</td>
</tr>
</tbody>
</table>

*Based on the Oxford Centre for Evidence-based Medicine Levels of Evidence.

GON, greater occipital nerve; SUNCT, short-lasting unilateral neuralgiform headache attacks with conjunctival injection and tearing; SUNA, short-lasting unilateral neuralgiform headache attacks with cranial autonomic features.

---

**Figure 1** Illustrative drawings of the anatomical course of the peripheral cranial nerves. These are used during the consent process to demonstrate the location of the nerves to be injected. Adapted with permission from Blumenfeld et al.
How to do it

**GON BLOCKS**

The GON arises as the medial branch from the dorsal primary ramus of the second cervical nerve. It emerges below the obliquus capitis inferior muscle and passes through the semispinalis muscle, before ascending to innervate the posterior scalp to the vertex. The GON can be localised superficially by identifying a point one-third (medially) of the way between the occipital protuberance (inion) and the mastoid process, approximately 2 cm lateral and 1.5–2.0 cm below the inion (figure 2). The occipital artery usually runs lateral to the GON, and this should be considered when injecting the nerve.

**Injection technique**

The patient should be comfortably seated on a chair with the head slightly flexed, and the clinician standing behind. Locate the GON, and if needed use topical anaesthetic cream before the injection. Use a 5 mL syringe with a 25-gauge needle. Gently insert the needle perpendicular to the skin, until meeting firm resistance, indicating the needle tip is at the periosteum. This helps to ensure there is no skull defect that might precipitate intracerebral infiltration of the anaesthetic. Withdraw the needle slightly and aspirate to confirm no arterial bleed. Then, redirect the needle slightly superiorly and gently inject the solution in a fanlike distribution (figure 3). Some practitioners inject the solution with the needle in the same position, which is sufficient if injecting a reasonable volume of anaesthetic. The patient may feel a burning sensation as the anaesthetic is infiltrated, but this should subside in a few minutes once the anaesthetic takes effect. Withdraw the needle and apply pressure to the site with gauze to minimise bleeding (online supplemental file 1). If the patient has had previous vasovagal episodes due to pain or previous injections, we recommend performing the injection with the patient in a lateral decubitus position, which will help avoid a sudden fall in blood pressure.

---

**Table 2 Constituents and volumes for individual nerve blocks**

<table>
<thead>
<tr>
<th>Nerve</th>
<th>Constituents per injection (volume injected, mL)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Greater occipital</td>
<td>Methylprednisolone 40 mg/mL (2 mL) Lidocaine 2% (1 mL) Bupivacaine 0.5% (1 mL) Total injection volume=4 mL</td>
</tr>
<tr>
<td>Lesser occipital</td>
<td>Lidocaine 2% (1 mL) Bupivacaine 0.5% (1 mL) Total injection volume=2 mL</td>
</tr>
<tr>
<td>Auriculotemporal</td>
<td>Supraorbital Lidocaine 2% (0.5 mL) Bupivacaine 0.5% (0.5 mL) Total injection volume=1 mL</td>
</tr>
<tr>
<td>Supratrochlear</td>
<td></td>
</tr>
</tbody>
</table>

* Methylprednisolone acetate maximum dose 160 mg per session.
† Lidocaine maximum dose: 4.5 mg/kg, not to exceed 300 mg per session (without vasoconstrictor).
‡ Bupivacaine maximum dose: 2.5 mg/kg, not to exceed 175 mg per session (without vasoconstrictor).
and hence a sudden decrease in cerebral perfusion pressure.

**LON BLOCKS**

The LON arises from the ventral primary rami of the second and third cervical nerves. It passes superiorly along the posterior border of the sternocleidomastoid muscle, dividing into cutaneous branches that innervate the lateral portion of the posterior scalp. The LON is localised by identifying a point two-thirds of the way between the inion and the mastoid process (figure 2). It is commonly injected along with the GON.

**Injection technique**

The procedure is broadly similar to the GON injection with the patient seated and clinician standing behind. Having located the LON, use a suitable 25-gauge needle, entering perpendicular to the skin, stopping once the periosteum is reached. After gentle aspiration to ensure no arterial entry, the injection is delivered (figure 3) (online supplemental file 2).

**SUPRATROCHLEAR NERVE BLOCKS**

The supratrochlear nerve is one of the terminal cutaneous branches of the frontal nerve, which in turn arises from the ophthalmic division of the trigeminal nerve (V1). The supratrochlear nerve exits the orbital cavity anteriorly and ascends the forehead to innervate the upper eyelid, forehead and anterior scalp. It is located superficially at the superomedial aspect of the supraorbital ridge, which is the injection site (figure 2).

**Injection technique**

Position the patient supine with their head in a neutral position. From here, the clinician, standing beside the patient, has easy access to the supratrochlear nerve. Use a 1.0 or 2.5 mL syringe with a 30-gauge needle. Locate the nasal bridge and the medial aspect of the supraorbital ridge. Gently insert the needle at the medial aspect of the corrugator muscle, just lateral to the procerus and above the eyebrow line to a depth of 4–5 mm. Gently aspirate to ensure no arterial flashback and then inject the solution, which will produce a small weal under the skin (figure 3) (online supplemental file 3).
SUPRAORBITAL NERVE BLOCKS
The supraorbital nerve is the larger of the terminal cutaneous branches of the frontal nerve and runs through the supraorbital notch to innervate the upper eyelid and conjunctiva. It then ascends the forehead, being closely associated medially with the supraorbital artery. The supraorbital nerve is located just above the supraorbital notch (figure 2).

Injection technique
With the patient supine and their head in a neutral position, palpate the supraorbital notch. Use a 1.0 or a 2.5 mL syringe with a 30-gauge needle. Insert the needle perpendicularly, just above supraorbital notch (avoid injecting into the supraorbital notch), to a depth of 4–5 mm. Gently aspirate to confirm no arterial entry and then inject the solution (figure 3) (online supplemental file 4).

AURICULOTEMPORAL NERVE BLOCKS
The auriculotemporal nerve arises as a posterior division of the mandibular branch of the trigeminal nerve. It innervates the temples and the temporomandibular joint. Its superficial branches innervate the tragus and the auricle of the ear; its proximal trunk is located superficially just anterior to the tragus (figure 2).

Injection technique
The positioning for this nerve injection can be with the patient seated and physician standing beside them, or with the patient supine and their head in a neutral position. At the point just anterior to the tragus, use a 5 mL syringe with a 30-gauge needle to infiltrate 1–2 mL into the subcutaneous tissue to a depth of about 4–6 mm. After gentle aspiration to exclude any arterial flashback, inject the solution (figure 3) (online supplemental file 5).

CASE STUDIES
Case 1
A 26-year-old woman was referred to the headache clinic when 4-weeks pregnant. She had a lifelong history of episodic migraine headaches with and without aura, which had transformed into daily headaches in the previous 3 weeks. She also described shooting pains over the right occipital region. The headaches were predominantly right sided associated with alloodynia over the right forehead and the right occipital region was tender. Her headaches had previously responded poorly to propranolol and she had stopped her current prophylactic, amitriptyline, on finding she was pregnant. Her Headache Impact Test (HIT6) score was 72 on presentation. The HIT6 score is a headache score between 36 and 78, which measures headache burden, with higher scores demonstrating a worse outcome.

After discussion, she opted for peripheral nerve blocks as a transitional migraine treatment. She underwent bilateral GON blocks with anaesthetic only, and right-sided supraorbital, supratrochlear and auriculotemporal blocks. There was sustained improvement in her migraine for 7 weeks, with an HIT6 score of 48 after 4 weeks. She continued the same combination of peripheral nerve blocks every 3 months throughout her pregnancy.

Case 2
A 40-year-old man was referred to the nerve block clinic with a 5-year history of seasonal stereotyped left hemicranial stabbing pains lasting about 2 hours, with occasional neuralgia affecting the left trigeminal nerve maxillary division (V2). There were associated left trigeminal autonomic symptoms including conjunctival injection, eye redness, lacrimation and nasal congestion. He was restless during these episodes, which occurred up to three times a day during the winter months, over a few weeks. Oxygen treatment and triptans had previously helped. He also had a history of left temporomandibular joint pain and previous episodic migraine. He had tried topiramate, verapamil and propranolol but had stopped this due to symptomatic bradycardia. Investigations were normal, including intracranial imaging looking for secondary causes. On examination, there was tenderness of the left greater and LON area, as well as left temporomandibular joint clicking and tenderness. We established the diagnoses of cluster headache and co-existing left temporomandibular joint dysfunction. The baseline HIT6 score was 78.

Due to his frequent cluster attacks and temporomandibular joint pain, we offered a left GON block with corticosteroid, and left lesser occipital, supraorbital, supratrochlear and auriculotemporal nerve blocks. On follow-up at 12 weeks, he reported a favourable response with only three further cluster attacks that had responded to sumatriptan injection rescue therapy.

CONCLUSION
Peripheral nerve blocks are effective in the acute and preventative management of several headache disorders. It is difficult to identify those who will respond best, but the procedures allow an interventional approach for those with troublesome and refractory headache. Neurologists can administer these blocks as a day procedure, in clinic or the emergency department, where quick pain relief can provide a satisfactory outcome.

How to do it
**How to do it**

**Key points**

- Peripheral nerve blocks have a role in acute and transitional treatment of acute migraine, chronic migraine, cluster headache and painful cranial neuralgias.
- Patient position and anatomical landmarks are key for their successful delivery.
- Corticosteroids are frequently used for greater occipital nerve blocks but may also be used for lesser occipital nerve blocks.
- Supraorbital, supratrochlear and auriculotemporal nerve blocks involve a combination of lidocaine and/or bupivacaine.
- Uncommon but important adverse effects include transient dizziness, light-headedness, transient headache exacerbation, and rarely localised lipoatrophy and alopecia with corticosteroids.

**FURTHER READING**


**Contributors** LF drafted the manuscript, edited and narrated the illustrative videos, and revised the manuscript for intellectual content. LI produced the illustrative videos, edited and revised the manuscript for intellectual content.

**Funding** The authors have not declared a specific grant for this research from any funding agency in the public, commercial or not-for-profit sectors.

**Competing interests** None declared.

**Patient consent for publication** Consent for patient illustration and supplemental videos has been obtained.

**Provenance and peer review** Commissioned. Externally peer reviewed by Nick Silver, Liverpool, UK.

**Data availability statement** All the content, figures, tables and videos in the manuscript are available to all the authors.

**Supplemental material** This content has been supplied by the author(s). It has not been vetted by BMJ Publishing Group Limited (BMJ) and may not have been peer-reviewed. Any opinions or recommendations discussed are solely those of the author(s) and are not endorsed by BMJ. BMJ disclaims all liability and responsibility arising from any reliance placed on the content. Where the content includes any translated material, BMJ does not warrant the accuracy and reliability of the translations (including but not limited to local regulations, clinical guidelines, terminology, drug names and drug dosages), and is not responsible for any error and/or omissions arising from translation or adaptation or otherwise.

**ORCID iDs**

Linford Fernandes http://orcid.org/0000-0002-1575-8776
Marc Randall http://orcid.org/0000-0002-3196-182X
Luis Idrovo http://orcid.org/0000-0003-2599-485X

**REFERENCES**