

Consultation

When consulting a patient, it is important to ensure you conduct a proper medical history. To do this thoroughly, there are a few important issues that need to be considered. Firstly, find out whether the patient in question has received a formal diagnosis of chronic migraine, by whom and when. Then, you should ask what previous treatments the patient has had – if any. We also often check which prophylactic medications they have had, their doses and duration of trial and side effects. It is important to look for drugs which might precipitate migraine such as oestrogen and nitrates as these might just need to be stopped to solve the problem.⁸ Liaison with the patient's GP at this point is important to ascertain the patient's main migraine triggers. For example, common migraine triggers include stress, dehydration, caffeine, alcohol, too much or too little sleep, or foods like chocolate, citrus fruits and cured meats, are all of which are implicated, as are hormonal changes.⁹ Because triggers are personal to each patient, it is essential for a diary to be kept as this provides vital clues which aid management. Even if you treat a patient with botulinum toxin, identifying triggers and reducing them will help to further reduce attacks. Finally, ensure to check for 'red flags' which might indicate that the problem is something else, especially if your patient has not been to a specialist prior to seeing you.¹⁰ For example, headaches that occur in patients after the age of 50 and have not been present before. Most migraine sufferers have had headaches since puberty or even before and follow a certain pattern which might change slowly over time. Worsening symptoms are also a sign that there may be something more sinister. In these cases migraines can be mistaken for thunderclap headaches, which are sudden onset severe headaches that raise suspicion of sub-arachnoid bleeds. Similarly, symptoms of temporal arteritis can appear alike to a migraine with sometimes an unilateral temporal tenderness, headache and visual disturbance. However, this generally occurs in over 50s and the headache pattern will be different to a patient's migraine or they may have no pre-existing migraine. Glaucoma symptoms which are predominantly eye pain and redness of the eye may also mimic a migraine, and any additional neurological symptoms including meningism are concerning.¹¹

Treatment approach

In 2009, pharmaceutical company Allergan conducted the seminal research into the use of botulinum toxin and migraines. The PREEMPT trials showed after 12 months of treatment of Botox (onabotulinum toxin A) at three monthly intervals, 70% of patients showed a greater than 50% reduction of headaches with minimal side effects.¹² The PREEMPT protocol is 31 injections in the head and neck, with a total of 155 units of Botox used. There is the option of a further 40 units to be administered in the occipitalis, temporalis and trapezius areas if required. Treatment protocols are generally based upon the PREEMPT trials which were methodologically robust.¹³ There are criticisms of the PREEMPT trials mainly concerning the marked placebo effect in both PREEMPT trials and the fact that most patients enrolled were overusing acute migraine medications.¹¹ A survey by Begasse de Dhaem *et al.* suggested that about 70% of clinicians varied the dose, sites and frequency of the toxin injections. Therefore, it would be interesting to explore the reasons behind this and develop new guidance as a result.¹⁴ There is a small sub-group of patients (around 10%) who fail to respond to treatment first time around, but research has shown that they do respond after two cycles of treatment at three months.¹⁵ Therefore, it is worth trying three cycles before considering ceasing treatment altogether. Dose should be at least 155 units of Botox, as earlier trials showed minimal difference from placebo below this dose.⁶ It is often worth treating trigger points and increasing the dose up to 195 units.¹¹ Side effects of this treatment are fairly minimal. Brow ptosis can occur, but this is less likely in aesthetic practices as clinicians should be aware of this complication and may adjust the distribution of the injections as a result. Anecdotally, we have not found an issue with altering the distribution of injections in this site. There have been trials comparing prophylactic migraine medications to Botox and found them to be comparable, but the studies are small.^{16,17} Other toxins such as Bocouture and Azzalure can be used off label, but different units will need to be used and adjusted, for example where 1U Botox/ Bocouture is equal to 2.5U Azzalure.¹⁸

Summary

Botox is licenced for use in chronic migraine. The key elements of optimal treatment are ensuring appropriate patient selection, correct technique, identifying and minimising triggers, good liaison with

the patient's GP and correct aftercare. It must be stressed that not all chronic migraineurs respond to Botox, so patient expectations need to be realistic and will need exploration at the initial consultation. Overall, treatment with Botox appears a safe method with long term tolerability.



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REFERENCES

- Steiner TJ *et al.*, 2003, Cephalgia: The International Journal of Headaches, The prevalence and disability burden of adult migraine in England and their relationships to age, gender and ethnicity
- John Hopkins Medicine, Migraine Headaches, <<https://www.hopkinsmedicine.org/health/conditions-and-diseases/headache/migraine-headaches>>
- National Institute for Health and Care Excellence, <<https://www.nice.org.uk/guidance/ta260/documents/migraine-chronic-botulinum-toxin-type-a-final-scope>>
- Dodick DW, 2018, A phase-by-phase review of migraine pathophysiology. Headache 58(Suppl 1):4–16
- Burstein R, Zhang X, Levy D, Aoki KR, Brin MF, 2014, Selective inhibition of meningeal nociceptors by botulinum neurotoxin type A: therapeutic implications for migraine and other pains
- National Institute for Health and Care Excellence, <<https://www.nice.org.uk/guidance/ta260/documents/migraine-chronic-botulinum-toxin-type-a-final-appraisal-determination3>>
- Ziegeler C, Brauns G, Jürgens T *et al.*, 2019, The Journal of Headache and Pain, Shortcomings and missed potentials in the management of migraine patients - experiences from a specialized tertiary care centre
- National Migraine Centre, Migraine and Headache Facts, <<https://www.nationalmigrainecentre.org.uk/migraine-and-headaches/migraine-and-headache-facts>>
- NHS, Migraine, <<https://www.nhs.uk/conditions/migraine/causes/>>
- Dr Suneeta Kochhar, 2018, <<https://www.gponline.com/red-flag-symptoms-headaches/neurology/neurology/article/1332134>>
- Gooriah R, Ahmed F, 2015, OnabotulinumtoxinA for chronic migraine: a critical appraisal. *Ther Clin Risk Manag.* 2015;11:1003-1013
- Dodick DW, Turkel CC, DeGryse RE *et al.*, 2010, Headache, OnabotulinumtoxinA for treatment of chronic migraine: pooled results from the double-blind, randomized, placebo-controlled phases of the PREEMPT clinical program
- Dodick DW, 2018, Headache, A phase-by-phase review of migraine pathophysiology.
- Begasse de Dhaem O *et al.* Headache, Modifications to the PREEMPT Protocol for OnabotulinumtoxinA Injections for Chronic Migraine in Clinical Practice
- Aurora SK, Winner P, Freeman MC *et al.* 2011, Headache, OnabotulinumtoxinA for treatment of chronic migraine: pooled analyses of the 56-week PREEMPT clinical program
- Cady RK, Schreiber CP, Porter JA, Blumenfeld AM, Farmer KU, 2019, Headache, A multi-center double-blind pilot comparison of onabotulinumtoxinA and topiramate for the prophylactic treatment of chronic migraine
- Magalhães E, Menezes C, Cardeal M, Melo A, 2010, Clinical Neurology and Neurosurgery, Botulinum toxin type A versus amitriptyline for the treatment of chronic daily migraine.
- C Blatchley, Treating Migraines, 2016, <<https://aestheticsjournal.com/feature/treating-migraines?authed>>