All headaches are not equal: A review of migraine as a state of brain dysfunction

De Wet Wolmarans,*1 Sarel J Brand,2 Linda Brand1

1 Centre of Excellence for Pharmaceutical Sciences, Division of Pharmacology, Faculty of Health Sciences, North West-University, Potchefstroom, South Africa
2 Unit for Environmental Sciences and Management, North-West University, Potchefstroom, South Africa

*Corresponding author, email: dewet.wolmarans@nwu.ac.za

Introduction and background

Individuals suffering from migraine headache, or migraineurs as they are often referred to, and the clinicians treating them are well-informed of the excruciating and debilitating nature of such episodes. However, the theoretical definition of migraine, i.e. a primary, recurrent and mostly unilateral headache characterised by throbbing pain,1 also seems to describe the presentation of often non-migraine headaches, thus somewhat failing to encompass the actual incapacitating nature of the condition. Importantly, migraine is diagnosed according to criteria stipulated by the International Headache Society (IHS)1 if patients experience 5 or more attacks of unilateral, pulsating, moderate to severe pain that last 4–72 hours, 2) if such attacks are associated with nausea and/or photophobia and, 3) if they are aggravated by, or prevent patients from engaging in normal routine physical activity.1 Interestingly, migraine is regarded as episodic or acute if it occurs less than 15 times per month, while chronic migraine is defined as more than 15 episodes per month continuing for more than 3 consecutive months.2 Considering that a single attack can last up to 72 hours, 1 and that clinicians often design therapeutic intervention based on accurate diagnoses, this classification seems somewhat counterintuitive. Indeed, patients will often regard a single migraine attack as one too many, with 15 migraine episodes per month essentially equating to permanent incapacity.1,4

Migraine as a state of brain dysfunction and neuronal hypersensitivity

Briefly, migraine results from aberrant activity in the trigeminovascular system in the meninges and can be divided into four phases that often, but not always, transpire in sequence.9

The premonitory phase – starting as early as 3 days before the actual pain10 – involves the sensitization of meningeal nociceptors and results from a complex interplay between homeostatic changes and hypothalamic hyperresponsiveness.9 During this phase, patients can already present with fatigue, food cravings, mood swings, yawning, myalgia, photophobia, nausea and rhinorrhoea.11 These symptoms tend to point to altered autonomic nervous system functioning and it is believed that changes in sympathetic and parasympathetic outflow during states of increased physiological or psychological stress may contribute to triggering peripheral pro-nociceptive signalling in the trigeminovascular neurons.12 That said, the subsequent cortical interpretation of such nociceptive signals originating from sensitised trigeminal neurons is modulated by the hypothalamic release of inhibitory or excitatory neurotransmitters.7 If this balance shifts towards being more excitatory, it will induce tonic firing of the trigeminovascular neurons resulting in pain transmission, with the opposite being true if the transmitters released are predominantly inhibitory.13 As the balance between excitatory and inhibitory neurotransmitter release is dependent on the brain state at the time of trigeminal sensitization, this phenomenon may explain why identical migraine triggers will not always result in headache. In summary, nociceptive trigeminovascular pathways are already sensitised during the premonitory phase following changes in the homeostatic balance, which paves the way for the propagation of nociceptive signalling.9

Second, nearly one third of migraineurs will experience sensory and neurological symptoms collectively referred to as migraine aura.14 While most patients who experience aura symptoms only report visual disturbances, transient motor and speech impairment as well as cognitive disability can also be present.15 Importantly, migraine aura is associated with a phenomenon...
broadly referred to as cortical spreading depression (CSD) that describes a slowly propagating wave of depolarization in neuronal cells that originates in the visual cortex and results in cortical inhibition for up to 30 minutes. Moreover, as the depolarizing wave proceeds, it is associated by sudden and massive changes in glutamate release as well as Na⁺ and Ca²⁺ flux across neuronal membranes that is accompanied first by hyperemia—or excessive blood supply—followed by a prolonged phase of oligemia. This is clinically important as central perfusion rates of < 20ml/100g/min is associated with ischemic neuronal cell death, which may pose a significant risk in patients already predisposed to ischemic stroke. In the case of stroke however, the onset of negative symptoms, e.g. numbness, often transpires suddenly as opposed to the slow progression of positive, e.g. visual disturbances, and then negative symptoms observed in migraine. Recently, it has been proposed that migraine with aura is founded in a distinct pathophysiological construct based on the differences in response of migraine with and without aura to oral sumatriptan intervention. Further, female patients presenting with aura migraine present with a twofold increased risk for ischemic stroke compared to non-aura migraineurs, pointing to a unique association between migraine with aura and aberrant vascular responses. Nevertheless, as migraine episodes will also occur in the majority of patients in the absence of an aura, the presence of an aura is neither necessary nor sufficient to trigger headache. While the pathophysiological role of aura in migraine episodes is still being investigated, clinicians must be attentive of the associated risks that this phenotype of migraine might possibly predict.

The third phase of a migraine episode involves a severe and throbbing headache that results from activation of prior sensitised trigeminovascular neurons. Briefly, the trigeminovascular pathway relays painful stimuli from the meninges to the cortex. Nociceptive neurons from the peripheral trigeminal ganglion converge with sensory neurons from surrounding skin and pericranial muscle, which then relays pain impulses via an already sensitised trigeminovascular system to the periorbital, occipital and cervical neck regions, accounting for descriptions of pain around the eyes and the back of the head and neck, respectively. Activation of the prior sensitised trigeminovascular pathway begins when peripheral nociceptors that innervate the dura mater are stimulated by—at the specific time—contextually relevant triggers, e.g. bright light, dietary constituents, water deprivation, and loud noise, among others. Such stimulation of peripheral nociceptors elicits the release of vasoactive peptides, e.g. calcitonin gene-related peptide (CGRP) and pituitary adenylate cyclase-activating polypeptide (PACAP) and other signalling molecules, e.g. nitric oxide (NO) and glutamate, which in turn sets in motion excessive pain signalling via the trigeminovascular neuronal pathway. At this stage of the migraine episode it is believed that nociceptive propagation is bolstered and extended by the release of CGRP and inflammatory mediators released from mast cells. Subsequent to this, sensitization and activation of central trigeminovascular neurons transpire over 30–60 minutes post peripheral nerve activation, reaching a maximum intensity after 2 hours. Central trigeminovascular activation is associated with severe cephalic muscle tenderness and sensitivity to touch.

Last, as the brain homeostasis gradually returns to normal, resolution and pain relief is often accompanied by asthenia, i.e. loss of energy, somnolence, and concentration difficulty. Importantly, a symptomatic resolution phase is more characteristic of aura migraine than of non-aura migraine, and normally demonstrates a positive correlation with pain severity.

**Four phases, multiple inputs**

Considering the four phases of migraine alluded to above, viz: sensitization and premonition, CSD and aura, headache, and resolution, the question can be raised as to which neurobiological factors may play a role in the initiation and propagation of nociceptive stimuli, as these may be current or future targets for therapeutic intervention. The answer to this question is complex and as such, only a brief overview will be provided.

During the last few decades, several significant advances have been made in our understanding of migraine. Historically, migraine was thought to be the result of dilating extracranial arteries and intracranial blood vessels that subsequently activated perivascular stretch receptors. In this regard for example, the 5HT₁₈ receptor agonists, i.e. ‘triptans’ have been developed as vasoconstrictors for the treatment of migraine. Today, however, we know that the triptans also act by means of their 5HT₁₂ and recently discovered 5HT₁₆ receptor targets to modulate trigeminovascular neurons, while also reducing the release of CGRP. In this section we will provide a brief overview of the major targets of investigation in current migraine research.

**Neuropeptides**

Since the discovery of CGRP and PACAP our understanding of the initiation and propagation of migraine pain has significantly broadened. In fact, the identification of these molecules, for the first time, sheds light on how triggers of migraine ultimately sensitise the trigeminal nociceptive pathway. This is important and valuable as we now have a better understanding of migraine as state of altered brain activity and nociceptive signalling, rather than being a condition of rapid vascular change.

Upon presentation with, or experience of a specific contextually relevant trigger, e.g. alcohol, caffeine, elevated oestrogen concentration, inflammation, or excessive stress, biological targets such as the transient receptor potential cation channel subfamilies A (TRPA1) and V (TRPV1) located in peripheral nociceptors are believed to be activated. Subsequently, CGRP is released to facilitate the propagation and intensity of trigeminovascular nociceptive transmission. In fact, CGRP is not only a potent vasodilator with proposed anti-inflammatory action, but it is also located in the afferents that innervate meningeal blood vessels. Moreover, CGRP has been shown to bolster glutamatergic neurotransmission and has been demonstrated to trigger migraine episodes in pain-free migraineurs, but not in healthy volunteers. This is significant as excessive glutamatergic signalling is associated with CSD, which is believed to underlie aura migraine. In addition, CGRP
also facilitates substance P release which may contribute to the hyperemia characteristic of the aura phase in migraine and intensifies signal transmission in the trigeminal ganglion, thereby contributing to the peripheral sensitization of trigeminal neurons.31

The other neuropeptide that needs mentioning here is PACAP which, although demonstrating fluctuating levels both within and between migraineurs,32 has been demonstrated to modulate circadian rhythm, a physiological system closely related to migraine in some individuals.33 Indeed, it has been shown that inhibiting PACAP signalling via its PAC, receptor has marked therapeutic potential in the treatment of migraine.27

Currently, investigations into the role neuropeptides play in processes of pain and central nervous system functioning are at the forefront of research. As relatively little is known about these molecules, the pathophysiological roles of neuropeptides, such as CGRP, PACAP and several others, e.g. orexins, oxytocin and neuropeptide Y, are continuously being studied as possible targets of intervention in the treatment of both migraine and other relevant conditions.

**Neurotransmitters**

Although a number of neurotransmitters, including noradrenalin and dopamine, have been implicated in the pathophysiology of migraine, emphasis will be placed on serotonin as this is the only neurotransmitter currently targeted by therapeutic intervention strategies. While glutamate also plays a major role in the pathophysiology of CSD and migraine, the effects of glutamate modulators, e.g. ketamine, are still being investigated.34 However, ketamine has been used as an abortive agent especially in migraine with aura.35 As indicated in Table 2, the anticonvulsants used as preventive treatment may also decrease glutamate neurotransmission.36

Since the development of the ‘triptan’ class of drugs, the involvement of serotonin in migraine has been believed to be restricted to its effect on extracranial and cranial vasoactive responses.37 Lately, however, the serotonergic system, originating in the raphe nuclei and innervating inter alia the basal ganglia, thalamus, cortex and the trigeminal ganglion, has been found to be intrinsically involved in the neural modulation of nociceptive signalling, independent of its actions on blood vessel responses. Initially migraine was proposed to be a condition characterised by low serotonergic activity with sudden increases during an episode. This hypothesis was based on findings that 5HT1B receptor antagonists, e.g. pizotifen and methysergide, are valuable prophylactics, while 5HT1D agonists, which function via auto-receptor induced reductions in serotonin release, are useful during the acute pain phase of migraine.38,39

Currently, though, our understanding of the role of serotonin in migraine involves a broader perspective in that it is now known that serotonin, depending on its concentration and the location of its specific receptors, is involved in the modulation of nociceptive pathways following trigeminal sensitization and that it can both inhibit or facilitate pain.40 In the central nervous system, serotonin mainly has an analgesic role as it inhibits nociceptive signal propagation from the peripheral to the central nociceptors.37 Therefore, low levels of central serotonin may disinhibit such nociceptive signals, resulting in excessive pain signalling. Serotonin also plays a significant and unique role in trigeminally sensitised migraineurs vs. non-sensitised controls in that low serotonin levels contribute to the hypersensitization often observed in the premonitory and aura phases of migraine episodes. Subsequently, sudden increases in serotonin release during the acute pain phase, contribute not to analgesia, but to pain via binding on its proalgesic 5HT2A receptors.41

In terms of the response of migraine pain to 5HT1B,D agonists, e.g. sumatriptan, several mechanisms have been proposed. First, sumatriptan induces more central nervous system adverse effects compared to placebo intervention, which somewhat undermines this argument.37 Whether vasoconstriction in fact plays any role in its analgesic effects remains highly debatable. A more plausible explanation for the therapeutic benefit of 5HT1B,D agonists in migraine is its modulation of nociceptive signalling via its actions on 5HT1F receptors, independent of its actions on vasoactive processes. In fact, sumatriptan, but not aspirin, has been shown to suppress signalling between the trigeminal ganglion and the cortex which abrogates the cortical interpretation of pain, while selective 5HT1F receptor agonists, e.g. lasmidiptan, that exert these effects in the absence of any vasoactive action, are currently being developed.37

**Migraine treatment: From bench to evidence-based bedside**

Migraine is treated with respect to acute and prophylactic intervention, while successful treatment of an acute migraine attack is defined by the following criteria:42

- The patient must be pain-free after 2 hours;
- The headache must improve from moderate to severe to mild or none in 2 hours;
- Efficacy must be consistent in two out of every three attacks; and
- Headache should not be recurrent and no further drug intake within 24 hours after successful treatment should be necessary.

Drugs for acute episodes (Table 1) can be divided in non-specific treatments such as paracetamol, non-steroidal anti-inflammatory drugs (including aspirin), opioids, and combinations of analgesics. These (except the opioids) usually constitute the first-line options for the treatment of mild to moderate migraine attacks. The second class of drugs indicated for acute intervention is the migraine-specific drugs (ergotamine tartrate and the triptans); these have historically been used only in ‘severe’ attacks, but given the intense and excruciating nature of most migraine episodes, these compounds enjoy significant favor among clinicians.43 With regard to the latter, triptans are preferred as first choice drugs and although ergotamine is very effective and demonstrates low relapse rates, its poor tolerability and relative risk of overdose headache (see Fact Box 1) rendered it to be preferred in a small group of migraineurs only, presenting with infrequent headaches of a long duration and complying
with dosing restrictions. All in all, a stratified care approach, where drugs are selected according to the degree of disability, is preferred to a step care approach, where treatment is initialised with safe and low-cost drugs.

The general principles for the prophylactic or preventive treatment of migraine have been summarised in Table 2. Drugs from diverse pharmacological classes are effective, although current evidence and clinical experience favor the beta-adrenoceptor antagonists, valproic acid, topiramate, and amitriptyline as first-line drugs. In fact, evidence that supports the use of other drugs, e.g. the calcium channel blockers, i.e. verapamil and flunarizine, is not as robust. Although multiple mechanisms of action may be involved in the prevention of migraine, it is suggested that all of the indicated prophylactics either inhibit cortical excitation or restore nociceptive dysmodulation. The anticonvulsants (and calcium channel blockers) reduce neuronal hyperexcitability in which glutamate dysfunction may play an important role. Further, as alluded to earlier, descending nociceptive inhibition may be restored by drugs with actions on serotoninergic 5HT1B,1D receptors.

While most studies and clinical trials do not differentiate between the management of episodic migraine with aura (classic migraine) and migraine without aura (common migraine), there are indications that the efficacy of drugs may vary according to the presence of an aura, although this is an inadequately studied topic. In addition to the differential effects of sumatriptan in migraineurs presenting with and without aura (see Section 2), it has also been found that in classic migraine, lamotrigine, daily aspirin and flunarizine may generally be effective prophylactic agents, and magnesium, furosemide and ketamine effective acute interventions.

Table 1: Treatment indicated for acute intervention in migraine episodes

<table>
<thead>
<tr>
<th>Active ingredient and trade name where applicable</th>
<th>Dosage interval (if repeated) and maximum daily dose</th>
<th>Pharmacokinetics</th>
<th>Mechanism of action and general comments</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Analgesics and Non-steroidal anti-inflammatory drugs</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Paracetamol Panado® and others</td>
<td>1g every 4 hours, max: 4g</td>
<td></td>
<td>MOA: Inhibits PG synthesis and inhibit neurogenically mediated inflammation in the trigeminovascular system; First-line treatment for mild to moderate and even some severe attacks; A combination of aspirin/paracetamol/caffeine is effective with a high speed of onset; All NSAIDs may be effective in treating migraine with/without aura and the failure of one does not exclude the use of another.</td>
</tr>
<tr>
<td>Aspirin Disprin®, Ecotrin, Dr. du Toit’s pain expeller, Myoprin and others</td>
<td>975 – 1000 mg every 4 – 6 hours, max 5.4 g (varies depending on indication)</td>
<td>T&lt;sub&gt;max&lt;/sub&gt; 0.5 – 1 hour, Elimination t&lt;sub&gt;½&lt;/sub&gt; 2 hours</td>
<td></td>
</tr>
<tr>
<td>Ibuprofen Brufen®, Nurofen® and others</td>
<td>400 mg every 4 hours, max 2400 mg</td>
<td>T&lt;sub&gt;max&lt;/sub&gt; 1 – 2 hours, Elimination t&lt;sub&gt;½&lt;/sub&gt; salicylate active metabolite 5 – 6 hours</td>
<td></td>
</tr>
<tr>
<td>Diclofenac Voltaren®, Diclofenac Biotech, Mylan Diclofenac and others Diclofenac powder Cataflam® and Catafast®</td>
<td>50 mg 3 – 4 times per day, max 150 mg, 50 mg single dose</td>
<td>T&lt;sub&gt;max&lt;/sub&gt; &lt; 1 hour, Elimination t&lt;sub&gt;½&lt;/sub&gt; 2 hours</td>
<td></td>
</tr>
<tr>
<td>Naproxen Synflex®, Aleve®, Nafasol® and others</td>
<td>500 mg orally/ rectally (although not clinically useful as monotherapy) 500 – 550 mg twice a day, max 1375 mg</td>
<td>T&lt;sub&gt;max&lt;/sub&gt; 2 hours Elimination t&lt;sub&gt;½&lt;/sub&gt; 14 hours</td>
<td></td>
</tr>
<tr>
<td><strong>Ergot alkaloids: Ergotamine tartrate tablets</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ergotamine tartrate 1 mg / Caffeine 100 mg Cafergot®</td>
<td>Oral, 2 mg at onset, may repeat twice in 30 min intervals, up to 6 mg in 24 hours. Total dose in any 7 days should not exceed 10 mg</td>
<td>Elimination t&lt;sub&gt;½&lt;/sub&gt; 2 hours, duration of action much longer. Caffeine increases absorption and increases plasma peak levels</td>
<td>MOA: Agonist on 5-HT&lt;sub&gt;1D&lt;/sub&gt; receptors; causes vasoconstriction of intracranial vessels and blocks the trigeminovascular pathway centrally. Contraindicated in vascular diseases, sepsis, uncontrolled hypertension, hepatic/renal dysfunction, porphyria, pregnancy. Nausea and vomiting in ≥ 10% patients, weakness, numbness and tingling of extremities, angina pain, brady/ tachycardia, ergotism. Ergotamine and triptans should not be used within 24 hours of each other. Metabolized by CYP450, concomitant use with CYP450 inhibitors may cause ergotism.</td>
</tr>
</tbody>
</table>

• The use of opioids is not recommended and has decreased (see paragraph 4.2)
All headaches are not equal: A review of migraine as a state of brain dysfunction

5-HT<sub>1D</sub> agonists

<table>
<thead>
<tr>
<th>Active ingredient</th>
<th>Trade name</th>
<th>Dosage</th>
<th>Mechanism of action</th>
<th>General comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sumatriptan</td>
<td>Imigran® 50 and 100 mg tablets, injection 6 mg/0.5 ml in a prefilled syringe and nasal spray 20 mg/0.1 ml in a unit-dose device.</td>
<td>Oral: initially 50 - 100 mg, dose may be repeated after 2 - 4 hours if symptoms recur, max 300 mg / 24 hours</td>
<td>Elimination t&lt;sub&gt;1/2&lt;/sub&gt; 2 hours Onset of action 10 min (s.c.), and 30 min (oral)</td>
<td>MOA Selective SHT&lt;sub&gt;1D&lt;/sub&gt; receptor agonists. Normalization of dilated intracranial arteries, inhibition of vasoactive peptide release, and inhibition of transmission through second-order neurons ascending to the thalamus.</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Extensively metabolised in liver.</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Migrex®, Triptom® 50 and 100 mg tablets</td>
<td>Intranasal 20 mg, may be repeated after 2 hours, max 40 mg / 24 hours</td>
<td>Slower onset of action, longer t&lt;sub&gt;1/2&lt;/sub&gt; (6 hours); less headache recurrence; may be used for a few days continuously for menstrual migraine, beginning 2 days ahead of menstrual period.</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Naratriptan</td>
<td>May be repeated once after 4 hours if needed. Max 5 mg/24 hours</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Hepatic and renal impairment: max 2.5 mg / 24 hours</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Eletriptan</td>
<td>Max: 160 mg / 24 hours</td>
<td>Eletriptan is contraindicated with CYP450 3A4 inhibitors.</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Relpax® 40 and 80 mg tablets</td>
<td>Max 30mg / 24 hours</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Maxalt® 5 and 10 mg tablets; 10mg wafers</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Zomig® 2.5 mg tablets and dispersible tablets</td>
<td>Max 15 mg / 24 hours</td>
<td>Elimination t&lt;sub&gt;1/2&lt;/sub&gt; 3 hours&lt;sup&gt;52&lt;/sup&gt; 2.5 mg three times per day for menstrual migraine prophylaxis</td>
<td></td>
</tr>
</tbody>
</table>

Table 2 - Migraine prophylaxis

Preventive (prophylactic therapy)

<table>
<thead>
<tr>
<th>Active ingredient and trade name where applicable</th>
<th>Dosage interval</th>
<th>Mechanism of action</th>
<th>General comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Beta adrenergic blockers</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Propranolol</td>
<td>Inderal® and others (Also timolol, metoprolol, atenolol and nadolol)</td>
<td>40–80 mg bd</td>
<td>Restore descending nociceptive inhibition&lt;sup&gt;56&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

General principles:<sup>46</sup>  
- Consider if frequent attacks require acute medication more than twice per week, the duration of an attack is longer than 24 hours, in case of medication failure or overuse and in uncommon migraine variants with the risk for permanent neurologic injury;  
- Administer daily to reduce the frequency, severity, and duration of attacks and to increase response to acute therapy;  
- May be given as intermittent therapy when headaches recur in a predictable pattern (e.g. exercise-induced/menstrual migraine);  
- Drug selection is primarily based on the side effect profiles and comorbid conditions; response is unpredictable and a 2–3 month trial is necessary to achieve benefit, while maximal benefit is typically observed following 6 months of treatment;<sup>59</sup>  
- Start with low doses, titrate until therapeutic effect or side-effects become intolerable;  
- Overuse of acute headache treatments will interfere with the effects of preventive treatment;<sup>59</sup>  
- Continue prophylaxis for at least 6–12 months after headache frequency and severity have diminished, gradually taper or discontinue;  
- Although different formulations of a specific triptan may be used in the same 24-hour period, only one triptan may be used during this time frame;  
- Triptans should not be used for more than 3 days of any given week due to the risk of overuse headache;<sup>57</sup>  
- Sumatriptan, rizatriptan and zolmitriptan should not be given within 2 weeks of MAOI therapy;  
- Concomitant SSRIs may cause serotonin syndrome, although the risk is very low and reconsideration of this FDA warning in 2006 is recommended;<sup>58</sup>  
- Side effects: paresthesia, fatigue, dizziness, flushing, warm sensations, and somnolence. “Triptan sensations,” e.g. tightness, heaviness, or pain in the chest, neck, or throat are reported by 25% of patients;<sup>59</sup>  
- Contraindicated in a history of ischemic heart disease and cerebrovascular disease. Patients at risk for unrecognized coronary artery disease should use triptans with caution;<sup>59</sup>  
- Triptans have traditionally been contraindicated in hemiplegic migraine and migraine with brain stem aura; however, this prohibition is being reconsidered as it is evidently safe to use in these conditions. <sup>35</sup>
Anticonvulsants

<table>
<thead>
<tr>
<th>Drug</th>
<th>Dose</th>
<th>Effect</th>
<th>Side-effects</th>
</tr>
</thead>
<tbody>
<tr>
<td>Valproic acid</td>
<td>300 mg/d, titrate slowly upwards to 600–900 mg/d</td>
<td>Reduce neuronal hyperexcitability</td>
<td>Nausea, tremor, somnolence, weight gain, hair loss, hepatotoxicity; Baseline liver function tests are required.</td>
</tr>
<tr>
<td>Topiramate</td>
<td>50–200 mg/d</td>
<td>AMPA/kainate glutamate antagonist, blocks neuronal excitability</td>
<td>50% of patients respond;</td>
</tr>
</tbody>
</table>

Antidepressants

<table>
<thead>
<tr>
<th>Drug</th>
<th>Dose</th>
<th>Effect</th>
<th>Side-effects</th>
</tr>
</thead>
<tbody>
<tr>
<td>Amitriptyline</td>
<td>10–25 mg at night, seldom &gt; 75–100 mg necessary</td>
<td>Effects are independent of antidepressant activity, probably related to down regulation of central 5-HT1A receptors, increased synaptic noradrenaline and enhanced opioid receptor actions</td>
<td>Anticholinergic effects may limit use especially elderly, increased appetite and weight gain, cardiovascular effects; Good tolerability, nausea, palpitations</td>
</tr>
<tr>
<td>Venlafaxine</td>
<td>75–150 mg/d</td>
<td>Calcium channel blocker reduces neuronal hyperexcitability; also dopamine, serotonin and histamine receptor antagonist</td>
<td>If no improvement after 2 months, discontinue use; Should be interrupted after 6 months and only reinstated if patient relapses;</td>
</tr>
<tr>
<td>Flunarizine</td>
<td>5–10 mg at night</td>
<td>Calcium channel blocker reduces neuronal hyperexcitability; also dopamine, serotonin and histamine receptor antagonist</td>
<td>Calcium channel blocker reduces neuronal hyperexcitability; also dopamine, serotonin and histamine receptor antagonist</td>
</tr>
<tr>
<td>Pizotifen</td>
<td>0.5–3 mg/d, max single dose 3 mg, max daily dose 4.5 mg</td>
<td>Histamine and serotonin receptor antagonist; weak anticholinergic</td>
<td>Drowsiness and weight gain</td>
</tr>
<tr>
<td>Clonidine</td>
<td>25 microgram bd, increase up to total of 150 micrograms/d in divided doses</td>
<td>Alpha 2 adrenoceptor agonist</td>
<td>Withdraw gradually to prevent rebound hypertension; Evidence of lack of efficacy in migraine prophylaxis</td>
</tr>
</tbody>
</table>

Fact Box 1 - Medication overuse headache

**Medication overuse headache (MOH; also known as analgesic rebound headache)**

All drugs used to treat headaches, even aspirin and paracetamol, can also cause headache when overused, often resulting in the worsening and/or changing of severity and symptoms previously observed or in chronicity of headaches. This poses a significant problem when considering that 3% of the population suffer from chronic headache, resulting in a MOH prevalence of more than 1% in the general population. Furthermore, patients who suffer from migraine are likely to develop MOH when using analgesics for other types of pain – an observation that is uncommon in patients free from migraine. The exact mechanisms responsible for MOH are still unclear, but various factors have been implicated including, but not limited to, changes in receptor expression and sensitivity (e.g. 5-HT1A receptors with the use of triptans), behavioural mechanisms (e.g. prophylactic use of analgesics), dependencies due to the psychotropic side-effects (e.g. opioids) or withdrawal symptoms (e.g. caffeine) and physical dependencies (e.g. opioids and ergot alkaloids).

MOH may result from the use of a single therapeutic agent or the concomitant use of several drugs or combination preparations and is described by the International Headache Society as headaches occurring on 15 or more days of a month in patients with pre-existing headache disorders who overuse one or more drugs that are used in the acute and/or symptomatic treatment of headache. Indeed, almost half of patients suffering from headache on 15 or more days during a single month (diagnosed as daily chronic headache; DCH) meet these criteria and physicians are encouraged to diagnose patients accordingly in addition to the pre-existing headache diagnosis. The most common treatment for MOH is withdrawal and the majority of patients improve after discontinuation while more serious cases may present with symptoms observed in drug addiction and should be referred and/or treated accordingly. With the exception of opioids, barbiturates and benzodiazepines, most drugs that lead to MOH can and should preferably be discontinued immediately. During discontinuation, patients may suffer from a variety of symptoms, including headache, nausea, tachycardia and anxiety for up to four weeks, although these symptoms more frequently last for an average of 3 to 4 days. While therapeutic protocols and guidelines for MOH do not exist, successful treatment is likely in outpatients who are well-instructed, motivated and treated symptomatically, e.g. with anti-emetics, ß-blockers, hydration. Mono-analgesics (naproxen 500 mg two to three times daily) may also be included while the use of corticosteroids, specifically prednisone (100 mg daily for three days), has been demonstrated to be effective in relieving withdrawal headache during detoxification. Patients remain at the greatest risk for relapse during the first year (41%) after withdrawal, where after this risk significantly decreases.

Novel drugs for migraine prophylaxis (FDA-approved, but not registered in SA for the indication)

**Botulinum toxin type A (BoNT-A; Botox; Onabotulinumtoxin A)**

Botulinum toxin type A (BoNT-A) is an FDA-approved treatment (2010) for the prevention of chronic migraine in adults older than 18 years. It has been demonstrated to be effective and safe by data from Phase III of the Research Evaluating Migraine Prophylaxis Therapy-program (PREEMPT). Treatment response, however, varies greatly among patients and therefore patient selection is crucial; this in turn depends on the site and frequency of pain, among others. Longer term treatments seem to increase its efficacy and while current guidelines recommend a cessation of treatment following 2 cycles in patients not responding to a single cycle of treatment, such early cessation may not be justified. It appears as though its benefit increases with additional treatment cycles, with 5 cycles of treatment sometimes being necessary to reach optimal effect. In responders, one BoNT-A treatment can be effective for approximately 3 months. BoNT-A not only inhibits the release of acetylcholine from presynaptic nerve endings, but may also modify the release of transmitters involved in pain transmission,
All headaches are not equal: A review of migraine as a state of brain dysfunction

e.g. substance P and CGRP, thereby preventing activation of central pain networks. Furthermore, BoNT-A is well tolerated and injection-related side effects are usually mild and transient with little or no systemic side effects.

CGRP antagonists

As alluded to earlier, CGRP is a vasoactive peptide released from activated trigeminal nerves which facilitates pain transmission from cerebral vessels to the central nervous system. In realization of the role CGRP plays in the pathophysiology of migraine, a range of monoclonal antibodies have recently been developed, including erenumab (Aimovig®; 2018), fremanezumab (AJOVY®; 2018), and galcanezumab (Emgality®; 2018). While erenumab targets the CGRP receptor, fremanezumab, galcanezumab and eptinezumab (still under regulatory review) bind and inactivate the CGRP molecule itself. The short-term side-effects of these drugs have been minimal and were mostly restricted to injection site reactions. However, fremanezumab, but not erenumab, may carry a risk for more severe hypersensitivity reactions, while general concern is expressed for possible long-term cardiovascular sequelae. These antibodies are usually self-administered once a month as a single subcutaneous injection. As may be expected, they are relatively expensive (approximately US $575/month) and patients would have to fail on treatment with at least two to three oral preventive drugs to become eligible for such treatment. While currently available preventive drugs offer relatively limited efficacy and are not free from side effects, CGRP monoclonal antibodies may offer hope to many migraine sufferers that remain unresponsive to current interventions.

Emergency treatment of migraine

When failing on customary acute migraine treatment (Table 1), patients may present to the emergency room during which time parenteral treatment is usually preferred. Recommendations for emergency treatment were compiled by the American Headache Society following a systematic review of clinical trials and a meta-analysis that included the following:

- Antidopaminergic drugs (metoclopramide, prochlorpromazine, haloperidol, and droperidol are commonly used) do not only relieve nausea, vomiting and gastroparesis associated with acute migraine, but are also analgesic, albeit for reasons still unknown. Akathisia (restlessness) and dystonic reactions may occur in as much as a third of patients receiving intravenous dopamine antagonists and to prevent this, intravenous diphenhydramine is often added; however, this practice is not supported by all clinicians.

- Subcutaneous sumatriptan is also effective although side effects include flushing, chest pain, and an occasional worsening of headache symptoms. Further, a higher incidence of headache recurrence within 24 hours is also noted. In general, antidopaminergic agents are usually better tolerated than sumatriptan.

- Nonsteroidal antinflammatory drugs, e.g. ketorolac and naproxen, alone or in combination with the antidopaminergics or sumatriptan may be of benefit.

- Antihistamines (monotherapy), ketamine, magnesium and propofol have not been found particularly effective.

- Dexamethasone (10 mg IV) can be added to counter the recurrence of headaches—which patients often experience within 24 hours of visiting the emergency department—but be mindful of giving steroids to patients with diabetes.

- Slow administration of parental drugs is necessary and may prevent some of the commonly reported adverse effects.

- While opioids were previously the mainstay of abortive treatment of acute migraine episodes, it is now recommended that they best be avoided due to the lack of evidence regarding their efficacy and possible long-term sequelae. In this regard, the decline in use of the emergency room opioids has also decreased the recurrence rate of acute episodes that would otherwise have necessitated more visits to an emergency care setting.

Conclusion

Migraine is a severe, debilitating and highly disabling condition that is founded in an altered state of brain function that results in a hypersensitive interpretation of sensory stimuli and exacerbated trigeminovascular nociceptive pain transmission. The condition is variably characterised by four phases, i.e. a premonitory phase, CSD and aura, acute headache and finally, resolution. Further, aura migraine may be associated with significant comorbidities related to epilepsy, ischemic stroke and cardiovascular pathology and should be regarded not only as a risk factor, but as a unique neurobiological disturbance.

References


