**DEFINITIONS**

**Primary headache disorders** - those not associated with an underlying pathology, eg migraine, tension-type, and cluster headache.

**Secondary headache disorders** - headache attributed to an underlying pathological condition. Includes any head pain of infectious, neoplastic, vascular or drug-induced origin.

**Chronic headache** occurs on more than 15 days per month for more than three months.

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**DIAGNOSIS OF PRIMARY HEADACHE**

An inadequate history is the probable cause of most misdiagnosis of headache type.

***D*** Practitioners should consider using headache diaries and appropriate assessment questionnaires to support the diagnosis and management of headache.

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**CHARACTERISTICS OF MIGRAINE**

*bold indicates the most helpful for distinguishing migraine from other headache*

- episodic moderate to severe headache that causes disability
- unilateral
- pulsating
- builds up over minutes to hours
- moderate to severe in intensity
- associated with nausea and/or vomiting and/or sensitivity to light and/or sensitivity to sound
- aggravated by routine physical activity
- **typical aura** (in 15–33% of patients with migraine)
- exacerbation by physical activity
- sensitivity to light between attacks
- positive family history of migraine.

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***C*** Patients who present with a pattern of recurrent episodes of severe disabling headache associated with nausea and sensitivity to light, and who have a normal neurological examination, should be considered to have migraine.

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**INVESTIGATIONS OF MIGRAINE**

***D*** Neuroimaging is not indicated in patients with a clear history of migraine, without red flag features for potential secondary headache, and a normal neurological examination.
The treatment of acute migraine attacks should be selected for each patient according to severity and frequency of attacks, other symptoms, patient preference and history of treatment.

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<thead>
<tr>
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<tbody>
<tr>
<td><strong>D</strong></td>
<td>Opioid analgesics should not be routinely used for the treatment of patients with acute migraine due to the potential for development of medication overuse headache.</td>
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</tr>
<tr>
<td><strong>A</strong></td>
<td>Aspirin 900 mg or ibuprofen 400 mg is recommended for acute treatment in patients with all severities of migraine.</td>
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<tr>
<td><strong>B</strong></td>
<td>Paracetamol 1,000 mg is recommended as acute treatment for mild to moderate migraine.</td>
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<tr>
<td><strong>A</strong></td>
<td>Oral triptans are recommended for acute treatment in patients with all severities of migraine if previous attacks have not been controlled using simple analgesics.</td>
<td></td>
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<tr>
<td><strong>A</strong></td>
<td>Almotriptan 12.5 mg, eletriptan 40-80 mg or rizatriptan 10 mg, are the preferred oral triptans for acute migraine.</td>
<td></td>
</tr>
<tr>
<td><strong>B</strong></td>
<td>If a patient does not respond to one triptan an alternative triptan should be offered.</td>
<td></td>
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<tr>
<td><strong>D</strong></td>
<td>Triptans should be taken at, or soon after, the onset of the headache phase of a migraine attack.</td>
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</tr>
<tr>
<td><strong>C</strong></td>
<td>A combination of sumatriptan 50-100 mg and naproxen sodium 500 mg may be helpful in acute migraine particularly in prolonged attacks which are associated with recurrence.</td>
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<tr>
<td><strong>D</strong></td>
<td>Oral and rectal anti-emetics can be used in patients with acute migraine attacks to reduce symptoms of nausea and vomiting and to promote gastric emptying.</td>
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<tr>
<td><strong>B</strong></td>
<td>A combination of aspirin and metoclopramide can be used for the treatment of patients with acute migraine attacks.</td>
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</tr>
<tr>
<td>✓</td>
<td>Practitioners should recognise that a patient’s standard therapy may not give a consistent response.</td>
<td></td>
</tr>
<tr>
<td>✓</td>
<td>When initiating acute treatment for migraine the risks of medication overuse headache should be discussed with the patient.</td>
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**NON-PHARMACOLOGICAL MANAGEMENT**

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<tbody>
<tr>
<td>B</td>
<td>Stress management should be considered as part of a combined therapies programme to help patients reduce the frequency and severity of migraine headaches.</td>
</tr>
<tr>
<td>B</td>
<td>Acupuncture should be considered for preventive management in patients with migraine.</td>
</tr>
</tbody>
</table>

**PROPHYLAXIS**

**Trials demonstrate that prophylaxis provides reduction in severity and frequency of headaches by 50%.**

### Betablockers

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<table>
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<tbody>
<tr>
<td>A</td>
<td>Propranolol 80-240 mg per day is recommended as first line therapy for prophylaxis in patients with migraine.</td>
</tr>
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</table>

### Antiepileptics

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<tbody>
<tr>
<td>A</td>
<td>In patients with episodic migraine and chronic migraine topiramate 50-200 mg per day is recommended to reduce headache frequency and severity.</td>
</tr>
<tr>
<td>A</td>
<td>In patients with episodic migraine sodium valproate 800-1,500 mg per day is recommended to reduce headache frequency and severity.</td>
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### Antidepressants

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<tr>
<td>B</td>
<td>Amitriptyline 25-150 mg per day is recommended for patients requiring prophylaxis of migraine.</td>
</tr>
<tr>
<td>B</td>
<td>Venlafaxine 75-150 mg per day is an effective alternative to tricyclic antidepressants for prophylaxis of migraine.</td>
</tr>
</tbody>
</table>
A diagnosis of tension-type headache should be considered in a patient presenting with bilateral headache that is non-disabling where there is a normal neurological examination.

**TREATMENT OF PATIENTS WITH TENSION TYPE HEADACHE**

**ACUTE TREATMENT**

A Aspirin and paracetamol are recommended for acute treatment in patients with tension-type headache.

**PROPHYLAXIS**

A Tricyclic antidepressants, particularly amitriptyline, 25-150 mg per day, are recommended as the agents of choice where prophylactic treatment is being considered in a patient with chronic tension-type headache.
- Cluster headache is the most common trigeminal autonomic cephalalgia, with a prevalence of 1 in 1,000.
- Characterised by attacks of severe unilateral pain in a trigeminal distribution.
- Associated with prominent ipsilateral cranial autonomic features.

### FEATURES DISTINGUISHING CLUSTER HEADACHE FROM MIGRAINE

<table>
<thead>
<tr>
<th>Headache type</th>
<th>Cluster headache</th>
<th>Migraine</th>
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</thead>
<tbody>
<tr>
<td>Duration</td>
<td>15 mins-3 hrs</td>
<td>4-72 hrs</td>
</tr>
<tr>
<td>Onset</td>
<td>rapid</td>
<td>gradual</td>
</tr>
<tr>
<td>Frequency</td>
<td>1 every other day-8/day</td>
<td>&lt;1/year-1/day (median 1-2/month)</td>
</tr>
<tr>
<td>Restlessness during an attack</td>
<td>100%</td>
<td>0%</td>
</tr>
<tr>
<td>Ipsilateral autonomic features</td>
<td>prominent</td>
<td>occasional</td>
</tr>
</tbody>
</table>

### ACUTE TREATMENT

A. Nasal sumatriptan or zolmitriptan is recommended for treatment of acute attacks of cluster headache in patients who cannot tolerate subcutaneous sumatriptan.

A. Subcutaneous injection of 6 mg sumatriptan is recommended as the first choice treatment for the relief of acute attacks of cluster headache.

### PROPHYLAXIS

B. Verapamil, 240-960 mg is recommended for the prophylaxis of cluster headache.
Consider the diagnosis of secondary headache in patients presenting with new onset headache or headache that differs from their usual headache.

**Red flag features which should prompt referral for further investigation:**
- new onset or change in headache in patients who are aged over 50
- thunderclap: rapid time to peak headache intensity (seconds to 5 mins)
- focal neurological symptoms (eg limb weakness, aura < 5 min or > 1 hr)
- non-focal neurological symptoms (eg cognitive disturbance)
- change in headache frequency, characteristics or associated symptoms
- abnormal neurological examination
- headache that changes with posture
- headache waking the patient up (NB migraine is the most frequent cause of morning headache)
- headache precipitated by physical exertion or valsalva manoeuvre (eg coughing, laughing, straining)
- patients with risk factors for cerebral venous sinus thrombosis
- jaw claudication or visual disturbance
- neck stiffness
- fever
- new onset headache in a patient with a history of human immunodeficiency virus (HIV) infection
- new onset headache in a patient with a history of cancer.

**Patients presenting with headache for the first time or with headache that differs from their usual headache should have a clinical examination, a neurological examination including fundoscopy, and blood pressure measurement.**

**Patients who present with headache and red flag features of potential secondary headache should be referred to an appropriate specialist for further assessment.**

**Brain CT should be performed in patients with headache who have unexplained abnormal neurological signs, unless the clinical history suggests MRI is indicated.**

**Clinicians requesting neuroimaging should be aware that both MRI and CT can identify incidental neurological abnormalities which may result in patient anxiety as well as practical and ethical dilemmas with regard to management.**
Patients with a first presentation of thunderclap headache should be referred immediately to hospital for same day specialist assessment.

In patients with thunderclap headache, unenhanced CT of the brain should be performed as soon as possible and preferably within 12 hours of onset.

In patients with thunderclap headache and a normal CT should have a lumbar puncture.

In patients who require a lumbar puncture for thunderclap headache, oxyhaemoglobin and bilirubin should be included in cerebrospinal fluid analysis.

Lumbar puncture in CT negative patients with suspected subarachnoid haemorrhage should be carried out as soon as possible after 12 hours has elapsed from the onset of symptoms.

In delayed presentations, lumbar puncture can be performed up to two weeks from onset of symptoms.

Described as:
- a high-intensity headache of rapid onset reaching maximum intensity in less than a minute in most, but can take a few minutes in some
- may be primary or secondary (no reliable differentiating features)
- subarachnoid haemorrhage is the commonest secondary cause, although a number of other conditions can also present with thunderclap headache (intracerebral haemorrhage, cerebral venous sinus thrombosis, arterial dissection, pituitary apoplexy).
**MEDICATION OVERUSE HEADACHE**

**Described as:**
headache which is present for 15 days or more per month and which has developed or worsened while taking regular symptomatic medication.

| D | Medication overuse headache must be excluded in all patients with chronic daily headache (headache ≥15 days/month for >3 months). |
| C | Clinicians should be aware that patients using any acute or symptomatic headache treatment are at risk of medication overuse headache. Patients with migraine, frequent headache and those using opioid-containing medications or overusing triptans are at most risk. |
| C | When diagnosing medication overuse headache, psychiatric comorbidity and dependence behaviour should be considered. |
| C | Patients with medication overuse headache who have psychiatric comorbidity or dependence behaviour should have these conditions treated independently. Referral to a psychiatrist or a clinical psychologist should be considered. |

**TREATMENT**

- Medication withdrawal should be attempted in all patients with medication overuse headache.

- **Patients with medication overuse headache caused by simple analgesics or triptans** should be advised to abruptly withdraw the overused medication. In the majority of patients this can be as an outpatient with structured advice.

- **Patients with medication overuse headache caused by opioids and opioid-containing analgesics** should be considered for gradual withdrawal of the overused medications.

- **If frequent headache persists after symptomatic medications have been withdrawn, prophylactic agents may be effective and should be considered.**

**NB** Abrupt withdrawal from medication initially results in worsening of headache.
PREGNANCY, CONTRACEPTION, MENSTRUATION AND THE MENOPAUSE

PREGNANCY

Where possible, the use of medication in pregnancy should be avoided, particularly in the first trimester. Paracetamol has been used routinely in all stages of pregnancy without apparent harmful effect, and, if drug treatment is essential then paracetamol is the analgesic of choice.

- Paracetamol 1,000 mg is the treatment of choice in pregnancy for all patients with migraine and tension-type headache when the pain is sufficient to require analgesia.

- If paracetamol provides insufficient analgesia aspirin 300 mg or ibuprofen 400 mg can be used in the first and second trimester of pregnancy.

Aspirin is contraindicated during the third trimester of pregnancy. Long term exposure or exposure to high doses of ibuprofen in late pregnancy is associated with an increased risk of fetal complications.

ORAL CONTRACEPTION

Women with migraine with aura using a COCP have a relative risk of 8.72 (95% CI 5.05 - 15.05) for developing stroke. Women over the age of 35 suffering from migraine without aura also have an increased risk of ischaemic stroke if they take COCP.

- Women with migraine with aura should not use a combined oral contraceptive pill.

- Patients with migraine without aura who are over the age of 35 should not use a combined oral contraceptive pill.

MENSTRUATION

The frequency and severity of migraine can increase around the time of menstruation.

- Patients with acute menstrual migraine can be treated with mefenamic acid or a combination of aspirin, paracetamol and caffeine.

- Sumatriptan, zolmitriptan, naratriptan and rizatriptan are recommended for the acute treatment of patients with menstrual migraine.

- Frovatriptan 2.5 mg/day or naratriptan 1 mg twice daily taken two days before day one of the menstrual cycle then for a further four or five days respectively is recommended for the prophylaxis of menstrual migraine.
This Quick Reference Guide provides a summary of the main recommendations in SIGN Guideline 107, *Diagnosis and management of headache in adults.*

Recommendations are graded A B C D to indicate the strength of the supporting evidence. Good practice points ☑️ are provided where the guideline development group wishes to highlight specific aspects of accepted clinical practice.

Details of the evidence supporting these recommendations can be found in the full guideline, available on the SIGN website: [www.sign.ac.uk](http://www.sign.ac.uk)