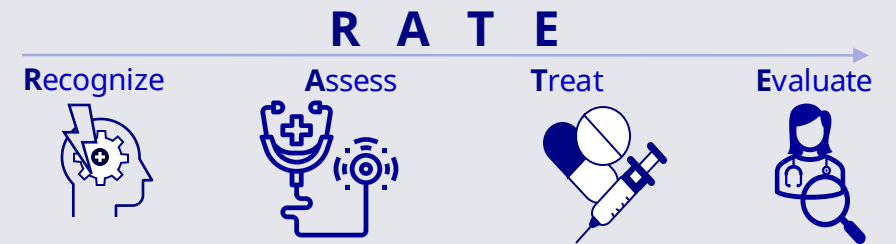




Migraine Management Toolkit: Diagnosis and Treatment Considerations

Start





Disclaimer

- The audience for this toolkit is US healthcare professionals.
- Consensus Treatment Guidelines or Clinical Practice Guidelines developed by medical societies may recommend therapies within their guidelines which do not have FDA approval for the indicated use. Prescription therapies noted within this presentation are limited to those with FDA approved indications for the treatment of acute migraine or prevention of migraine.

Agree and proceed





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Toolkit objectives

Introduction

- ▶ Migraine is a common neurological disease worldwide
- ▶ Migraine is highly disabling and impacts patients' quality of life
- ▶ Migraine is associated with substantial economic burden
- ▶ Migraine is associated with reduced work productivity
- ▶ Migraine is underdiagnosed
- ▶ Patients seek treatment across many providers: Highest in primary care
- ▶ Variable and non-specific symptoms may lead to delayed diagnosis
- ▶ Recognizing, Assessing, Treating, and Evaluating patients with migraine

Recognize

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- ▶ Migraine subtypes and association with a range of symptoms
- ▶ Stages of a migraine attack
- ▶ Knowing when to suspect migraine is important
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- ▶ Migraine in pregnancy
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- ▶ Preventive migraine treatment aims to reduce the frequency, severity, and duration of attacks
- ▶ Unique considerations for special populations experiencing migraine
- ▶ Summary: an individualized treatment management plan should be implemented following diagnosis

Evaluate

- ▶ Appraisal of treatment plans is important to ensure ongoing efficacy and safety
- ▶ Treatment plans and long-term follow-up

Summary

Introduction



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Toolkit objectives

- To describe the unmet medical need in the diagnosis and treatment of migraine in non-headache specialties
- To review ICHD-3 diagnostic criteria and differential diagnoses of migraine
- To provide an overview of current acute and preventive treatments, and guidance on the clinical management of patients with migraine



Migraine is a common neurological disease worldwide

Introduction



Recognize



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Migraine prevalence: more than 1 billion people live with migraine globally, with approximately ~40 million in the US¹⁻³



2nd

most prevalent **neurological disease** in the US^{a,4}



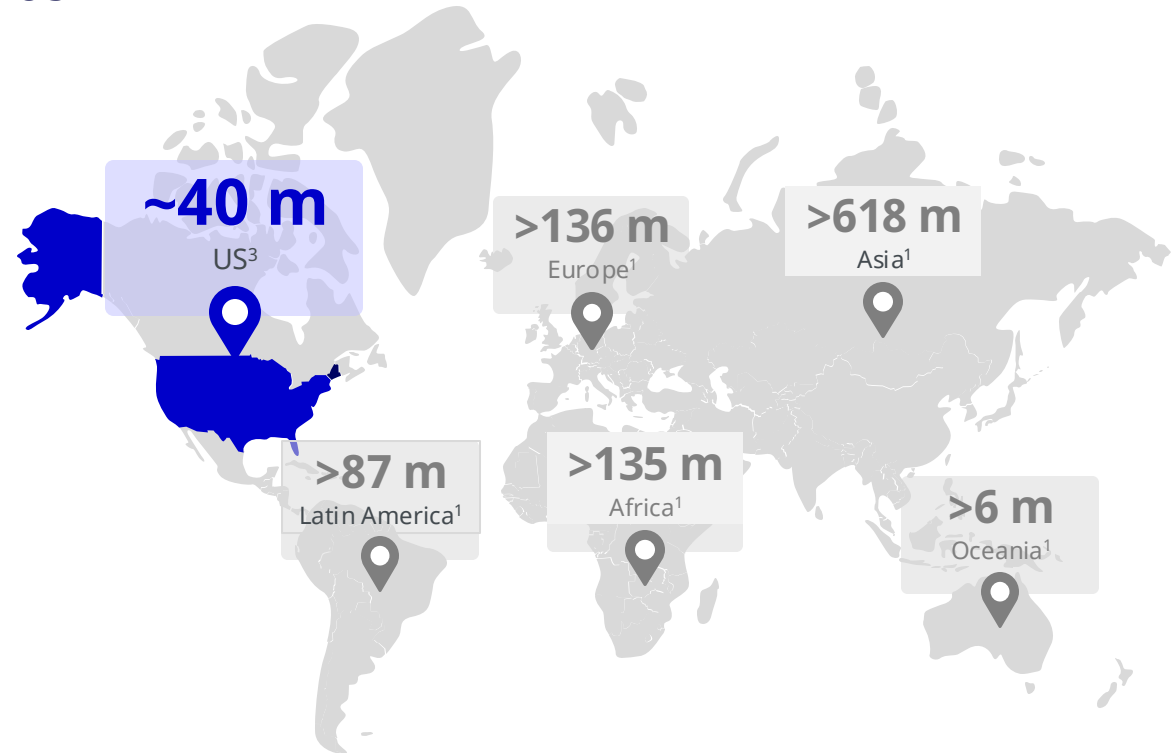
~3x

more common in females than males^{5,6}



25-55 years

Most common in the **productive years**⁶



^aTension-type headache is the most prevalent neurological disease.

1. GBD 2016 Headache Collaborators. Lancet Neurol 2018;17:954-76; 2. Stovner LJ, et al. J Headache Pain 2022;23:34; 3. Law HZ, et al. Plast Reconstr Surg Glob Open 2020;8:e2790;

4. GBD 2017 US Neurological Disorders Collaborators. JAMA Neurol 2021;78:165-76; 5. Lipton RB, et al. Headache 2022;62:122-40; 6. Lipton RB, et al. Headache 2018;58:1408-26.



Migraine is highly disabling and impacts patients' quality of life

Among the **top 3 most debilitating neurological diseases in the US**^{1,a}

Migraine impacts multiple aspects of individuals' lives



report being very or extremely limited in completing daily activities during headache phase²



report impaired sleep²



feel fatigued often or always³



need help with daily tasks²



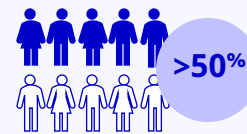
worry about long-term financial security⁴



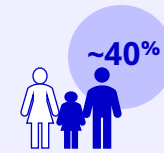
report that they would get more enjoyment from their free time without headaches⁴



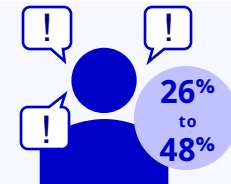
indicate that migraine has contributed to relationship problems⁴



report feeling helpless at least some of the time⁵



believe that they would be a better parent if they did not have headaches⁴



experience migraine-related stigma often or very often⁶

^aBased on systematic analysis of the Global Burden of Disease 2017 study. Data on incidence, prevalence, mortality, and DALY in the US of 14 major neurological disorders were analyzed; migraine (2.40 million absolute DALYs) ranked third behind stroke (3.58 million) and Alzheimer disease and other dementias (2.55 million).¹

DALY, disability-adjusted life years; US, United States.

1. GBD 2017 US Neurological Disorders Collaborators. JAMA Neurol 2021;78:165-76; 2. Gibbs SN, et al. Headache 2020;60:1351-64 3. Martelletti P, et al. J Headache Pain 2018;19:115. 4. Buse DC, et al. Headache 2019;59:1286-99; 5. Hubig LT, et al. J Headache Pain 2022;23:9; 6. Shapiro RE, et al. Neurology 2024;102:e208074.



Migraine is associated with substantial economic burden

High economic burden due to direct and indirect costs^{1,2}

Episodic migraine costs

>\$2600

per person per year in the US¹

Chronic migraine costs

>\$8200

per person per year in the US¹

Estimated cost of productivity losses due to migraine in the US²



60,000–686,000

annual workdays affected by absenteeism and presenteeism across different industries



Costs of lost productive time ranging between

\$18 million and **\$155 million**



Annual indirect costs estimated to be **~6–9× higher** than direct costs

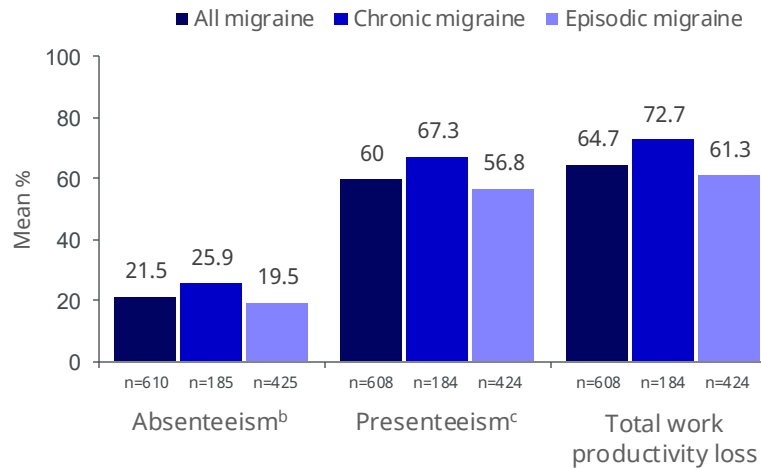




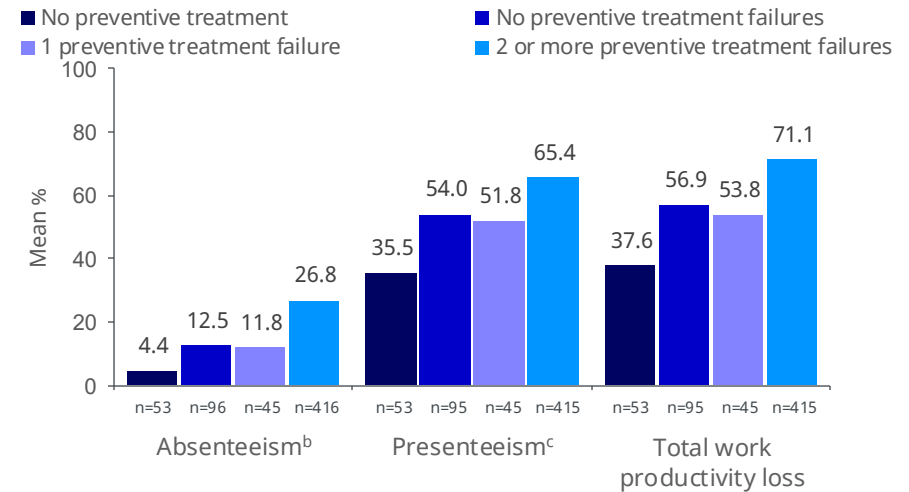
Migraine is associated with reduced work productivity

Productivity losses are greater among those with chronic migraine and those who have had two or more preventive treatment failures^{1a}

Work productivity impairment by migraine frequency



Work productivity impairment by preventive treatment history



A headache education and evaluation program² in Japan at Fujitsu Co. Ltd. (81,159 employees)^d consisting of an e-learning program, video seminars, and online consultation with a staff physician led to:

- Annual productivity gains of ~15.2 days per employee per year after completion of program (1.17 days for absenteeism, 14.0 days for presenteeism)
- Annual productivity savings of US \$4,531 per employee

^aWeb-based survey of 1101 people with self-diagnosed migraine in the United States (My Migraine Voice); ^bWork time missed; ^cReduced on-the-job effectiveness; ^dEmployee e-learning participation rate was 90.5%; ~3 in 5 participants had a headache disorder (16.7% migraine; 40.7% tension-type headache, 0.5% cluster headache).

1. Gibbs SN, et al. Headache 2020;60:1351-64; 2. Sakai F, et al. Cephalalgia 2023;43:1-14.



Migraine is underdiagnosed

Introduction



Recognize



Assess



Treat



Evaluate



Despite advances in our understanding of the pathophysiology and management of migraine, it remains widely underdiagnosed¹



Average time between onset
and diagnosis is **~3.3 years**¹



~20-70%
of patients with migraine
do not consult an HCP^{1,2}



~30%
do not receive an
accurate diagnosis^{2,a,b}



Patients with chronic migraine are
57% less likely to receive an accurate
diagnosis than those with episodic migraine^{2,a,c}

7% of patients consult with HCPs who may not be well suited for ongoing headache care; the most frequently reported HCPs providing headache care who are not customary providers of headache care were ENTs, allergists, psychiatrists, ER/urgent care physicians, and dentists.²

^aBased on findings from the longitudinal internet-based survey from the CaMEQ study (N=16,789; among the 9,184 patients with MIDAS scores ≥ 6 and valid health insurance data, 86.3% of patients met criteria for episodic migraine and 13.7% met criteria for chronic migraine)
^bCommon misdiagnoses for migraine include sinus headache, stress headache and tension-type headache; ^cAmong patients under care (management or treatment) of a healthcare practitioner well suited for ongoing headache care (n=2,699)
CaMEQ, Chronic Migraine Epidemiology and Outcomes; ENT, ear nose and throat specialist; ER, emergency room; MIDAS, Migraine Disability Assessment
1. Lipton RB, et al. Headache 2022;62:122-40; 2. Buse DC, et al. Headache 2021;61:628-41



Patients seek treatment across many providers: Highest in primary care

Introduction



Recognize



Assess



Treat

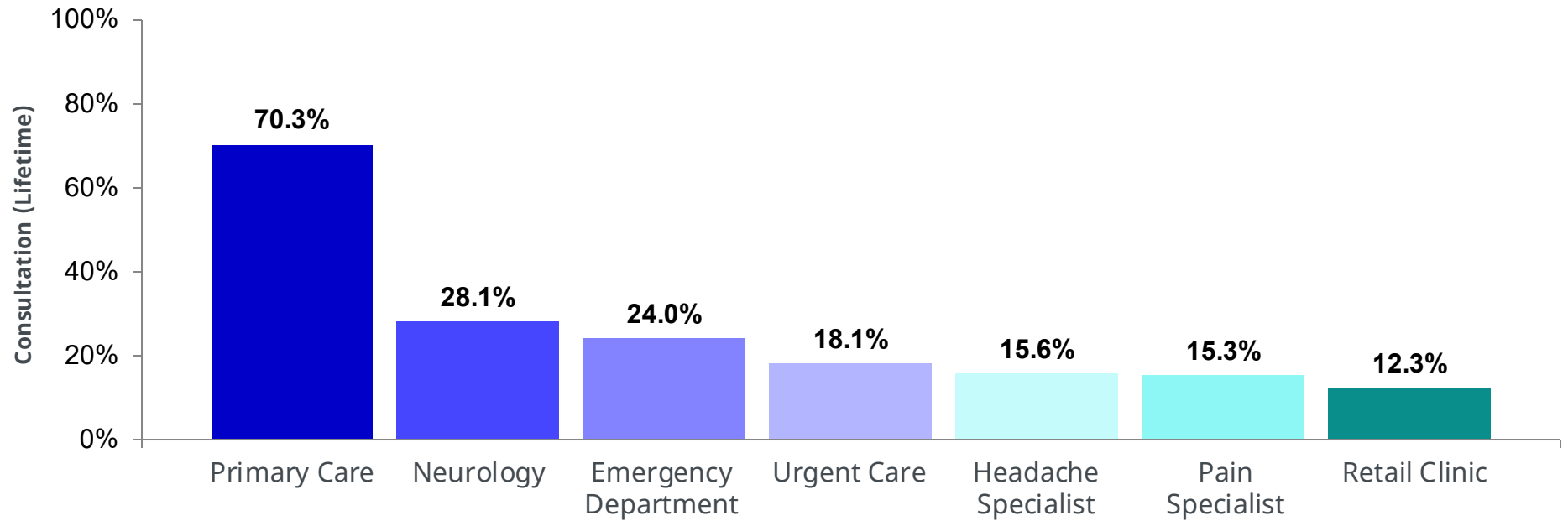


Evaluate



Lifetime Migraine or Headache Consultation by Specialty

n=21,143



Introduction



Recognize



Assess



Treat



Evaluate



Variable and non-specific symptoms may lead to delayed diagnosis

Patients with migraine can experience variable and non-specific symptoms



A **range of symptoms** are associated with migraine, and these can **differ among patients** and **between headache attacks** within the same patient



Migraine diagnostic criteria (ICHD-3) require a combination of common symptoms for migraine diagnosis, but some characteristics may not be present such as **aura, throbbing,** and **severe pain**



Overlapping symptoms with other conditions such as **sinus, tension-type, and cluster headaches** can lead to misdiagnosis



Patients' **failure to describe** all of the **relevant symptoms** they experience during attacks may also lead to underdiagnosis of migraine



Recognizing, Assessing, Treating, and Evaluating patients with migraine

Introduction



Recognize



Assess



Treat



Evaluate



Introducing the mnemonic **RATE** (**R**ecognize, **A**ssess, **T**reat, **E**valuate) as a potential approach that HCPs can utilize to help guide the diagnosis and management of patients presenting with migraine^a



Recognize

When to suspect migraine in patients¹:

- Recurrent headache of moderate to severe intensity
- Visual aura, nausea, and vomiting
- Family history of migraine



Assess

Use a combination of¹:

- Detailed history taking
- Physical examination
- Screening for secondary headache and considering differential diagnoses
- Validated diagnostic and screening tools

Treat

Acute treatment^{1,2}

- For patients with a confirmed diagnosis of migraine

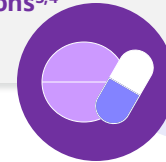
Preventive treatment^{1,2}

- For patients whose attacks significantly interfere with daily routines despite optimized acute treatment
- For those who have frequent attacks, intolerance or contraindication(s) to acute treatments, or failure or overuse of acute treatments

- Based on patient preference²

Special populations^{3,4}

- Pregnancy
- Breastfeeding



Evaluate

- Primary care practitioners should play a role in the long-term management of migraine¹
- It is important to evaluate the impact of the treatment plan on migraine¹
- Treatment plans should be revised and optimized for each patient, as appropriate¹



HCPs, healthcare professionals; RATE, Recognize, Assess, Treat, and Evaluate.

^aThe RATE program described here for patients with migraine follows a similar theme to the proposed RATE strategy for the diagnosis and treatment of patients with chronic pain.⁵

1. Eigenbrodt AK, et al. Nat Rev Neurol 2021;17:501–14; 2. Ailani J, et al. Headache 2021;61:1021–39; 3. Lucas S. Obstet Gynecol 2019;134:211; 4. ACOG Committee on Clinical Practice Guidelines–Obstetrics. Obstet Gynecol 2022;139:944–72; 5. Gebke KB, et al. Postgrad Med 2023;135:244–253.



Recognize: primary versus secondary headache

Recognize

Introduction



Recognize



Assess



Treat



Evaluate



Most patients presenting with headache in clinical practice have a primary headache disorder¹

Primary Headache²

- No known underlying cause
- The most common primary headache disorders include:
 - **Migraine**
 - **Tension-type headache**
 - **Cluster headache**
- Most patients with a chief complaint of headache who present to their HCP for an evaluation will have primary headache

Secondary Headache²

- The result of another condition causing activation of pain-sensitive structures
- Common secondary headaches include those related to:
 - Infection
 - Vascular disease
 - Trauma
- Certain "red flags" or warning signs indicate a need for evaluation

HCP, healthcare practitioner.

1. Dodick DW. Continuum (Minneapolis, MN) 2021;27(3):572-85. 2. Rizzoli P, Mullally WJ. Am J Med. 2018;131:17-24.





Recognize: migraine subtypes and association with a range of symptoms

Recognize

Introduction



Recognize



Assess



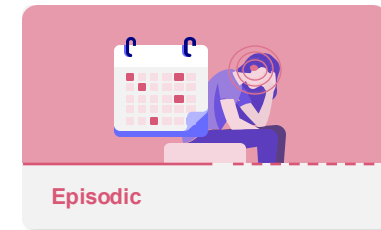
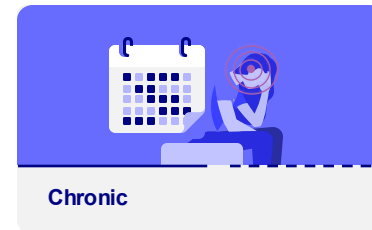
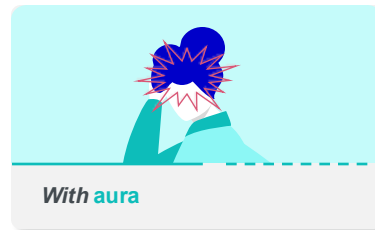
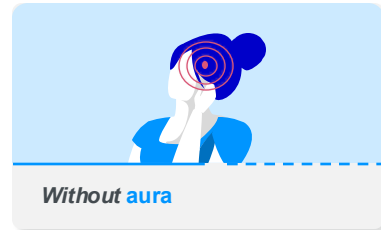
Treat



Evaluate



Migraine is a primary headache disorder that can be categorized into subtypes based on symptoms and headache frequency¹



Migraine is associated with a range of symptoms and pain can be unilateral or bilateral^{2,3}

Migraine is characterized by recurrent headache attacks of moderate to severe pain often accompanied by other symptoms²:



Photophobia



Nausea



Phonophobia



While pain is typically unilateral, **~4 in 10 patients** report bilateral pain during migraine attacks³

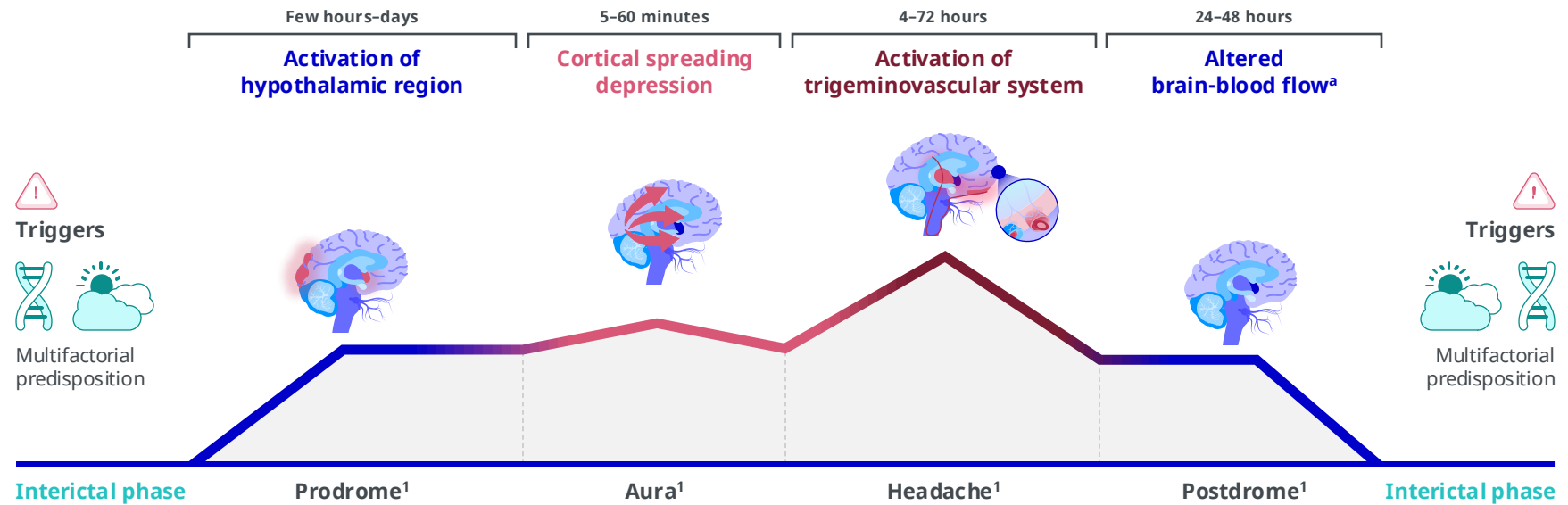




Recognize: stages of a migraine attack

Recognize

Functional changes in different areas of the brain produce an array of symptoms at different stages of an attack¹⁻³



Stages can occur sequentially or overlap, and some may not occur at all for some patients¹

Only ~1 in 3 people with migraine experience aura with some or every attack⁴

^aPostdrome is the least studied and understood phase.

1. Ferrari MD, et al. Nat Rev Dis Primers 2022;8:2; 2. Dodick DW. Lancet 2018;391:1315-30; 3. Andreou AP, Edvinsson L. J Headache Pain 2019;20:117; 4. Lipton RB, et al. Neurology 2002;58:885-94.

Introduction



Recognize



Assess



Treat



Evaluate



Introduction



Recognize



Assess



Treat



Evaluate



Recognize: knowing when to suspect migraine is important

Recognize

When to suspect migraine in patients presenting with headache^{1,2}



Recurrent headache of moderate to severe intensity



Visual aura



Nausea



Vomiting



Family history of migraine



Recognize: common comorbidities associated with migraine

Recognize

Introduction



Recognize



Assess



Treat

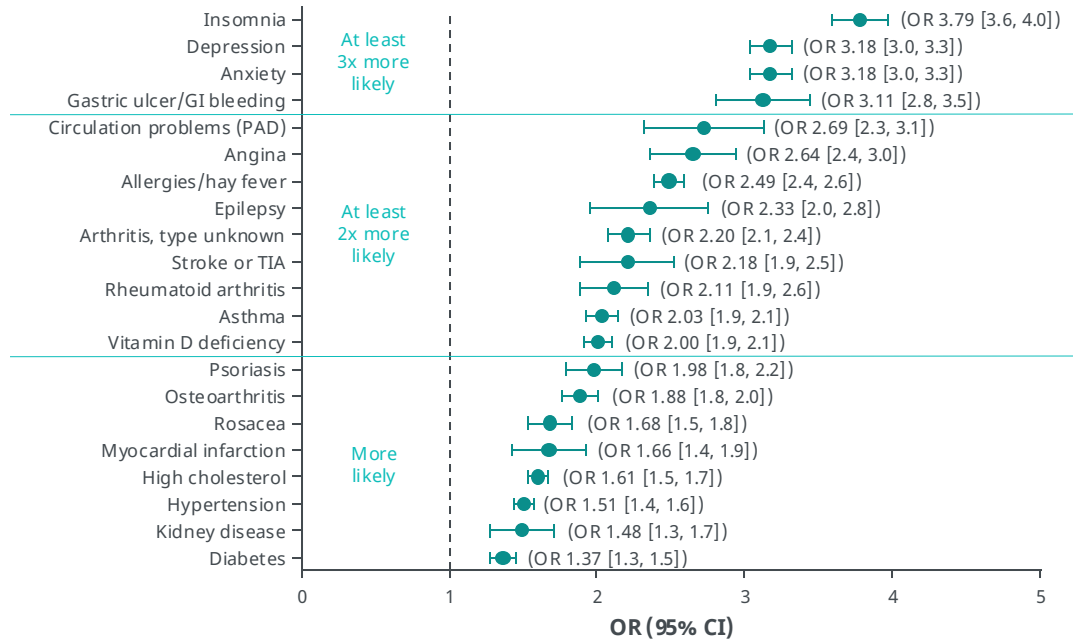


Evaluate



People with migraine experience a wide range of comorbid health conditions

Relative odds for migraine among comorbid conditions (vs migraine-free controls)¹



Comorbidities and chronification



~3% of individuals with episodic migraine progress to chronic migraine each year²



Risk factors for chronification include female sex, lifestyle factors (e.g. high caffeine consumption), ineffective treatment, medication overuse and untreated comorbidities³



Conversely, risk of nearly all comorbidities increases with headache frequency^{1,4}



Risk of sleep and psychiatric comorbidities also increases with pain intensity¹

CI, confidence interval; GI, gastrointestinal; OR, odds ratio; PAD, peripheral artery disease; TIA, transient ischemic attack.

Data from prospective web-based survey of US population samples with migraine (N=15,133). Data adjusted for sociodemographic characteristics.

1. Buse DC, et al. J Headache Pain 2020;21:23; 2. Dodick DW. Lancet 2018;391:1315-30; 3. Torres-Ferrús M, et al. J Headache Pain 2020;21:42; 4. Lipton RB, et al. Neurology 2019;93:e2224-36



Recognize: migraine in pregnancy

Recognize

Introduction



Recognize



Assess



Treat



Evaluate



Some patients experience migraine for the first time during pregnancy¹

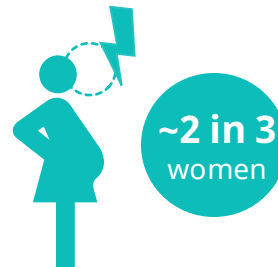
Migraine in pregnancy

Prevalence:



Frequency of migraine attacks tends to decrease during pregnancy; however, **~3–7%** of women experience **new-onset** migraine in pregnancy during the first trimester¹

Migraine attacks tend to **recur**, often within the first **6 months** after delivery¹



~2 in 3
women

Screen²

ACOG recommends careful screening of migraine during pregnancy²



Patients should be screened **pre-pregnancy** and at their **initial pregnancy visit** for:

- History of migraine
- Frequency of attacks
- Associated symptoms
- Intervention and/or medication history



The **appearance** or **worsening** of migraine during pregnancy warrants attention³

Migraine symptoms, when accompanied by **high blood pressure**, may increase the risk of developing vascular complications such as **preeclampsia**³

Introduction



Recognize



Assess



Treat



Evaluate



Recognize: patient understanding of migraine attack progression may help them anticipate and identify symptoms

Recognize

Identifying symptoms associated with the earliest stages of migraine can serve as warning signs and indicate a need to consult a HCP^{1,2}



Patients may report **prodromal symptoms** up to **48 hours before** the onset of headache¹

These may include:

hyperactivity

hypoactivity

depression

specific food cravings

yawning

fatigue

difficulty concentrating

neck stiffness / pain

blurred vision

sensitivity to light and sound



- Patients with migraine should consider maintaining a **headache diary** to **recognize their symptoms** and phases they experience before and after each headache²



Identifying these symptoms **early** can impact treatment initiation and is key to **reducing the severity** of or **prevention of headache**²

HCP, healthcare practitioner.

1. Headache Classification Committee of the International Headache Society. The International Classification of Headache Disorders, 3rd edition. Cephalalgia 2018;38:1–211; 2. American Migraine Foundation. The timeline of a migraine attack, 2018: <https://americanmigrainefoundation.org/resource-library/timeline-migraine-attack/>.



Recognize: perceived migraine triggers are unique and specific to each patient¹⁻³

Recognize

Introduction



Recognize



Assess



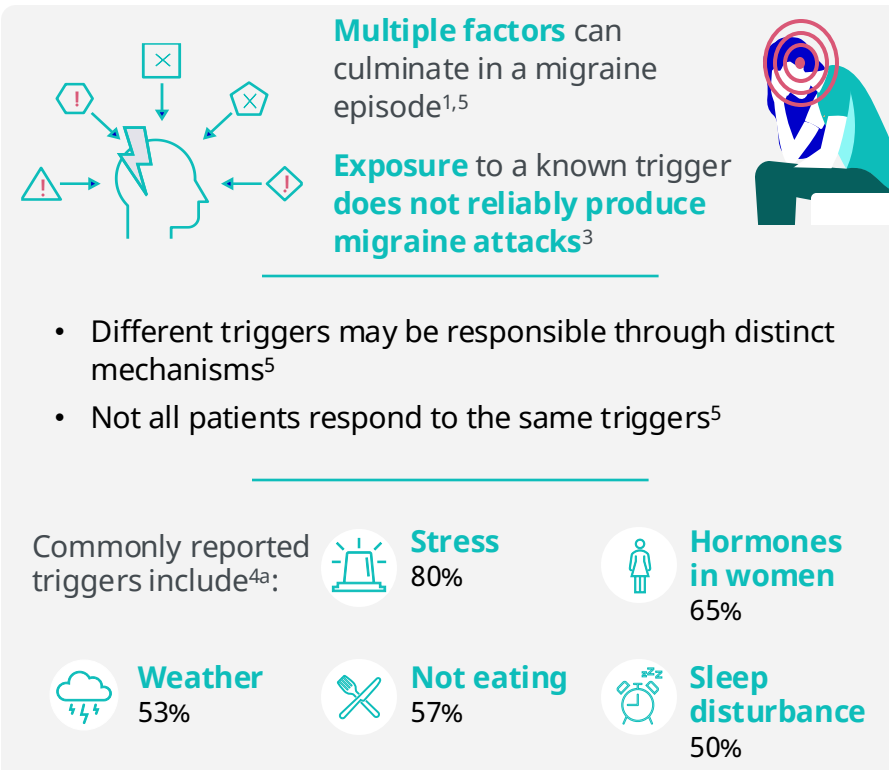
Treat



Evaluate



Patients report a variety of factors or triggers for migraine⁴



Patient management strategies for identification and mitigation of triggers¹

- While prodromal symptoms can make it difficult to identify triggers, patients should consider the following:



RECORD: Keeping a diary or calendar may help patients to identify triggers



ACT: Trigger avoidance is often unsuccessful and may disrupt participation in social and leisure activities. Patients should consider healthy lifestyle choices, improvement in diet and sleep



FIND OUT ABOUT FEAR: Worrying about the next attack may lead to patients avoiding or reducing their activities. This is an important predictor of migraine chronification and medication overuse, meaning identification and treatment are crucial



Identification of specific triggers may not always be possible; HCPs should encourage positive health behaviors and shared decision-making for a preventive approach⁶

^aBased on a retrospective study of 1750 patients with migraine. HCPs, healthcare providers

1. Marmura MJ. Curr Pain Headache Rep 2018;22:81; 2. Lipton RB, et al. Headache 2014;54:1661-9; 3. Martin VT and Behbehani MM. Med Clin North Am 2001;85:911-41; 4. Kelman L. Cephalalgia 2007;27:394-402; 5. Hoffmann J, et al. Ann Clin Transl Neurol 2015;2:22-8; 6. Sauro KM, Becker WJ. Headache 2009;49:1378-86.



Recognize: knowing when to suspect migraine is important

Recognize

Introduction



Recognize



Assess



Treat



Evaluate



Summary



Suspect migraine when there is the following:^{1,2}

- Recurrent headache of moderate to severe intensity
- Visual aura
- Family history of migraine



- Psychiatric, pain, sleep, cardiovascular, neurologic, and respiratory disorders are the most common comorbidities associated with migraine³



- Patients should be screened pre-pregnancy and at their initial pregnancy visit for history and frequency of attacks, associated symptoms, intervention and/or medication history⁴



- Patient identification of triggers and early symptoms may help to reduce the severity of, or in some cases, prevent migraine attack^{5,6}

1. Eigenbrodt AK, et al. Nat Rev Neurol 2021;17:501–14; 2. Headache Classification Committee of the International Headache Society. The International Classification of Headache Disorders, 3rd edition. Cephalalgia 2018;38:1–211; 3. Buse DC, et al. J Headache Pain 2020;21:23; 4. ACOG Committee on Clinical Practice Guidelines—Obstetrics. Obstet Gynecol 2022;139:944–72; 5. American Migraine Foundation. The timeline of a migraine attack, 2018: <https://americanmigrainefoundation.org/resource-library/timeline-migraine-attack/>; 6. Graf M, et al. Headache 2023;63:506–16.



Assess: patient history taking and physical examination as key first steps for diagnosis

Assess

Introduction



Recognize



Assess



Treat



Evaluate



Patient history



A thorough **headache history** is the **most** important tool in diagnosing **migraine**¹



To aid **accurate diagnosis**, it is necessary to ask patients a series of **questions** on **symptoms** experienced and **severity** of migraine attacks²

Physical examination

A physical examination can determine other factors that may exacerbate migraine and include **fundoscopy**, **palpation of head and neck**, **cardiovascular screening**, and **neurologic examination**²

Example questions when conducting a patient history of suspected migraine^{2,3}

 Pain pattern	<ul style="list-style-type: none"> • When and how do headaches begin? • Are they episodic or chronic? • How long do headaches last? • How frequent is headache pain? • Are there factors that trigger an attack? 	 Medical history	<ul style="list-style-type: none"> • Has there been any experience of depression, anxiety and/or sleep disorders? • Are there any comorbidities such as asthma, hypothyroidism, or hypertension?
 Associated symptoms	<ul style="list-style-type: none"> • Do other symptoms precede the attack? • Are there any symptoms that accompany the attack (e.g., nausea, photophobia, phonophobia)? 	 Treatment history	<ul style="list-style-type: none"> • What previous and current treatments have been taken and when? • To exclude medication overuse headache,^a what acute headache medications have been taken? • What non-headache specific treatments have been taken? • If treatment was discontinued, why?
 Nature of pain	<ul style="list-style-type: none"> • What is the location of the pain? • How severe is the pain? 	 Family and social history	<ul style="list-style-type: none"> • Is there a family history of migraine? • How have the headaches impacted your lifestyle? • Have there been any sleep disturbances?

^aMedication overuse: ≥15 days/month use of simple (non-opioid) analgesics for >3 months; ≥10 days/month use of opioids, triptans, ergotamines, or their combination analgesics for >3 months⁴

1. Martin VT, et al. Ann Med;2021;53:1979–1990; 2. Weatherall MW. Ther Adv Chronic Dis;2015;6:115–123; 3. Ravishankar K. Ann Indian Acad Neurol 2012;15:7–14; 4. Headache Classification Committee of the International Headache Society. The International Classification of Headache Disorders, 3rd edition. Cephalalgia 2018;38:1–211.



Assess: primary headache disorders

Assess

Introduction



Recognize



Assess



Treat



Evaluate



Differential diagnoses: characteristics of primary headache disorders

Migraine¹

Headache characteristics:



Usually
unilateral in
location



**4-72
hours**

Moderate
or severe
intensity



Usually
pulsating pain



Accompanying symptoms:



Photophobia, phonophobia,
nausea, vomiting

Often aggravated by routine physical activity

Tension-type headache¹

Headache characteristics:



Typically bilateral or
circumferential



**Hours
to days^a**

Mild or
moderate
intensity



Usually pressing or
tightening pain



Accompanying symptoms:



Often none; sometimes
photophobia or phonophobia (but
not both); sometimes mild nausea
in chronic tension-type headache

Not aggravated by routine physical activity

Cluster headache¹

Headache characteristics:



Strictly unilateral
and orbital,
supraorbital,
and/or temporal



**15-180
minutes**

Severe
or very
severe



Overwhelming
pain



Accompanying symptoms:



Ipsilateral to the headache:
cranial autonomic symptoms, e.g.,
conjunctival injection,^b lacrimation
(tearing) and nasal congestion

Associated with restlessness or agitation

^aOr unremitting; ^bIncreased redness of the eye due to enlargement of conjunctival vessels.²

1. Eigenbrodt AK, et al. Nat Rev Neurol 2021;17:501-14; 2. Park JK, et al. Invest Ophthalmol Vis Sci 2013;54:5249-57.



Assess: apply ICHD-3 diagnostic criteria to determine migraine category

Assess

Introduction



Recognize



Assess



Treat



Evaluate



A patient's clinical history should be evaluated against diagnostic criteria to aid migraine diagnosis

The ICHD-3 provides diagnostic criteria for three main categories of migraine¹

Migraine categories				
<p>Migraine without aura History of ≥ 5 attacks</p>	<p>Headache attacks lasting 4–72 hours <i>(when untreated or unsuccessfully treated)</i></p>	<p>With ≥ 2 of the following characteristics:</p> <ul style="list-style-type: none"> • Unilateral location • Pulsating quality • Moderate or severe pain intensity • Aggravation by or causing avoidance of routine physical activity^a 	<p>Accompanied by ≥ 1 of the following symptoms:</p> <ul style="list-style-type: none"> • Nausea and/or vomiting • Photophobia AND phonophobia 	<p>Symptoms not better accounted for by another ICHD-3 diagnosis</p>
<p>Migraine with aura^b History of ≥ 2 attacks</p>	<p>Recurrent migraine aura symptoms lasting 5–60 minutes that accompany or are followed within 60 minutes, by headache which may not fulfil criteria for migraine with aura</p>	<p>With ≥ 3 of the following characteristics:</p> <ul style="list-style-type: none"> • ≥ 1 aura symptom that spreads gradually over ≥ 5 minutes • ≥ 2 aura symptoms that occur in succession • Individual aura symptoms of 5–60 minutes • ≥ 1 unilateral^c aura symptom • ≥ 1 positive aura symptom e.g., scintillations/pins and needles • Aura with/preceded by headache within 60 min 	<p>≥ 1 of the following fully reversible aura symptoms:</p> <ul style="list-style-type: none"> • Visual • Sensory • Speech and/or language • Motor • Brainstem • Retinal 	<p>Symptoms not better accounted for by another ICHD-3 diagnosis</p> <p>Transient ischemic attack excluded</p>
<p>Chronic migraine Attacks on ≥ 15 days/month for >3 months</p>	<p>≥ 5 headache attacks meeting criteria for migraine with or without aura</p>	<p>With ≥ 1 of the following characteristics on ≥ 8 days/month for >3 months:</p> <ul style="list-style-type: none"> • Features of migraine without aura • Features of migraine with aura • Believed by the patient to be migraine at onset and relieved by a triptan or ergot derivative 		<p>Symptoms not better accounted for by another ICHD-3 diagnosis</p>

ICHD-3, International Classification of Headache Disorders (3rd edition).

^aE.g., walking or climbing stairs; ^bMigraine with aura can be subcategorized as typical aura, brainstem aura (≥ 2 brainstem symptoms, e.g., dysarthria, vertigo, tinnitus, hypacusis, diplopia, ataxia, decreased consciousness), hemiplegic migraine (motor weakness) or retinal migraine (monocular visual disturbances); ^cAphasia is always unilateral while dysarthria may or may not be.

Headache Classification Committee of the International Headache Society. The International Classification of Headache Disorders, 3rd edition. Cephalalgia 2018;38:1–211.



Assess: assessing secondary headaches

Assess

Introduction



Recognize



Assess



Treat



Evaluate



Secondary headaches are those in which the headache is a symptom of another disorder recognized as a potential underlying cause¹; patients with suspected secondary headache require a diagnostic workup that depends on the red flag identified and suspected underlying disease²

SNOOP4 and **SNNOOP10+** are **red flag** detection tools reported in the literature that are used to identify potentially life-threatening secondary headaches²⁻⁴

SNOOP4				
Red flag sign or symptom	History and examination findings	Secondary headache causes		Diagnostic workup
S ystemic	<ul style="list-style-type: none"> Signs of infection (e.g., fever, chills, weight loss) Secondary risk factors include HIV or systemic cancer⁵ 	<ul style="list-style-type: none"> Infection Malignancy 	<ul style="list-style-type: none"> Rheumatic disease Giant cell arteritis 	<ul style="list-style-type: none"> Neuroimaging Lumbar puncture
N eurologic	<ul style="list-style-type: none"> Abnormal neurologic examination^a Change in behavior or personality 	<ul style="list-style-type: none"> Malignancy Inflammatory disorder 	<ul style="list-style-type: none"> Infection 	
O nset (sudden)	<ul style="list-style-type: none"> Headache reaching peak intensity in <1 minute (thunderclap, often triggered by subarachnoid hemorrhage or RCVS^{6,b}) 	<ul style="list-style-type: none"> Subarachnoid hemorrhage Stroke 	<ul style="list-style-type: none"> RCVS 	<ul style="list-style-type: none"> Head CT Lumbar puncture (if CT negative)
O lder age at onset	<ul style="list-style-type: none"> New onset headache at age >50 years 	<ul style="list-style-type: none"> Malignancy Infection 	<ul style="list-style-type: none"> Giant cell arteritis 	<ul style="list-style-type: none"> MRI
P attern change	<ul style="list-style-type: none"> Change in headache pattern or characteristics Progressive headache^c 	<ul style="list-style-type: none"> Malignancy Inflammatory or vascular disorder 		
P recipitated by Valsalva maneuver	<ul style="list-style-type: none"> Headache precipitated by Valsalva maneuver, sneezing, coughing, or exercise 	<ul style="list-style-type: none"> Chiari malformation type 1 Posterior fossa lesions Intracranial hypertension or hypotension 	<ul style="list-style-type: none"> Malignancy Arachnoid cysts Subdural hematoma 	<ul style="list-style-type: none"> Neuroimaging
P ostural	<ul style="list-style-type: none"> Headache precipitated or aggravated by postural change 	<ul style="list-style-type: none"> Intracranial hypertension Intracranial hypotension 		<ul style="list-style-type: none"> Neuroimaging Lumbar puncture MRI with gadolinium^d
P apilledema	<ul style="list-style-type: none"> Papilledema, visual obscurations, diplopia, or field defects 	<ul style="list-style-type: none"> Intracranial hypertension Inflammatory disorder 	<ul style="list-style-type: none"> Malignancy 	<ul style="list-style-type: none"> Thorough fundoscopic examination

^aE.g. confusion, impaired consciousness.^{2,7} ^bFactors associated with RCVS include: sexual activity, postpartum state, vasoactive drugs, autoimmune disorders, temperature differences, air travel, and COVID-19 infection.⁸ ^cLoss of headache-free periods. ^dTo rule out dural enhancement with suspected CSF leak.²

CT, computed tomography; HIV, human immunodeficiency virus; MRI, magnetic resonance imaging; RCVS, reversible cerebral vasoconstriction syndrome; SNOOP, systemic symptoms, neurological signs, onset, older age at onset, and prior medical history. 1. Ravishanker K. Headache 2016;56:1685-1697; 2. Martin VT, et al. Ann Med;2021;53:1979-1990; 3. Do TP, et al. Neurology 2019;92:134-144; 4. Dodick DW. Semin Neurol 2010;30:74-81; 5. Smith JH. Practical Neurology, March/April 2018; 40-49; 6. Schwedt TJ. Continuum (Minneap Minn) 2015;21:1058-71; 7. Micieli A and Kingston W. Front Public Health 2019;7:52; 8. Ribas MZ, et al. Egypt J Neurol Psychiatric Neurosurg 2023;59:5.





Assess: assessing secondary headaches

Assess

Introduction



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Treat

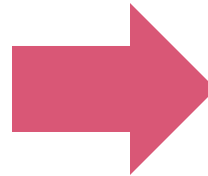


Evaluate



SNNOOP10 retains the red flags of SNOOP4 and adds additional signs or symptoms of related secondary headaches^{1,2}

SNOOP4
Red flag sign or symptom
S ystemic
N eurologic
O nset (sudden)
O lder age at onset
P attern change
P recipitated by Valsalva maneuver
P ostural
P apilledema



SNNOOP10	
Additional red flag sign or symptom	Notes
N eoplasm in history	Relevant accompanying symptoms include emesis, headache duration ≤ 10 weeks, atypical headache pattern, pulsating quality and moderate to severe intensity, gait instability, and extensor plantar response
P rogressive headache and atypical presentations	Not commonly described and defined in the literature; may be the only signs of serious underlying pathology
P regnancy or puerperium	Headaches during this period have a higher risk of severe pathology due to physiologic changes and interventions
P ainful eye with autonomic features	Can be due to a structural lesion; neuroimaging is recommended
P osttraumatic onset of headache	Headache occurring directly in relation to a trauma is a red flag
P athology of the immune system (such as HIV)	The risk of severe pathology is dependent on the degree of immunosuppression
P ainkiller overuse or new drug at onset of headache	Medication overuse is the most common cause of a secondary headache; onset of headache due to a new drug can be a sign of incompatibility with the given drug

1. Dodick DW. Semin Neurol 2010;30:74–81; 2. Do TP, et al. Neurology 2019;92:134–144.



Assess: when to consider referring for neuroimaging

Assess

Neuroimaging is occasionally required to confirm or reject suspected secondary headache¹

AHS guidelines recommend *against* neuroimaging in patients with headaches consistent with migraine who have a normal neurologic examination and there are no atypical features or **red flags** present²

Strong recommendation, high quality evidence



AHS guidelines note that **neuroimaging may be considered** for presumed migraine for the following reasons^{2a}:

Strong recommendation, low quality evidence

- Unusual, prolonged, or persistent aura
- Increasing frequency, severity, or change in migraine clinical features
- First or worst migraine
- Migraine with brainstem aura
- Confusional migraine
- Hemiplegic migraine
- Late-life migrainous accompaniments
- Migraine aura without headache
- Side-locked migraine
- Post-traumatic migraine

MRI is preferred to CT as it provides a higher resolution and avoids harmful exposure to ionizing radiation¹

- **MRI has its limitations** as it can also **detect clinically insignificant** findings (such as white matter lesions, arachnoid cysts, and meningiomas) which can alarm patients and lead to further unnecessary evaluations

^aMost of these are consensus based with little or no literature support?

AHS, American headache society; CT, computed tomography; MRI, magnetic resonance imaging.

1. Eigenbrodt AK, et al. Nat Rev Neurol 2021;17:501–14; 2. Evans RW, et al. Headache 2020;60:318–336.



Assess: assessing migraine in pregnancy

Assess

Introduction



Recognize



Assess



Treat



Evaluate



Pregnancy can increase the risk of some secondary headache disorders¹

ACOG recommends careful screening and assessments of headache during pregnancy²



Secondary headaches should be **carefully evaluated**²:

- History
- Physical examination
- Other tests (e.g., imaging or spinal fluid examination), as appropriate

Strong recommendation;
low-quality evidence



Features of secondary headache that warrant prompt attention²:

- “Thunderclap” headache
- Rapid onset
- Visual changes
- High blood pressure
- Vomiting
- Fever
- Neurologic deficits or altered consciousness

Good practice point



Primary headache disorders typically improve during pregnancy; therefore, HCPs should have¹:



Lower threshold for further investigation

Higher clinical suspicion for secondary headache (particularly in the third trimester when vascular risk increases)

ACOG, American College of Obstetricians and Gynecologists; AHS, American Headache Society; HCP, healthcare practitioners.

^aFor additional information, please refer to the full guideline.

1. Negro A, et al. J Headache Pain 2017; 18:106; 2. Adapted from: ACOG Committee on Clinical Practice Guidelines—Obstetrics. Obstet Gynecol 2022; 139:944–72;

Introduction



Recognize



Assess



Treat



Evaluate



Assess: diagnostic screening tools aid migraine diagnosis

Assess

Various screening tools are available to aid in the diagnosis of migraine



Screening tools



Headache diary



Can assist with re-evaluation of diagnosis at follow-up



Headache calendar



Records temporal occurrence of headaches and related events at follow-up



ID Migraine™ screener



Identifies individuals who are likely to have migraine based on their answers to three questions regarding headache-associated nausea, photophobia, and disability



Migraine Disability Assessment (MIDAS)



A brief, self-administered questionnaire designed to quantify headache-related disability over a 3-month period; the MIDAS score has moderately high test-retest reliability in headache sufferers and is correlated with clinical judgment regarding the need for medical care



Assess: diagnostic screening tools aid migraine diagnosis

Assess

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Evaluate



Headache diary can assist with re-evaluation of diagnosis at follow-ups¹⁻³



Name (first name(s)): _____ (family name) _____

Record in this diary for 1 week. Start date: _____ / _____ / _____ Finish date: _____ / _____ / _____

Please read **the instructions** carefully. Then complete the following **every** evening at bedtime (where indicated, tick the most appropriate box or circle the most appropriate answer).

Please remember to bring this diary to your next appointment

1. Date (day/month)	/	/	/	/	/	/	/
2. Did you have a headache today? (circle one) (If no, the diary is completed today)	yes / no	yes / no	yes / no	yes / no	yes / no	yes / no	yes / no
3. At what time today did your headache start? (hours and minutes) (if It began yesterday and was still there today, draw a left arrow ←)							
4. When did it go away? (hours and minutes) (if it is still there when you go to bed for the night, draw a right arrow →)							
5. On which side is/was the headache (circle one)	right left both	right left both	right left both	right left both	right left both	right left both	right left both

1. Eigenbrodt AK. Nat Rev Neurol. 2021;17(8):501-514. 2. Stewart WF et al. Pain. 2000;88(1):41-52. 3. European Headache Federation. Accessed June 20, 2024. https://static-content.springer.com/esm/art%3A10.1186%2Fs10194-018-0899-2/MediaObjects/10194_2018_899_MOESM16_ESM.pdf.





Assess: diagnostic screening tools aid migraine diagnosis

Assess

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Headache calendar records temporal occurrence of headaches and related events at follow-ups^{1,2}



YEAR:

NAME:

DOB:

Symptomatic drugs:

Daily prophylactic drugs:

Hormones:

Other regular medication:

	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31
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1. Eigenbrodt AK. Nat Rev Neurol. 2021;17(8):501-514. 2. European Headache Federation. Accessed June 20, 2024. https://static-content.springer.com/esm/art%3A10.1186%2Fs10194-018-0899-2/MediaObjects/10194_2018_899_MOESM17_ESM.pdf.



Assess: diagnostic screening tools aid migraine diagnosis

Assess

Introduction



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The ID Migraine™ screener can help increase the recognition of migraine in previously undiagnosed individuals^{1,2}

The 3-item **ID Migraine screener** is a validated and reliable tool that may help rapidly diagnose migraine in the primary care setting:

	1. Has a headache limited your activities for a day or more in the last 3 months?
	2. Are you nauseated or sick to your stomach when you have a headache?
	3. Does light bother you when you have a headache?
	Total (Questions 3)

If the patient responds affirmatively to 2 or all 3 items, a migraine diagnosis is likely^a

^aTesting positive ≥2 times yields a sensitivity of 0.81 (95% CI, 0.77–0.85).

1. Lipton RB et al. Neurology. 2003;61(3):375-382. 2. de Mattos ACM, et al. Arq Neuro-Psiquiatr. 2017;75 (7):446-450.



Assess: diagnostic screening tools aid migraine diagnosis

Assess

Introduction



Recognize



Assess



Treat



Evaluate



The Migraine Disability Assessment (MIDAS)^{1,2}

The **MIDAS** (Migraine Disability Assessment) questionnaire was put together to help you measure the impact your headaches have on your life.

INSTRUCTIONS: Please answer the following questions about ALL of the headaches you have had in the last 3 months. Select your answer in the box next to each question. Select zero if you did not have the activity in the last 3 months. Please take the completed form to your healthcare professional.

1. On how many days in the last 3 months did you miss work or school because of your headaches?
2. How many days in the last 3 months was your productivity at work or school reduced by half or more because of your headaches? (Do not include days you counted in question 1 where you missed work or school)
3. On how many days in the last 3 months did you not do household work (such as housework, home repairs and maintenance, shopping, caring for children and relatives) because of your headaches?
4. How many days in the last 3 months was your productivity in household work reduced by half or more because of your headaches? (Do not include days you counted in question 3 where you did not do household work.)
5. On how many days in the last 3 months did you miss family, social or leisure activities because of your headaches?

Total (Questions 1-5)

What your Physician will need to know about your headache:

- A. On how many days in the last 3 months did you have a headache? (If a headache lasted more than 1 day, count each day.)
- B. On a scale of 0-10, on average how painful were these headaches? (where 0=no pain at all, and 10=pain as bad as it can be.)

MIDAS Grade	Definition	MIDAS Score
I	Little or No Disability	0-5
II	Mild Disability	6-10
III	Moderate Disability	11-20
IV	Severe Disability	21+

1. Stewart WF et al. *Pain*. 2000;88(1):41-52. 2. Lipton RB, et al. *Handb Clin Neurol*. 2010;97:23-32



Introduction



Recognize



Assess



Treat



Evaluate



Assess: referring to a specialist

Assess

Reasons for specialist referral for patients with migraine, TTH, or MOH:



A migraine diagnosis cannot be confirmed



Diagnosis of any of the following, which are best managed by specialists:

- migraine with aura including motor weakness
- chronic migraine
- cluster headache
- trigeminal neuralgia
- persistent idiopathic facial pain



Serious secondary headache or serious etiology are suspected



Unusual migraine aura, especially:

- prolonged aura (duration >1 hour)
- aura featuring brainstem symptoms and/or motor weakness
- new aura without headache in a patient >50 years and in the absence of a prior history of migraine



• Persistent management failure



• Comorbid disorders requiring specialist management



Assess: patient centricity and education have key roles in the diagnosis of migraine

Assess

Introduction



Recognize



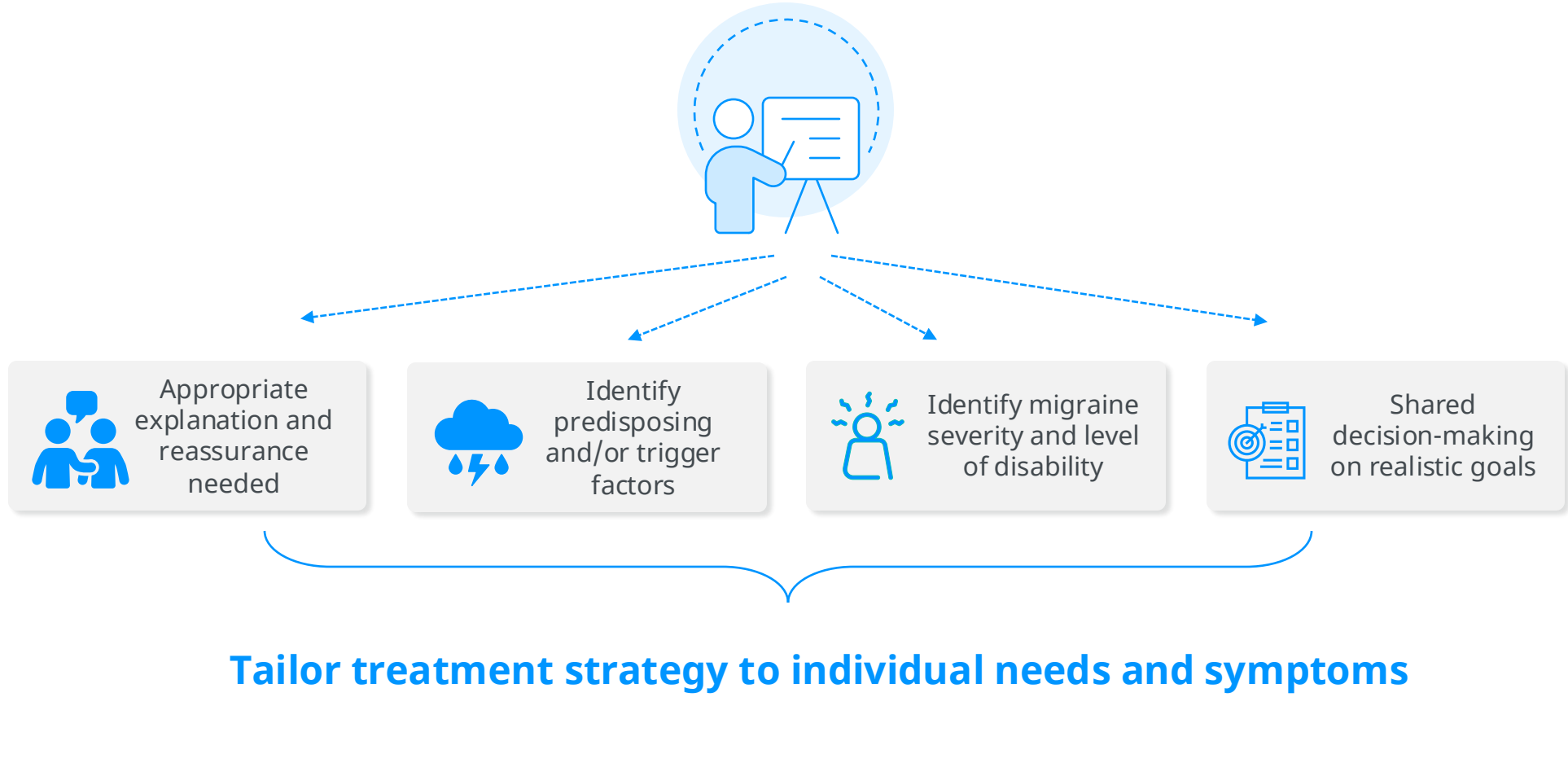
Assess



Treat



Evaluate





Assess: diagnosis and assessment of migraine

Assess

Introduction



Recognize



Assess



Treat



Evaluate

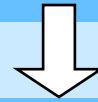


Summary

Key recommendations for diagnosing migraine



History and physical exam



Exclude secondary headache

Screen for presence of red flags (SNOOP4 or SNNOOP10)

−

+

→ Secondary headache



Identify primary headache disorder

ID Migraine™ screen

−

+

→ If necessary, refer to headache specialist



Diagnose primary headache disorder such as tension-type or cluster headaches using ICHD-3 criteria

Diagnose migraine



Introduction



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Treat: an individualized treatment management plan should be implemented following diagnosis

Treat

Summary of Recommendations from the AHS 2021 Consensus Statement



Acute migraine treatment^{1,2}

- For patients with a confirmed diagnosis of migraine



Preventive migraine treatment¹

- For patients whose attacks significantly interfere with daily routines despite acute treatment
- For those who have frequent attacks, intolerance or contraindication(s) to acute treatments, or failure or overuse of acute treatments
- Based on patient preference



Migraine treatment in special populations^{3,4}

- Pregnancy
- Breastfeeding

AHS, American Headache Society.

1. Ailani J, et al. Headache 2021;61:1021–39; 2. Headache Classification Committee of the International Headache Society. Cephalalgia 2018;38:1–211; 3. Lucas S. Obstet Gynecol 2019;134:211; 4. ACOG Committee on Clinical Practice Guidelines–Obstetrics. Obstet Gynecol 2022;139:944–72.



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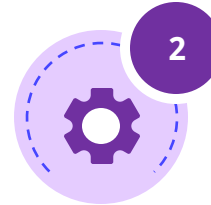
Treat: **acute** migraine treatment aims to stop attacks, or reduce headache severity and other associated symptoms

Treat

Summary of goals for **acute** migraine treatment from the **2021 AHS Consensus Statement:**



Provide rapid symptomatic relief without recurrence



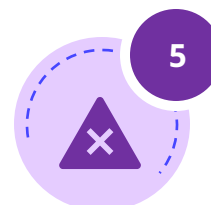
Restore function



Minimize the need for rescue medications or repeat dosing



Optimize self-care and reduce healthcare utilization



Minimize adverse events



Reduce overall treatment costs

Acute treatments



AHS, American Headache Society.

The AHS Consensus Statement provides timely recommendations for clinicians and is not intended to be, and should not be understood or applied as, a Clinical Practice Guideline. Adapted from: Ailani J, et al. Headache 2021;61:1021-39.





Treat: **acute** migraine treatment aims to stop attacks, or reduce headache severity and other associated symptoms

Treat

Introduction



Recognize



Assess



Treat



Evaluate



Migraine-Specific Treatments¹

- Triptans
- Ergotamine derivatives
- Ditans
- CGRP receptor antagonists (gepants)
- Neuromodulation devices

Nonspecific Treatments¹

- NSAIDs^a
- Acetaminophen
- Others (combination analgesics)

FDA Approved Acute Migraine Treatments:

Ergotamine
Derivative³

1946

Triptan²

1992

Ditan⁴ Gepant⁵

2019

2019

Year of approval of initial product within class

^aCelecoxib oral solution is approved as a migraine-specific treatment¹

CGRP, calcitonin gene-related peptide; FDA, Food and Drug administration; NSAIDs, non-steroidal anti-inflammatory drugs.

1. Ailani J et al. *Headache*. 2021;61:1021-1039. 2. Sumatriptan. Package insert. GlaxoSmithKline. 3. FDA. https://www.accessdata.fda.gov/drugsatfda_docs/nda/97/020148_migranal_toc.cfm. Accessed April 6, 2022. 4. FDA

https://www.accessdata.fda.gov/drugsatfda_docs/nda/2019/211280Orig1s000TOC.cfm. Accessed April 6, 2022. 5. FDA <https://www.accessdata.fda.gov/scripts/cder/daf/index.cfm?event=overview.process&ApplNo=211765>. Accessed April 6, 2022.

AHS Consensus





Treat: **acute** migraine treatment aims to stop attacks, or reduce headache severity and other associated symptoms

Treat

Introduction



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Evaluate



Acute treatment of migraine



The AHS 2021 Consensus Statement recommends that all patients with a confirmed diagnosis of migraine should be offered acute pharmacological and/or nonpharmacological treatment¹

The AHS 2021 Consensus Statement¹

Nonpharmacologic interventions include counseling patients on the benefits of:

- Proper nutrition
- Regular exercise
- Adequate hydration
- Proper sleep
- Stress management
- Maintaining a migraine diary

Mild-to-moderate attacks

NSAIDs, nonopioid analgesics, acetaminophen, or caffeinated analgesic combinations

Moderate-to-severe attacks

OR

Mild-to-moderate attacks that respond poorly to nonspecific medications

Migraine-specific agents (triptans, ergotamine derivatives, gepants^a, ditans)

Criteria for initiating gepants, ditans or neuromodulatory devices for acute treatment¹:

- Contraindication or inability to tolerate triptans or
- Inadequate response to ≥ 2 oral triptans

^aAmong the available gepants: rimegepant, ubrogepant, and zavegepant are indicated for treatment of acute migraine.
NSAID, nonsteroidal anti-inflammatory drug.
1. Ailani J, et al. Headache 2021;61:1021-39.



Approval Timeline





Treat: preventive migraine treatment aims to reduce the frequency, severity, and duration of attacks

Treat

Introduction



Recognize



Assess



Treat



Evaluate



2021 AHS Consensus Statement on Preventive Treatments for Migraine

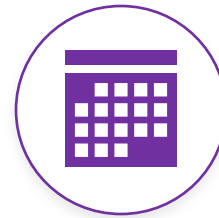
Patients with migraine should be considered for preventive treatment if:



Attacks significantly interfere with patient daily routines despite acute treatment



Patient has contraindications to and/or failures with acute treatments



Patient experiences ≥ 2 headache days per month based on the degree of disability^a



Patient overuses acute treatments



Patient experiences tolerability issues with acute treatments



Patient preference for prevention

Prevention should be	Headache days/month	Degree of disability
Offered	6 or more 4 or more 3 or more	None Some Severe
Considered	4 or 5 3 2	None Some Severe

Overuse defined as follows:
Ten or more days per month for ergotamine derivatives, triptans, opioids, combination analgesics, and a combination of drugs from different classes that are not individually overused
Fifteen or more days per month for nonopioid analgesics, acetaminophen, and NSAIDs

^aAs measured by the MIDAS, Migraine Physical Function Impact Diary, or Headache Impact Test.
AE, adverse event; AHS, American Headache Society; HRQOL, health-related quality of life.
Ailani J et al. *Headache*. 2021;61:1021-1039.

Introduction



Recognize



Assess



Treat



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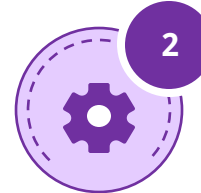
Treat: preventive migraine treatment aims to reduce the frequency, severity, and duration of attacks

Treat

Summary of goals for **preventive** migraine treatment from the **2021 AHS Consensus Statement:**



1
Reduce attack frequency, severity, duration, and disability



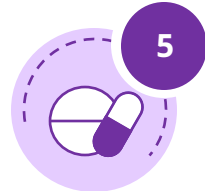
2
Improve function and reduce disability



3
Improve responsiveness to and avoid escalation in use of acute treatment



4
Improve health-related quality of life



5
Reduce reliance on unwanted, poorly tolerated, or ineffective acute treatments



6
Optimize self-care and enhance sense of personal control



7
Reduce psychological symptoms and headache-related distress



8
Reduce overall treatment costs

Preventive treatments +

AHS, American Headache Society.

The AHS Consensus Statement provides timely recommendations for clinicians and is not intended to be, and should not be understood or applied as, a Clinical Practice Guideline.
Adapted from: Ailani J, et al. Headache 2021;61:1021-39.





Treat: preventive migraine treatment aims to reduce the frequency, severity, and duration of attacks

Treat

Introduction



Recognize



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Evaluate



Interventions with evidence of efficacy in migraine prevention^{1a}

- Anti-CGRP monoclonal antibodies
- Anticonvulsants
- Beta-Blockers
- CGRP receptor antagonists (gepants)
- Neuromuscular blocking agent

Non-pharmacological interventions¹

- Non-invasive neuromodulation
- Nutraceuticals
- Biobehavioral therapies
- Lifestyle modifications

FDA Approved Preventive Migraine Treatments:



Year of approval of initial product within class

AHS Consensus



CGRP, calcitonin gene-related peptide; mAb, monoclonal antibody; SSRI, selective serotonin reuptake inhibitor.

^aThe AHS 2021 Consensus Statement identifies additional agent classes with evidence of efficacy in migraine prevention which do not possess FDA approval for that use.¹

1. Ailani J et al. Headache. 2021;61:1021-1039. 2. FDA. https://www.accessdata.fda.gov/drugsatfda_docs/label/2011/016418s080,016762s017,017683s008lbl.pdf. Accessed April 9, 2025. 3. FDA.

<https://www.accessdata.fda.gov/scripts/cder/daf/index.cfm?event=overview.process&ApplNo=018723>. Accessed May 6, 2022. 4. FDA. https://www.accessdata.fda.gov/drugsatfda_docs/bla/2010/103000orig1s5215.pdf. Accessed May 6, 2022. 5. FDA. <https://www.fda.gov/news-events/press-announcements/fda-approves-novel-preventive-treatment-migraine>. Accessed May 6, 2022. 6. FDA. <https://www.accessdata.fda.gov/scripts/cder/daf/index.cfm?event=overview.process&varApplNo=212728>. Accessed May 6, 2022.





Treat: **preventive** migraine treatment aims to reduce the frequency, severity, and duration of attacks

Treat

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Preventive treatment of migraine



The AHS 2021 Consensus Statement recommends that preventive treatment should be considered for patients whose attacks significantly interfere with daily routines despite acute treatment; for those who have frequent attacks, intolerance or contraindication(s) to acute treatments, or failure or overuse of acute treatments; or based on patient preference^{1a}

The AHS 2021 Consensus Statement¹ and the 2024 Updated Position Statement²



Use evidence-based treatments^{1,2}

2024 Update: CGRP-targeting migraine therapies are a first-line option for migraine prevention. Initiation of these therapies should not require trial and failure of non-specific migraine preventive medication approaches.²

Classes of agents with an FDA-approved indication for preventive treatment of migraine^b include:

Oral treatments: anticonvulsants, beta-blockers, gepants

Intramuscular injection: neuromuscular blocking agent

Subcutaneous injection: anti-CGRP monoclonal antibodies

Intravenous infusion: anti-CGRP monoclonal antibodies



Allow an adequate trial before switching¹

Oral treatments:
≥8 weeks at target therapeutic dose or usual effective dose

Neuromuscular blocking agent:
After ≥2 quarterly injections (6 months)

Anti-CGRP mAbs:
≥3 months (monthly administration) or ≥6 months (quarterly administration)



Approval Timeline

CGRP, calcitonin gene-related peptide; HIT, Headache Impact Test; IV, intravenous; mAb, monoclonal antibody; MIDAS, Migraine Disability Assessment; MHD, monthly headache day; MMD, monthly migraine day.

^aOveruse defined as use of ergotamine derivatives, triptans, opioids or combination analgesics on ≥10 days/month for >3 months, or use of nonopioid analgesics, nonsteroidal anti-inflammatory drugs or simple analgesics on ≥15 days/month for >3 months. "Frequent attacks" includes ≥3 monthly headache days with severe disability, ≥4 monthly headache days with some disability or ≥6 monthly headache days without disability; ^bThe AHS 2021 Consensus Statement identifies additional agent classes with evidence of efficacy in migraine prevention which do not possess FDA approval for that use!

1. Ailani J, et al. Headache 2021;61:1021-39; 2. Charles AC, et al. Headache 2024; 64:333-41.





Treat: unique considerations for special populations experiencing migraine

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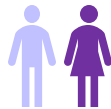


Evaluate



Migraine affects women during key stages in their lifecycle

Migraine is more common and disabling in women than in men



~3× more common
in women than
men^{1,2}

The largest prevalence
difference occurs at **30.2
years of age**³

Migraine is the **leading cause of disability** among women of
reproductive age (15–49 years) worldwide⁶

Compared with
men with
migraine,
women with
migraine
experience:^{4,5}



More frequent headache

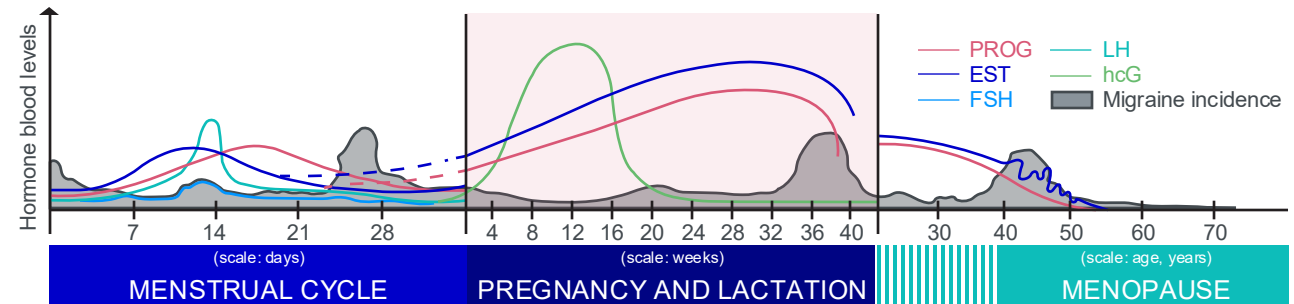
More intense headache

Higher risk of **chronification**

Greater headache-related **disability**

Migraine can be triggered by fluctuations in hormones throughout a woman's lifetime⁷

Incidence of migraine rises steeply at
puberty, peaks during the reproductive
years and subsides after menopause⁸



**Menstrually-related
migraine**

1. Andreou AP, et al. J Headache Pain. 2019;20:117:1–17; 2. Dodick DW. Migraine. Lancet. 2018;391:1315–30; 3. Victor TW, et al. Cephalalgia. 2010;30:1065–72; 4. Buse DC, et al. Headache. 2013; 53(8):1278–99; 5. Labastida-Ramirez A, et al. Cephalalgia. 2019;39:435–44; 6. Steiner TJ, et al. J Headache Pain. 2020; 21(1):4–7; 7. Sacco S, et al. J Headache Pain. 2012;13:177–189; 8. Krause DN, et al. Nat Rev Neurol 2021;17:621–33.



Treat: unique considerations for special populations experiencing migraine

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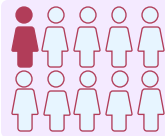


Evaluate



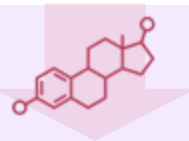
Menstrually-related migraine

Estrogen withdrawal hypothesis: drop in estrogen during menses is thought to trigger migraine attacks



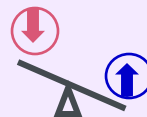
10% of women with migraine report migraine onset **at menarche**¹

Among women who have **migraine without aura**, the greatest risk of an attack is around **menses**²

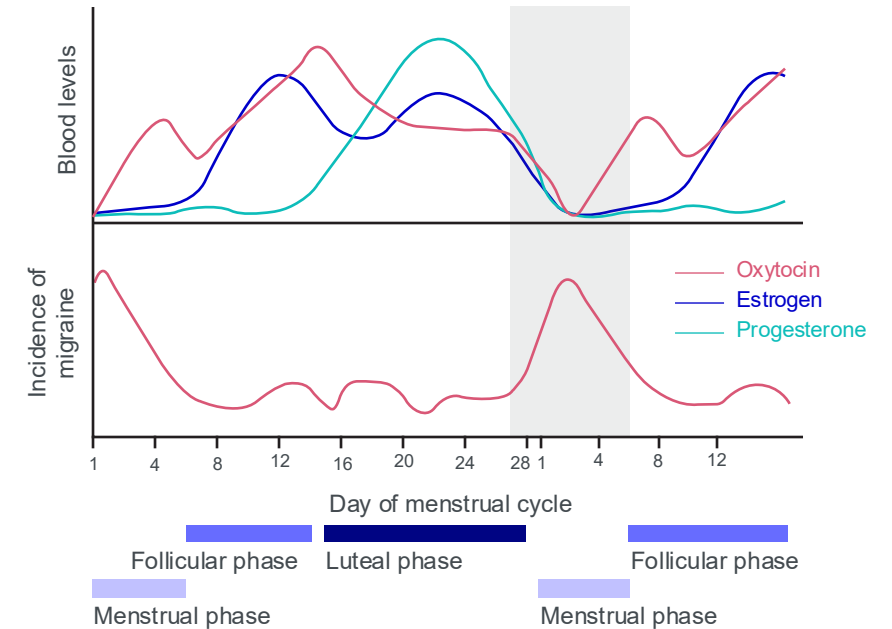


Estrogen withdrawal is thought to trigger the attacks, though the drop in oxytocin may also contribute²

Sharp decline in estrogen **shifts the balance toward a pro-migraine state** through increased CGRP signaling²



Fluctuations in the incidence of migraine and hormone blood levels over the menstrual cycle²



There is currently no specific treatment for menstrually-related migraine³

Figure adapted from Krause D, et al. 2021.
CGRP, calcitonin gene-related peptide.

1. Victor TW, et al. Cephalalgia 2010;30:1065-72; 2. Krause DN, et al. Nat Rev Neurol 2021;17:621-33; 3. Determan M. Pharmacy Times July 28, 2023.

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Treat: an individualized treatment management plan should be implemented following diagnosis

Treat

Summary of Recommendations from the 2021 AHS Consensus Statement



Acute treatment of migraine¹

- Mild-to-moderate attacks:
 - NSAIDs
 - Acetaminophen
 - Nonopioid analgesics
 - Caffeinated analgesic combinations
- Moderate-to-severe attacks or mild-to-moderate attacks that respond poorly to nonspecific medications:
 - Migraine-specific agents (triptans, ergotamine derivatives, gepants, ditans)



Preventive treatment of migraine^{1,2,a}

- Oral:
 - Anticonvulsants
 - Beta-blockers
 - Gepants
- Intramuscular:
 - Neuromuscular blocking agent^b
- Subcutaneous injection
 - Anti-CGRP mAbs
- Intravenous infusion
 - Anti-CGRP mAbs

^aThe AHS 2021 Consensus Statement identifies additional agent classes with evidence of efficacy in migraine prevention which do not possess FDA approval for that use.¹ ^bFor chronic migraine only. AHS, American Headache Society; CGRP, calcitonin gene-related peptide; mAbs, monoclonal antibodies; NSAID, nonsteroidal anti-inflammatory drug.

1. Ailani J, et al. Headache 2021; 61: 1021-39; 2. Charles AC, et al. Headache 2024; 64:333-41.



Evaluate: appraisal of treatment plans is important to ensure ongoing efficacy and safety

Evaluate

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Recommendations for the management of migraine from the EHF/EAN Consensus statement



Clinical management & follow-up



Evaluating response

- Assess effectiveness and tolerability regularly^a
- **Suboptimal outcomes:** Review diagnosis, treatment strategy, dosing and adherence
- **Treatment failure:** Conclude only after thorough review of underlying reasons
- **Specialist referral:** Only if diagnostically challenging, difficult to treat or complicated by comorbidities



Managing complications

- Discourage or stop medication overuse
- **MOH:** Withdraw overused medications (ideally abruptly, except in case of opioids)
- **Chronic migraine:** Specialist referral and use of preventive treatment indicated



Recognizing comorbidities

- Identify comorbid conditions
- Select migraine drugs and adjust dose according to comorbidities
- Alleviate comorbidities if possible



Long-term follow-up

- **Primary care should be responsible for the long-term management of patients with migraine, maintaining stability of effective treatment and reacting to change**
- Timely transition from specialist care back to primary care with comprehensive treatment plans in place
- Patients can be referred to primary care once sustained efficacy with preventive therapy for up to 6 months is achieved with no substantial treatment-related adverse effects

CGRP, calcitonin gene-related peptide; EAN, European Academy of Neurology; EHF, European Headache Federation; MOH, medication overuse headache.

^aEvaluate treatment responses shortly after initiation (after 2–3 months) or a change in treatment, and regularly thereafter (every 6–12 months). Note that the efficacy of CGRP antibodies and onabotulinumtoxinA should only be assessed after 3–6 months or 6–9 months, respectively.

Adapted from: Eigenbrodt AK, et al. Nat Rev Neurol 2021;17:501–14.

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Evaluate: treatment plans and long-term follow-up

Evaluate



- **Engage** in long-term management in primary care to maintain stability of effective treatment and to react to change¹



- **Manage** complications and discourage medication overuse¹



- **Identify** and alleviate comorbidities if possible¹



- **Appraise** treatment response to assess effectiveness and tolerability regularly¹
 - Patient-oriented, validated outcome measures can help to verify that patients have experienced a meaningful response and identify the need for therapy adjustments²



Evaluate: treatment plans and long-term follow-up

Evaluate

Various validated tools are available to measure response in treatment



Migraine Disability Assessment (MIDAS)^{1,2}

Five scorable questions on school, social, and employment impact, frequency of headache and intensity of headache pain



Migraine Treatment Optimization Questionnaire (MTOQ-5)³

Five questions on functioning, rapid relief, consistency of relief, risk of occurrence, tolerability (to evaluate acute treatment response)



Migraine Assessment of Current Therapy (Migraine-ACT)⁴

Four questions on consistency of response, global assessment of relief, headache impact, emotional response (to evaluate acute treatment response)



Work Productivity and Activity Impairment Questionnaire: Migraine (WPAI:Migraine)⁵

Six questions on absenteeism, presenteeism, work productivity loss, activity impairment



Summary

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Evaluate



- Migraine is a primary headache disorder that can be categorized into subtypes (migraine with or without aura, chronic migraine, episodic migraine) based on symptoms and headache frequency¹
- It has a high socioeconomic and personal impact² and yet is widely underdiagnosed³

Recognize

- Early identification of symptoms, triggers, and comorbidities may help to plan treatment plan and reduce the severity of or prevent migraine attack⁴

Assess

- Key recommendations for diagnosing migraine include:⁵⁻⁷
 - History and physical exam
 - Screen for presence of red flags
 - Identify primary headache disorder using diagnostic tools

Treat

- Acute treatments include: NSAIDs, nonopioid analgesics, acetaminophen, or caffeinated analgesic combinations, triptans, ergotamine derivatives, gepants, ditans⁸
- Preventive treatments include: anticonvulsants, beta-blockers, anti-CGRP mAbs, gepants, neuromuscular blocking agent⁸

Evaluate

- Providers should engage in long-term follow-up to evaluate response to treatment, manage complications, and recognize comorbidities in patients
- Validated tools can assess disability, effectiveness of acute medications and quality of life⁹⁻¹³

NSAIDs, nonsteroidal anti-inflammatory drugs; CGRP, calcitonin gene-related peptide; mAbs, monoclonal antibodies.

1. IHS Headache Classification Committee. The International Classification of Headache Disorders, 3rd edition. Cephalgia 2018;38:1-211; 2. Ashina M et al. Lancet. 2021;397:1485-98; 3. Buse DC, et al. Headache 2021;61:628-41; 4. American Migraine Foundation. The timeline of an attack, 2018: <https://americanmigrainefoundation.org/resource-library/timeline-migraine-attack/>; 5. Martin VT, et al. Ann Med;2021;53:1979-1990; 6. Weatherall MW. Ther Adv Chronic Dis;2015;6:115-123; 7. Eigenbrodt AK. Nat Rev Neurol. 2021;17(8):501-514; 8. Ailani J, et al. Headache 2021;61:1021-39; 9. Lipton RB, et al. Headache 2001;41:854-61; 10. Stewart WF, et al. Pain 2000;88:41-52; 11. Lipton RB, et al. Cephalgia 2009; 29:751-9; 12. Dowson AJ, et al. Curr Med Res Opin 2004;20:1125-35; 13. Ford JH, et al. J Patient Rep Outcomes 2023;7:34

