

More Than Just a Headache

Introduction

Migraine is a common and debilitating neurologic disease affecting more than 10% of the world's population (approximately 1 billion people globally). In the United States, the prevalence of migraine is estimated to be around 12%. Migraine is 2 to 3 times more common in women than in men, and its prevalence peaks in midlife (30–49 years of age), impacting individuals in their prime working years. 2,3

Though a common disease with substantial impact, migraine is underdiagnosed and undertreated. The American Migraine Prevalence and Prevention study of 18,968 people found that 44% of subjects who met the International Classification of Headache Disorders 2nd edition (ICHD-2) criteria for migraine had never received a medical diagnosis of migraine. Furthermore, in an observational study of 2991 subjects who had a medical or self-diagnosis of sinus headache, 88% met the International Headache Society (IHS) criteria for migraine-type headache.

Pathophysiology

Many brain regions and systems are

thought to be involved in migraine pathophysiology, including the cerebral cortex, hypothalamus, trigeminocervical complex, thalamus, and meningeal nerves and vessels.6-10 Additionally, a range of neurotransmitters (eg, serotonin) and neuropeptides (eg, somatostatin) may contribute to the underlying biology of migraine. 11,12 Of these, the calcitonin gene-related peptide (CGRP), a 37-amino acid neuropeptide, is produced in central and peripheral neurons.^{8,13,14} CGRP is now widely considered to play a central role in migraine pathophysiology through its interaction with the CGRP receptor, a G-protein coupled receptor expressed in the trigeminovascular system.8,11,13,14

Current clinical data suggest that the interaction between CGRP and this receptor regulates key events that underlie migraine pathophysiology, including trigeminovascular neuron sensitization and neuropeptide release. ^{15–17} Most recently, clinical evidence has demonstrated that monoclonal antibodies disrupting the interaction between CGRP and its receptor are effective preventive treatments for migraine. ^{18–21}

Diagnosing Migraine

Migraine diagnosis is described in the third edition of the ICHD (ICHD-3), developed by the IHS.²² To meet ICHD-3 diagnostic criteria for migraine, a person must experience at least 5 headache attacks that each fulfill specific criteria for duration, quality, and severity, and are not better accounted for by a different diagnosis (**Figure 1**).²²

The Course of a Migraine Attack

Migraine attacks occur over hours to days and consist of several phases. **Figure 2** illustrates potential symptoms that may be associated with each phase and indicates how the symptoms may vary in intensity and duration over the course of a migraine attack.

During the prodromal phase, which may last for up to 48 hours, fatigue, food cravings, sensitivity to light and sound, nausea, neck discomfort, and cognitive symptoms have been reported.^{22–26} The aura phase may be shorter (5–60 minutes) and is often characterized by changes in vision, skin sensations, and/or language problems.²² The ictal (headache) phase is defined by moderate-

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to-severe head pain, but may also include sensitivity to light and sound, nausea and/or vomiting, sensitivity to touch, neck discomfort, cranial autonomic symptoms, and cognitive symptoms.^{22,25–27} This phase is the most debilitating and may last up to 72 hours.²² During the postdrome, which follows headache during a migraine attack, patients may experience sensitivity to light and sound, nausea, fatigue, cognitive symptoms, and neck discomfort for up to 48 hours.^{25,28,29} Some of these non-headache pain symptoms may continue into the interictal phase even in the absence of head pain.^{27,30–32} Not every migraine patient will experience all symptoms and phases; for example, approximately 30% of patients report aura symptoms, and 60%–70% experi-

The characteristic symptoms of migraine help facilitate differential diagnosis from other primary headache disor- Diagnostic Tools and Migraine ders including tension-type and cluster headache.²² **Table 1** summarizes the An accurate diagnosis of migraine heading migraine.^{36,37} pre-specified criteria that may be used to ache depends heavily on obtaining an identify each headache type.²² Key dis- accurate patient history.³⁶ Validated di- ment that was validated in primary care

Figure 2. Phases of Migraine^{22-32,46}

Aura

Changes in vision²²

Language problems

Skin sensations/

tingling²²

Prodrome

Food cravings^{22–25}

Fatigue^{22–25}

Nausea^{22,24,25}

Symptoms^{23–25}

discomfort²²⁻²⁶

and sound

Sensitivity to light

phonophobia)^{22–25}

Cognitive

Neck

ence sensitivity to touch.^{17,22,33–35}

Figure 1. ICHD-3 Criteria for Migraine Diagnosis²²

- □ ≥5 headache attacks
- ☐ Each attack lasting 4–72 hours
- \square Each attack including ≥ 2 of the following:
 - Unilateral location
 - Pulsating quality
 - O Moderate-to-severe pain intensity
- O Aggravated by or causing avoidance of routine physical activity
- \square Each attack including ≥ 1 of the following:
 - Nausea and/or vomiting
 - O Photophobia and phonophobia
- □ Not better accounted for by another ICHD-3 diagnosis

Abbreviation: ICHD, International Classification of Headache Disorders.

eral location, long duration (4–72 hours), proach to recording and assessing posfrequency, associated symptoms such as sible migraine symptoms and facilitate nausea and/or vomiting, and sensitivity diagnosis.^{36,37} Both general headache to light and sound or to touch.^{22,33–35}

Classification

Headache (ictal)

Moderate-to-severe head pain²²

Sensitivity to light and sound²²

Sensitivity to touch (allodynia)

~ 4–72 hours²²

Time

Nausea, vomiting²²

Neck discomfort^{25,26}

Cognitive symptoms

Cranial autonomi

and migraine-specific tools are available, which may be used concurrently to help exclude secondary headache and diagnose primary headache, includ-

Interictal

Sensitivity to light

and sound30-32

symptoms^{27,30}

Cognitive

Fatigue³⁰

Postdrome

Sensitivity to light

Nausea^{25,28}

Fatigue^{25,28,29}

symptoms^{25,28,29}

discomfort^{25,28,29}

≤ 48 hours²²

Cognitive

Neck

ID Migraine[™] is a 9-question assesseficial.³⁶ tinctive features of migraine are unilat- agnostic tools provide a systematic ap- practices and is considered the gold

when migraine is suspected, as it does cause is not overlooked.³⁷ not screen for other headache types.³⁶

cation use, or trigger identification).³⁶ specific ID Migraine[™] tool.⁴⁰ The headache diary is beneficial because

has a positive predictive value of 93.3.38 ing this screening takes only a few min-may impact treatment decisions.42 The ID Migraine[™] tool may be used utes and can help ensure a secondary

The Brief Headache Screen (BHS) is a Migraine is one of the most burdensome

the patient can record their symptoms established, it can be further refined as social obligations.^{2,44} Migraine also in real time instead of recalling them based on the monthly frequency of imposes a substantial economic burden after a time delay.^{36,39} The diary is used headache days.^{22,41} Episodic migraine is on society through increased health care to assess general headache symptoms, defined as fewer than 15 headache days costs and lost days of work.⁴⁵ On averto help determine whether migraine- per month, whereas chronic migraine age, patients with migraine lose 2-4 specific tests and screening may be ben- is characterized by 15 or more headache work days each month (not including SNOOP is a diagnostic screener that or tension-type headache) for more than graine who do go to work).⁴⁵ may be used to help rule out secondary 3 months, with headaches fulfilling the headaches, based on systemic and neu- diagnostic criteria for migraine on at **Conclusion**

standard for diagnosing migraine.^{36,38} A rologic symptoms, onset, other associleast 8 of the headache days.²² While simplified 3-item version assesses nau- ated conditions, and prior headache his- diagnosis is assumed to be stable and is sea, light sensitivity, and headache-re- tory.³⁷ The elements of SNOOP are red used to guide treatment decisions and lated disability.³⁸ A "yes" response to 2 flags for further investigation: **S**ystem-gauge outcomes, some patients have of the 3 items has high specificity and ic symptoms, **N**eurological symptoms, natural fluctuations in the severity of the sensitivity for diagnosis of migraine.³⁸ In **O**nset sudden, **O**lder (Onset after age disease (eg, they may transition between the primary care setting, ID Migraine[™] 50), and **P**attern change.^{36,37} Complet- chronic and episodic migraine), which

Burden of Migraine

For a general record of symptoms, a 7-item self-administered questionnaire neurologic diseases.¹ Headache disorheadache diary may be used by the pa- to help distinguish between different ders, including migraine, are the second tient to document the frequency, intensity per of headaches, including episodic leading cause of years lost to disability ty, duration, and other characteristics of headache syndromes and daily headache worldwide, after low back pain.^{1,43} More headaches.³⁶ Diaries can be used in digi-syndromes. Within the daily syndromes, than half of migraine patients experience tal or paper format, are simple to use, and it can also identify medication overuse severe impairment of daily activities or may be customized based on the data headache.³⁶ The BHS has demonstrated a need for bed rest during a migraine of interest (eq., symptom history, mediatrong agreement with the migraine- attack.² Many patients also report that their family activities are disrupted, and Once a migraine diagnosis has been that they miss everyday activities, such days per month (including migraine-like reduced productivity in people with mi-

Migraine affects more than 1 in 10 individuals worldwide¹ and can have a severe and disabling impact on everyday life.^{2,44} Migraine is underdiagnosed, undertreated, and frequently misdiagnosed as other common headache disorders.⁵ The substantial direct and indirect costs associated with migraine impose a considerable societal burden.⁴⁵ While much remains unknown, there is an evolving understanding of the pathophysiology of migraine. A growing body of evidence suggests that the interaction between CGRP and the CGRP receptor and activation of the trigeminovascular system may play a central role.^{8,13,14} In sum, migraine is more than just a headache. It is one of the most common and burdensome neurologic diseases.^{1,2} Understanding the characteristics and features of migraine and the symptoms that differentiate it from other headache disorders may help facilitate diagnosis and treatment in the primary care setting.

Table 1. Distinguishing Migraine From Other Primary Headaches²²

	Tension-type	Migraine	Cluster
Intensity	Mild to moderate	Moderate to severe	Severe or very severe
Location (of head pain)	Often bilateral	Often unilateral	Unilateral, usually be- hind or around one eye
Physical activity	Not aggravated by routine physical activity	Aggravated by routine physical activity	
Duration	30 min–1 week	4–72 hours	15–180 min
Frequency	Infrequent to daily	Recurrent, variable frequency	Once every other day to 8 times per day during clusters
Symptoms (in addition to head pain)	Light OR sound sensitivity (not both) Pericranial tenderness No nausea or vomiting	Light and sound sensitivity Nausea/vomiting Aura ¹⁷ Sensitivity to touch ^{33–35}	Eye redness or water- ing and constricted pupils Nasal congestion and facial sweating Eyelid swelling or drooping
Demographics	Higher prevalence in women than in men ⁴⁷	Prevalence in women 2–3 times higher than in men ²	Prevalence in men 3 times higher than in women

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Migraine patients may not experience all phases and listed symptoms, and not all possible symptoms are listed.

Adapted from Blau JN. Lancet. 1992;339:1202-1207.

≤ 48 hours²² ~ 5–60 min²²

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References

- GBD 2017 Disease and Injury Incidence and Prevalence Collaborators. Lancet. 2018;392(10159):1789-1858.
- Lipton RB, Bigal ME, Diamond M, Freitag F, Reed ML, Stewart WF. Neurology. 2007;68(5):343-349.
- 3. Martelletti P, Quintana R, Carboni V, Schwedt TJ, Lanteri-Minet M. *Cephalalgia*. 2018;38(1S):1-115. MTIS2018-088.
- 4. Diamond S, Bigal ME, Silberstein S, Loder E, Reed M, Lipton RB. *Headache*. 2007;47(3):355-363.
- 5. Schreiber CP. Arch Intern Med. 2004;164:1769-1772.
- 6. Charles A. Lancet Neurol. 2018;17(2):174-182.
- 7. Tedeschi G, Russo A, Conte F, et al. Cephalalgia. 2016;36(2):139-147.
- 8. Russo AF. Annu Rev Pharmacol Toxicol. 2015;55:533-552.
- 9. Maniyar FH, Sprenger T, Monteith T, Schankin C, Goadsby PJ. *Brain*. 2014;137(1):232-241.
- 10. Noseda R, Kainz V, Jakubowski M, et al. *Nat Neurosci*. 2010;13(2):239-245.
- 11. Tajti J, Szok D, Majláth Z, Tuka B, Csáti A, Vécsei L. *Neuropeptides*. 2015;52:19-30.
- 12. D'Andrea G, Leon A. Neurol Sci. 2010;31(suppl 1):S1-7.
- 13. Edvinsson L. Br J Clin Pharmacol. 2015;80(2):193-199.
- 14. Raddant AC, Russo AF. Expert Rev Mol Med. 2011;13:1-18.
- 15. Edvinsson L, Haanes KA, Warfvinge K, Krause DiN. *Nat Rev Neurol*. 2018;14(6):338-350.
- 16. Goadsby PJ, Edvinsson L. Ann Neurol. 1993;33(1):48-56.
- 17. Goadsby PJ, Holland PR, Martins-Oliveira M, Hoffmann J, Schankin C, Akerman S. *Physiol Rev.* 2017;97(2):553-622.
- 18. Silberstein SD, Dodick DW, Bigal ME, et al. *N Engl J Med*. 2017;377(22):2113-2122.
- 19. Goadsby PJ, Reuter U, Hallström Y, et al. *N Engl J Med*. 2017;377(22):2123-2132.
- 20. Stauffer VL, Dodick DW, Zhang Q, Carter JN, Ailani J, Conley RR. JAMA Neurol. 2018;75(9):1080-1088.
- 21. Sun H, Dodick DW, Silberstein S, et al. *Lancet Neurol*. 2016;15(4):382-390.
- 22. Headache Classification Committee of the International Headache Society (IHS). *Cephalalgia*. 2018;38(1):1-211.

- 23. Laurell K, Artto V, Bendtsen L, et al. Cephalalgia. 2016;36(10):951-959.
- 24. Schulte LH, Jürgens TP, May A. J Headache Pain. 2015;16(1):1-5.
- 25. Giffin NJ, Ruggiero L, Lipton RB, et al. *Neurology.* 2003;60(6):935-940.
- 26. Lampl C, Rudolph M, Deligianni CI, Mitsikostas DD. *J Headache Pain*. 2015;16:80.
- 27. Vuralli D, Ayata C, Bolay H. J Headache Pain. 2018;19(1):109.
- 28. Giffin NJ, Lipton RB, Silberstein SD, Olesen J, Goadsby PJ. *Neurology*. 2016;87(3):309-313.
- 29. Kelman L. Cephalalgia. 2006;26(2):214-220.
- 30. Houtveen JH, Sorbi MJ. PLoS One. 2013;8(8):1-10.
- 31. Ashkenazi A, Mushtaq A, Yang I, Oshinsky ML. *Cephalalgia*. 2009;29(10):1042-1048.
- 32. Main A, Dowson A, Gross M. Headache. 1997;37(8):492-495.
- 33. Baykan B, Ekizoglu E, Karli N, et al. Clin J Pain. 2016;32(7):631-635.
- 34. Misra UK, Kalita J, Bhoi SK. Clin J Pain. 2013;29(7):577-582.
- 35. Bigal ME, Ashina S, Burstein R, et al. *Neurology*. 2008;70(17):1525-1533.
- 36. Buse DC, Sollars CM, Steiner TJ, Jensen RH, Al Jumah MA, Lipton RB. Curr Pain Headache Rep. 2012;16(3):237-254.
- 37. Dodick D. Semin Neurol. 2010;30(1):74-81.
- 38. Lipton RB, Dodick D, Sadovsky R, et al. *Neurology*. 2003;61(3):375-382
- 39. Jensen R, Tassorelli C, Rossi P, et al. Cephalalgia. 2011;31(15):1549-
- 40. Maizels M, Houle T. Headache. 2008;48(3):385-394.
- 41. American Headache Society. *Headache*. 2019;59(1):1-18.
- 42. Serrano D, Lipton RB, Scher Al, et al. *J Headache Pain*. 2017;18(1):1-
- 43. Institute for Health Metrics and Evaluation (IHME). Seattle, WA: IHME, 2018.
- 44. Vo P, Paris N, Bilitou A, et al. Neurol Ther. 2018;7(2):321-332.
- 45. Bonafede M, Sapra S, Shah N, Tepper S, Cappell K, Desai P. *Headache*. 2018;58(5):700-714.
- 46. Blau JN. Migraine: *Lancet*. 1992;339(8803):1202-1207.
- 47. Russell MB. *J Headache Pain*. 2007;8(2):71-76.