


RESEARCH SUBMISSIONS

Diagnostic accuracy of an artificial intelligence online engine in migraine: A multi-center study

Robert P. Cowan MD¹ | Alan M. Rapoport MD² | Jim Blythe PhD³ | John Rothrock MD⁴ | Kerry Knievel DO⁵ | Addie M. Peretz MD¹ | Elizabeth Ekpo MD⁶ | Bharati M. Sanjanwala MSc¹ | Yohannes W. Woldeamanuel MD¹ 

¹Division of Headache and Facial Pain, Department of Neurology and Neurological Sciences, Stanford University School of Medicine, Stanford, California, USA

²Neurology, University of California, Los Angeles, California, USA

³Information Sciences Institute, University of Southern California, Los Angeles, California, USA

⁴Neurology, The George Washington University School of Medicine and Health Sciences, Washington, District of Columbia, USA

⁵Neurology, Barrow Neurological Institute, Phoenix, Arizona, USA

⁶Neurology, University of California Davis, Davis, California, USA

Correspondence

Yohannes W. Woldeamanuel, Division of Headache and Facial Pain, Department of Neurology and Neurological Sciences, Stanford University School of Medicine, Stanford, California, USA.
Email: ywoldeam@stanford.edu

Funding information

The authors are thankful to The SunStar Foundation for financially sponsoring this study. YWW is a recipient of the National Institute of Neurological Disorders and Stroke of the National Institutes of Health under award number 1K01NS124911-01. The funding sources had no role in the design of this study, its execution, analyses, interpretation of the data, or decision to submit results.

Abstract

Objective: This study assesses the concordance in migraine diagnosis between an online, self-administered, Computer-based, Diagnostic Engine (CDE) and semi-structured interview (SSI) by a headache specialist, both using International Classification of Headache Disorders, 3rd edition (ICHD-3) criteria.

Background: Delay in accurate diagnosis is a major barrier to headache care. Accurate computer-based algorithms may help reduce the need for SSI-based encounters to arrive at correct ICHD-3 diagnosis.

Methods: Between March 2018 and August 2019, adult participants were recruited from three academic headache centers and the community via advertising to our cross-sectional study. Participants completed two evaluations: phone interview conducted by headache specialists using the SSI and a web-based expert questionnaire and analytics, CDE. Participants were randomly assigned to either the SSI followed by the web-based questionnaire or the web-based questionnaire followed by the SSI. Participants completed protocols a few minutes apart. The concordance in migraine/probable migraine (M/PM) diagnosis between SSI and CDE was measured using Cohen's kappa statistics. The diagnostic accuracy of CDE was assessed using the SSI as reference standard.

Results: Of the 276 participants consented, 212 completed both SSI and CDE (study completion rate = 77%; median age = 32 years [interquartile range: 28–40], female:male ratio = 3:1). Concordance in M/PM diagnosis between SSI and CDE was: $\kappa = 0.83$ (95% confidence interval [CI]: 0.75–0.91). CDE diagnostic accuracy: sensitivity = 90.1% (118/131), 95% CI: 83.6%–94.6%; specificity = 95.8% (68/71), 95% CI: 88.1%–99.1%. Positive and negative predictive values = 97.0% (95% CI: 91.3%–99.0%) and 86.6% (95% CI: 79.3%–91.5%), respectively, using identified migraine prevalence of 60%. Assuming a general migraine population prevalence of 10%, positive and

Abbreviations: AI, artificial intelligence; CDE, Computer-based, Diagnostic Engine; CI, confidence interval; ICHD-1, International Classification of Headache Disorders, 1st edition; ICHD-2, International Classification of Headache Disorders, 2nd edition; ICHD-3, International Classification of Headache Disorders, 3rd edition; IQR, interquartile range; LR, likelihood ratio; ML, machine learning; M/PM, migraine/probable migraine; NPV, negative predictive value; PPV, positive predictive value; SSI, semi-structured interview; STARD, Standards for Reporting of Diagnostic Accuracy Studies.

This is an open access article under the terms of the [Creative Commons Attribution-NonCommercial](https://creativecommons.org/licenses/by-nc/4.0/) License, which permits use, distribution and reproduction in any medium, provided the original work is properly cited and is not used for commercial purposes.

© 2022 The Authors. *Headache: The Journal of Head and Face Pain* published by Wiley Periodicals LLC on behalf of American Headache Society

negative predictive values were 70.3% (95% CI: 43.9%–87.8%) and 98.9% (95% CI: 98.1%–99.3%), respectively.

Conclusion: The SSI and CDE have excellent concordance in diagnosing M/PM. Positive CDE helps *rule in* M/PM, through high specificity and positive likelihood ratio. A negative CDE helps *rule out* M/PM through high sensitivity and low negative likelihood ratio. CDE that mimics SSI logic is a valid tool for migraine diagnosis.

KEYWORDS

artificial intelligence, diagnosis, diagnostic accuracy study, migraine, online engine, semi-structured interview

INTRODUCTION

Migraine is an underdiagnosed and undertreated disabling disease worldwide.^{1,2} Several studies have shown that patients face significant delay of up to 12–17 years in obtaining an accurate migraine diagnosis.^{3–6} An inadequate number of headache-trained professionals and time-consuming in-person diagnostic interviews by undertrained clinicians contribute to a significant bottleneck at the initial visit.^{2,7} Currently, there are only 688 headache specialists in the United States for more than 40 million people experiencing migraine, or one headache specialist for 58,000 individuals with migraine.⁸ Globally, the rising migraine prevalence affects nearly one billion people—most of whom are not receiving appropriate headache care.⁹ Diagnostic delay increases the risk of chronic migraine, treatment refractoriness, comorbidities, and medication overuse.² Of nine headache-related variables in a 7-year follow-up study, a 6-month delay in migraine diagnosis was the only factor differentiating headache freedom from persistent headache.¹⁰ Another 10-year longitudinal study showed a 2-fold increased risk of persistent migraine in children diagnosed after age 12.¹¹ A separate longitudinal study demonstrated that a 5-year diagnostic delay of migraine increased consultations and unnecessary investigations by 40% and 35%, respectively.¹² A US study showed that only 25% of patients with chronic migraine who consulted a health-care professional received an accurate initial diagnosis, and a mere 1.8% received optimal migraine management.⁵

The diagnostic approach in migraine primarily relies on patient history¹³—making it well suited to self-administered digital health tools. Accurate web-based algorithms may improve scalability of accurate migraine diagnoses and might thereby facilitate access to appropriate treatment. Digital health tools can reduce migraine care costs and expedite health-care delivery by restructuring the live clinical encounter.^{14–16} The integration of machine learning in digital diagnostic tools can approximate the decision-making used in in-person screening and triaging patients.^{17,18} The utility of digital health tools lies at the intersection of increasing burden of chronic conditions (e.g., migraine) and patient-centered care.^{14–17} Patients improve self-efficacy and self-management when using digital health tools.^{14–17} Artificial intelligence (AI)-powered diagnoses can facilitate disease management when collaborative health care is not readily available.¹⁹ Digital health tools can enhance the efficiency of large population studies and clinical trials.^{19,20}

Computer-assisted and computer-based diagnostics have long been posited as a potential solution to the shortage of physicians.^{18,21} Examples of digital health tools validated for diagnosis or self-management include MindDoc for depression,²² SleepAp for obstructive sleep apnea,²³ and EncephalApp for covert hepatic encephalopathy.²⁴ Computerized headache diagnosis is a timely topic given the prevalence of headache disorders, self-report-based diagnosis, and paucity of trained providers.

Review of computerized migraine diagnostic tools

We conducted a systematic review of all published studies that evaluated computerized migraine diagnostic tools (41 studies since 1960).²⁵ In 1991, the first computer diagnosis based on the 1988 International Classification of Headache Disorders (ICHD-1)²⁶ and the 1962 Ad Hoc criteria²⁷ was compared to interview-based diagnosis by headache physicians and psychologists.²⁸ A 95.9% concordance rate was found between the interviewers and the computerized diagnosis²⁸—both using the Ad Hoc criteria. The concordance between computerized diagnosis based on the Ad Hoc criteria and computerized diagnosis based on ICHD-1 criteria was 77% for migraine.²⁸ This study²⁸ did not compare the computerized and interviewers' diagnosis using ICHD-1. In 2005, the first ICHD-2-based computerized tool performed with 68% (345/500) concordance in diagnosing primary headache types compared to interview-based diagnosis by headache-trained clinicians.²⁹ Missing data and clinicians' errors contributed to 29% non-concordance while computer error accounted for 3% non-concordance.²⁹ Both studies^{28,29} did not report concordance rates between computerized and interview migraine diagnosis. These early digital tools required physicians to complete the forms,^{28,29} reducing their utility and scalability.

In the last 5 years, researchers^{30–32} have evaluated different algorithms and expert systems in diagnosing migraine and other headache types, reporting 30%–95% accuracy. However, several of these tools were developed based on non-ICHD criteria, retrospective analysis, and were not tested against live interviews; others contained contamination (i.e., the diagnostician was exposed to diary data prior to interviews). The coronavirus disease 2019 pandemic has contributed to the increase in digital health research and care in headache.³³

Today, the ground truth of headache diagnosis is the ICHD-3. In statistics and machine learning, the “ground truth” is similar to the

“gold standard” in diagnostic accuracy studies. However, in the hands of clinicians, the ICHD-3 serves as a guide, rather than a rigid diagnostic template. It is the semi-structured interview (SSI) that is the vehicle used by the diagnostician to achieve an ICHD-3 diagnosis. Previous studies have found inconsistencies in headache diagnostic accuracy between self-administered questionnaires (sans machine learning [ML]) and clinical interviews.^{34,35} Sophisticated SSI when substituted by simplified tools such as ID Migraine³⁶ might lead to misdiagnosing an exceptional patient. Diagnostic disparities due to non-ML self-administered questionnaires may be obviated by ML that approximates the cognitive processes at play in the clinician’s diagnostic analytics. This is because both ML and clinician’s diagnostic reasoning involve iterative processes relying on repetitive exposures to clinical cases^{37,38} and both require expert feedback for development.^{37,38}

In this study, we determined the concordance in migraine diagnosis between an online, self-administered Computer-based Diagnostic Engine (CDE) and a headache specialist (SSI), developed by headache fellowship-trained clinicians. Both use the ICHD-3. We hypothesized that the CDE could diagnose migraine as accurately as the headache specialist. Furthermore, we examined areas in which discrepancies occurred between the two approaches and discussed efforts to improve concordance.

METHODS

Study design

This was a cross-sectional study that enrolled participants in a two-part assessment involving the SSI and CDE. The order of the assessments was randomly assigned. For the SSI assessment, participants took part in a phone interview conducted by headache specialists involving an SSI that was developed by the headache specialists using ICHD-3 criteria. For the CDE assessment, every participant completed the CDE—a web-based algorithm constructed by tying ICHD-3 diagnostic criteria for all primary headache disorders and the more common secondary headache disorders to specific questions presented using a version of mixed chaining and case-based reasoning.³⁹ Participants were recruited between March 2018 and August 2019. The first-part assessment was followed a few minutes later by the second-part assessment in all participants. Our questionnaires were designed to minimize the occurrence of respondent fatigue by utilizing well-recognized approaches such as branching features that help reroute some questions, forced entry, non-compounded directed questions, and clear communications of length of time needed to complete questionnaire.⁴⁰ Both participants and clinicians were blinded to results.

PARTICIPANT RECRUITMENT, INCLUSION AND EXCLUSION CRITERIA

Participant recruitment was carried out at three centers and nearby communities using convenience sampling: Stanford University

Headache Center, Jan and Tom Lewis Migraine Treatment Program at Barrow Neurologic Institute, and George Washington University Headache Center. Inclusion criterion: adults aged 18 years or older. Exclusion criterion: children aged younger than 18 years.

Development of the CDE

The digital diagnostic tool used in this study was developed by authors RPC, AMR, and JB based on a detailed decision tree designed to ask sufficient questions to diagnose all ICHD-3 primary headaches as well as several secondary headaches such as medication overuse headache and post-traumatic headache. The CDE uses the National Institutes of Health⁴¹ and American Medical Association⁴² recommended level of 6th grade for preparing health information materials including diagnostic questionnaires. Medical jargon was avoided, and easy-to-understand phrases were used. The CDE is compatible with any internet-connected computer, smart phone, or tablet and utilizes a forced choice format for filling responses. The CDE contains 10 questions on demographics and 168 questions on headache assessment which can increase depending on the number of the participant’s headache types. The CDE headache assessment questions broadly involve headache and headache-related disability history, headache treatment history, personal and family history of headache as well as familial headache treatments, emotions, and habits in relation to headache.

CDE development and testing was continuous over a period of several years before the study was conducted. The CDE diagnostic rule set involves considering each question in turn and dynamically recomputing, relative to the current answer set and the diagnostic rules, whether the question may logically affect the emerging diagnostic impression. The question is only asked if this is true. This approach minimizes the number of questions required to reach a diagnostic impression *dependent on the initial question ordering*, which is chosen to create a coherent, conversational experience, and assuming no prior probabilities for diagnostic outcomes. Utilizing a rule-based engine, responses were tied to ICHD-3 diagnostic criteria and a version of mixed chaining and case-based reasoning³⁹ analysis designed to identify the fewest questions that would lead to a definitive diagnosis and rule out others. This design was intended to emulate the diagnostic process used in the SSI. A simplified definition of technical terms related to artificial intelligence (adopted from references^{43,44}) used in our study is provided in Table S1.

Development of the SSI questionnaire

The SSI questionnaire was prepared by copying the migraine criteria from the ICHD-3 and rephrasing it in a question format. The SSI was used by five headache specialists from the three headache centers who performed a phone interview with each participant and made a diagnosis of type of headache or no headache. The

SSI contains seven questions on demographics and a minimum of 65 detailed questions on headache assessment that can branch up to 135 questions depending on responses and the number of the participant's headache types—with the caveat that there may be many more questions that can be asked at the interviewer's discretion (see File S1). The SSI allowed the headache specialists to probe further using their own interviewing approaches and additional questions in cases of unclear responses or perceived inconsistencies. Data on these additional questions was not collected. Neither the CDE nor the SSI utilized a physical or neurologic examination.

Outcome measures

Outcome measures included “migraine/probable migraine” (M/PM) as a positive CDE test and “no migraine” as a negative CDE test diagnosis using the SSI as reference standard. A second analysis was done using “migraine” as a positive CDE test and “no migraine” as a negative CDE test. A third analysis was done using “migraine” as a positive CDE test and “no migraine/probable migraine” as a negative CDE test. These methods allowed us to compare the accuracy of CDE in migraine diagnosis as well as its precision in discerning probable migraine from definitive migraine. Participants' age and sex were recorded. We interpreted the CDE index test without knowledge of the SSI reference standard. All analyses were preplanned. While other headache types were identified in both the CDE and SSI, there were too few to assign significance in a subset analysis.

Sample size estimation

Assuming a migraine prevalence of 35% in headache clinics⁴⁵⁻⁴⁷ and a sample sensitivity of 80% for CDE, the sample size needed for a two-sided 85% sensitivity confidence interval (CI) with a width of at most 0.15, is 203.^{48,49} Assuming a migraine prevalence of 35% in headache clinics⁴⁵⁻⁴⁷ and a sample specificity of 80% for CDE, the sample size needed for a two-sided 85% specificity CI with a width of at most 0.15, is 110.^{48,49} The whole table sample size required so that both CIs have widths less than 0.15, is 203, the larger of the two sample sizes.^{48,49} We adjusted the final sample size by accounting for an estimated 20% to 25% of participants with missing/incomplete data.^{50,51} Hence, we enrolled a total of 266 participants to ensure we had 203 evaluable participants' data. The CI was based on binomial distribution (Clopper-Pearson exact method⁵²). Sample size calculation was performed using PASS 2020 software (NCSS, LLC).

Statistical analysis

This is the primary analysis of these data. Descriptive statistics (i.e., median and interquartile range [IQR]) were used to describe age and sex ratio. The concordance in migraine diagnosis between the SSI

and CDE was measured using unweighted Cohen's kappa (κ) statistics; unweighted κ was selected because the outcomes are nominal variables. Kappa values were interpreted using Cohen's recommendations as “no agreement” for $\kappa \leq 0$, “none to slight agreement” for $\kappa = 0.01-0.20$, “fair agreement” for $\kappa = 0.21-0.40$, “moderate agreement” for $\kappa = 0.41-0.60$, “substantial agreement” for $\kappa = 0.61-0.80$, and “almost perfect agreement” for $\kappa = 0.81-1.00$.⁵³ The diagnostic accuracy of the CDE was assessed using the SSI as the reference standard. The sensitivity, specificity, positive predictive value (PPV), negative predictive value (NPV), accuracy, as well as the positive and negative likelihood ratios (LRs) were used to measure the performance of the CDE and if an accurate diagnosis was made. For PPV, NPV, and accuracy estimations, both the migraine prevalence in our study population determined by the SSI results and a standard estimate of migraine prevalence in the general population⁵⁴ were used to determine approximate boundaries on these parameters. This allowed us to compare CDE's utility between clinical and community settings. Accuracy was defined as the overall probability that a patient is correctly classified and was calculated as: sensitivity \times prevalence + specificity \times (1 - prevalence). The two-step Fagan's nomogram⁵⁵ based on Bayes' Theorem⁵⁶ was used to examine pre- to post-test probability changes in migraine diagnosis using CDE. It is noteworthy to distinguish between diagnostic test performance in the current sample (e.g., sensitivity, specificity, PPV, NPV, LR) and theoretical post-test probability in future samples (i.e., Fagan's nomogram). Agreement rates between CDE and SSI among nine migraine-related symptoms (i.e., presence of unilateral headache, moderate/severe head pain intensity, aura, nausea and/or vomiting, headache duration of 4–72 h, pulsating headache, photophobia, phonophobia, aggravation by or avoidance of routine physical activity) were further analyzed to identify the symptom domains with high and low discrepancy. The results are reported in accordance with the Standards for Reporting of Diagnostic Accuracy Studies (STARD).⁵⁷ Statistical analyses were performed using MedCalc for Windows, version 20.022 (MedCalc Software) and Microsoft Excel 2021.

RESULTS

Characteristics of included participants

A total of 266 participants were recruited to the study from the three headache centers: 143 participants from Stanford University Headache Center, 43 participants from Jan and Tom Lewis Migraine Treatment Program at Barrow Neurologic Institute, and 80 participants from George Washington University Headache Center. Of the 266 recruited participants, 202 participants completed both the CDE and SSI (study completion rate = 76%). The remaining 64 (24%) participants were excluded due to incomplete or missing data. Of the 202 participants, 102 (50.5%) were newly diagnosed (i.e., diagnosis based on SSI without a prior diagnosis) while the remaining 100 (49.5%) participants were known cases with confirmed diagnoses of different headache types. Responders had a median age of

32 years (IQR: 28, 40), female:male ratio of 3:1, 59% White, and 28% were recruited from headache clinics while 72% came from the local communities. The age and female:male ratio of patients recruited from the three headache centers is displayed in Table 1. The racial demographics of participants is available in Table S2. Participants with headache had a median monthly headache day frequency of 3 (IQR = 1–13). Use of headache medication classes and frequency of monthly headache medication consumption is available in Table S3. The duration of the SSI interview as well as the time needed to complete the CDE lasted from 5 min in participants with no headache history up to 45 min in participants reporting multiple headache types. The mean time to complete the SSI was 30 min, while the mean time to complete the CDE was 48 min.

Diagnostic accuracy performance

There was almost perfect concordance in M/PM diagnosis between CDE and SSI, $\kappa = 0.82$ (95% CI: 0.74–0.90; Figures 1 and 2). The CDE performed with an overall diagnostic accuracy of 91.6% (95%

TABLE 1 Demographic characteristics of patients recruited from the three headache centers

	Recruitment headache centers			
	Stanford (n = 143)	GWU (n = 80)	Barrow (n = 43)	Total (n = 202)
Median age (IQR), years	32 (29, 41)	33 (26, 39)	36 (28, 45)	32 (28, 40)
Female, n (%)	81 (57%)	69 (86%)	35 (81%)	152 (75%)

Abbreviations: GWU, George Washington University; IQR, interquartile range.

CI: 86.9%–95.0%), sensitivity of 89.0% (95% CI: 82.5%–93.7%), and specificity of 97.0% (95% CI: 89.5%–99.6%). Positive and negative predictive values were 98.4% (95% CI: 93.9%–99.6%) and 81.0% (95% CI: 72.5%–87.3%), respectively, using the identified M/PM prevalence of 67% (95% CI: 60.4%–73.7%). The age and sex ratio of SSI-based diagnosis is shown in Table 2. A 2 × 2 contingency table allowing calculations of the diagnostic performance of the CDE is displayed in Table 3. Assuming a general migraine population prevalence of 10%,⁵⁴ the positive and negative predictive values were 76.5% (95% CI: 45.4%–92.8%) and 98.8% (95% CI: 98.0%–99.2%), respectively. The positive and negative LR_s were 29.4 (95% CI: 7.5–115.1) and 0.11 (95% CI: 0.07–0.18), respectively. Based on Fagan's nomogram, a M/PM diagnosis on the CDE increases a 50% pre-test probability of having M/PM to a 97% post-test probability (Figure 2). Similarly, a negative result on CDE ("no migraine") decreases a 50% pre-test probability of having "no migraine" to a 10% post-test probability (Figure 2). If a patient from a high-risk population (i.e., headache clinic setting with a 67% M/PM prevalence) tests positive, the post-test probability that the patient truly has M/PM will be 98%. Alternatively, if the high-risk patient tests negative, the post-test probability that she or he truly has M/PM will only be 18%. For a patient from a low-risk population (e.g., community migraine prevalence of 10%⁵⁴) who tests positive on CDE, the post-test probability that the patient truly has M/PM will be 76%. On the other hand, if the low-risk patient tests negative, the post-test probability that she or he truly has M/PM will decrease to 1%. On stratified analysis, the diagnostic accuracy of CDE for M/PM diagnosis was 87%, 86%, and 82% in the subgroups of participants recruited from community, newly diagnosed participants, and known cases with confirmed diagnoses, respectively.

For the second analysis using "migraine" as a positive CDE and "no migraine" as a negative CDE (excluding probable migraine), there

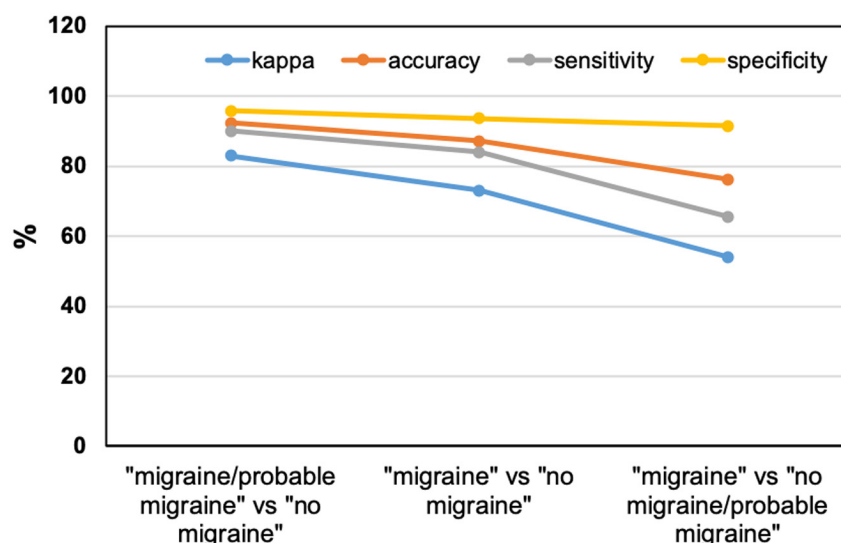


FIGURE 1 Diagnostic accuracy performance of the CDE. The diagnostic accuracy performance (measured by kappa, accuracy, sensitivity, specificity LR+) of the CDE increased in the following order: "migraine" vs. "no migraine/probable migraine," "migraine" vs. "no migraine," "migraine/probable migraine" vs. "no migraine." CDE, Computer-based Diagnostic Engine; LR+, positive likelihood ratio [Color figure can be viewed at wileyonlinelibrary.com]

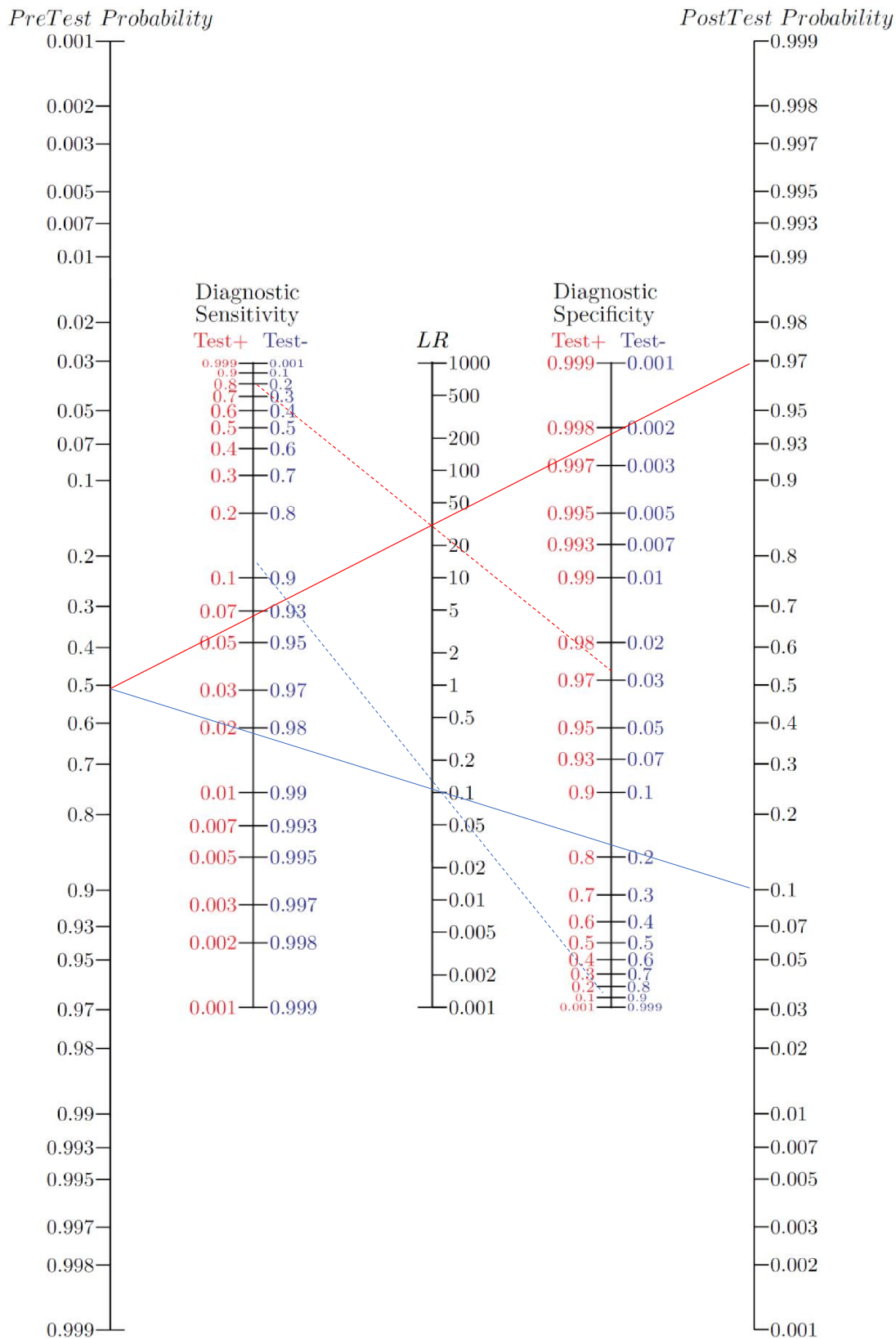


FIGURE 2 The two-step Fagan's nomogram. A migraine/probable migraine diagnosis on the CDE increases a 50% pre-test probability of having migraine/probable migraine to a 97% post-test probability (red solid line). With a negative CDE result ("no migraine"), a 50% pre-test probability of having "no migraine" lowers to a 10% post-test probability (blue solid line). The dotted lines indicate the sensitivity and specificity of 89% and 97%, respectively; as well as the positive (red dotted line) and negative (blue dotted line) likelihood ratios of 29.4 and 0.11, respectively. CDE, Computer-based Diagnostic Engine; LR, likelihood ratio [Color figure can be viewed at wileyonlinelibrary.com]

was substantial concordance in migraine diagnosis between CDE and SSI, $\kappa = 0.73$ (95% CI: 0.62–0.84). The CDE performed with an overall diagnostic accuracy of 87.2% (95% CI: 80.6%–92.3%), sensitivity of 84.0% (95% CI: 75.1%–90.8%), and specificity of 93.6% (95% CI: 82.5%–98.7%). Positive and negative predictive values were 96.3% (95% CI: 89.8%–98.8%) and 74.6% (95% CI: 64.7%–82.4%), respectively, using the identified migraine prevalence of 67% (95% CI: 58.2%–74.4%). Assuming a general migraine population prevalence of 10%,⁵⁴ the positive and negative predictive values were 59.4% and 98.1%, respectively. The positive and negative LR_s were 13.2 (95% CI: 4.39–39.5) and 0.17 (95% CI: 0.11–0.27), respectively. Based on Fagan's nomogram, a positive CDE increases a 50% pre-test probability of having migraine to a 92.9% post-test probability. Similarly,

a negative result on CDE ("no migraine") decreases a 50% pre-test probability of having "no migraine" to a 14.5% post-test probability.

For the third analysis using "migraine" as a positive CDE and "no migraine/probable migraine" as a negative CDE, there was moderate concordance in migraine diagnosis between CDE and SSI, $\kappa = 0.67$ (95% CI: 0.57–0.77). The CDE performed with an overall diagnostic accuracy of 83.2% (95% CI: 77.3%–88.1%), sensitivity of 75.6% (95% CI: 66.9%–83.0%), and specificity of 94.0% (95% CI: 86.5%–98.0%). Positive and negative predictive values were 94.7% (95% CI: 88.4%–97.7%) and 72.9% (95% CI: 66.1%–78.8%), respectively, using the identified migraine prevalence of 58.9% (95% CI: 51.8%–65.8%). Assuming a general migraine population prevalence of 10%,⁵⁴ the positive and negative predictive values were 46.3% and 96.0%, respectively. The positive and negative LR_s were 12.6 (95% CI: 5.33–29.6) and 0.26 (95% CI: 0.19–0.36), respectively. Based on Fagan's nomogram, a positive CDE increases a 50% pre-test probability of having migraine to 93% post-test probability. Similarly, a negative result on CDE ("no migraine/probable migraine") decreases a 50% pre-test probability of having "no migraine" to a 21% post-test probability.

The summary of the diagnostic accuracy results is displayed in Table 4.

The agreement rate between CDE and SSI (Figure 3) among nine migraine-related symptoms was 47% for phonophobia, 47%

TABLE 2 Demographic characteristics of SSI-based diagnosis

	SSI diagnosis	
	Migraine/probable migraine (n = 131)	No migraine/probable migraine (n = 71)
Median age (IQR), years	34 (28, 41)	31 (28, 37)
Female-to-male ratio	94 (71%)	28 (40%)

Abbreviations: IQR, interquartile range; SSI, semi-structured interview.

TABLE 3 A 2 × 2 contingency table for calculations of diagnostic accuracy performance of the CDE (Computer-based Diagnostic Engine) using the SSI (semi-structured interview) as a gold standard

		SSI		
		Migraine/probable migraine	No migraine/probable migraine	Total
CDE	Migraine/probable migraine	121 (true positive)	2 (false positive)	123 (true positive + false positive)
	No migraine/probable migraine	15 (false negative)	64 (true negative)	79 (false negative + true negative)
Total		136 (true positive + false positive)	66 (false negative + true negative)	202

TABLE 4 Diagnostic accuracy performance of the CDE

Diagnostic accuracy	"Migraine/probable migraine" vs. "no migraine"	"Migraine" vs. "no migraine"	"Migraine" vs. "no migraine/probable migraine"
Kappa % (95% CI)	82% (74%–90%)	73% (62%–84%)	67% (57%–77%)
Accuracy % (95% CI)	92% (87%–95%)	87% (81%–92%)	83% (77%–88%)
Sensitivity % (95% CI)	89% (83%–94%)	84% (75%–91%)	76% (67%–83%)
Specificity % (95% CI)	97% (90%–100%)	94% (83%–99%)	94% (87%–98%)
PPV % (95% CI)	98% (94%–100%)	96% (90%–99%)	95% (88%–98%)
NPV % (95% CI)	81% (73%–87%)	75% (65%–82%)	73% (66%–79%)
LR ₊ , ratio (95% CI)	29 (8–115)	13 (4–39)	13 (5.33–29.6)
LR ₋ , ratio (95% CI)	0.11 (0.07–0.18)	0.17 (0.11–0.27)	0.26 (0.19–0.36)

Note: Except for LR₋, all values are rounded off to the nearest whole number.

Abbreviations: CDE, Computer-based Diagnostic Engine; CI, confidence interval; LR, likelihood ratio; NPV, negative predictive value; PPV, positive predictive value.

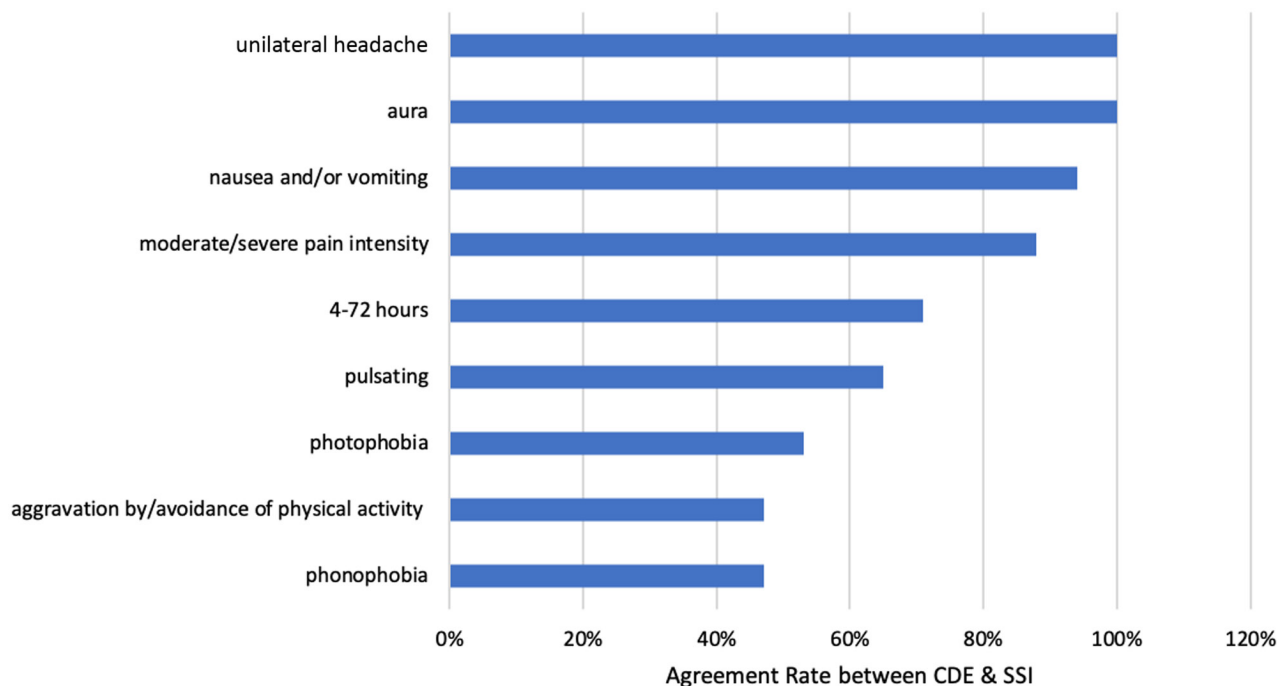


FIGURE 3 Agreement rates between CDE and SSI among nine migraine-related symptoms. The agreement rate between CDE and SSI among nine migraine-related symptoms was 47% for phonophobia, 47% for aggravation by/avoidance of routine physical activity, 53% for photophobia, 65% for pulsating headache, 71% for 4–72 h headache duration, 88% for headache pain intensity, 94% for nausea and vomiting, 100% for aura, and 100% for unilateral headache, ascendingly. These agreement rates were based on the 17 participants that were either false negative or false positive in migraine/probable migraine diagnosis in which the CDE performed with an overall diagnostic accuracy of 91.6% (95% CI: 86.9%–95.0%). $\kappa = 0.82$ (95% CI: 0.74–0.90). CDE, Computer-based Diagnostic Engine; CI, confidence interval; SSI, semi-structured interview [Color figure can be viewed at wileyonlinelibrary.com]

for aggravation by/avoidance of routine physical activity, 53% for photophobia, 65% for pulsating headache, 71% for 4–72 h headache duration, 88% for headache pain intensity, 94% for nausea and vomiting, 100% for aura, and 100% for unilateral headache, ascendingly. These agreement rates were based on the 17 participants that were either false negative or false positive in M/PM diagnosis in which the CDE performed with an overall diagnostic accuracy of 91.6% (95% CI: 86.9%–95.0%), $\kappa = 0.82$ (95% CI: 0.74–0.90).

DISCUSSION

By virtue of being a disruptive digital health technology, the CDE has enormous utility in addressing the unmet need of diagnostic delay, under-/misdiagnosis and under-/mismanagement of migraine in both clinical as well as community settings. By providing accurate migraine diagnosis, the CDE can be a step closer toward addressing the rising headache burden—especially when accompanied by improvement in optimum headache care. Besides accelerating remote telemedicine, the CDE can allow self-diagnosis thereby improving patients' self-efficacy. The CDE creates an opportunity for enhancing data-driven clinical research by enabling improved data collection for headache outcomes in addition to validating personalized treatment delivery. With the use of CDE for patient triaging, referral of patients with migraine can be streamlined to provide efficient

use of primary and tertiary care settings. Given the present study's validation in headache clinics with interviews being conducted by headache specialists, the ideal users would be patients attending headache clinics. In the future, we plan to validate this study in primary care setting with interviews being conducted by primary care providers.

Our results show a high level of agreement between the self-administered CDE and the SSI phone interviews conducted by headache specialists. A positive CDE will help to *rule in* a migraine diagnosis, driven by its near-perfect specificity and high positive LR. A negative CDE will aid to *rule out* migraine diagnosis because of its high sensitivity and low negative LR. The reason that the CDE's sensitivity (90.08%) was slightly lower than its specificity (95.77%) may be because computerized diagnostic tools are more prone to false positive errors compared to traditional interviewing.^{58,59} The computer may be limited to branch adequately and generate additional or follow-up questions to clarify and refine responses.^{58,59} Unfiltered responses are more common in computerized diagnostic tools than in traditional interviewing;^{58,59} in the latter, the physician can redirect the patient to focus on important questions while reassuring the patient on trivial complaints.^{60,61} This clearly gave the SSI an advantage over the CDE where the questions had to be understood and answered the best way possible. Similarly, nonverbal behavior can be identified by the physician.⁵⁹ Nonverbal behavior includes visual cues

(e.g., facial expression, body language) in face-to-face interviews as well as conversational/auditory cues (e.g., intonation, hesitation, sighs, pressured speech, annoyance, sarcasm) in telephone interviews.⁶²⁻⁶⁴ Conversational cues play a role during telephone-based diagnostic interviews, can help to create rapport, and facilitate probing for in-depth interviews.^{62,63,65,66} Compared to in-person patient interviews, telephone-based interviews have the advantage of making participants feel at ease, make them feel “on their own turf” and relaxed to disclose sensitive information (e.g., stigmatizing conditions such as migraine).^{66,67} In addition, interviewers have the opportunity to engage in informal dialogue prior to the formal interview—this can further improve rapport.⁶⁵ The CDE has a ML component, which is currently training on reducing unfiltered responses, inadvertent errors, and false positivity that lower sensitivity outcomes. Currently, the CDE does not contain advanced AI such as a neural network. In the near future, we plan to upgrade and test the accuracy of a next-generation CDE that will involve a neural network with continued training in classification decisions based on its internally generated representation to supplement and enhance the existing rule-based system.

The CDE was estimated to perform variably with higher PPVs and lower NPVs for high-prevalence migraine settings (tertiary headache clinics), and vice versa for low-prevalence (primary care or community) settings. This variance in diagnostic accuracy may be due to patients presenting at tertiary headache centers exhibiting a more protracted headache history than those attending primary care. Such difference in presentation may introduce recall and/or response bias in patient-reported headache symptoms. These biases can influence the CDE performance, particularly by reducing its sensitivity, NPV, and negative LR. In contrast, patients who are found to have no migraine in a tertiary setting usually present with comorbidities of other headache types—reducing the specificity, PPV, and positive LR of the CDE test.

The agreement rates between CDE and SSI among migraine-related symptoms showed that distinctly memorable experiences which can be worded with brief and straight forward questions such as the presence of aura or unilateral headache were perfectly consistent. Alternatively, symptoms that require verbal rephrasing such as phonophobia, photophobia, aggravation by or avoidance of routine physical activity were found to be highly discrepant between the CDE and SSI (Figure 3). Identification of these specific symptom domains with high discrepancy will help us develop a more robust and accurate next-generation CDE.

In general, predictive values are useful to answer “What is the probability that migraine will be present or absent in the context of a positive or negative CDE result?”⁶⁸ However, LRs tell us how much more likely a CDE result is in patients with migraine than it is in patients without migraine. The advantage of LRs over predictive values is their transferability and applicability beyond our study population.⁶⁹ Likelihood ratios can be beneficial directly at the individual patient level because they allow the clinician to quantitate the probability of migraine for any individual patient. By virtue of combining prevalence (pre-test probability) and LRs, the results from the

Fagan’s nomogram (Figure 2) provide the most useful and the most robust outcome measures.

In the simplest terms, if the ICHD-3 is transformed directly into a decision tree and uses a rule-based engine, then a history collected online should be both 100% sensitive and specific to a semi-structured interview diagnosis strictly adherent to the same ICHD-3 diagnostic criteria. The accuracy performance of the CDE was lower in distinguishing definite from probable migraine compared to discerning migraine from non-migraine. This discrepancy may be attributed to inconsistencies in patient responses to the same questions when asked by interview compared to completing self-response questionnaires. Headache specialists often rephrase the same question in different formats and approaches to ensure a consistent response is elicited—similar to building a patient’s case history. Embedding daily headache diaries within the CDE and SSI may also help reduce recall and/or response bias.

For example, a patient may initially deny the presence of photophobia or light sensitivity accompanying their headache attacks; however, the same patient may respond “Yes” when the interviewer rephrases the question if the patient prefers a darker room during a headache attack. This same patient may quickly click “No” and pass on to the next question without giving it a second thought when issued a self-administered question. These differences can create discrepancies in migraine diagnostic accuracy performance between the CDE and SSI. The CDE can address this through ML to reformulate and rephrase a question when the response does not appear consistent with the building data set.

Variations are expected when comparing psychometric properties from two modes of questionnaire administration.⁷⁰ Previous studies have shown interview-based diagnosis may be more accurate than self-administered questions due to lower cognitive demand for respondents, lower recall bias, better comprehension of the question, and the option of asking the interviewer to clarify the question.^{59,70-72} However, interviews may exhibit interviewer bias (e.g., interviewer-respondent rapport, communication style), acquiescence (yes-saying) bias, question order bias, as well as social desirability bias and lower willingness to disclose sensitive information compared to self-administered tools.^{59,70-72} Also, the channel of questionnaire presentation (auditory, oral, visual) impacts results.⁷⁰⁻⁷² The CDE uses a forced choice format, which helps gather a complete data set. Questionnaires are known to provide more complete data than traditional interviews.^{59,73,74} Patients can respond to the CDE at their own pace, which allows them to ask family members or relatives to contribute to some aspects of the responses, to check medicines, and to take breaks. The stigma and anxiety surrounding migraine diagnosis may make digital health tools more appealing to some patients than face-to-face interviews. The Head-HUNT study validating telephonic SSI versus face-to-face interview found an agreement kappa of 0.79 (95% CI = 0.66–0.92) in 172 participants with headache; there was a lapse of 2–4 weeks between the telephonic SSI and face-to-face interview.⁷⁵ Although not a validation study, Russell et al. reported no significant differences between telephonic interview

and face-to-face diagnosis in 219 patients with self-reported migraine.⁷⁶ The differences in accuracy and psychometric property between telephonic SSI and face-to-face interview could emanate from the reasons mentioned above.

The CDE and the SSI should accurately reflect the ICHD-3 diagnostic criteria. The live interviewer can check the profile being created against the larger data set in the SSI; that is, the years of experience the expert brings to the interview. The computer equivalent of this body of experience is a larger data set against which to compare the responses of an incoming data set, identify a best fit, and then ask additional questions to confirm/reject that fit. By virtue of having an algorithm that benefits from continuous training, the next generation CDE is expected to progressively improve with a robust ML platform and refine its precision—similar to other ML diagnostic tools.^{77,78} The CDE is currently upgrading to incorporate this type of artificial intelligence.

The present study was not sufficiently powered to comment on headache types other than migraine. The CDE fits with ICHD-3 fifth-digit hierarchical classification. The same rule-based engine is applied to all ICHD-3 diagnostic criteria/CDE questions. In the absence of an adequately powered study looking at other headache diagnoses, this approach represents a significant step beyond presently available diagnostic instruments for headache diagnosis. Both the SSI and CDE contain important “red flags” screening questions for secondary headaches, as shown in File S1 for the SSI. Including these headache diagnostic elements is vital for generalizability and validity, particularly in a community or primary care setting, to assess accuracy of the CDE in relation to these elements.

Given the large discrepancy between the number of people with migraine compared to the number of headache specialists, a computer-based diagnostic tool can be implemented within the health-care system to aid primary care or emergency department providers in ascertaining accurate diagnoses thereby lowering the burden on neurology/headache providers. An accurate computer-based diagnostic tool can reduce inefficiencies in headache care and enable remote provision of headache management, leading to reduction in health-care-related cost, shortening diagnostic delay, and improving access to care.^{79–83} Accurate computer-based migraine diagnosis can decrease rate of misdiagnosis (false positive/negative cases or over/underdiagnoses) and improve effectiveness of triage systems for headache consultations.^{25,79–81} Reduction in false negative cases will be crucial for an early migraine diagnosis, avoiding diagnostic delay thereby minimizing the risk for progression to chronic migraine and medication overuse headache, which require more aggressive and expensive treatment.^{25,80,81} Additionally, it will help avoid anxiety of patients and their families about a missed migraine diagnosis.⁸⁴ On the other hand, reduction in false positive cases will lessen the financial burden associated with unnecessary referral and costly medications,^{79,81} lower the health-care resource waste,^{79–81} as well as alleviate unnecessary cyberchondria (anxiety from misdiagnosis by digital tools) from patients and their families.^{84–86}

This study conforms with Class I evidence for diagnostic accuracy studies as per the American Academy of Neurology classification scheme for the following reasons: by virtue of being a cross-sectional

study with prospective data collection; by having disease status determination (CDE) without knowledge of the diagnostic test result (SSI); by having clearly defined exclusion/inclusion criteria; and by having both the diagnostic test (CDE) and disease status (SSI) measured in at least 80% of participants.⁸⁷ According to QUADAS (Quality Assessment of Diagnostic Accuracy Studies)⁸⁸ and STARD,⁵⁷ it is recommended to avoid excluding “difficult-to-diagnose” patients to prevent overoptimistic diagnostic accuracy results. Likewise, it is recommended to avoid excluding “confirmed cases” to reduce underestimating diagnostic accuracy performance. Hence, our enrollment of participants with nearly equal chance of being undiagnosed and diagnosed cases reduces the potential risk of participant selection bias.

It is true that the “gold standard” of migraine diagnosis is based on a complete patient history accompanied by a physical examination.¹³ Usually, imaging or additional laboratory investigations are not necessary in primary headache diagnosis—unless rarely indicated following suspicion of a secondary headache disorder.¹³ This process, which requires a patient visit to a headache clinic, can take up to 1 h per patient, excluding time to travel and waiting times. The significance of our study is to ultimately shorten the delay in migraine diagnosis. Migraine diagnostic delay is due to the time-consuming traditional headache care delivery approach involving a clinic visit, shortage of headache-trained providers, and the growing burden of primary headache disorders worldwide.^{2,12,89,90} Given the increase in the global burden of migraine estimated to affect a billion people,⁹ it would not be possible to capture every patient with migraine seeking the traditional in-person clinical visits. Hence, the SSI phone interview of patient history is the closest approach analogous to the traditional method of migraine diagnosis—making it our preferred “gold standard” for remote diagnosis in our study. However, we anticipate that future versions of SSI-based “gold standard” references will include virtual neurological examination and possible actigraphy/wearable biosensors to have some level of objective assessment of the patient.^{91–93}

The limitations of our study include its generalizability to settings other than tertiary headache clinics and the community. Our study helped us to see how the CDE performed with complex patients from academic headache centers as well as patients from the general population with milder headache. Our convenience sampling method is another study limitation as it can create selection bias. Probability (e.g., random) sampling can avoid sampling bias and provide better statistical inferences than convenience sampling; however, the median age group and female preponderance in our study population offer some degree of representativeness of the general migraine population. We are currently conducting studies to evaluate additional psychometric properties of the SSI (e.g., inter-rater reliability) and CDE (e.g., test-retest reliability). To our knowledge, there are no prior studies published that measured inter-rater reliability rate for an SSI in adult migraine diagnosis. There are two pediatric headache studies with percent agreement range of 61%–83%.^{94,95} The CDE diagnosis for migraine needs to be tested for intraindividual test-retest reliability, for example, within a 6-month period. However, the longer the test-retest period gap, the lower the test-retest reliability can be for episodic and chronic migraine—because migraine is an unstable condition that fluctuates

over time.⁹⁶ To our knowledge, there is no headache-specific published study that measured test–retest reliability of a computerized headache diagnostic tool. An example of test–retest correlation coefficients in seven computer-administered neurobehavioral measure scores ranged from 0.60 to 0.92.⁹⁷ The CDE is limited to anglophone patients; voice command and translations to other languages (currently underway) can help its implementation in diverse patient populations. Another study limitation is the potential for respondent fatigue in relation to time-to-complete the CDE questions, given that it can take more than 40 min on average to complete the CDE. We did not compare data quality and reliability in relation to time-to-complete the CDE. Our sample size may not be adequate to accommodate the stratified analysis results shown for community, newly diagnosed participants, and known cases with confirmed diagnoses; we have not conducted post hoc power analysis to examine the stratified analysis.

Our study featured low risk of bias⁸⁸ in the flow and timing of participants; that is, most participants received both the index and reference tests within an appropriate interval. The CDE index test showed low risk of bias (index test was interpreted without knowledge of reference test result) and low concern of applicability (its conduct or interpretation). Similarly, our reference test (SSI) exhibited low risk of bias. Given the fact that the SSI interviewers had the option to introduce new questions, there may be some concern about conduct or interpretation of the SSI reference test—particularly in terms of inter-rater agreement. In the absence of inter-rater reliability statistics, it is difficult to rule in/out concern about conduct or interpretation of the SSI reference.

That both the CDE and SSI were developed based on the standard headache criteria (i.e., ICHD-3), and that our reference standard was interview based are strengths of our study. This study provides initial testing of the CDE, which we plan to further validate in a larger, randomly sampled population as well as in headache patients presenting at primary care settings. Moreover, the logic and dataset of the CDE are expected to improve with “experience”; and because it is searchable, it opens the door for systematic subset analyses within diagnostic categories.

CONCLUSION

This study demonstrates that a novel computer-based algorithm utilizing ML can reliably and consistently apply the logic of a semi-structured interview, executed by a trained headache specialist, in a way that is scalable and capable of refinement with experience.

AUTHOR CONTRIBUTIONS

Study concept and design: Robert P. Cowan, Alan M. Rapoport, Jim Blythe, Yohannes W. Woldeamanuel. *Acquisition of data:* Robert P. Cowan, John Rothrock, Kerry Kniewel, Addie M. Peretz, Elizabeth Ekpo, Bharati M. Sanjanwala, Yohannes W. Woldeamanuel. *Analysis and interpretation of data:* Robert P. Cowan, Jim Blythe, Yohannes W. Woldeamanuel. *Drafting of the manuscript:* Yohannes W. Woldeamanuel. *Revising it for intellectual content:* Robert P. Cowan, Alan M. Rapoport, Jim Blythe,

John Rothrock, Kerry Kniewel, Addie M. Peretz, Elizabeth Ekpo, Bharati M. Sanjanwala, Yohannes W. Woldeamanuel. *Final approval of the completed manuscript:* Robert P. Cowan, Alan M. Rapoport, Jim Blythe, John Rothrock, Kerry Kniewel, Addie M. Peretz, Elizabeth Ekpo, Bharati M. Sanjanwala, Yohannes W. Woldeamanuel.

CONFLICT OF INTEREST

Robert P. Cowan, Alan M. Rapoport, Jim Blythe hold equity in the company that developed the CDE (Computer-based Diagnostic Engine). John Rothrock, Kerry Kniewel, Addie Peretz, Elizabeth Ekpo, Bharati M. Sanjanwala, Yohannes W. Woldeamanuel: no conflict.

ETHICAL CLEARANCE

Ethical clearance from institutional review boards of the three centers and written informed consent from all participants was collected prior to study enrollment.

DATA AVAILABILITY STATEMENT

Qualified researchers may obtain access to all anonymized data used for this study.

ORCID

Yohannes W. Woldeamanuel  <https://orcid.org/0000-0003-4879-6098>

REFERENCES

1. Vos T, Lim SS, Abbafati C, et al. Global burden of 369 diseases and injuries in 204 countries and territories, 1990–2019: a systematic analysis for the Global Burden of Disease Study 2019. *Lancet*. 2020;396(10258):1204–1222.
2. Buse DC, Armand CE, Charleston L, et al. Barriers to care in episodic and chronic migraine: results from the Chronic Migraine Epidemiology and Outcomes Study. *Headache*. 2021;61(4):628–641.
3. Viticchi G, Falsetti L, Bartolini M, et al. Migraine: incorrect self-management for a disabling disease. *Neurol Int*. 2018;10(1):7510.
4. Buse DC, Rupnow MFT, Lipton RB. Assessing and managing all aspects of migraine: migraine attacks, migraine-related functional impairment, common comorbidities, and quality of life. *Mayo Clin Proc*. 2009;84(5):422–435.
5. Adams AM, Serrano D, Buse DC, et al. The impact of chronic migraine: the Chronic Migraine Epidemiology and Outcomes (CaMEO) study methods and baseline results. *Cephalalgia*. 2015;35(7):563–578.
6. Peres MFP, Swerts DB, de Oliveira AB, Silva-Neto RP. Migraine patients' journey until a tertiary headache center: an observational study. *J Headache Pain*. 2019;20(1):88.
7. Steiner TJ, Antonaci F, Jensen R, Lainez MJA, Lanteri-Minet M, Valade D. Recommendations for headache service organisation and delivery in Europe. *J Headache Pain*. 2011;12(4):419–426.
8. www.ucns.org. Accessed June 3, 2021.
9. GBD 2016 Disease and Injury Incidence and Prevalence Collaborators. Global, regional, and national incidence, prevalence, and years lived with disability for 328 diseases and injuries for 195 countries, 1990–2016: a systematic analysis for the Global Burden of Disease Study 2016. *Lancet*. 2017;390(10100):1211–1259.
10. Kienbacher C, Wöber C, Zesch HE, et al. Clinical features, classification and prognosis of migraine and tension-type headache in children and adolescents: a long-term follow-up study. *Cephalalgia*. 2006;26(7):820–830.

11. Galinski M, Sidhoum S, Cimerman P, Perrin O, Annequin D, Tourniaire B. Early diagnosis of migraine necessary in children: 10-year follow-up. *Pediatr Neurol*. 2015;53(4):319-323.
12. Viticchi G, Silvestrini M, Falsetti L, et al. Time delay from onset to diagnosis of migraine. *Headache*. 2011;51(2):232-236.
13. Headache Classification Committee of the International Headache Society (IHS). The International Classification of Headache Disorders, 3rd edition. *Cephalalgia*. 2018;38(1):1-211.
14. Finkelstein J, Knight A, Marinopoulos S, et al. Enabling patient-centered care through health information technology. *Evid Rep Technol Assess (Full Rep)*. 2012;206:1-1531.
15. Petersen CL, Weeks WB, Norin O, Weinstein JN. Development and implementation of a person-centered, technology-enhanced care model for managing chronic conditions: cohort study. *JMIR Mhealth Uhealth*. 2019;7(3):e11082.
16. Granström E, Wannheden C, Brommels M, Hvitfeldt H, Nyström ME. Digital tools as promoters for person-centered care practices in chronic care? Healthcare professionals' experiences from rheumatology care. *BMC Health Serv Res*. 2020;20(1):1108.
17. Slevin P, Kessie T, Cullen J, Butler MW, Donnelly SC, Caulfield B. Exploring the potential benefits of digital health technology for the management of COPD: a qualitative study of patient perceptions. *ERJ Open Res*. 2019;5(2):00239-2018.
18. Mesko B, Györffy Z. The rise of the empowered physician in the digital health era: viewpoint. *J Med Internet Res*. 2019;21(3):e12490.
19. Orton M, Agarwal S, Muhoza P, Vasudevan L, Vu A. Strengthening delivery of health services using digital devices. *Glob Health Sci Pract*. 2018;6(suppl 1):S61-S71.
20. Marra C, Chen JL, Coravos A, Stern AD. Quantifying the use of connected digital products in clinical research. *NPJ Digit Med*. 2020;3(1):50.
21. Meskó B, Drobni Z, Bényei É, Gergely B, Györffy Z. Digital health is a cultural transformation of traditional healthcare. *mHealth*. 2017;3:38.
22. Burchert S, Kerber A, Zimmermann J, Knaevelsrud C. Screening accuracy of a 14-day smartphone ambulatory assessment of depression symptoms and mood dynamics in a general population sample: comparison with the PHQ-9 depression screening. *PLoS One*. 2021;16(1):e0244955.
23. Behar J, Roebuck A, Shahid M, et al. SleepAp: an automated obstructive sleep apnoea screening application for smartphones. *IEEE J Biomed Health Inform*. 2015;19(1):325-331.
24. Duarte-Rojo A, Allampati S, Thacker LR, et al. Diagnosis of covert hepatic encephalopathy: a multi-center study testing the utility of single versus combined testing. *Metab Brain Dis*. 2019;34(1):289-295.
25. Woldeamanuel YW, Cowan RP. Computerized migraine diagnostic tools: a systematic review. *Ther Adv Chronic Dis*. 2022;13:204062232110652.
26. Headache Classification Committee of the International Headache Society. Classification and diagnostic criteria for headache disorders, cranial neuralgias and facial pain. *Cephalalgia*. 1988;8(suppl 7):1-96.
27. Classification of headache. *Arch Neurol*. 1962;6(3):173.
28. Penzien DB, Andrew ME, Knowlton GE, et al. Computer-aided system for headache diagnosis with the IHS headache diagnostic criteria: development and validation. *Cephalalgia*. 1991;11(suppl 11):325-326.
29. Gallai V, Sarchielli P, Alberti A, et al. Application of the 1988 International Headache Society diagnostic criteria in Nine Italian Headache Centers using a computerized structured record. *Headache*. 2002;42(10):1016-1024.
30. Çelik U, Yurtay N, Koç ER, Tepe N, Güllüoğlu H, Ertaş M. Diagnostic accuracy comparison of artificial immune algorithms for primary headaches. *Comput Math Methods Med*. 2015;2015:1-8.
31. Simić S, Banković Z, Villar JR, Simić D, Simić SD. A hybrid fuzzy clustering approach for diagnosing primary headache disorder. *Log J IGPL*. 2021;29(2):220-235.
32. Roesch A, Dahlem MA, Neeb L, Kurth T. Validation of an algorithm for automated classification of migraine and tension-type headache attacks in an electronic headache diary. *J Headache Pain*. 2020;21(1):75.
33. Bentivegna E, Tassorelli C, de Icco R, Sances G, Martelletti P. Telehealthcare in migraine medicine: from diagnosis to monitoring treatment outcomes. *Expert Rev Neurother*. 2022;22(3):237-243.
34. Samaan Z, MacGregor EA, Andrew D, McGuffin P, Farmer A. Diagnosing migraine in research and clinical settings: the validation of the Structured Migraine Interview (SMI). *BMC Neurol*. 2010;10(1):7.
35. Rasmussen BK, Jensen R, Olesen J. Questionnaire versus clinical interview in the diagnosis of headache. *Headache*. 1991;31(5):290-295.
36. Lipton RB, Dodick D, Sadovsky R, et al. A self-administered screener for migraine in primary care. *Neurology*. 2003;61(3):375-382.
37. Pelaccia T, Forestier G, Wemmer C. Deconstructing the diagnostic reasoning of human versus artificial intelligence. *CMAJ*. 2019;191(48):E1332-E1335.
38. Pelaccia T, Tardif J, Tribi E, Charlin B. An analysis of clinical reasoning through a recent and comprehensive approach: the dual-process theory. *Med Educ Online*. 2011;16:5890.
39. Hayes-Roth F, Waterman D, Lenat D. *Building Expert Systems*. Addison-Wesley Longman Publishing Co., Inc.; 1983.
40. Lavrakas PJ. Respondent fatigue. In: *Encyclopedia of Survey Research Methods*. Vol. 2. SAGE Publications; 2008:742-743. Accessed January 9, 2022. <https://link.gale.com/apps/doc/CX3073300494/GVRL?u=stan90222&sid=bookmark-GVRL&xid=9ceeb36a>
41. *Clear & Simple*. National Institutes of Health; 2021. Accessed December 12, 2021. <https://www.nih.gov/institutes-nih/nih-office-director/office-communications-public-liaison/clear-communication/clear-simple>
42. Weiss B. *Health Literacy and Patient Safety: Help Patients Understand. Manual for Clinicians*. 2nd ed. American Medical Association; 2007. Accessed December 12, 2021. <http://www.partnershiphp.org/Providers/HealthServices/Documents/Health%20Education/CandLToolkit/2%20Manual%20for%20Clinicians.pdf>
43. Raynor W. *The International Dictionary of Artificial Intelligence*. Glendale Publishing Company, Ltd.; 1999.
44. *50 AI Terms Every Beginner Should Know*; 2021. Accessed April 1, 2022. <https://www.telusinternational.com/articles/50-beginner-ai-terms-you-should-know>
45. Sanin L, Mathew N, Bellmeyer L, Ali S. The International Headache Society (IHS) headache classification as applied to a headache clinic population. *Cephalalgia*. 1994;14(6):443-446.
46. Fejes E, Feher G, Gurdan Z, et al. Characteristics of patients referred to a specialized headache clinic. *Sci Rep*. 2020;10(1):1146.
47. Dong Z, Di H, Dai W, et al. Application of ICHD-II criteria in a headache clinic of China. *PLoS One*. 2012;7(12):e50898.
48. Hajian-Tilaki K. Sample size estimation in diagnostic test studies of biomedical informatics. *J Biomed Inform*. 2014;48:193-204.
49. Newcombe RG. Two-sided confidence intervals for the single proportion: comparison of seven methods. *Stat Med*. 1998;17(8):857-872.
50. Chow S-C, Shao J, Wang H, Lokhnygina Y. *Sample Size Calculations in Clinical Research*, 3rd ed. Chapman and Hall/CRC; 2017.
51. Julious SA. *Sample Sizes for Clinical Trials*, 1st ed. Chapman and Hall/CRC; 2009.
52. Clopper CJ, Pearson ES. The use of confidence or fiducial limits illustrated in the case of the binomial. *Biometrika*. 1934;26(4):404-413.
53. Cohen J. A coefficient of agreement for nominal scales. *Educ Psychol Measur*. 1960;20(1):37-46.
54. Woldeamanuel YW, Cowan RP. Migraine affects 1 in 10 people worldwide featuring recent rise: a systematic review and meta-analysis of community-based studies involving 6 million participants. *J Neurol Sci*. 2017;372:307-315.

55. Caraguel CGB, Vanderstichel R. The two-step Fagan's nomogram: ad hoc interpretation of a diagnostic test result without calculation. *Evid Based Med*. 2013;18(4):125-128.
56. Fagan T. Nomogram for Bayes's theorem. *N Engl J Med*. 1975; 293(5):257.
57. Bossuyt PM, Reitsma JB, Bruns DE, et al. STARD 2015: an updated list of essential items for reporting diagnostic accuracy studies. *BMJ*. 2015;351:h5527.
58. Mayne JG. A health questionnaire based on paper-and-pencil medium individualized and produced by computer. *JAMA*. 1969;208(11):2060.
59. Bachman JW. The patient-computer interview: a neglected tool that can aid the clinician. *Mayo Clin Proc*. 2003;78(1):67-78.
60. Beckman HB. The effect of physician behavior on the collection of data. *Ann Intern Med*. 1984;101(5):692. doi:10.7326/0003-4819-101-5-692
61. Lilford RJ, Bourne G, Chard T. Comparison of information obtainable by computerized and manual questionnaires in an antenatal clinic. *Med Inform*. 1982;7(4):315-320.
62. Sturges JE, Hanrahan KJ. Comparing telephone and face-to-face qualitative interviewing: a research note. *Qual Res*. 2004;4(1):107-118.
63. Opendakker R. Advantages and disadvantages of four interview techniques in qualitative research. *Forum Qual Soc Res*. 2006;7(4):1-13.
64. Tausig JE, Freeman EW. The next best thing to being there: conducting the clinical research interview by telephone. *Am J Orthopsychiatry*. 1988;58(3):418-427.
65. Burnard P. The telephone interview as a data collection method. *Nurse Educ Today*. 1994;14(1):67-72.
66. Kavanaugh K, Ayres L. "Not as bad as it could have been": assessing and mitigating harm during research interviews on sensitive topics. *Res Nurs Health*. 1998;21(1):91-97.
67. Novick G. Is there a bias against telephone interviews in qualitative research? *Res Nurs Health*. 2008;31(4):391-398.
68. Altman DG, Bland JM. Statistics notes: diagnostic tests 2: predictive values. *BMJ*. 1994;309(6947):102.
69. Deeks JJ, Altman DG. Diagnostic tests 4: likelihood ratios. *BMJ*. 2004;329(7458):168-169.
70. Bowling A. Mode of questionnaire administration can have serious effects on data quality. *J Public Health*. 2005;27(3):281-291.
71. Tourangeau R, Rips LJ, Rasinski K. *The Psychology of Survey Response*. Cambridge University Press; 2000.
72. Bolarinwa O. Principles and methods of validity and reliability testing of questionnaires used in social and health science researches. *Niger Postgrad Med J*. 2015;22(4):195.
73. Hall GH. Experiences with outpatient medical questionnaires. *BMJ*. 1972;1(5791):42-45.
74. Inui TS, Jared RA, Carter WB, et al. Effects of a self-administered health history on new-patient visits in a general medical clinic. *Med Care*. 1979;17(12):1221-1228.
75. Hagen K, Zwart J-A, Vatten L, Stovner L, Bovim G. Head-HUNT: validity and reliability of a headache questionnaire in a large population-based study in Norway. *Cephalgia*. 2000;20(4):244-251.
76. Russell MB, Rasmussen BK, Thorvaldsen P, Olesen J. Prevalence and sex-ratio of the subtypes of migraine. *Int J Epidemiol*. 1995;24(3):612-618.
77. Deodhar A, Rozycki M, Garges C, et al. Use of machine learning techniques in the development and refinement of a predictive model for early diagnosis of ankylosing spondylitis. *Clin Rheumatol*. 2020;39(4):975-982.
78. Richens JG, Lee CM, Johri S. Improving the accuracy of medical diagnosis with causal machine learning. *Nat Commun*. 2020;11(1):3923.
79. Müller KI, Alstadhaug KB, Bekkelund SI. Acceptability, feasibility, and cost of telemedicine for nonacute headaches: a randomized study comparing video and traditional consultations. *J Med Internet Res*. 2016;18(5):e140.
80. Müller KI, Alstadhaug KB, Bekkelund SI. A randomized trial of telemedicine efficacy and safety for nonacute headaches. *Neurology*. 2017;89(2):153-162.
81. Stubberud A, Linde M. Digital technology and mobile health in behavioral migraine therapy: a narrative review. *Curr Pain Headache Rep*. 2018;22(10):66.
82. Qubty W, Patniyot I, Gelfand A. Telemedicine in a pediatric headache clinic: a prospective survey. *Neurology*. 2018;90(19):e1702-e1705.
83. Davis LE, Harnar J, LaChey-Barbee LA, Pirio Richardson S, Fraser A, King MK. Using teleneurology to deliver chronic neurologic care to rural veterans: analysis of the first 1100 patient visits. *Telemed J E Health*. 2019;25(4):274-278.
84. Meyer AND, Giardina TD, Spitzmueller C, Shahid U, Scott TMT, Singh H. Patient perspectives on the usefulness of an artificial intelligence-assisted symptom checker: cross-sectional survey study. *J Med Internet Res*. 2020;22(1):e14679.
85. White RW, Horvitz E. Cyberchondria: studies of the escalation of medical concerns in web search. *ACM Trans Inform Syst*. 2009;27(4):1-37.
86. Vismara M, Caricasole V, Starcevic V, et al. Is cyberchondria a new transdiagnostic digital compulsive syndrome? A systematic review of the evidence. *Compr Psychiatry*. 2020;99:152167.
87. Gronseth G, Cox J, Gloss D, on behalf of the Guideline Development D and IS of the American Academy of Neurology. *Clinical Practice Guideline Process Manual*. The American Academy of Neurology; 2017.
88. Whiting PF. QUADAS-2: a revised tool for the quality assessment of diagnostic accuracy studies. *Ann Intern Med*. 2011;155(8):529.
89. Viticchi G, Falsetti L, Bartolini M, et al. Migraine: incorrect self-management for a disabling disease. *Neurol Int*. 2018;10(1):24-27.
90. Woldeamanuel YW, Cowan RP. Migraine affects 1 in 10 people worldwide featuring recent rise: a systematic review and meta-analysis of community-based studies involving 6 million participants. *J Neurol Sci*. 2017;372:307-315.
91. Moccia M, Lanzillo R, Brescia Morra V, et al. Assessing disability and relapses in multiple sclerosis on tele-neurology. *Neurol Sci*. 2020;41(6):1369-1371.
92. Roy B, Nowak RJ, Roda R, et al. Teleneurology during the COVID-19 pandemic: a step forward in modernizing medical care. *J Neurol Sci*. 2020;414:116930.
93. Boes CJ, Leep Hunderfund AN, Martinez-Thompson JM, et al. Primer on the in-home teleneurologic examination. *Neurol Clin Pract*. 2021;11(2):e157-e164.
94. Hershey AD, Winner P, Kabbouche MA, et al. Use of the ICHD-II criteria in the diagnosis of pediatric migraine. *Headache*. 2005;45(10):1288-1297.
95. Tsze DS, Cruz AT, Mistry RD, et al. Interobserver agreement in the assessment of clinical findings in children with headaches. *J Pediatr*. 2020;221:207-214.
96. Serrano D, Lipton RB, Scher AI, et al. Fluctuations in episodic and chronic migraine status over the course of 1 year: implications for diagnosis, treatment and clinical trial design. *J Headache Pain*. 2017;18(1):101.
97. Campbell KA, Rohlman DS, Storzbach D, et al. Test-retest reliability of psychological and neurobehavioral tests self-administered by computer. *Assessment*. 1999;6(1):21-32.

SUPPORTING INFORMATION

Additional supporting information may be found in the online version of the article at the publisher's website.

How to cite this article: Cowan RP, Rapoport AM, Blythe J, et al. Diagnostic accuracy of an artificial intelligence online engine in migraine: A multi-center study. *Headache*. 2022;62:870–882. doi:10.1111/head.14324