A Systematic Review of Instruments to Identify Mental Health and Substance Use Problems Among Children in the Emergency Department

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ABSTRACT

Objective: Specialized instruments to screen and diagnose mental health problems in children and adolescents are not yet standard components of clinical assessments in emergency departments (EDs). We conducted a systematic review to investigate the psychometric properties, accuracy, and performance metrics of instruments used in the ED to identify pediatric mental health and substance use problems.

Methods: We searched seven electronic databases and the gray literature for psychometric validation studies, diagnostic studies, and cohort studies that assessed any instrument to screen for or diagnose mental illness, emotional or behavioral problems, or substance use disorders. Studies had to include children and adolescents with mental health presentations or positive screens for substance use. Two reviewers independently screened studies for relevance and quality. Diagnostic study quality was assessed with the four QUADAS-2 domains. Psychometric study quality was assessed with published criteria for instrument reliability, validity, and usability. We present a descriptive analysis of the reported psychometric properties and diagnostic performance of instruments for each study.

Results: Of the 4,832 references screened, 14 met inclusion criteria. Included studies evaluate 18 instruments for identifying suicide risk (six studies), alcohol use disorders (six studies), mood disorders (one study), and ED decision making (need for assessment, admission; one study). Nine studies include a psychometric focus but quality varies, with no studies fully meeting criteria for reliability, validity, and usability. Seven studies examine diagnostic performance of an instrument, but no study has a low risk of bias for all QUADAS-2 domains. The HEADS-ED instrument has good inter-rater reliability (r = 0.785) for identifying general mental health problems and modest evidence for ruling in patients requiring hospital admission (positive likelihood ratio [LR+] = 6.30). Internal consistency (reliability) varies for instruments to screen for suicide risk ($\alpha = 0.46-0.97$), and no instruments have both high sensitivity and high specificity. The Ask Suicide-Screening Questions (ASQ) is highly sensitive (98%) and has strong evidence for ruling out risk (negative likelihood ratio [LR-] = 0.04). Among screening instruments for alcohol use disorders, internal consistency is high for the consumption subscale of the Alcohol Use Disorders Identification Test ($\alpha = 0.83-0.88$) and the Adolescent Drinking Index ($\alpha = 0.92$). Both instruments also had sound internal validity. Diagnostically, a two-item instrument based on DSM-IV criteria is the most accurate in identifying patients

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while considering whether the instruments can address setting-related barriers and professional concerns.

with a disorder (area under the curve = 0.89) and has modest evidence for ruling in and out risk (LR+ = 8.80, LR - = 0.13). Conclusions: From available evidence, we recommend that ED clinicians use 1) the HEADS-ED to rule in ED

admission among pediatric patients with visits for mental health care, 2) the ASQ to rule out suicide risk among pediatric patients with any visit type, and 3) the DSM-IV two-item instrument to rule in/rule out alcohol use disorders among pediatric patients currently using alcohol. These instruments require minimal to no training or time commitment. We also recommend that clinicians become familiar with each instrument's psychometric properties to understand the guality of the evidence base. In this review, however, we identify methodologic limitations in the evidence base. To develop a robust evidence base, additional research is necessary.

M ental health care visits by children and adoles-cents to general and pediatric emergency departments (EDs) have increased significantly over the past 10 years in the United States.^{1,2} These children and adolescents may or may not have been previously diagnosed with a mental disorder or problem.³ Using specialized instruments to identify mental health problems among children and adolescents during ED care is supported by the American Academy of Pediatrics (AAP) and the Committee on Pediatric Emergency Medicine.³ However, we do not know whether such instruments are a standard component of clinical assessments by ED clinicians in general and pediatric EDs. Research suggests that using instruments to identify mental health problems among pediatric ED patients is not widespread⁴⁻⁶ even if the ED visit is mental health related.⁶

Two setting-related barriers to using specialized instrument in screening for mental health problems are absence of validated instruments and time limitations.⁶ ED physicians and nurses also raise professional concerns about their lack of knowledge, lack of training, and high discomfort with mental health care.^{6,7} In 2011, the AAP recognized several instruments, including those for assessing suicide risk, as addressing both setting-related barriers and professional concerns.³ These instruments are evidencebased, have sound psychometric and/or diagnostic properties, and can be rapidly deployed at the bedside and by nonspecialists. More recently, several other instruments have been evaluated with pediatric mental health patients to identify concerns related to mental health during an ED visit (e.g., Cappelli et al.,⁸ Gipson et al.⁹).

This systematic review has two objectives: 1) to evaluate the psychometric properties and accuracy of instruments intended for children and adolescents who present to the ED with a mental health concern and 2) to provide a more definitive knowledge base on the performance metrics of available instruments

METHODS

Study Design

A protocol for the review was developed and registered with PROSPERO (Registration # CRD42016033708). Reporting of the review adheres to the PRISMA statement checklist.¹⁰ Funding for the review was provided by the Emergency Strategic Clinical Network of Alberta Health Services (Alberta, Canada). In-kind support of research librarian services was provided by the Alberta SPOR SUPPORT Unit, Knowledge Translation platform. These funding organizations had no involvement in any aspect of the conduct, analysis, and manuscript preparation of this review.

Search Strategy

A research librarian developed the search strategies for the following databases: MEDLINE, Embase, CEN-TRAL, EBM Reviews (Cochrane Database of Systematic Reviews, Health Technology Assessment Database and Database of Abstracts of Reviews of Effects), PsycINFO, CINAHL, Social Services Abstracts, and Pro-Quest Theses and Dissertations. Restrictions were language (English) and date (2000–2015, including literature published up to October 1-14, 2015). The gray literature was searched using Google Scholar, Health Services Research Projects in Progress, Health Services/Sciences Research Resources, and Health Services/Technology Assessment Texts. Reference lists of relevant reviews were searched as well. Clinical trials were searched using www.clinicaltrials.gov. Also searched were conference proceedings of the past two vears (2014 and 2015) from the American College of Emergency Physicians, Canadian Association of Emergency Physicians, and Emergency Medicine Academy. Data Supplement S1 (available as supporting information in the online version of this paper) provides the search terms developed for the MEDLINE database.

Criteria for Considering Studies for This Review

This review aimed to include a broad range of mental health instruments that have been evaluated with pediatric mental health patients in the ED. Studies that recruited both nonpsychiatric and psychiatric patients were included because the study instrument was still used with mental health patients. Mental health instruments were defined as any instruments that could identify mental illness, emotional or behavioral disorders, substance use problems (substance use disorders or harmful/hazardous use; problems that have been associated with mental disorders¹¹), or suicide risk. To be included in the review, a study had to evaluate an instrument with children and adolescents (6-18 years) presenting to an ED with a mental health concern (psychiatric) and/or recent substance use. Studies of young adult and pediatric mental health patients were included if the mean age of the entire group was <20 years. Eligible study designs included psychometric, diagnostic, and cohort studies. Diagnostic studies compared an instrument to a reference standard, but a comparator was not required for psychometric studies to be included in this review. The primary outcomes of interest were 1) instrument validity and reliability for psychometric studies and 2) positive and negative likelihood ratios (LR+ and LR-, respectively) for diagnostic and cohort studies. Secondary outcomes of interest were 1) factor solution or factor loading (if factor solution was not available) and model fit for psychometric studies and 2) area under the curve for diagnostic and cohort studies.

Screening for Eligibility

References were organized and screened using End-Note X7.2.1. Two reviewers (AS, SWK) independently screened title and abstract for the first 100 articles in the EndNote library and then calculated inter-rater agreement with the kappa statistic.¹² Once a sufficiently high kappa was reached (≥ 0.8), the remaining references in the library were divided in two and each reviewer screened one half using title and abstract. Two reviewers (AS, SWK) independently reviewed the full text of studies that were identified as potentially relevant using our predetermined inclusion and exclusion criteria. Any discrepancies were discussed between the reviewers and taken to a third party (ASN) if no agreement could be reached.

Data Abstraction

Data were extracted using a standardized form that assessed study characteristics (e.g., language of publication, country), characteristics of the study population, study setting, instrument description and reference standard (diagnostic studies only), and results. Data were extracted by one reviewer (SWK) and reviewed for completeness and accuracy by another reviewer (AS). Discrepancies were resolved by discussion or by contacting corresponding authors of included studies.

Quality Assessment

Methodologic quality for all studies was assessed independently by two reviewers (SWK, AS). Disagreements were resolved by consensus or by involving a third reviewer (ASN) as required. The quality of instrument development reported in psychometric studies (appropriateness of analytic techniques and psychometric statistics) was assessed using a modified version of an evaluation tool for early childhood social-emotional screening and assessment measures (developed by Gokiert et al.).¹³ That tool includes criteria to assess instrument reliability, validity, and usability, with each category being assessed on multiple factors (see Data Supplement S2, available as supporting information in the online version of this paper). We modified the tool by excluding criteria that were not relevant to psychometric studies (e.g., sensitivity and specificity) or that were not relevant to an ED-based instrument (e.g., size of the early childhood standardization sample). Each study was evaluated on the psychometric evidence provided (e.g., no evidence provided, criterion not met, criterion met). When a study did not fully meet a criterion, the overall study quality was downgraded (see Data Supplement S2).

The methodologic quality and applicability of diagnostic studies was assessed using QUADAS-2.¹⁴ The QUADAS-2 reports on four domains: patient selection, index test, reference standard, and flow (of patients through the study) and timing (of the index test and reference standard). Signaling questions for each domain were answered to indicate whether a study had a low, high, or unclear risk of bias. If all signaling questions in a domain were answered "yes," then risk of bias was judged as low. If any signaling question in a domain was answered "no," the study was flagged as having a potential for bias. If any signaling question in a domain was answered "unclear" due to insufficient information, risk of bias was judged as unclear. Assessing applicability for each QUADAS-2 domain involved determining whether the study matched the objective of the systematic review. Applicability was judged as high (a match between study and review), low (no match), or unclear (insufficient evidence to determine a match).

Data Analysis

Evidence tables were developed to describe the studies, including information on study design, methodologic quality, study population and setting, sample size, instruments, and comparators. Psychometric outcome data were extracted from studies to determine each instrument's psychometric properties and compare them against set criteria (see Data Supplement S2).

Diagnostic data were extracted from articles to assess the sensitivity and specificity of each instrument. LR+ and LR- likelihood ratios were extracted to provide estimates of how instrument cut scores change the odds of a diagnosis. If sensitivity and specificity data were provided but LR+ and LR- were not, we calculated LR+ and LR- as LR+ = sensitivity/(1 - specificity) and LR = (1 - sensitivity)/specificity. LR+ expresses the strength of evidence available to "rule in" a diagnosis, whereas LR- indicates whether a diagnosis could be "ruled out." General guidelines for interpreting LR+ values to rule in a disorder are >10, strong evidence; 5 to 10, modest evidence; 2 to 5, weak evidence; and 0.5 to 2, no significant change in the likelihood of a disorder. General guidelines for interpreting LR- values to rule out a disorder are 0.2 to 0.5, weak evidence; 0.1 to 0.2, modest evidence; and <0.1, strong evidence.¹⁵ In this review we aimed to identify diagnostic instruments with large LR+ and small LR- values.

We examined area under the receiver operating characteristic curve to determine instrument accuracy. We used general guidelines for interpreting the values:



0.90–1 (excellent accuracy), 0.80–0.90 (good accuracy), 0.70–0.80 (fair accuracy), 0.60–0.70 (poor accuracy), and 0.50–0.60 (fail).¹⁶

We were unable to assess for publication bias due to the small numbers of studies for any given instrument. Heterogeneity in patient populations, clinical instruments, and outcome reporting limited our ability to conduct a meta-analysis. Instead, we conducted a descriptive analysis of the psychometric and diagnostic results of each study.

RESULTS

Literature Search and Selection

Our search strategy identified 7,425 citations, with 4,832 citations remaining after removing duplicates. Of these, 168 were considered potentially relevant based on their title and abstract (Figure 1). Full-text review identified 14 studies that met inclusion criteria; 10 provided psychometric data and seven provided diagnostic data.

Description of Included Studies

Table 1 presents 18 instruments that were evaluated in the 14 included studies. Instruments for identifying alcohol use disorders^{17–22} and suicide risk^{9,23–28} were assessed in six and seven studies, respectively, making these two clinical focuses the most common conditions assessed. Characteristics of included studies, stratified by clinical focus, are presented in Table 2. Three studies^{19,21,26} report early findings of later published studies.^{18,20,25}

Study Quality

Details on the quality of the studies that assess instrument psychometrics are provided in Data Supplement S3 (available as supporting information in the online version of this paper). Data Supplement S4 (available as supporting information in the online version of this paper) provides details on the quality of the studies that report diagnostic evaluations.

General Screening

One study presents a psychometric and diagnostic assessment of an instrument for general use.⁸ The instrument, the HEADS-ED, meets criteria for content and predictive validity and partially meets the criterion for convergent/concurrent validity. The criterion for inter-rater reliability is partially met. The instrument

Clinical Instruments Used in the Studies for the Identification of Mental Health and Alcohol Use Problems Among Children in the ED

General screening	
HEADS-ED	Home, Education, Activities/Peers, Drug/Alcohol, Suicidality, Emotions/ Behavior, Discharge Resources
Suicide risk	
ASQ	Ask Suicide-Screening Questions
C-SSRS	Columbia-Suicide Severity
0 00110	Bating Scale
RSQ	Risk of Suicide Questionnaire
Composite: SIQ-JR.	Suicide Ideation Questionnaire for
AUDIT-C. BADS-2	patients aged 13 or 14 years.
	Alcohol Use Disorders Identification
	Test-Consumption subscale.
	Revnolds Adolescent
	Depression Scale
SQS	Single-Question Screen
TQS	Two-Question Screen
Alcohol use disorders	
ADI	Adolescent Drinking Index
AUDIT	Alcohol Use Disorders Identification
AUDIT-C	Alconol Use Disorders Identification
	Lest-Consumption subscale
CAGE	Cut down, Annoyed, Guilty,
ODALET	Eye-opener
GRAFFI	Trouble
DSM-IV two-item scale	(1) Alcohol abuse (drinking in
	hazardous situations);
	(2) Alcohol dependence
FAST	Fast Alcohol Screening Test
RAFFT	Relax, Alone, Friends, Family, Trouble
RAPS4-QF/RAPS-QF	Remorse, Amnesia/blackouts,
	Perform, Starter/eye-opener,
	Quantity, Frequency
RUFT-Cut	Riding with a drinking driver,
	Unable to stop, Family/Friends,
	Trouble, Cut down
TWEAK	Tolerance, Worried, Eye-opener,
	Amnesia, Kut-down

requires minimal training in assessment and interpretation. The QUADAS-2 assessment shows a low risk of bias for the approach to patient selection and for flow and timing in the diagnostic study of the HEADS-ED. In the study, the risk of bias is unclear for how the index test and reference standard were used.⁸

Suicide Risk

Four studies provide psychometric data on three instruments that assess for risk of suicide: C-SSRS,⁹ RSQ,^{25,26} and a composite of several instruments (SIQ-JR, AUDIT-C, RADS-2).²⁸ The C-SSRS meets criteria for content validity and internal consistency and partially meets the criterion for predictive validity. No evidence is provided for other forms of reliability and validity. The C-SSRS has an administration time of less than 15 minutes and requires minimal training in assessment and interpretation. The RSQ meets the

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Overview of Included Studies

				Clinical U	Jsability
Author, Year, Country	Study Design (Enrollment)	Participants: Inclusion and Exclusion Criteria	Instrument	Administrator During the Study	Time to Complete
Assessment of G Cappelli et al., 2012, ⁸ Canada	eneral Mental Health S Prospective cohort (consecutive)	Status and Well-being Inclusion: Patients aged < 18 y with a mental health concern. Exclusion: Patients with a need for immediate medical or surgical care.	HEADS-ED	Crisis intervention worker, research assistant	Varies
Gipson et al., 2015, ⁹ US	Prospective cohort (convenience)	Inclusion: Patients aged 13–17 y seeking emergency psychiatric services.	C-SSRS	Social worker, physician	Range = 1–2 to 5–10 min
Horowitz et al., 2012, ²⁴ US	Prospective cohort (convenience)	Inclusion: Patients aged 10–21 y with either medical/surgical or psychiatric concerns. Exclusion: Patients with any disorder preventing an ability to comprehend questions or relay answers, triage level 1 suggesting physiologic instability, or parent/guardian unavailable or did not speak English	ASQ	Research assistant	<2 min
Folse and Hahn, 2009, ²⁵ US Folse et al., 2006, ²⁶ US	Prospective cohort (convenience)	Inclusion: Patients aged > 12 y who were seeking psychiatric or nonpsychiatric care, who were medically stable and could understand English. Exclusion: Patients for whom privacy conditions supported a discussion without risk of being overheard by others in the ED	RSQ	Registered nurse	90 s
Horowitz et al., 2001, ²⁷ US	Prospective cohort (consecutive)	Inclusion: Patients aged < 18 y seeking emergency care primarily for psychiatric reasons. Exclusion: Patients with impairments that would prevent completion of the assessment tools or missing data on the tool or criterion standard	RSQ	Triage nurse	<2 min
King et al., 2009, ²⁸ US	Prospective cohort (convenience)	Inclusion: Patients aged 13–17 y seeking pediatric or psychiatric emergency services. Exclusion: Patients with severe cognitive impairment, abnormal vital signs, or parent/guardian unavailable or did not speak English	Composite: SIQ-JR, AUDIT-C, RADS-2	Self-report	5–10 min
Rutman et al., 2008, ²³ US	Prospective cohort (convenience)	<i>Inclusion:</i> Patients aged 12–17 y with injury, illness, and/or psychiatric concerns. <i>Exclusion:</i> Patients with critical illness or injury, developmental delay, or intoxication or if patient or parent/ guardian did not speak English.	SQS TQS	Research assistant, study physician	<1 min
Assessment of A Hernandez et al., 2014, ¹⁷ US	Iconol Use Disorders Prospective cohort (convenience)	<i>Inclusion:</i> Patients aged 13–17 y who reported consuming alcohol within 6 h of their ED admission, had a positive blood alcohol test or breathalyzer reading, and/or scored four or above on the AUDIT. <i>Exclusion:</i> Patients who were suicidal, in police custody, requiring hospitalization, or whose medical care interrupted the screening process.	ADI	Self-report	<5 min

Table 2 (continued)

				Clinical	Usability
Author, Year, Country	Study Design (Enrollment)	Participants: Inclusion and Exclusion Criteria	Instrument	Administrator During the Study	Time to Complete
Kelly et al., 2009, ¹⁸ US Kelly et al.,	Prospective cohort (nonconsecutive)	Inclusion: Patients aged 18–20 y, who received medical clearance, currently used alcohol. Adolescents could be alcohol positive at the time of ED presentation. Exclusion: Patients with a Glasgow Coma Score < 15, not accompanied by a parent/guarding if < 10 u	AUDIT-C, CRAFFT, RAPS4-QF, FAST, RUFT- Cut, DSM-IV 2- item scale AUDIT, CAGE,	Self-report	1–5 min*
2004, 05		by a parent/guardian if < 18 y, admitted due to suspected child abuse or neglect, unable to read and complete a self-report questionnaire, or intoxicated and not oriented to time and place/able to provide informed consent.	RAPS-QF, RUFT-Cut		
Kelly et al., 2002, ²⁰ US Kelly and Donovan, 2001, ²¹ US	Prospective cohort (nonconsecutive)	Inclusion: Patients aged 12–20 y with no serious head injury, not seriously ill or critically injured, accompanied by a parent or legal guardian if < 18 y, able to read and complete a self-report questionnaire, who currently used alcohol and properly completed the questionnaire (e.g., provided usable data). Adolescents could be alcohol positive at the time of ED presentation. <i>Exclusion:</i> Patients too ill or critically	AUDIT, TWEAK, CAGE AUDIT	Self-report	1–2 min†
Bastiaens et al., 2000, ²² US	Prospective cohort (consecutive)	Injured to approach. Inclusion: Patients aged 13–18 y referred for psychiatric assessment. Exclusion: Patients who presented by themselves or who did not have a parental figure who had lived with them for at least 6 mo.	RAFFT	Triage nurse	<2 min

†2 minutes for AUDIT; <2 minutes for TWEAK; <1 minute for CAGE.

criterion for content validity, partially meets the criterion for criterion validity, and does not meet the criterion for internal consistency. No evidence is provided for other forms of reliability and validity. The RSQ has an administration time of less than 15 minutes and requires no training. The composite tool meets criteria for content and concurrent validity and for internal consistency. No evidence is provided for other forms of reliability and validity. The composite tool has an administration time of less than 15 minutes and requires minimal training in assessment and interpretation.

Three studies evaluated the diagnostic properties of four instruments: the ASQ,²⁴ RSQ,²⁷ SQS,²³ and TQS.²³ The ASQ shows a high risk of bias for methods of patient selection, an unclear risk of bias for how the index test and reference standard were used in the diagnostic evaluation and a low risk of bias for

flow and timing of instrument delivery. The RSQ shows a low risk of bias for all four domains: methods of patient selection, how the index test and reference standard were used in the diagnostic evaluation, and flow and timing of instrument delivery. The SQS and TQS show a high risk of bias for methods of patient selection, a low risk of bias for use of the index and reference standard, and a low risk of bias for flow and timing.

Alcohol Use Disorders

Five studies report on the psychometric properties of 11 different instruments to assess for alcohol use disorders: the ADI, AUDIT, AUDIT-C, CAGE, CRAFFT, RAPS4-QF/RAPS-QF, FAST, RUFT-Cut, TWEAK, and a DSM-IV two-item scale. The criterion for content validity is met by all instruments, and the criterion for construct validity is achieved for the ADI and AUDIT.^{17,21} The criterion for internal consistency is partially met for all 11 instruments.^{17–21} The criterion for criterion validity is met for the ADI and partially met for the AUDIT.^{17,20} One study evaluates concurrent validity of the AUDIT, TWEAK, and CAGE, with the instruments partially meeting the criterion.²⁰ All the instruments have an administration time of less than 15 minutes and require limited to no training.

Three studies evaluate the diagnostic properties of nine of the 11 instruments.^{18,19,22} One study²² considered risk of bias low for methods of patient selection and two studies^{18,19} considered the risk high. All three studies consider the risk of bias high for the index test or its conduct or interpretation. All studies consider the risk of bias low for the reference standard, its conduct, or its interpretation. Risk of bias for flow and timing is considered low²² or unclear.^{18,19}

Reliability and Validity of Instruments

The reliability and validity of instruments for use in the ED are presented in Table 3.

General Screening

The HEADS-ED is a reliable instrument when used by multiple ED clinicians (r = 0.785), with reliability of the instrument's seven domains ranging from r = 0.57 (Emotions and behaviors domain) to r = 0.90 (Drugs and alcohol domain). The instrument has also been shown to predict which pediatric mental health patients are in need of a full psychiatric assessment and admission to hospital.⁸

Suicide Risk

The C-SSRS is a reliable instrument ($\alpha = 0.81$ for the five-item intensity scale) with the ability to predict: 1) ED revisits for suicide attempts by adolescents who seek emergency mental health care (intensity scale score; odds ratio [OR] = 1.09; 95% confidence interval [CI] = 1.01–1.17) and 2) ED revisits for mental health care by adolescents who report suicidal ideation during their index ED visit (OR, 1.09; 95% CI = 1.00–1.19).⁹

The RSQ has been studied with adolescent patients in mental health ED visits, but psychometric properties largely reflect data from patients in nonpsychiatric visits.^{25,26} Two studies found modest reliability ($\alpha = 0.64$ and $\alpha = 0.65$) for the two-item RSQ rather than the four-item version. Correlations of RSQ items with psychiatric and suicide-related diagnoses vary. Question 1 ("Are you here because you tried to hurt yourself?") has the strongest correlations with psychiatric (r = 0.87 and r = 0.76) and suicide-related (r = 0.72 and r = 1.00) diagnoses.^{25,26} The composite instrument of suicide, alcohol-, and mood-related questions has excellent reliability, ranging from 0.89 to 0.97. Adolescents who screen at elevated risk of suicide using the composite instrument score higher on hopelessness than psychiatrically hospitalized adolescents.²⁸

Alcohol Use Disorders

The AUDIT and its consumption subscale, the AUDIT-C, are the most extensively studied instruments.^{18–21} The reliability of the full instrument is excellent ($\alpha = 0.83$ –0.88) as is reliability for the subscale ($\alpha = 0.81$). The full instrument has validity in distinguishing between adolescents with hazardous and nonhazardous drinking (p < 0.001); breathalyzer-positive and breathalyzer-negative adolescents (p < 0.02); and age, sex, and racial differences in alcohol consumption (all statistically significant). The underlying factor structure of the AUDIT is also sound (construct validity).

The 24-item ADI also demonstrates excellent reliability ($\alpha = 0.92$).¹⁷ Its underlying factor structure is sound, and it can distinguish between various alcohol and other drug use outcomes.¹⁷ Modest reliability has been reported for other alcohol use disorder instruments: the CAGE, TWEAK, CRAFFT, RAPS4-QF, FAST, DSM-IV two-item, and RUFT-Cut. The TWEAK and CAGE are able to distinguish between adolescents with hazardous and nonhazardous drinking (p < 0.001 and p < 0.05, respectively).²⁰ The TWEAK has also been shown to distinguish age-related drinking differences.²⁰

Diagnostic Accuracy of Instruments

Our findings on the accuracy of mental health instruments for use in the ED are presented in Table 4.

General Screening

If used as part of a general mental health evaluation, the HEADS-ED has good accuracy in identifying pediatric mental health patients who require hospital admission (area under the curve = 0.82).⁸ Risk of admission is sixfold more likely if patients receive a HEADS-ED score > 7 and a suicidal risk score of 2 (range = 0 to 2).

Suicide Risk

In the three studies that examine suicide risk instruments in pediatric psychiatric patients,^{23,24,27} none of

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	Instrument tremint	Partici	ipants	Findings	
Author	(No. Items or Domains)	Sample, <i>n</i> (% Female)	Age (y), Range (Mean)	Validity	Reliability
General Screening Cappelli et al. ^s	HEADS-ED (7 domains)	313 (58.1)	4-17 (14.3)	 Concurrent validity (correlation with CANS-MH 3.0 items), range r = 0.32-0.89 Concurrent validity (correlation with CDI items), range r = 0.17-0.43 Predictive validity Predictive of consultation for a full psychiatric assessment and subsequent admission to inpatient care, mean difference in scores for those admitted vs. discharged (p < 0.000), and those admitted vs. discharged (p < 0.000). 	Inter-rater reliability • r = 0.785 Single-item reliability, HEADS-ED items, range • r = 0.57–0.90
Suicide Risk Gipson et al. ⁹	C-SSRS (3 subscales)	178 (55.6)	13–17 (15.25)	 Predictive validity of intensity scale score Not predictive of a return psychiatric emergency visit, but predictive of a suicide attempt at return psychiatric emergency visit (OR = 1.09; 95% CI = 1.01–1.17); among subgroup of adolescents who reported suicidal ideation at visit, predictive of a collescents who reported suicidal ideation at visit, predictive of a collescents who reported suicidal ideation at visit, predictive of a collescents who reported suicidal ideation at visit. Predictive validity of intensity scale attempt at return visit. The duration item (how long suicidal thoughts last) was the only item-level predictor of return psychiatric emergency visits (OR = 1.67; 95% CI = 1.06–3.04). Predictive validity of severity scale score Not predictive of a return psychiatric emergency visit. Score categorization (intent vs. no intent) did not predict a return visit. 	Internal consistency reliability • 5-item intensity scale, α = 0.81 • 1-item severity and behavior scales, NA
					(Continued)

Table 3 Overview of Psychometric Studies That Evaluated Instruments to Identify Mental Health Problems and Alcohol Use Disorders Among Patients Presenting with Mental Health or Alcohol-related

Table 3 (continued)					
	linstrument triemi	Partici	ipants	Findings	
Author	(No. Items or Domains)	Sample, <i>n</i> (% Female)	Age (y), Range (Mean)	Validity	Reliability
Folse and Hahn ²⁵	RSQ (4 items)	59* (61.0)	12-24 (19.3)	 Criterion validity (criterion: positive/hegative screens significantly correlate with presence/absence of postevaluation diagnoses indicating imminent suicide risk) A positive screen for question 1 of the RSQ was correlated with a postevaluation primary diagnosis (r = 0.87) and suicide-related diagnosis (r = 0.72; p < 0.01). A positive screen for question 2 of the RSQ was correlated with postevaluation primary diagnosis (r = 0.49) and suicidewith postevaluation primary diagnosis (r = 0.40) 	Internal consistency reliability • 4-item instrument, $\alpha = 0.46$ • 2-item instrument (questions 1 and 2), $\alpha = 0.64$
Folse et al ²⁶		39 [°] (71.8)	12–24 (18)	 diagnosis (r = 0.57; p < 0.01). Questions 3 and 4 of the RSQ did not contribute to validity. Criterion validity (criterion: positive/negative screens significantly correlate with presence/absence of postevaluation diagnoses indicating imminent suicide risk) 	Internal consistency reliability • 4-item instrument,
				 A positive screen for question 1 of the RSQ was correlated with a psychiatric-related discharge diagnosis (r = 0.76) and suicide-related discharge diagnosis (r = 1.00; p < 0.01). A positive screen for question 2 of the RSQ was correlated with a psychiatric-related discharge diagnosis (r = 0.34) and suicide diagnosis (r = 0.48; p < 0.01). A positive screen for question 3 of the RSQ was correlated with a psychiatric-related discharge diagnosis (r = 0.28; p < 0.01). 	$\alpha = 0.63$ 2-item instrument (questions 1 and 2), $\alpha = 0.65$
King et al. ²⁸	Composite: SIQ-JR (15 items), AUDIT-C (3 items), RADS-2 (30 items)	298 (50.3)	13–17 (15.02)	• Question 4 of the RSQ did not contribute to validity. Concurrent validity (difference between total scores, mean \pm SD) • Adolescents who screened positive for elevated suicide risk with the composite instrument had results on the BHS [‡] (11.7 \pm 5.3) comparable to or higher than those previously reported for psychiatrically hospitalized adolescents (6.9 \pm 4.9).	Internal consistency reliability • SIQ-JR, $\alpha = 0.97$ • AUDIT-C, $\alpha = 0.89$ • RADS-2, $\alpha = 0.95$
					(Continued)

Table 3 (continued)					
	Instrument	Partici	ipants	Findings	
Author	(No. Items or Domains)	Sample, <i>n</i> (% Female)	Age (y), Range (Mean)	Validity	Reliability
Alcohol Use Disorders Hernandez et al. ¹⁷	ADI (24 items)	740 (41.5)	13–17 (15.26)	 Construct validity, model fit 24-item, 4-factor model⁶: RMSEA = 0.090, CFI = 0.812, TLI = 0.789 8-item, 4-factor model⁶: RMSEA = 0.050, CFI = 0.984, TLI = 0.967 8-item, 4-factor model of adolescents who had a negative alcohol screen: RMSEA = 0.079, CFI = 0.968, TLI = 0.936 8-item, 4-factor model of adolescents who had a negative alcohol screen: RMSEA = 0.074, CFI = 0.968, TLI = 0.936 8-item, 4-factor model of adolescents who had a negative alcohol screen: RMSEA = 0.074, CFI = 0.964, TLI = 0.936 8-item, 4-factor model of adolescents who had a positive alcohol screen (0.22 vs. 0.30, p< 0.001) Hangover (0.02 vs. 0.17, p < 0.001) Positive alcohol screen (0.20 vs. 0.30, p < 0.001) Ciorrette use (0.13 vs. 0.16, p < 0.001) 	Internal consistency reliability • 24-tem scale, $\alpha = 0.92$ • 8-item adapted scale, $\alpha = 0.79$
Kelly et al. ¹⁸	AUDIT-C (3 items), CRAFFT (6 items), RAPS4-QF (6 items), FAST ^{II} (3 items), RUFT- Cut (5 items), DSM-IV 2-item scale	181 (43)	18-20 (NR)	 Cocaine use (0.28 vs. 0.25, p < 0.01) Other substance use (0.27 vs. 0.25, p < 0.001) 	Internal consistency reliability • AUDIT-C, $\alpha = 0.81$ • RAPS4-QF, $\alpha = 0.68$ • FAST, $\alpha = 0.64$ • RUFT-Cut, $\alpha = 0.64$
Kelly et al. ¹⁹	AUDIT (10 items), CAGE (4 items), CRAFFT (6 items), RAPS-QF (6 items), RUFT-Cut (5 items)	93 (45)	18-20 (19.0)	β	DSM-IV 2-item, $\alpha = 0.41$ Internal consistency reliability AUDIT, $\alpha = 0.83$ RAPS4-QF, $\alpha = 0.63$ CAGE, $\alpha = 0.56$ CAGE, $\alpha = 0.56$ RUFT-Cut, $\alpha = 0.64$
					(Continued)

lable 3 (continuea)					
	Instrument tremut	Partic	cipants	Findings	
Author	(No. Items or Domains)	Sample, <i>n</i> (% Female)	Age (y), Range (Mean)	Validity	Reliability
Kelly et al. ²⁰	AUDIT (10 items), TWEAK (5 items), CAGE (4 items)	103 (46.6)	12-20 (17.5)	Concurrent validity (correlation between total scores) • AUDIT and TWEAK (r = 0.83, p < 0.001) • AUDIT and CAGE (r = 0.43, p < 0.001) • TWEAK and CAGE (r = 0.43, p < 0.001) • TWEAK and CAGE (r = 0.53, p < 0.001) Criterion validity (criterion: hazardous vs. nonhazardous drinking, mean score \pm SD) • AUDIT (15.9 \pm 7.3 vs. 4.3 \pm 3.2, p < 0.001) • D-item AUDIT (14.3 \pm 6.4 vs. 3.8 \pm 2.8, p < 0.001) • D-item AUDIT (14.3 \pm 6.4 vs. 3.8 \pm 2.8, p < 0.001) • TWEAK (3.93 \pm 1.5 vs. 1.93 \pm 1.3, p < 0.001) • CAGE (1.1 \pm 1.3 rs. 0.56 \pm 0.65, p < 0.05) • CAGE (1.1 \pm 1.3 rs. 0.56 \pm 0.65, p < 0.05) • CAGE (1.1 \pm 1.3 rs. 0.56 \pm 0.05) • CAGE (1.1 \pm 1.3 rs. 0.56 \pm 0.05) • CAGE (1.1 \pm 1.3 rs. 0.56 \pm 0.05) • CAGE (1.1 \pm 1.5 vs. 1.93 \pm 1.3, p < 0.001) • CAGE (1.1 \pm 1.5 vs. 1.93 \pm 1.3, p < 0.001) • CAGE (1.1 \pm 1.5 vs. 1.93 \pm 1.3, p < 0.001) • AUDIT (15.6 \pm 9.6 vs. 6.8 \pm 6.2, p < 0.02) TWEAK and CAGE (NS) Construct validity (group differences for alcohol use, mean \pm SD) AUDIT (5.02 \pm 4.9 vs. 10.39 \pm 8.0, p < 0.001) • AUDIT (5.02 \pm 4.9 vs. 10.39 \pm 8.0, p < 0.001) • CAGE (NS) • CAUDIT • CAGE (NS) • CAGE (NS)	Internal consistency reliability • AUDIT, $\alpha = 0.88$ • TWEAK, $\alpha = 0.50$ • CAGE, $\alpha = 0.66$
Kelly et al. ²¹	AUDIT (10 items)			 TWEAK (3.9 ± 1.5 vs. 2.1 ± 1.8, p < 0.009) AUDIT and CAGE (NS) Race/ethnicity (Gaucasian vs. African American adolescents) AUDIT (8.56 ± 7.6 vs. 5.07 ± 3.9, p < 0.02) TWEAK and CAGE (NS) TWEAK and CAGE (NS) Construct validity, model fit 9-item, 1-factor model: RMSEA = 0.21, CFI = 0.80, NNFI = 0.76, GFI = 0.75 10-item, 3-factor model*: RMSEA = 0.04, CFI = 0.99, NNFI = 0.98, GFI = 0.94 	Internal consistency reliability • Total scale, α = 0.86
CANS-MH 3.0 = Ch NR = not reported; *91.5% of the adole †92.1% of the adole †92.1% of the adole the adole figer Hopeless Sci §Goodness-of-fit str Modified version. **Analysis allowed fr	iid and Adolescent Needs an NS = not statistically significa scent sample presented with secent sample presented with ale scores have shown predic atistics for first-order confirma drinking removed to see if dif or correlated error variances.	d Strengths-Mental H unt, RMSEA = root me nonpsychiatric compl nonpsychiatric compl ritve validity for suicide tory factor analysis. fferences were inflated	lealth instrument, vei san square error of a laints. e attempts in adolesi d.	sion 3.0; CDI = Children's Depression Inventory; CFI = Comparative Fit Inc pproximation; TLI = Tucker Lewis index. cents. ²⁹	sx; NA = not applicable;

Table 4

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		Partic	cipants				Findings					
		Sample. <i>n</i>	Ade (v). Bande		Recommended Score/ Cut-score (Score	Sensitivity	Specificity	Predic Value	tive es	LR		
Author	Instrument	(% Female)	(Mean)	Reference	Range)	(95% CI)	(95% CI)	+	I	+		Accuracy
General Scree Cappelli et al. ⁸	ning HEADS-ED	313 (58.1)**	4–17 (14.3)	CANS-MH	HEADS-ED score > 7 (0–14) and suicidal risk score of 2 (0–2)	0.82 (NR)	0.87 (NR)	NR	NR	6.30	0.21	0.82
Suicide Risk Horowitz et al. ²⁴	ASQ	180 (58.3)**	10–21 (14.4)	SIQ/SIQ-JR*	A positive response to at least 1/4 of the	0.98 (91.7–99.7)	0.66 (55.2–75.0)	71.3	96.9	2.8	0.04	0.83
Horowitz et al. ²⁷	RSQ	144 (54)**	10–21 (13.6)	SIQ/SIQ-JR*	A positive response to at least 1/4 of the	0.98 (NR)	0.37 (NR)	55	67	1.56	0.05	0.87
Rutman et al. ²³	SQS TQS	212 [*] (50.0)	12–17 (14.7)	CES-D	A positive response to the 1 item A positive response to at least 1/2 of the items	0.56 (50%–63%) 0.78 (73%–84%)	0.93 (90%–96%) 0.82 (77%–87%)	83 72	78 87	8.0	0.47 0.27	K K
Alcohol Use L Kelly et al. ¹⁸	%sorders AUDIT-C CRAFFT RAPS4-QF FAST [∥] RUFT-Cut DSM-IV DSM-IV	181 [‡] (43)	18-20 (NR)	DSM-IV AUD	$\begin{array}{l} Cut\text{-score} = 6^{\$} \ (0-12) \\ Cut\text{-score} = 3 \ (0-6) \\ Cut\text{-score} = 3 \ (0-6) \\ Cut\text{-score} = 3 \ (0-16) \\ Cut\text{-score} = 3 \ (0-11) \\ Cut\text{-score} = 1 \ (0-2) \end{array}$	0.74 (NR) 0.69 (NR) 0.79 (NR) 0.83 (NR) 0.80 (NR) 0.88 (NR)	(RN) 77.0 (RN) 70.0 (RN) 7	N N N N N N N N N N N N N N N N N N N N	K K K K K K K K K K K K	3.22 2.56 3.07 8.80 8.80	0.34 0.42 0.29 0.23 0.13	0.83 0.79 0.80 0.83 0.83 0.83
Kelly et al. ¹⁹	2-item AUDIT CAGE CRAFFT RAPS-QF PUIET Cut	93 [*] (45)	18-20 (19.0)	DSM-IV AUD	Cut-score = 10 (0-40) Cut-score = 1 (0-4) Cut-score = 3 (0-6) Cut-score = 3 (0-6)	0.82 (NR) 0.66 (NR) 0.82 (NR) 0.82 (NR) 0.82 (NR) 0.82 (NR) 0.82 (NR)	0.78 (NR) 0.58 (NR) 0.67 (NR) 0.54 (NR) 0.54 (NR)	71 52 55 MP	88 84 81 81	3.73 1.57 2.48 1.78	0.23 0.59 0.33 0.33	0.85 0.68 0.79 0.76
Bastiaens et al. ²²	RAFFT	226 (45)	13–18 (15.4)	DSM-IV SUD	2 positive answers	0.89 (NR)	(NN) 63.0	E N N	L H	2.87	0.16	NR PN
AUD = alcohc CES-D = Cen SUD = substa *For adolescei	I use disorder ter for Epidemic nce use disord nts < 14 years (diagnosis; CAN ⁶ ological Studies er diagnosis. old.	S-MH = Child and Depression Scale	d Adolescent Nee s; DSM = Diagnos	eds and Strengths-Mental stic and Statistical Manual	Health instrument; C I of Mental Disorders	D = cannabis diagnc ; LR = likelihood rati	o; NA = r	 children not appli 	en's dep icable; N	R = not	nventory; reported;

§Cut-point of 6 recommended for females.
*Percentage of patients with psychiatric ED visits not reported.
*Percentage of patients who were alcohol positive at the time of study enrollment not reported.
**pediatric psychiatric population

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the instruments demonstrate both high sensitivity and specificity. Instruments that are more sensitive are less specific (ASQ, RSQ), and those that are more specific are less sensitive (SQS, TQS). The ASQ and RSQ provide strong evidence to rule out risk. In terms of ruling in risk, an adolescent with a positive response to at least one of the four ASQ items has an almost threefold higher risk for suicide (LR+ = 2.8).²⁴

Alcohol Use Disorders

Diagnostic accuracy ranges widely among the 12 instruments evaluated with adolescent ED patients. Most effective for detecting an alcohol use disorder (area under the curve = 0.89) was using two diagnostic items based on DSM-IV criteria ("In the past year, have you sometimes been under the influence of alcohol in situations where you could have caused an accident or gotten hurt?" and "Have there often been times when you had a lot more to drink than you intended to have?").¹⁸ Adolescents who answer yes to at least one of the two items are eightfold more likely to be diagnosed with a disorder (LR+ = 8.80), but evidence for ruling out a disorder with these items is modest (LR- = 0.13).¹⁸

DISCUSSION

Using specialized instruments to screen and diagnose mental health problems among children who present to the ED is supported by the AAP and the Committee on Pediatric Emergency Medicine.³ However, actual use of evidence-based, mental health screening practices among ED physicians has been documented at less than 10%.⁶ Physician screening can benefit children who present to the ED with mental health and/ or substance use concerns but have yet to receive a diagnosis. Such screening can inform consultation with mental health professionals and in-house assessments by them. If mental health resources are not available in the ED, physician screening becomes an integral step in comprehensive discharge planning and referral to mental health services for more detailed psychiatric assessment. For children who present with known mental health and/or substance use concerns, physicians can use instruments to identify immediate risks (e.g., risk of suicide) and guide disposition decision making. This systematic review identifies three instruments, the HEADS-ED, the ASQ, and the DSM-IV, as evidence-based screening options for ED physicians. These instruments require minimal (HEADS-ED) to no training (ASQ, DSM-IV two-item instrument) before use.

Using the recommended cut scores, the HEADS-ED provides reasonable sensitivity and specificity for the ED setting. The instrument will detect 82% of pediatric mental health patients requiring admission (18% who require admission will go undetected) and will correctly report 87% of patients who do not need admission (13% who do not require admission will be identified as needing admission).

Recent research has found screening for suicide risk to be acceptable to clinicians, parents, and pediatric patients.³⁰ The ASO provides strong evidence to rule out risk and is a highly sensitive instrument for detecting risk of suicide in our population of interest, pediatric psychiatric patients (98% sensitivity and 66% specificity), as well as general pediatric ED population (97% sensitivity and 88% specificity). While ASQ instrument specificity is modest in our population of interest, pediatric psychiatric patients, high sensitivity is most essential in evaluating risk of suicide. The falsepositive rate of 34% in a pediatric psychiatric patient population and 12% in a general pediatric patient population may be best addressed by ED physicians regarding the instrument as an initial screen. Patients who screen positive (whether true or false positive) should be assessed further for risk of suicide. Whether this subsequent assessment is conducted by a mental health professional or by a physician may depend on the resources available in the ED at the time of the visit. Any decision to not screen a child because of concerns over a potential false-positive result, or discomfort with what clinical decisions should follow screening outcomes, should be tempered by the potential benefit of identifying a child in need. One-third of young people with suicidal ideation will go on to develop a plan, and of these young people, approximately 60% will attempt suicide.³¹

Pediatric patients who answer yes to at least one of the two items on the DSM-IV instrument are at eightfold greater risk of having an alcohol use disorder (modest evidence to rule in a disorder). The instrument also provides ED physicians with good sensitivity and specificity. It will detect 88% of patients with a disorder (12% with a disorder will go undetected) and will correctly report 90% of patients who do not have a disorder (10% who do not have a disorder will be identified as having one). The DSM-IV instrument can be used by physicians as an initial screen with patients whose presenting complaints or history indicate the need for assessment. Children who screen positive could be discharged with a recommendation to follow up with more specialized addictions services for assessment and potential treatment. This approach can reduce the potential impact on ED patient flow and resources.

The different diagnostic approaches and varied findings in this review suggest that the field of EDbased mental health instrumentation requires maturation and refinement. Additional research must address methodologic limitations identified in this review. This will develop a robust evidence base for the instruments identified as psychometrically and diagnostically sound. Part of this research must define minimally acceptable thresholds for instrument sensitivity/specificity and LRs to rule in and rule out disorders. In the ED, instruments that have maximum sensitivity and specificity and strong evidence to rule in/rule out disorders are most desirable. Easy-to-use instruments with those characteristics, and with minimum training requirements and implementation time, can provide ED physicians and other ED clincians with critical information rapidly. Such information is valuable in guiding further assessment, consultation, and discharge planning.

LIMITATIONS

Our findings in this review have several limitations. First, some psychometric studies that we included lacked evidence or reporting on reliability and validity. We were therefore unable to comprehensively evaluate inter-rater and test-retest reliability or convergent/concurrent and construct validity of the screening instruments. Second, the diagnostic studies commonly enrolled a convenience sample of patients and did not use a prespecified threshold for the index test. This creates potential for bias in patient selection or interpretation of the index test, which could have influenced the screening results. Future studies should avoid using the tool to decide on outcomes (independence). Third, methodologic heterogeneity among the studies was significant, specifically in the screening instruments used. Instruments such as the HEADS-ED were assessed in only one study and therefore tests of the reliability, validity, and diagnostic accuracy of many of these instruments have not been replicated. Further, the limited number of trials available for each instrument prevented us from assessing the risk of publication bias. Finally, as with any systematic review, selection bias is possible. Although we conducted an

extensive search of the electronic and gray literature, the search was limited by language (English only) and date of publication (2000–2015). Without these restrictions, our search might have identified some additional studies.

CONCLUSION

Reliable, valid, and accurate instruments are available for use in pediatric mental health visits to the ED. However, these instruments need a more robust evidence base through additional research that addresses the methodologic and psychometric limitations we identify in this review. From available evidence, we recommend that emergency care clinicians use the HEADS-ED to rule in ED admission among children with visits for mental health care, the ASQ to rule out suicide risk among children with any visit type, and DSM-IV two-item instrument to rule in/rule out alcohol use disorders among children who currently use alcohol. We also recommend that clinicians familiarize themselves with instrument validity and reliability, and the evidence base for this information, to understand current instrument strengths and limitations. With this understanding, clinicians can make best use of these valuable instruments in assessing child and adolescent mental health during ED visits.

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Supporting Information

The following supporting information is available in the online version of this paper:

Data Supplement S1. Search strategy developed for Ovid MEDLINE(R) In-Process & Other Non-Indexed Citations and Ovid MEDLINE(R).

Data Supplement S2. Modified quality criteria from Gokiert et al.¹³

Data Supplement S3. The quality of instruments reported in psychometric studies using quality criteria from Gokiert et al.¹³

Data Supplement S4. Quality assessment of diagnostic studies using the QUADAS-2.