

Quality indicators for the assessment and management of pain in the emergency department: A systematic review

Antonia S Stang MDCM MBA MSc^{1,2,3}, Lisa Hartling PhD^{4,5}, Cassandra Fera BHSc⁶,
David Johnson MD^{1,3,7}, Samina Ali MDCM^{4,8,9}

AS Stang, L Hartling, C Fera, D Johnson, S Ali. Quality indicators for the assessment and management of pain in the emergency department: A systematic review. *Pain Res Manag* 2014;19(6):e179-e190.

BACKGROUND: Evidence indicates that pain is undertreated in the emergency department (ED). The first step in improving the pain experience for ED patients is to accurately and systematically assess the actual care being provided. Identifying gaps in the assessment and treatment of pain and improving patient outcomes requires relevant, evidence-based performance measures.

OBJECTIVE: To systematically review the literature and identify quality indicators specific to the assessment and management of pain in the ED.

METHODS: Four major bibliographical databases were searched from January 1980 to December 2010, and relevant journals and conference proceedings were manually searched. Original research that described the development or collection of data on one or more quality indicators relevant to the assessment or management of pain in the ED was included.

RESULTS: The search identified 18,078 citations. Twenty-three articles were included: 15 observational (cohort) studies; three before-after studies; three audits; one quality indicator development study; and one survey. Methodological quality was moderate, with weaknesses in the reporting of study design and methodology. Twenty unique indicators were identified, with the majority (16 of 20) measuring care processes. Overall, 91% (21 of 23) of the studies reported indicators for the assessment or management of presenting pain, as opposed to procedural pain. Three of the studies included children; however, none of the indicators were developed specifically for a pediatric population.

CONCLUSION: Gaps in the existing literature include a lack of measures reflecting procedural pain, patient outcomes and the pediatric population. Future efforts should focus on developing indicators specific to these key areas.

Key Words: *Emergency department; Pain assessment and management; Quality indicators; Systematic review*

Pain is the most common reason for seeking health care in the Western world (1-3). Multiple national and international bodies, including the WHO, have focused on the importance of pain management (4-6). Research has shown that inadequate pain treatment can have detrimental effects; it can result in extended length of hospitalization, slower healing, altered pain processing, depression, anxiety, and substantial social and economic costs to society (5,7,8). Although pain is a common reason for visiting the emergency department (ED), evidence consistently indicates that pain is undertreated in the ED (9-15).

Les indicateurs de qualité pour évaluer et prendre en charge la douleur à l'urgence : une analyse systématique

HISTORIQUE : D'après les données probantes, la douleur n'est pas assez traitée en salle d'urgence (SU). La première étape pour atténuer l'expérience de la douleur chez les patients en SU consiste à évaluer les soins administrés de façon précise et systématique. Pour déterminer les lacunes dans l'évaluation et le traitement de la douleur et améliorer le résultat des patients, il faut des mesures de rendement pertinentes et fondées sur des faits probants.

OBJECTIF : Faire l'analyse systématique des publications et déterminer les indicateurs de qualité propres à l'évaluation et à la prise en charge de la douleur en SU.

MÉTHODOLOGIE : Les chercheurs ont interrogé quatre grandes bases de données bibliographiques entre janvier 1980 et décembre 2010 et fouillé manuellement les revues et délibérations de congrès ou colloques pertinents. Ils ont inclus les recherches originales qui décrivaient la mise sur pied ou la collecte de données sur au moins un indicateur de qualité propre à l'évaluation ou à la prise en charge de la douleur en SU.

RÉSULTATS : La recherche a permis d'extraire 18 078 citations. Vingt-trois articles ont été retenus : 15 études d'observation (cohortes), trois études avant-après, trois vérifications, une étude d'élaboration d'indicateurs de qualité et une enquête. La méthodologie était de qualité modérée, comportant des faiblesses dans la déclaration de la conception et de la méthodologie. Vingt indicateurs uniques ont été relevés, la majorité (16 sur 20) mesurant les processus de soins. Dans l'ensemble, 91 % des études (21 sur 23) rendaient compte d'indicateurs pour l'évaluation ou la prise en charge de la douleur à la présentation, en opposition à la douleur causée par une intervention. Trois études incluaient les enfants. Cependant, aucun indicateur n'a été élaboré expressément pour la population pédiatrique.

CONCLUSION : Le peu de mesures reflétant la douleur liée à l'intervention, les résultats des patients et la population pédiatrique font partie des lacunes des publications actuelles. Les futurs efforts devraient porter sur l'élaboration d'indicateurs axés sur ces secteurs clés.

Policy makers, researchers and health care providers use quality indicators, or performance measures, to measure and improve the quality of care provided to patients. Previous research and experience has shown that quality indicators and performance measurement improve health care outcomes (16,17). The first step in improving the treatment of pain for ED patients is to accurately and systematically assess the actual care being provided. Identifying gaps in the assessment and treatment of pain and improving patient outcomes requires relevant, evidence-based performance measures. Quality indicators have previously

¹Department of Pediatrics; ²Community Health Sciences, University of Calgary; ³Alberta Children's Hospital Research Institute, Calgary;

⁴Department of Pediatrics, University of Alberta; ⁵Alberta Research Centre for Health Evidence, Edmonton, Alberta; ⁶School of Physical and Occupational Therapy, McGill University, Montreal, Quebec; ⁷Departments of Physiology and Pharmacology, University of Calgary, Calgary;

⁸Department of Emergency Medicine, Faculty of Medicine and Dentistry, University of Alberta; ⁹Women and Children's Health Research Institute, Edmonton, Alberta

Correspondence: Dr Antonia S Stang, Departments of Pediatrics and Community Health Sciences, University of Calgary, Alberta Children's Hospital Research Institute, 2888 Shaganappi Trail, Calgary, Alberta T3B 6A8. Telephone 403-955-7493, fax 403-955-7552, e-mail antonia.stang@albertahealthservices.ca



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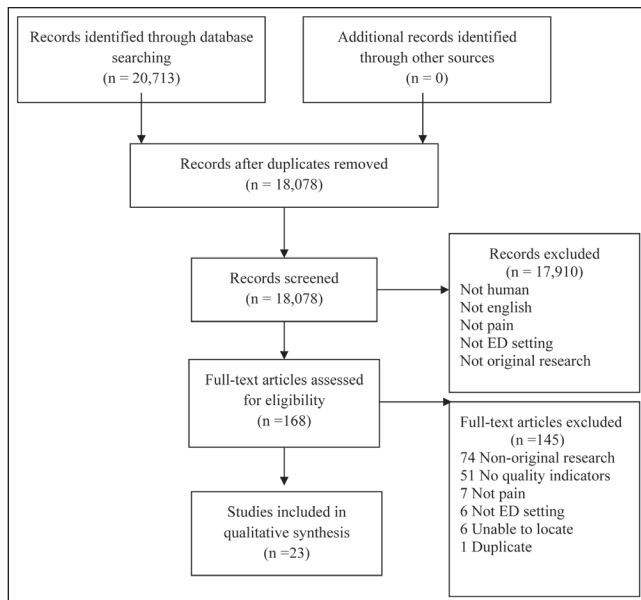


Figure 1 Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) diagram. ED Emergency department

been developed and applied to a number of clinical conditions and health care settings, including the ED management of asthma, acute myocardial infarction and pneumonia (18-22). The objective of the present study was to systematically review the literature and identify quality indicators for the assessment and management of pain in the ED setting. To create health care system improvements in the ED that meet the needs of patients with presenting (eg, illness or injury-related) and procedural (eg, medical test or other procedure-related) pain, an accurate summary of existing performance measures is required.

METHODS

Literature search and selection

The search strategy was developed by a medical research librarian (AM), in consultation with the research team and preidentified content experts. A commonly accepted definition of quality indicators that has been employed in previous research was used: "explicitly defined and measurable items referring to the structures (staff, equipment, and facilities), processes (prescribing, investigations, interactions between professionals and patients) or outcomes (such as mortality, morbidity or patient satisfaction) of care" (23-25). A systematic literature search of journals published between January 1980 and December 2010 was conducted using the following bibliographic databases: Medline, Cochrane Library, EMBASE and CINAHL. Search terms included those corresponding to quality indicators and quality improvement combined with terms describing the assessment and treatment of pain in the ED (Appendix 1). Due to resource constraints, only English-language publications were considered.

All titles and abstracts generated by the search were independently screened by two of three authors (AS, SA and CF) and any articles team members believed were likely to provide potential indicators were included for full-text review. Using predefined eligibility criteria, the full text of screened articles were independently assessed for inclusion by two authors (AS, CF). Original research that described the development or collection of data for one or more quality indicators relevant to the assessment or management of pain in the ED setting was included. Studies that were conducted outside of the ED, including the prehospital setting, inpatient setting or operating room were excluded, as were studies conducted in the ED that described the epidemiology, assessment or management of pain but did not measure the quality of care using quality indicators or a related term such as performance measure, audit filter, indicator, care indicator, benchmark, clinical path expected outcome or quality

measure. To identify additional studies, reference lists of included articles and review papers were screened, and relevant journals and proceedings of key scientific meetings were hand searched (Appendix 2). Consensus was required for final exclusion of screened articles and disagreements were resolved through involvement of a third author (SA). Before implementation, ethics approval for this study was obtained from the University of Calgary's Conjoint Health Research Ethics Board (Calgary, Alberta).

Data extraction

Data were abstracted from each study by one of two authors (AS, CF) using a standardized, piloted form, with accuracy of extraction verified by the second author (AS, CF). Abstracted data included: study methodology (design, setting, population), indicator definition and indicator measurement properties (reliability, validity, data source) and the degree to which the quality indicator was judged to be operational (described explicitly and in sufficient detail to be potentially implemented by a reader) (26).

Assessment of methodological quality

Methodological quality was rated using the Newcastle-Ottawa Quality Assessment Scale (NOS) (27) for observational studies and a modified version of the NOS for before-after studies (28). To the authors' knowledge, there is no validated tool to measure the methodological quality of indicator development studies. The indicator development process used in included studies was compared with the steps recommended by the Agency for Healthcare Research and Quality for quality indicator measure development, implementation, maintenance and retirement (29) (Appendix 3).

Analysis

The results of the present review are primarily described in a qualitative manner, with the use of counts and proportions where relevant. The primary outcome was the count of existing quality indicators for the assessment and treatment of pain in the ED, including the type of indicator (structure [such as staff, equipment, and facilities], process [prescribing, investigations, interactions between professionals and patients] or outcome [such as mortality, morbidity or patient satisfaction]) (25) and the aspect of pain measured (assessment or management). Secondary outcomes included the indicator data source, the proportion of indicators that are operational, the results of indicator measurement, and the type of pain, either presenting (eg, musculoskeletal injury, abdominal pain, sickle cell crisis) or procedural (eg, fracture reduction, intravenous insertion, lumbar puncture).

RESULTS

Study selection

The search identified 18,078 citations (Figure 1). Review of titles and abstracts resulted in the retrieval of 168 full-text articles. No additional articles were identified through the manual search of references, journals or conference proceedings. Twenty-three articles met the inclusion and exclusion criteria for the present review (Appendix 4).

Study characteristics

The studies included in the review are described in Table 1; 15 of the studies were observational (cohort) studies, three were before-after studies, three were audits, one was a quality indicator development study and one was a survey. The number of patients enrolled in each of the studies ranged from 18 (30) to 156,729 (31), with a median of 302 patients. The majority of participants were adults. One study was performed in a pediatric ED (32), while three studies focused on pain in the elderly (33-35). The majority (n=10) of studies were performed in EDs in the United States, four in Australia, five in the United Kingdom, one in France and one in Canada; one study focused on indicator development, and was not health care facility-based.

Overall, there were weaknesses in the reporting of study design and methodology, which made assessment of methodological quality challenging for the present review. Methodological quality was

TABLE 1
Summary of included studies

Study	Study design	Setting (annual visits)	n	Age, years	Objective
Arendts and Fry (69), 2006	Retrospective cohort	ED, trauma centre, Australia (46,000)	857	Median 47 (range 0–101)	To determine: the proportion of patients that require parenteral opiate analgesia for pain in an ED and who receive the opiate in <60 min; and whether any factors are predictive for the first dose of analgesia being delayed beyond 60 min
Chu and Brown (44), 2009	Analytic observational	ED, tertiary referral hospital, Australia (70,000)	72	Median 47 (IQR 35–56)	To provide exploratory data seeking an association between access block and time to parenteral opioid analgesia in patients presenting to the ED with renal colic
Eder et al (41), 2003	Retrospective	ED, United States (NR)	261	Mean \pm SE 40 \pm 15, median 40	To evaluate ED documentation of patient pain in light of the Joint Commission of Accreditation of Healthcare Organization's emphasis on pain assessment and management
Forero et al (45), 2008	Retrospective cohort	ED, referral hospital, Australia (>40,000)	13,449	23% 0–14, 41% 15–44, 26% 45–75, 9% \geq 76	To explore the association of morphine use with factors influencing time to initial analgesia
Goodacre and Roden (39), 1996	Audit	ED, United Kingdom (NR)	200	NR	To evaluate a protocol for intervention used to improve on shortcomings in the use of analgesia in an accident and ED
Grant et al (42), 2006	Retrospective analysis of patient records	ED, United Kingdom (NR)	473	NR (<18 excluded)	To evaluate the effectiveness of analgesia delivery, in the ED setting, to patients presenting with acute pain, with regard to guidelines from the British Association of Accident and Emergency Medicine
Guru and Dubinski (49), 2000	Prospective cohort	ED, Canada (NR)	71	Mean \pm SE 35.5 \pm 15.7	To assess how well pain was evaluated and treated in accordance with recommended guidelines in the Canadian Association of Emergency Physicians consensus document
Hawkes et al (32), 2008	Before-after	ED, third level trauma, Ireland (50,000, 10,000 pediatric)	240	Before, median 8 (IQR 4–12); after, 10 (IQR 4–13)	To describe the provision of analgesia for pediatric patients in a large Irish ED and to analyse and modify the protocol in this hospital with a view to improving the quality of care provided for children
Hwang et al (43), 2008	Retrospective observational	ED, urban, academic tertiary care ED, United States (NR)	1068	Mean \pm SD 47 \pm 19	To evaluate the association of three ED crowding factors with the quality of general ED pain care
Hwang et al (33), 2006	Retrospective cohort	ED, tertiary care teaching facility, United States (70,000)	158	Mean 83 (range 52–101)	To evaluate the effect of ED crowding on the assessment and treatment of pain in older adults
Jackson (34), 2010	Before-after (educational intervention)	ED, United States (NR)	302	Mean 83 (range 65–99)	To explore the results of a staff educational intervention with evaluation of medical record documentation before and after implementation
Kuan et al (50), 2009	Before-after (educational intervention)	ED, tertiary referral centre, Ireland (40,000)	151	Audit 1; median 38 (range 15–90) audit 2, 37 (16–83) audit 3, 38 (15–82)	To evaluate the impact of a brief educational intervention on prompt recognition and treatment of pain in the ED
Mitchell et al (46), 2009	Observational retrospective	ED, metropolitan teaching hospital, Australia (32,000)	436	Median 57	To determine whether overcrowding negatively impacts on the timeliness of pain management, an indicator of ED quality of care
Odesina et al (30), 2010	Retrospective medical record review	ED, suburban university health care centre, United States (29,200)	18	Mean \pm SE 29.9 \pm 5.61	To examine the current sickle cell disease pain management practice patterns, explore evidence-based sickle cell disease pain management clinical guidelines, and develop and implement an adapted ED sickle cell pain management clinical pathway
Pines et al (54), 2008	Retrospective	ED, urban tertiary care, United States (55,000)	13,758	Mean \pm SD 39 \pm 16	To study the impact of ED crowding on both treatment and delays in treatment in a broad cohort of ED patients who presented at triage with a complaint of severe pain
Pines and Hollander (48), 2010	Retrospective cohort	ED, United States, multicentre, academic tertiary care (57,000), community hospital (35,000)	5616	Mean \pm SD 44 \pm 17	To study the association between ED crowding and the use of, and delays in, analgesia in patients with back pain in two EDs
Pletcher et al (31), 2008	Analysis of national database	ED, United States, multicentre (NR)	156,729	Reported by race: white, mean \pm SE 39.0 \pm 22; black, 34.3 \pm 19; Hispanic, 31.9 \pm 20; Asian/other, 36.7 \pm 21	To determine whether opioid prescribing in EDs has increased, whether non-Hispanic white patients are more likely to receive an opioid than other racial/ethnic groups, and whether differential prescribing according to race/ethnicity has diminished since 2000

Continued on next page

TABLE 1 – CONTINUED
Summary of included studies

Study	Study design	Setting (annual visits)	n	Age, years	Objective
Ritsema et al (40), 2007	Retrospective cohort	ED, multicentre, United States (NR)	2064	1998–2000, mean \pm SE 33 \pm 1.2, 2001–2003, 35 \pm 1.2	To compare the quality of ED pain management before and after implementation of the Joint Commission on the Accreditation of Healthcare organizations standards in 2001
Ricard-Hibon et al (36), 2004	Survey	ED and prehospital EMS, France, multicentre (23% <15,000; 44% 15,000–30,000; 33% >30,000)	363 responses	NR	To evaluate the existence of a quality control program for acute pain management in the ED and in the prehospital EMS, and the needs in training and support to implement these procedures.
Shah and Lendrum (37), 2004	Audit	ED, United Kingdom (NR)	25	NR	To compare the practice in the Chesterfield Royal hospital ED with the standards set by The British Association of Accident and Emergency Medicine, and critically evaluate the results.
Tanabe et al (47), 2007	Retrospective cohort	ED, United States, academic medical centres, multicentre (NR)	159	Mean \pm SE 32 \pm 10	To characterize the initial management of patients with sickle cell disease and an acute pain episode, to compare these practices with the American Pain Society Guideline for the Management of Acute and Chronic Pain in Sickle-Cell Disease in the ED, and to identify factors associated with a delay in receiving an initial analgesic.
Terrell et al (35), 2009	Task force (quality indicator development)	N/A	N/A	\geq 65	To develop ED-specific quality indicators for older patients
Vega-Stromberg et al (38), 2002	Audit	Acute care, United States, multicentre (NR)	NR	NR	To describe an interdisciplinary model for process improvement within an integrated health care system.

ED Emergency department; EMS Emergency medical services; IQR Interquartile range; NR Not recorded; SE Standard error; N/A Not applicable

TABLE 2
Summary of quality indicators and results of measurement

Quality indicators	Type of indicator	Studies reporting indicator, n	Results of indicator measurement, including range of results when similar indicators were measured in multiple studies
Pain assessment			
Patients with any documented pain assessment	Process	5	57% to 94%
Patients with documented pain assessment using validated pain score	Process	1	23%
Patients with physician-documented pain assessment	Process	1	85% to 86%
Timeliness of pain assessment	Process	2	Mean 40 min to 174 min (from arrival/triage)
Patients with documented pain reassessment after treatment	Process	4	32% to 50%
Timely reassessment of pain relief after treatment	Process	4	0% to 55.3% of patients, mean 113 min
Pain assessment documented before discharge from ED	Process	2	56% (of sites)
Patients with pain rated 0/10 at discharge	Process	1	8%
Pain management			
Patients administered any analgesia	Process	9	6% to 79%
Patients with analgesia offered at triage	Process	3	18% to 83%
Timely access to any analgesia	Process	12	Mean or median >60 min in 6 of 9 studies, % of patients with delay \geq 1 h 14% to 81% (from arrival/triage)
Timely access to parenteral opioid analgesia	Process	2	Median 0.8 h to mean 67.5 min from arrival/triage
Elderly patients treated with meperidine	Process	2	32.8%
Patients receiving appropriate analgesic dose	Process	1	92%
Patients receiving analgesic by appropriate route	Process	1	55%
Provision of bowel regimen with opioid analgesia prescription	Process	1	NR
Sites with training specifically for pain management for physicians	Structure	1	56% of sites
Sites with training specifically for pain management for nurses	Structure	1	68% of sites
Sites with pain therapeutics protocols	Structure	1	69% of sites
Patients satisfied with pain management	Outcome	1	34% to 39%

ED Emergency department; NR Not reported

moderate (Appendix 3). The NOS was applied to studies described by the study authors or classified by data extractors as observational studies. Application of the NOS identified weaknesses in comparability of study groups and outcome ascertainment, particularly in the description of subjects lost to follow-up. For the studies identified by authors or data extractors as before-after studies, there were weaknesses identified in the selection of the pre- and postintervention sample as well as in outcome reporting and pre- and postintervention comparability with respect to the timeframe of data collection. The quality indicator development study demonstrated weaknesses in the assessment of candidate indicators. A quality assessment could not be completed for four studies due to study design, namely survey (36) and audit (37-39) methodology.

Primary outcome

Table 2 provides a summary of the quality indicators, the number of studies reporting each indicator and the results of data collected on the indicators. The most commonly measured indicators reflected the documentation and timeliness of pain assessment and reassessment, and the receipt and timeliness of analgesia. Table 3 provides details on the quality indicators extracted from each of the individual studies, including the type of indicator, data source, results of indicator measurement, extent to which indicator as reported is operational and type of pain. There were a total of 20 indicators measured, with considerable overlap in the indicators reported across studies. Eighty percent (16 of 20) of the indicators measured care processes, 15% (three of 20) measured structure and 5% (one of 20) measured patient outcome. For the purposes of categorizing indicator type, indicators measuring pain assessment were considered process indicators. Sixty percent (12 of 20) of the indicators reflected pain management and 40% (eight of 20) reflected pain assessment.

Secondary outcomes

Within the included studies, data for the specific indicators were abstracted from chart review (14 of 23), administrative databases (six of 23) and surveys (three of 23). Ninety-one percent of the studies (21 of 23) reported indicators reflecting presenting pain complaints. Some of the included studies evaluated the quality of pain assessment and management for patients with all types of presenting pain (10 of 23), while others focused on specific conditions including back pain, fractures, sickle cell disease, hip fracture and renal/biliary colic (11 of 23). The remaining studies did not specify presenting or procedural pain (two of 23). For 19 of 23 of the studies, our study team rated the indicators as operational (Table 3).

Data collected on the indicators in the individual studies are summarized in Tables 2 and 3. The proportion of patients with documentation of pain assessment ranged from 57% (40) to 94% (41) with 23% of assessments in the latter study measured using a validated pain scale. The proportion of patients with pain reassessment ranged from 32% (42) to 50% (43). For the studies that provided data on indicators reflecting time to analgesia (from arrival or triage), the mean or median was >60 min in six of the nine studies (30,32,33,43-48). The receipt and timing of analgesia varied according to pain severity and type of presenting pain. For example, in one study (49) the proportion of patients offered analgesia ranged from 6% for mild pain (score one to three of 10) to 68% for severe pain (score seven to 10 of 10). With respect to timing of analgesia, the proportion of patients in severe pain (score seven to 10 of 10) who received analgesia within 20 min of arrival or triage ranged from 24% (42) to 83% (50). In another study, 59% of patients with severe pain had a delay >1 h in the time to first analgesic, when measured from triage presentation (48). With respect to specific types of presenting pain, the time to analgesia included a mean of 141 min (from triage) for hip fracture (33), a median of 48 min (from arrival) for renal colic (44) and a median of 90 min (from triage) for sickle cell pain crisis (47). One study measured patient satisfaction, with 34% to 39% of patients reporting 'excellent' satisfaction with pain management (38).

DISCUSSION

The increasing awareness of the prevalence and impact of untreated pain in the ED setting, and the growing body of evidence on this topic,

precludes a comprehensive summary and review of all studies describing the epidemiology, assessment and management of pain in the ED. As such, we chose to focus on studies that provided quality indicators for the assessment and management of pain in the ED. Our approach enabled us to provide data on performance using existing indicators and to identify important gaps in measurement and quality improvement that are relevant to clinicians, administrators and researchers. Our comprehensive review has identified a total of 20 indicators. While previous work has focused on reviewing existing quality indicators for trauma care (26,51), pediatric emergency medicine (52) and palliative care (53), to the best of our knowledge, this is the first publication to systematically review quality indicators specific to the assessment and management of pain in the ED.

Clinical implications

The clinical implications of the present review are directly relevant to nurses, physicians and administrators alike. Although a significant proportion of patients in the studies reviewed had documentation of pain assessment, few patients were assessed using a validated pain scale. An even smaller proportion of patients had documentation of pain reassessment. Given the negative implications of delays in pain treatment, these findings highlight the importance of encouraging the use of a validated pain scale, and conducting and documenting pain reassessments. It is imperative that pain not only be measured with a valid, objective tool, but also be frequently reassessed to optimize pain management. Our results further suggest that there are delays in the receipt of analgesia, even for patients in severe pain (48) or with clinical conditions known to be associated with significant pain (47). A number of the included studies identified an association between ED crowding and delays in pain management (33,43,45,48,54). An awareness of the impact of crowding on pain management may help to further focus the attention of individual caregivers and hospital administrators on the importance of adequate and timely pain management.

Research implications

A key finding of the present review is that 91% of the identified indicators were specific to presenting pain only. The importance of procedural pain management and the harmful effects of such pain, when left untreated, have long been recognized (8,55-59). Poor management of procedural pain has been documented in multiple care settings including hospitalized children (60), children and adults in acute and critical care settings (61), and adults in the ED (13,62). The relative lack of indicators for procedural pain highlights a significant gap in measurement and a potential missed opportunity for quality improvement.

Previous research has documented variation in pain management based on patient demographic characteristics including race, age and sex (63-68). Results from one of the studies included in the present review indicate that although opioid prescribing in the United States for patients with a pain-related ED visit increased after national quality improvement efforts in the 1990s, differences according to race and ethnicity have not diminished (31). With respect to sex, one of the included studies demonstrated a longer time to analgesia for female patients with acute pain compared with men (47). Other identified factors associated with delays in analgesia that deserve further exploration included language barriers (46) and insurance status (Medicaid) (40). Delays in analgesia were also more likely among children (69) and elderly patients (40,69), suggesting that those at the extremes of the age spectrum are suboptimally assessed and treated for acute pain. Although many of the reviewed studies included or focused on elderly patients, few of the studies included children and none of the indicators were specific to the pediatric population. Given the particular vulnerability of neonates, infants and children to both the short- and long-term consequences of untreated pain (8,55-57), the lack of pediatric-specific pain indicators for the ED represents another important gap in the existing quality measurement and improvement literature.

A final gap identified in the present review was the lack of measures reflecting patient-focused outcomes. Previous work involving

TABLE 3
Detailed summary of results of data collection on quality indicators

Study	Quality indicator	Type*	Data source	n†	Result	Operational‡	Type of pain§
Arendts and Fry (69), 2006	Delay to analgesia (% of patients with time from arrival to analgesia ≥ 60 min)	Process	Administrative database	857	47%	Yes	Presenting, all types
Chu and Brown (44), 2009	Time to parenteral opioid analgesia (time of arrival to parenteral opioid)	Process	Chart review	69	Median 0.80 h (IQR 0.37 h to 1.37 h)	Yes	Presenting, renal colic
	Time to parenteral opioid analgesia (% ≤ 1 h)	Process	Chart review	69	60.9%	Yes	Presenting, renal colic
Eder et al (41), 2003	% patients with initial pain assessments	Process	Chart review	261	94%	Yes	Presenting, all types
	% of patients with initial pain assessments measured using pain scale	Process	Chart review	261	23%	Yes	Presenting, all types
	% of patients with documented pain assessment subsequent to therapy	Process	Chart review	261	39%	Yes	Presenting, all types
	% of patients with documented pain assessment subsequent to therapy measured using pain scale	Process	Chart review	261	19%	Yes	Presenting, all types
Forero et al (45), 2008	Time to analgesia (time from triage to administration of morphine)	Process	Chart review	1097	Median 79 min, 95% CI 71–85	Yes	Presenting, all types
Goodacre and Roden (39), 1996	Percent of fracture clinic referrals with no analgesia offered in ED	Process	Chart review	200	Initial audit 91% repeat audit 69%	No	Presenting, MSK, fracture
	% of orthopedic admissions with no analgesia offered in ED	Process	Chart review	200	Initial audit 39% repeat audit 22%	No	Presenting, MSK, fracture
Grant et al (42), 2006	% of patients in severe pain (score 7 to 10/10) who receive appropriate analgesia within 20 min of arrival or triage (whichever is earlier)	Process	Administrative database	213	24%	Yes	Presenting, all types
	% of patients in moderate pain (score 4 to 6/10) offered analgesia at triage	Process	Administrative database	105	18%	Yes	Presenting, all types
	% of patients who have documented re-evaluation of analgesia requirements	Process	Administrative database	NR	32%	Yes	Presenting, all types
Guru and Dubinsky (49), 2000	% patients offered analgesia for mild pain (score 0 to 3/10)	Process	Survey	71	6%	Yes	Presenting, all types
	% analgesia offered for moderate pain (score 4 to 6/10)	Process	Survey	71	18%	Yes	Presenting, all types
	% analgesia offered for severe pain (score 7 to 10/10)	Process	Survey	71	68%	Yes	Presenting, all types
	% rating no pain (score 0/10) on discharge	Process	Survey	71	8%	Yes	Presenting, all types
Hawkes et al (32), 2008	Time from triage to analgesia major fracture (long bone, rib, clavicle)	Process	Chart review	36	Pre median 54 min (IQR 25 to 90); Post median 7 min (IQR 4 to 12); P=0.0004	Yes	Presenting, major fracture
	% receiving analgesia major fracture (long bone, rib, clavicle)	Process	Chart review	36	Pre 55.6%; post 61.1% P=0.735	Yes	Presenting, major fracture
	Time from triage to analgesia other diagnoses	Process	Chart review	183	Pre median 14 min (IQR 4 to 45); Post median 6 min (IQR 6 to 61) P=0.794	Yes	Presenting, all types
	% receiving analgesia other diagnoses	Process	Chart review	183	Pre 34.7%; post 39.8%	Yes	Presenting, all types
Hwang et al (33), 2006	% of patients with documentation of pain assessment	Process	Chart review	158	72.8%	Yes	Presenting, hip fracture
	Time to pain assessment by a physician (from triage)	Process	Chart review	128	Mean 40 min (range 0 min to 600 min)	Yes	Presenting, hip fracture
	% of patients with documentation of administration of pain medication	Process	Chart review	128	64.1%	Yes	Presenting, hip fracture
	Time to pain treatment (from triage)	Process	Chart review	128	Mean 141 min range (10 min to 525 min)	Yes	Presenting, hip fracture
	% of patients receiving opioid who were treated with meperidine	Process	Chart review	73	32.8%	Yes	Presenting, hip fracture
Hwang et al (43), 2008	% of patients with physician documented pain assessment	Process	Chart review	1068	Census low: 86%, Census high: 85%	Yes	Presenting, all types
	% of patients with any documented pain assessment	Process	Chart review	1068	Census low: 90% Census high: 90%	Yes	Presenting, all types

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TABLE 3 – CONTINUED
Detailed summary of results of data collection on quality indicators

Study	Quality indicator	Type*	Data source	n†	Result	Operational‡	Type of pain§
Hwang et al (43), 2008	% of patients with follow-up pain assessment	Process	Chart review	961	Census low: 50%, Census high: 47%	Yes	Presenting, all types
	% of patients who received analgesic medication	Process	Chart review	642	Census low: 65%, Census high: 55%	Yes	Presenting, all types
	Time to first clinician pain assessment	Process	Chart review	899	Census low: mean 106 min, Census high: mean 174 min	Yes	Presenting, all types
	Time to first analgesic medication ordering	Process	Chart review	604	Census low: mean 104 min, Census high: mean 136 min	Yes	Presenting, all types
	Time to first analgesic medication administration	Process	Chart review	590	Census low: mean 125 min, Census high: mean 167 min	Yes	Presenting, all types
Jackson (34), 2010	Time to first pain treatment after assessment (% <60 min)	Process	Chart review	220	Pre 41.8%; post 50%	Yes	Presenting, hip fracture
	Time to reassessment documentation after treatment (% <60 min)	Process	Chart review	149	Pre 30.9%; post 55.3%	Yes	Presenting, hip fracture
Kuan et al (50), 2009	% of patients with analgesia offered at triage	Process	Chart review	151	43%	Yes	Presenting, all types
	% of patients with analgesia offered at triage for severe pain (≤20 min of arrival or at triage for severe pain [score 7 to 10/10])	Process	Chart review	24	83%	Yes	Presenting, all types
	% of patients with timely reassessment of pain relief (within ≤30 min for 90% of patients with severe pain [score 7 to 10/10])	Process	Chart review	19	0%	Yes	Presenting, all types
	% of patients with timely treatment of pain (≤60 min of arrival for >75% of patients with moderate pain [score 4 to 6/10])	Process	Chart review	64	14%	Yes	Presenting, all types
Mitchell et al (46), 2009	Time to analgesia (from ED arrival)	Process	Chart review	232	Median 53 mins (IQR 30.5 to 114.5)	Yes	Presenting, fracture, renal/biliary colic
Odesina et al (30), 2010	Time from registration to receiving first opioid	Process	Chart review	44	Mean ± SD 67.5±48.1 min	Yes	Presenting, sickle cell disease
	Time to reassessment for pain relief after first opioid administration	Process	Chart review	44	Mean ± SD 113±118.4 min	Yes	Presenting, all types
Pines and Hollander (54), 2008	% of patients with severe pain (score 9 to 10/10) with no analgesia in the ED	Process	Administrative database	13,758	51%	Yes	Presenting, all types
	% of patients with severe pain (score 9 to 10/10) with delay >1 h in time to analgesia from triage	Process	Administrative database	6746	59%	Yes	Presenting, all types
	% of patients with severe pain (score 9 to 10/10) with delay >1 h in time to analgesia from placement in room	Process	Administrative database	1319	20%	Yes	Presenting, all types
Pines et al (48), 2010	% of patients with administration of any analgesia	Process	Administrative database	5616	79%	Yes	Presenting, back pain
	Time from triage to first analgesia	Process	Administrative database	4425	Median 130 min (IQR 73 to 217)	Yes	Presenting, back pain
	% of patients with delay of ≥1 h from triage to analgesia	Process	Administrative database	4425	81%	Yes	Presenting, back pain
	Time from room placement to analgesia	Process	Administrative database	4425	Median 86 min (IQR 51 to 135)	Yes	Presenting, back pain
	% of patients with delay of ≥1 h from room placement to analgesia	Process	Administrative database	4425	67%	Yes	Presenting, back pain
Pletcher et al (31), 2008	% of patients receiving opioid analgesic prescription for pain-related visits	Process	Administrative database	156,729	29% (95% CI 28% to 30%)	Yes	Presenting, all types
Ricard-Hibon et al (36), 2004	% of sites with training specifically for pain management for physicians	Process	Survey	356	56%	No	Presenting, all types
	% of sites with training specifically for pain management for nurses	Process	Survey	356	68%	No	Presenting, all types
	% of sites with pain therapeutics protocols	Process	Survey	356	69%	No	Presenting, all types

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TABLE 3 – CONTINUED
Detailed summary of results of data collection on quality indicators

Study	Quality indicator	Type*	Data source	n†	Result	Operational‡	Type of pain§
	% of sites where pain intensity was evaluated 'systematically or often' at the beginning of patient management	Process	Survey	356	64%	No	Presenting, all types
	% of sites where pain intensity was evaluated 'systematically or often' at the end of patient management	Process	Survey	356	56%	No	Presenting, all types
Ritsema et al (40), 2007	Proportion of patients whose pain was assessed	Process	Administrative database	2064	1998 to 2000 57%, 2001 to 2003 74% (P<0.01)	Yes	Presenting, long bone fracture
	Proportion of patients receiving analgesia	Process	Administrative database	2064	Any analgesia 1998 to 2000 68% 2001 to 2003 76%	Yes	1, long bone fracture
Shah and Lendrum (37), 2004	% of patients with moderate pain (score 4 to 6/10) with analgesia offered at triage	Process	Chart review	13	24%	Yes	Presenting, MSK, abdominal pain
	% of patients severe pain (score 7 to 10/10) who receive appropriate analgesia within 20 min of arrival or triage (whichever is earlier)	Process	Chart review	12	42%	Yes	Presenting, MSK, abdominal pain
	% of patients with moderate pain (score 4 to 6/10) with documented re-evaluation and action within 60 min of receiving first dose of analgesic	Process	Chart review	13	0%	Yes	Presenting, MSK, abdominal pain
	% of patients with severe pain (score 7 to 10/10) with documented re-evaluation and action within 30 min of receiving first dose of analgesic	Process	Chart review	13	0%	Yes	Presenting, MSK, abdominal pain
Tanabe et al (47), 2007	Time from triage to analgesia	Process	Chart review	529	Median 90 min (IQR 54–149)	Yes	Presenting, sickle cell disease
	% of patients who received morphine or hydromorphone as initial analgesic	Process	Chart review	529	87%	Yes	Presenting, sickle cell disease
	% of patients who received recommended dose of analgesic	Process	Chart review	529	92%	Yes	Presenting, sickle cell disease
	% of patients who received analgesic by either intravenous or subcutaneous route	Process	Chart review	529	55%	Yes	Presenting, sickle cell disease
Terrell et al (35), 2009	If an older person remains in the ED for >6 h, then a second pain assessment should be documented within 1 h of arrival to the ED	Process	Chart review	N/A	N/A	No	Presenting, all types
	If an older person received pain treatment while in the ED, then a pain assessment should be documented before discharge home from the ED	Process	Chart review	N/A	N/A	No	Not specified
	If an older person presents to the ED and has moderate to severe pain (score ≥4/10) then pain treatment should be initiated	Process	Chart review	N/A	N/A	No	Presenting, all types
	If an older person receives analgesic medication while in the ED, then meperidine should be avoided	Process	Chart review	N/A	N/A	No	Not specified
	If an older person receives an opioid analgesia prescription on discharge from the ED, then a bowel regimen should also be provided	Process	Chart review	N/A	N/A	No	Not specified
Vega-Stromberg et al (38), 2002	Patient reports of satisfaction with pain management (% excellent rating)	Outcome	Survey	NR	34% quarter A, 39% quarter B	No	Not specified

*Type of indicator: Process, Structure, Outcome; †Number of subjects; ‡Operational: described explicitly and in sufficient detail to be potentially implemented by a reader; §Type of pain: presenting or procedural pain, condition/illness. IQR Interquartile range, MSK Muskuloskeletal; N/A Not applicable, NR Not recorded

performance measurement has highlighted the importance of outcome indicators or, at a minimum, evidence of a link between the process of care and patient outcome (70,71). There was only one outcome indicator identified in the present review (patient satisfaction) (38). Research on the link between pain management and patient satisfaction has generated conflicting results (72-74) and the utility of isolated measurement of patient satisfaction with pain management as a quality improvement tool has been questioned (14). Other potentially

relevant and complementary outcomes that warrant further exploration include achievement of pain reduction to the level desired by the patient, quality of life measures, absence from work/school due to pain and pain-related health care visits/admissions. Were these measures to be considered alongside patient satisfaction, we may then have a more comprehensive understanding of patient-level outcomes.

Although many studies have suggested that oligoanalgesia in the ED is both a common and long-standing problem (10-14,75,76), a

recent commentary has questioned whether this is true, citing the retrospective nature of the majority of these studies (77). Retrospective data cannot account for patient preference, pain perception and appropriate medical justifications for avoiding analgesia. Furthermore, chart abstraction has variable reliability, and documentation (or lack thereof) does not necessarily reflect analgesia administration (77). Most of the studies included in the present review are subject to the same criticisms. Despite this recent controversy, it is clear that awareness and documentation of pain and its management are, at best, sub-optimal. Given that modification of clinical behaviour begins with recognition of the problem, improvements in documentation of patient care and objective measures of quality of care will likely result in better patient outcomes. In addition to further prospective research on pain management, the development of patient-centred outcome indicators and work linking process measures to these patient outcomes would help to clarify the degree to which our current analgesia practices impact the patient.

Strengths and limitations

The strengths of the present review include its comprehensive search strategy, clearly defined eligibility criteria and standardized data extraction. The main weakness of the present review was the variability in study design of included articles, and weaknesses in their description of methodology, outcome assessment and reporting. This made it difficult to compare performance on the indicators among studies. Furthermore, there is no validated quality assessment tool for before-after studies or quality indicator development studies. We mitigated this limitation by using a version of the NOS adapted for before-after studies and rated the quality indicator development study against the indicator development and assessment process, as outlined by the Agency for Healthcare Research and Quality. Additionally, much of the data on the quality indicators is based on chart review and administrative databases. The reliability and accuracy of these data sources for measures of true analgesia provision has, as previously discussed, been raised (77). Finally, our search did not include any literature after December 2010; however, given the large number of screened articles identified, resource limitations prevent further updating of the search.

CONCLUSION

The present review provides a summary of existing quality indicators for pain assessment and management in the ED setting. From a clinical perspective, the results highlight the importance of pain assessment and reassessment using a validated pain score and the potential for delays in analgesia administration, particularly at times when the ED is busy. We also identified gaps in existing performance measures, including a lack of measures for procedural pain and those specific to the pediatric population. Finally, the present review highlights the need for further work linking process measures to patient outcomes. These results can serve as the foundation for future work aimed at generating a complete list of quality indicators that will comprehensively measure the assessment and management of pain in the ED setting and establish a link between measurement of the indicators and patient outcomes.

DISCLOSURES: All authors listed have contributed substantially to the manuscript. This project was supported by an Alberta Innovates Health Solutions (AIHS) Knowledge Transfer/Innovation Research Grant secured by Drs A Stang and S Ali. Dr L Hartling holds a New Investigator Salary Award from the Canadian Institutes of Health Research (CIHR). The funding source (AIHS) had no involvement in data collection, analysis or interpretation. None of the authors were paid to write this article. The corresponding author (Dr A Stang) had full access to all the data and had final responsibility for the decision to submit for publication. All authors have no financial relationships with any organizations that might have an interest in the submitted work. All authors have no other relationships or activities that could appear to

have influenced the submitted work. This work has not been submitted or accepted for previous publication.

ACKNOWLEDGEMENTS: The authors thank Andrea Milne, who conducted the literature search.

DISCLOSURES: All authors have no financial relationships with any organizations that may have an interest in the submitted work. All authors have no other relationships or activities that could appear to have influenced the submitted work.

APPENDIX 1: MEDLINE* SEARCH STRATEGY

MeSH and key words for quality indicators:

1. exp Quality Indicators, Health Care/
2. exp "Quality of Health Care"/
3. exp Quality Assurance, Health Care/
4. exp Clinical Competence/
5. exp Guideline Adherence/
6. exp "Outcome and Process Assessment (Health Care)"/
7. exp Peer Review, Health Care/
8. exp Program Evaluation/
9. exp Practice Guideline/
10. exp Practice Guideline as Topic/
11. exp Patient Care Management/
12. exp Benchmarking/
13. exp "Outcome Assessment (Health Care)"/
14. exp "Process Assessment (Health Care)"/
15. (adherence adj3 guideline*).tw
16. quality performance measure.tw
17. (performance adj3 guideline*).tw
18. (program adj 3 evaluat*).tw
19. (performance adj3 evaluat*).tw
20. (quality adj3 assess*).tw
21. (quality adj3 improv*).tw
22. OR/1-23

MeSH and keywords for emergency care:

23. exp Emergency Medicine/
24. exp Emergency Medical Services/
25. exp Emergency Service, Hospital/
26. exp Emergencies/
27. exp Ambulatory Care/
28. exp Critical Care/
29. exp Trauma Centers/
30. exp Emergency Nursing/
31. exp Emergency Treatment/
32. emergency department*.tw
33. (emergency adj3 admission*).tw
34. (emergency adj3 admit*)
35. emergency room*.tw
36. emergency ward*.tw
37. acute care.tw
38. (ED or ER).tw
39. (EDs or ERs).tw
40. (emergenc\$ adj5 (department\$ or ward\$ or unit\$ or room\$ or hospital\$ or care or patient\$ or physician\$ or doctor\$ or treatment\$)).tw
41. emergency unit*.tw
42. emergency care.tw
43. (trauma adj (centres or centers)).tw
44. (emergency or emergencies).jn
45. OR/23-44
46. 22 (quality indicator terms) AND 45 (emergency terms)

MeSH and key words terms for pain:

- 47. exp Pain/
- 48. exp Neuralgia/
- 49. exp Nociceptors/
- 50. pain.tw
- 51. exp Pain Measurement/
- 52. exp Pain threshold/
- 53. (pain adj2 management).tw
- 54. (pain adj 2 assessment).tw
- 55. (pain adj (scale or score)).tw
- 56. exp Analgesia/
- 57. exp Anesthesia/
- 58. exp "Anesthesia and Analgesia"/
- 59. analgesi\$.tw
- 60. an?esthes\$.tw
- 61. OR/50-60
- 62. 46 (quality indicator and emergency terms) AND 61 (pain terms)
- 63. Limit year to "1980-Current"

*The search strategy was translated as appropriate for the other databases

**APPENDIX 2
Journals and conference proceedings**

Journals	Conference proceedings
Academic Emergency Medicine	American Academy of Pediatrics
Annals of Emergency Medicine	American College of Emergency Physicians
BMJ Quality and Safety	
Canadian Journal of Emergency Medicine	Canadian Association of Emergency Physicians
Canadian Medical Association Journal	
The Journal of Emergency Medicine	Pediatric Academic Societies
Journal of the American Medical Association	Society for Academic Emergency Medicine
New England Journal of Medicine	
Pediatrics	

**APPENDIX 3: SUMMARY OF METHODOLOGICAL QUALITY ASSESSMENT: TABLE 1
Observational studies (Newcastle Ottawa Scale)†**

Article	Selection‡	Comparability§	Outcome¶
Arendts and Fry (69), 2006	****	**	**
Chu and Brown (44), 2009	****	*	*
Eder et al (41), 2003	***	*	*
Forero et al (45), 2008	****	**	**
Grant et al (42), 2006	***		**
Guru and Dubinsky (49), 2000	***		*
Hwang et al (43), 2008	****	**	**
Hwang et al (33), 2006	****	**	*
Mitchell et al (46), 2009	****	**	*
Odesina et al (30), 2010	***		*
Pines and Hollander (54), 2008	****	**	**
Pines et al (48), 2010	****	**	**
Pletcher et al (31), 2008	****	*	**
Ritsema et al (40), 2007	****	*	**
Tanabe et al (47), 2007	****	**	*
Maximum	****	**	***

†Not completed for four studies due to study design (one survey [Ricard-Hibon 2004 (56)] and three audits [Goodacre 1996 (25), Shah 2004 (62), Vega-Stromberg 2002 (76)]). ‡Maximum of four stars for: representativeness of the exposed cohort; selection of the nonexposed cohort; ascertainment of exposure; and demonstration that outcome of interest was not present at start of study; §Maximum of two stars for: comparability of cohorts on the basis of the design or analysis; ¶Maximum of three stars for: assessment of outcome; was follow-up long enough for outcome to occur; and adequacy of follow-up of cohorts

**APPENDIX 3: TABLE 2
Before-after studies (adapted from Newcastle Ottawa Scale)**

Article	Selection†	Comparability‡	Outcome§	Intervention¶	Pre- and post inter-vention periods††
Hawkes et al (32), 2008	*		**		
Jackson (34), 2010				**	
Kuan et al (50), 2009	*	*	***	**	
Maximum	***	*	*****	**	**

†Maximum of three stars for: representativeness of the postintervention group; representativeness of the preintervention group; pre-post intervention groups drawn from the same source. ‡Maximum of one star for: comparability of the pre- and postintervention groups on the basis of design or analysis. §Maximum of five stars for: validity of outcome assessment; reliability/accuracy of outcome assessment; method of outcome assessment the same for pre-and postintervention groups. ¶Maximum of two stars for: reporting point in time that intervention occurs; clearly describing intervention. ††Maximum of two stars for: pre and postintervention data collected during similar time frame

**APPENDIX 3: TABLE 3
Quality assessment of included indicator development study†**

Article	Identification of candidate indicators‡	Assessment of candidate indicators§
Terrell et al (35), 2009	**	*
Maximum	***	*****

†Based on the Agency for Healthcare Research and Quality (AHRQ) Document on Quality Indicator Measure Development, Implementation, Maintenance, and Retirement; ‡Maximum of three stars for: literature review to identify candidate indicators; development of conceptual model; expert engagement. §Maximum of seven stars for: initial specifications; literature review on evidence base for candidate indicators; panel review using modified Delphi or Nominal Group process; risk adjustment; empirical analysis; finalization of specifications; summary of evidence for each recommended indicator

APPENDIX 4: INCLUSION CRITERIA FOR FULL TEXT REVIEW

Objective: To identify existing quality indicators for the assessment and management of pain in the ED and acute care setting.

Definitions

Quality of care is defined by the Institute of Medicine (IOM) as the "degree to which health services for individual's increases the likelihood of desired health outcomes and are consistent with current professional knowledge".

Quality indicators are explicitly defined and measurable items referring to the structures (staff, equipment, and facilities), processes (prescribing, investigations, interactions between professionals and patients) or outcomes (such as mortality, morbidity or patient satisfaction) of care.

Alternate terms for quality indicators: performance measure, audit filter, indicator, care indicator, benchmark, clinical path expected outcome, quality measure.

Screening Questions.

1. Is the article in English?
Yes: go to question 2

- No: Exclude and mark as 1-Not English
2. Is the complete article available?
Yes: go to question 3
No: Exclude and mark as 2-Abstract only (we will find full references for these)
 3. Are the subjects of the article human?
Yes: go to question 4
No: Exclude and mark as 3-Not Human
 4. Is this article a Duplicate?
Yes: Exclude and mark as 4-Duplicate and link to duplicate article by highlighting both in same color
No: go to question 5
 5. Is the article original research?
Yes: go to question 6
No: Exclude and mark as 5-Not original research
Examples of original research include:
Systematic Reviews and Meta-analysis
Experimental studies (randomized controlled trials, cluster randomized controlled trials, nonrandomized trials)
Observational studies (cohort study, case-control study, case report, case-series, ecological study, survey, audit, time-series analysis)
Consensus methodologies (systematic methods to combine expert

opinion and medical evidence including consensus development conference, Guideline Based, Nominal Group Technique, Delphi (or modified Delphi) Technique, Rand/UCLA appropriateness method)

Examples of non-original research include: letters to the editor, commentaries, nonsystematic reviews, book chapters, guideline, news items

6. Does the article include the development or collection of data on one or more quality indicators?
Yes: Go to question 7
No: Exclude and mark as 6-No Quality Indicators
7. Does the article contain quality indicators relevant to the assessment or treatment of pain?
Yes: Go to question 8
No: Exclude and mark as 7-Not Pain
8. Does the article contain quality indicators relevant to the assessment or management of pain in the ED setting?
Yes: Go to question 9
No: Exclude and mark as 8-Not ED setting
9. Is there another reason for exclusion?
Yes: Exclude and mark as 9 with explanation for reason
No: Include

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