

ORIGINAL RESEARCH

Clinical Decision Support Systems for Opioid Prescribing for Chronic Non-Cancer Pain in Primary Care: A Scoping Review

Sheryl Spithoff, MD, MSc, Stephanie Mathieson, PhD,
Frank Sullivan, FRSE, FRCP, FRCGP, Qi Guan, MSc, Abhimanyu Sud, MD, CCFP,
Susan Hum, MSc, and Mary Ann O'Brien, PhD

Background and Objectives: Clinical decision support systems (CDSSs) may help clinicians prescribe opioids for chronic noncancer pain (CNCP) more appropriately. This scoping review determined the extent and range of the current evidence on CDSSs for opioid prescribing for CNCP in primary care, and whether investigators followed best evidence and current guidance in designing, implementing and evaluating these complex interventions.

Methods: We searched 9 electronic databases and other data sources for studies from January 1, 2008 to October 11, 2019. Two reviewers independently screened the citations. One reviewer extracted data and a second verified for accuracy. Inclusion criteria: study of a CDSS for opioid prescribing for CNCP in a primary care clinical setting. We reported quantitative results in tables and qualitative results in narrative form.

Results: Our search yielded 5068 records, of which 14 studies met our inclusion criteria. All studies were conducted in the United States. Six studies examined local (eg, health center) CDSSs and 8 examined prescription drug monitoring program CDSSs. Three CDSSs incorporated evidence-based components. Study aims were heterogeneous and study designs included both quantitative and qualitative methodologies. No studies assessed patient health outcomes. Few studies appeared to be following guidance for evaluating complex interventions.

Conclusions: Few studies have rigorously assessed the use of CDSSs for opioid prescribing for CNCP in primary care settings. Going forward, investigators should include evidence-based components into the design of CDSSs and follow guidance for the development and evaluation of complex interventions. (J Am Board Fam Med 2020;33:529–540.)

Keywords: Biomedical Technology Assessment, Chronic Pain, Clinical Decision-Making, Clinical Decision Support Systems, Electronic Health Records, Information Technology, Opioid-Related Disorders, Outcomes Assessment, Prescription Drug Monitoring Programs, Translational Medical Research

Introduction

Two countries at the epicenter of the opioid crisis, Canada and the United States,^{1–4} recently released clinical practice guidelines for opioid prescribing for

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From the Department of Family and Community Medicine, University of Toronto, Toronto, Canada (SS); Institute for Musculoskeletal Health, Sydney School of Public Health, Faculty of Medicine and Health, University of Sydney; Sydney, Australia (SM); University of St Andrews; St Andrews, United Kingdom (FS); Department of Family and Community Medicine, University of Toronto; Toronto, Canada (FS); Institute of Health Policy, Management and Evaluation, University of Toronto; Toronto, Canada (QG); Department of Family and Community Medicine; Medical Psychiatry Alliance; Institute of Health Policy, Management and Evaluation, University of Toronto; Toronto, Canada (AS); Department of Family and Community Medicine, Women's College Hospital; Toronto, Canada (SH); Department of Family and Community Medicine, University of Toronto; Women's College Research Institute; Toronto, Canada (MAO).

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chronic noncancer pain (CNCP).^{5,6} These guidelines recommend against using opioid analgesics for CNCP as a first-line medication because the harms frequently outweigh the benefits.^{7–10} When opioids are prescribed for CNCP, the guidelines recommend risk mitigation strategies and opioid dose tapering. Both guidelines target primary care providers (PCPs), since they write about half of all opioid analgesic prescriptions in North America.^{11–13} However, evidence shows that PCPs may have difficulty adopting recommended clinical practices.^{14–21} Clinical decision support may provide assistance.

Clinical decision support systems (CDSSs) are electronic systems that assist health care providers in clinical decision-making, by providing patient-specific data at the point-of-care.^{14–16} Studies show that CDSSs lead to improvements in clinician performance (a care process measure), such as ordering appropriate tests and safer prescribing.^{17–25} Some CDSS design components are evidence-based, including requiring a reason for an override, activating automatically (ie, the CDSS runs without requiring provider initiation), integrating into the electronic medical record (EMR), and providing advice to patients (eg, written materials), as well as clinicians.^{14,20,26–28} These components lead to improvements in care process outcomes. Studies in which the CDSS evaluators are also the developers tend to show positive impact on process outcomes.^{26,27}

However, the impact of CDSSs on important patient or population health outcomes is unclear,^{17–20} and widespread adoption is often limited by implementation issues.^{29–34} In addition, CDSSs can be difficult to develop and evaluate because they are complex interventions that seek to change the functioning of a complex adaptive system, such as a primary care clinic.³⁵ The Medical Research Council in the United Kingdom (UK) recommends that researchers design and evaluate complex interventions through a carefully staged series of studies targeting key uncertainties, as well as a definitive evaluation.^{35,36} All steps should include process evaluations and assess for unintended consequences.³⁷

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Corresponding author: Sheryl Spithoff, MD, MSc, Department of Family and Community Medicine, Women's College Hospital, 76 Grenville Street, Toronto, ON, Canada M5S 1B2. (E-mail: sheryl.spithoff@wchospital.ca).

CDSSs can have a variety of roles in improving adherence to opioid prescribing guidelines for CNCP. They can be used to reduce the number of new opioid prescriptions for acute pain³⁸ and to reduce the initiation of opioid prescribing for CNCP. They can also be used to improve prescribing and other measures, like risk mitigation strategies for patients already receiving opioids for CNCP. This is the most challenging role for a CDSS as these patients are at high risk of harms and changing prescribing is very difficult.^{39,40}

Several studies have evaluated CDSSs for opioid prescribing for CNCP in primary care settings.^{41–44} These studies report that the use of a CDSS led to a reduction in opioid prescribing or improved adherence to clinical practice guidelines.^{41–44} Several studies have also evaluated prescription drug monitoring program (PDMP) CDSSs for opioid prescribing for CNCP in primary care settings. PDMP CDSSs are large, centralized, government-run databases that prescribers can access for point-of-care information on a patient's opioid prescriptions.^{45,46} While 1 PDMP CDSS study found that physicians wrote fewer opioid prescription in 61% of cases,⁴⁷ another study reported no association between PDMP implementation status and requirement levels (from no requirements to a mandatory requirement to check the PDMP before prescribing) and physicians' opioid prescribing for CNCP.⁴⁸ Four other PDMP CDSS studies examined PCPs' use of, and views on PDMPs.^{49–52} To date, however, the literature in this emerging field has not been systematically summarized and analyzed so the benefits and risks of implementing a CDSS are unclear.

This scoping review determined the extent and range of the current evidence on CDSSs for opioid prescribing for CNCP in primary care. Our secondary aim was to determine whether researchers followed best evidence for the design of the CDSSs and current guidance for the evaluation of complex interventions.

Methods

We conducted a scoping review using the frameworks^{53,54} described by Colquhoun et al,⁵⁵ and the methods outlined by The Joanna Briggs Institute.⁵⁶ We followed the reporting guidelines from the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) Extension for Scoping reviews.⁵⁷ We created an a priori protocol

and used an iterative approach. Modifications included a secondary research aim and a change to the data extraction plan.

Study Eligibility

We included peer-reviewed and nonpeer-reviewed studies that used quantitative, qualitative, and mixed-methods methodologies. We excluded non-systematic reviews, letters, opinion articles, analysis articles, clinical practice guidelines, and policy documents. We included all studies where the population was PCPs (ie, family physicians, emergency medicine physicians, nurse practitioners, and primary care internists) working in a primary care setting. Studies that reported less than 50% PCPs or did not report the percentage of PCPs were excluded, unless results were reported by subgroup. We included all studies that assessed a CDSS that sought to improve opioid prescribing for CNCP patients in a primary care clinical setting. We excluded studies where PCPs were working in a secondary and tertiary settings, such as a pain clinic or addiction clinic. We excluded primary care pediatric clinics. We defined a CDSS as an electronic system that assisted health care providers in clinical decision-making, by providing patient-specific data at the point-of-care.^{14–16} We included studies where the CDSS was integrated into the EMR, or functioned independently (eg, Web-accessed), or was embedded within a larger intervention. We excluded studies where CDSS use was not specified, where it was used for another reason, or where it was not implemented in clinical settings.

Data Sources and Searches

We searched electronic databases (MEDLINE [via OVID], EMBASE, CINAHL, CENTRAL, PsycINFO and International Pharmaceutical Abstracts [via OVIDSP]) from January 1, 2008 to October 11, 2019. CDSSs developed before this period likely evolved or became obsolete.⁵⁹ We built a comprehensive search strategy, including the terms “opioid,” and “clinical decision support systems.” Since studies used a large number of different keywords and medical subject headings for a CDSS, we had to conduct a broad search using a large variety of terms, including computer systems, health informatics, clinical decision-making (Appendix 1 Medline search strategy). The Medline strategy (Appendix 1) was adapted for the other databases. We used the Canadian Agency

for Drugs and Technologies approach to our gray literature search (Appendix 2 Gray literature search).⁶⁰ We also searched trial registries (ClinicalTrials.gov, World Health Organization International Clinical Trials Registry Platform), checked reference lists of additional eligible studies and contacted experts (ie, lead authors on included studies, registered protocols and systematic reviews of CDSSs).

Screening and Selection

Two researchers independently screened abstracts to determine whether they met inclusion criteria. Two researchers then independently screened the full text of all relevant articles. For both steps, after we screened 10 to 15 titles and articles, we checked inter-reviewer agreement to ensure it was at least 80% before continuing further. When there were disagreements, a third researcher (MAO) assisted in making the final decision. We contacted authors for more information when full text was not available online.⁵⁸

Data Extraction

We created and pilot tested a data extraction form to record the following items: study population and setting, description of the intervention and implementation process, type of CDSS, inclusion of evidence-based CDSS components (components that the literature has consistently found to have an impact on outcomes: requiring a reason for an override; activating automatically; integrating into the EMR; and providing advice to patients and clinicians),^{14,20,26–28} study aims, methodology and design, study outcomes, funding information, conflicts of interest, and adherence to guidance for complex interventions (eg, study was part of a stepped approach to development and evaluation), assessment for unintended consequences, planned process evaluation, process and outcome measures, theoretical approach to guide implementation and/or evaluation. One reviewer extracted data and another researcher reviewed their work (SMS, MAO, QG, SM, SH). This was a modification from our protocol that specified that 2 researchers would independently extract the data.

Data Synthesis

We used a flow diagram to report on study selection. We reported quantitative data in tabular format. We wrote narrative summaries using contextual and process-oriented data. We did not conduct a detailed

assessment of study quality, assess for reporting bias, or risk of bias consistent with current guidance on conducting scoping reviews.^{55–57}

Results

Our literature search identified 5068 citations from which 14 were included in the scoping review (Figure 1). Six studies examined local CDSSs (eg, specific health system, center or clinic)^{41,43,44,61–63} while 8 examined state-run, Web-based, central PDMP CDSSs^{47,49–52,64–66}. Results using these 2 typologies are summarized in Table 1. Study descriptions are detailed in Appendix 3.

CDSS Description

Types of CDSSs included protocols (ie, forms that guide clinical management) in the EMR, Intranet dashboards, EMR alerts, data repositories and Web-based clinical tools. Four local CDSSs were integrated into the EMR^{43,44,62} and 2 automatically activated.^{44,62} The other 2 required the PCP to activate the CDSS. Studies assessing PDMP CDSSs did not report any evidence-based design components.

Study Characteristics

All studies occurred in the United States and practice settings were mostly primary care clinics. Three were set in the emergency department.^{44,47,49} All the local CDSSs, and 3 of the PDMP CDSS studies^{47,64,66} were designed to assess whether a CDSS alone or incorporated into a multi-faceted intervention improved prescribing or adherence to guidelines. The remaining PDMP CDSS studies determined providers' behavior, knowledge of, attitudes toward and use of CDSSs. Local CDSS study designs included 4 prepost interventions, a cluster randomized controlled trial (RCT) and a mixed-methods evaluation. The 8 PDMP CDSS studies included a wide variety of study designs including: 3 prepost interventions, a cross-sectional survey, 2 qualitative, 1 mixed-methods, and 1 retrospective cohort. Study aims and designs are summarized in Table 2 and described in detail in Appendix 3. One study was part of a stepped approach in evaluating a complex intervention.⁶³ About half of the studies that assessed the impact of an intervention included a process evaluation

Figure 1. PRISMA Flow Diagram.

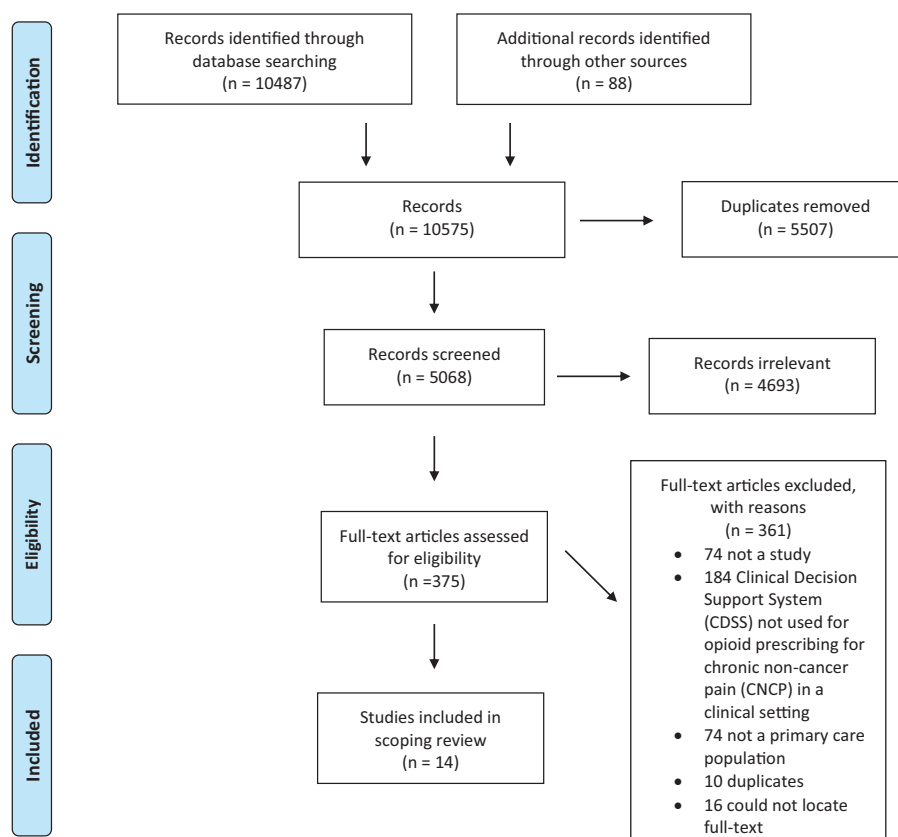


Table 1. Study Setting, Participants, Clinical Decision Support System (CDSS) Type, and Inclusion of Evidence-Based Components

Characteristic		Local CDSS* N (%)	PDMP CDSS** N (%)
Country	United States	6/6 (100%)	8/8 (100%)
Practice settings	Primary care clinic	5/6 (83%)	6/8 (75%)
	Emergency department	1/6 (17%)	2/8 (25%)
Types of PCPs	Physicians	6/6 (100%)	7/8 (88%)
	NPs	6/6 (100%)	4/8 (50%)
CDSS type	Dashboard	2/6 (33%)	0/8 (0%)
	Protocol	2/6 (33%)	0/8 (0%)
	Alert	1/6 (17%)	0/8 (0%)
	Clinical tool	1/6 (17%)	0/8 (0%)
	Data repository	0/6 (0%)	8/8 (100%)
Evidence-based CDSS components ^o	Integrated into EMR	3/6 (50%)	0/5 (0%)***
	Automatically activates	2/6 (33%)	0/5 (0%)***
	Requires a reason for over-ride	0/6 (0%)	0/5 (0%)***
	Provides advice to patients and providers	0/6 (0%)	0/5 (0%)***

Abbreviations: CDSS, clinical decision support system; EMR, electronic medical record; N/A, Not applicable; NP, nurse practitioner; PDMP, prescription drug monitoring program; PCPs, primary care providers.

*Local CDSSs are used locally within a specific health center, health system or clinic.

**PDMP CDSSs are large, centralized, government-run databases.

***We excluded 3 studies because they included multiple PDMP CDSSs, and did not provide information on a specific CDSS.^{45,47,49}

Note: Unless a study stated a component was included (e.g. automatic activation), we assumed it was not.

(measures assessing if program components had been implemented as intended).^{41,43,47,49,62–64} Two studies reported using a theoretical approach in implementation and evaluation processes.^{61,63}

Implementation Processes

All the studies on local CDSSs described their implementation process, but provided little detail. None of the PDMP CDSS studies described implementation processes.

Study Findings

Local CDSSs

Anderson et al⁴¹ found that the CDSS and summary reports improved compliance with guidelines; Canada et al⁴³ reported that a CDSS plus monetary incentives improved adherence to guidelines; Downes et al⁶² found that a CDSS and electronic reports reduced opioid prescribing and increased urine drug testing and use of pain

Table 2. Aims and Designs of Included Studies

Aims	Design	Local CDSS* N (%)	PDMP CDSS** N (%)
To determine if a multi-faceted intervention improved prescribing/guideline adherence	Cluster RCT***	1/6 (17%)	0/8 (0%)
	Pre-post	4/6 (33%)	0/8 (0%)
To determine whether a CDSS improved prescribing/guideline adherence	Pre-post	0/6 (0%)	1/8 (13%)
To determine if PCPs used a CDSS	Retrospective cohort	0/6 (0%)	1/8 (13%)
	Cross-sectional survey	0/6 (0%)	1/8 (13%)
To determine if an intervention affected provider knowledge, behavior, attitudes and/or use related to CDSS	Mixed-methods	0/6 (0%)	1/8 (13%)
	Pre-post	0/6 (0%)	2/8 (25%)
To learn about factors affecting opioid prescribing for CNCP, including use of CDSS	Qualitative	0/6 (0%)	2/8 (25%)
To pilot a multi-component intervention, including a CDSS	Mixed-methods	1/6 (17%)	0/8 (0%)

Abbreviations: CDSS, clinical decision support system; CNCP, chronic non-cancer pain; N/A, not applicable; PDMP, prescription drug monitoring program; RCT, randomized controlled trial; PCPs, primary care providers.

*Local CDSSs are used locally within a specific health center, health system or clinic.

**PDMP CDSSs are large, centralized, government-run databases.

***CDSS included in both study arms.

Table 3. Funding and Relationship between Developers and Evaluators

		Local CDSS* N (%)	PDMP CDSS** N (%)
Funding for CDSS development	Public/Non-profit	3/6 (50%)	0/8 (0%)
	Industry	0/6 (0%)	0/8 (0%)
	Not sponsored	0/6 (0%)	0/8 (0%)
	Unclear or not reported	3/6 (50%)	8/8 (100%)
Funding for evaluation	Public/non-profit	4/6 (67%)	5/8 (63%)
	Industry	0/6 (0%)	0/8 (0%)
	Not sponsored	0/6 (0%)	2/8 (25%)
	Unclear or not reported	2/6 (33%)	1/8 (13%)
Relationship between developers and evaluators	Same person, group, or organization	4/6 (67%)	0/8 (0%)
	Different person, group, or organization	0/6 (0%)	0/8 (0%)
	Unclear or not reported	2/6 (33%)	8/8 (100%)

Abbreviations: CDSS, clinical decision support system; PDMP, Prescription drug monitoring program.

*Local CDSSs are used locally within a specific health center, health system or clinic.

**PDMP CDSSs are large, centralized, government-run databases.

contracts; Gugelmann et al⁴⁴ found that the CDSS reduced opioid prescribing; Liebschutz et al⁶¹ reported that a multi-faceted intervention that included a CDSS in both study arms also reduced opioid prescribing; and Seal et al⁶³ found in a multi-component intervention (with CDSS in both arms) that providers “abandoned use” of the CDSS.

PDMP CDSSs

Baehren et al⁴⁷ found that physicians who used PDMP data wrote fewer opioid prescriptions in 61% of cases and more opioid prescriptions in 39% of cases; Binswanger et al⁶⁴ found that a multi-component intervention improved adherence to guidelines; Chaudhary et al⁵² found that most PCPs reported always checking the PDMP before prescribing opioids to new patients. Click et al⁵⁰ found that providers have positive views about PDMPs, but reported barriers in using them. Coleman⁵¹ found that in 5 of 7 records of patient prescribed opioids, providers accessed the PDMP. Freeman et al⁶⁵ reported that PDMPs are key tools for PCPs and that barriers included a lack of integration; Kohlbeck et al⁴⁹ reported that an educational intervention increased providers’ knowledge of, behavior, and attitudes toward PDMP CDSSs; Patchett et al⁶⁶ reported that a multi-component intervention increased use of a PDMP and led to a reduction in opioid prescribing.

Funding and Conflict of Interest

All but 2 local CDSS studies reported on funding for CDSS evaluation;^{44,62} and 3 others were missing information on funding for CDSS development.^{44,63}

All PDMP studies except 1⁶⁶ provided information on funding for evaluation, but none provided information on funding for development. For all 6 local CDSS studies, the developers were also the evaluators or the relationship was unclear or not stated. No evaluators of PDMPs provided information on their relationship to the PDMP developer (Table 3).

Discussion

We identified 14 studies published between 2009 and 2019 that examined CDSSs for opioid prescribing for CNCP in primary care clinical settings. Six of the studies examined local CDSSs (that were used locally within a specific health center, health system or clinic) and 8 examined PDMP CDSSs. Studies evaluating CDSS impact found that the CDSS (alone or more commonly, part of a dual or multi-component intervention) led to more appropriate prescribing practices and/or adherence to guidelines. Several PDMP CDSS studies assessed providers’ views on, and/or their use of PDMP CDSSs. These studies reported frequent use of the PDMP CDSS and positive views toward the CDSS with some acknowledgment of the barriers and limitations. These findings are similar to a recent qualitative rapid review that asked providers about the use of PDMPs.⁶⁷ No study in our review, however, contained an assessment of patient health outcomes or assessed for unintended consequences. In addition, in 4 studies the evaluators were also the CDSS developers, a potentially useful situation, but one that presents a potential conflict of interest,^{26,27}

that was not addressed by the investigators. We also found that few CDSSs included evidence-based components and that in only 1 study investigators reported following current guidance for development and evaluation of complex interventions.^{35,36}

Our finding that there were only 14 studies, and only 1 RCT, which met our inclusion criteria is surprising. In contrast, a 2015 systematic review found 7 RCT studies of CDSSs for antibiotic prescribing by primary care providers.²⁸ There may be several contributing factors. The prescription opioid crisis only gained widespread attention in the last decade,⁶⁸ and it takes time to develop a complex intervention like a CDSS.³⁶ It is also possible that some CDSSs failed to show promise early on and development was subsequently stalled or halted. Accordingly, there are a number of reports on the development of a CDSSs for opioid prescribing for CNCP where clinical outcomes have not been reported yet.^{69–72} And finally, it is possible that CDSSs are being used without an evaluation plan, as has occurred with many PDMP CDSSs.⁷³ This may be because of a demand for immediate solutions to the opioid crisis and an evaluation of a CDSS takes significant time and money. However, since CDSSs frequently do not improve patient outcomes,^{17–20} and may lead to unintended consequences, a comprehensive evaluation is essential.⁷⁴

Most studies in our review that assessed the impact of the CDSS reported an improvement in prescribing or better adherence to clinical practice guidelines. This aligns with previous research in other fields: CDSSs have a modest impact on clinician performance (a care process outcome).^{17–25} However, these results need careful interpretation. Most studies were prepost, nonrandomized control or observational designs. Although—consistent with guidance for scoping reviews^{55,56}—we did not conduct a quality assessment; these types of study designs have greater threats to validity.⁷⁵ In addition, in most of the studies, the CDSS was part of a larger intervention, so its specific impact was unclear. Another reason for caution is that no studies assessed patient health outcomes, such as quality of life, morbidity, and mortality.^{76–78} Reductions in opioid prescribing and better adherence to guidelines may have unintended consequences.³⁶ For example, studies report that patients often turn to illicit sources of opioids when they have reduced access to prescribed opioids, increasing their risk of overdose.^{79–84} Several studies in a systematic review

found that heroin overdoses increased after a PDMP CDSS was implemented.⁷⁴ A more recent systematic review, however, found no consistent association between population-level opioid-related harms (including heroin use and overdoses) and PDMP CDSSs.⁸⁵ We also noted a conflict of interest in some studies where the developers were also the evaluators. Systematic reviews in other fields have demonstrated that when the CDSS evaluator is also the developer, outcomes are better.^{26,27} It is possible that developers achieve better outcomes because they design effective implementation plans,²⁶ but it is possible that the conflict of interest leads to conscious or unconscious bias.^{26,86–92} Interestingly, none of the studies reported funding from or involvement of for-profit entities. It is possible that CDSSs developed by for-profit entities are not undergoing a publicly reported evaluation. This is problematic, and as a recent criminal case demonstrated, can lead to potential harm to patients.⁹³

We found that few of the CDSSs incorporated evidence-based design components. In only 1 study did researchers follow guidance for designing and evaluating complex interventions. Developers may not have incorporated evidence-based components because of the lag time between development and evaluation: when the CDSS was created the developers may not have had access to systematic reviews on the various components. The developer may also feel that the evidence does not apply to this particular subspecialty or setting.⁹⁴ Another reason may be a general excitement and overconfidence in e-health technologies.⁹⁵ Funders and developers may be too eager to solve the problem of unsafe opioid prescribing using eHealth technologies and are not ensuring that developers are building on information from the medical literature.⁹⁵ Changes are occurring. Between 2012 and 2016, the Substance Abuse and Mental Health Services Administration funded 9 projects to integrate PDMP data into EMRs.⁹⁶ Investigators might not follow guidance for complex evaluations because it is a lengthy and expensive iterative process before a definitive evaluation.^{35–37,97} This is a widespread issue—few complex interventions seem to undergo modeling, pilot and feasibility testing,⁹⁸ and many lack process evaluations.^{99,100} This is problematic. If researchers conduct a trial without testing components, possible causal pathways, uncertainties, contextual factors, and implementation approaches, they risk wasting

resources on an expensive trial and perhaps causing harm.^{35,37,101} Conversely, if the evaluation takes too long, the technology could become obsolete before it gains widespread uptake.⁵⁹ Adopting rapid, concurrent and iterative pilot and feasibility studies may be the best approach.^{102–104}

Limitations

There are 2 main limitations in our review. In the gray literature search we may have missed non-English-language studies, as we conducted the searches only in English. Second, several of the studies included both PCPs and other provider types (we excluded those with less than 50% PCPs), and, as these studies only reported aggregate outcomes, they may not accurately reflect the PCP population.

Conclusion and Next Steps

Our review reveals that few studies have rigorously assessed the use of CDSSs in the context of opioid prescribing for CNCP in the primary care setting. More high-quality studies are needed. Going forward, investigators should include evidence-based components into the design of CDSSs and follow guidance for the development and evaluation of complex interventions, including pilot studies, process evaluations, and an assessment for unintended consequences.

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References

1. Fischer B, Keates A, Bühringer G, Reimer J, Rehm J. Non-medical use of prescription opioids and prescription opioid-related harms: why so markedly higher in North America compared to the rest of the world?. *Addict Abingdon Engl* 2014;109:177–81.
2. Report of the International Narcotics Control Board for 2017 [Internet]. International Narcotics Control Board. 2017. Available from: <https://www.incb.org/incb/en/publications/annual-reports/annual-report-2017.html>.
3. Seth P. Overdose Deaths Involving Opioids, Cocaine, and Psychostimulants—United States, 2015–2016. *MMWR Morb Mortal Wkly Rep* [Internet]. 2018 [cited 2018 Apr 23];67. Available from: <https://www.cdc.gov/mmwr/volumes/67/wr/mm6712a1.htm>.
4. Government of Canada. National Report: Apparent opioid-related deaths in Canada [Internet]. 2018 [cited 2018 Apr 23]. Available from: www.canada.ca/en/public-health/services/publications/healthy-living/national-report-apparent-opioid-related-deaths-released-march-2018.html.
5. Dowell D, Haegerich TM, Chou R. CDC Guideline for prescribing opioids for chronic pain—United States, 2016. *JAMA* 2016;315:1624–45.
6. Busse J. The 2017 Canadian guideline for opioids for chronic non-cancer pain [Internet]. National pain center. 2017. Available from: <http://www.cmaj.ca/content/suppl/2017/05/03/189.18.E659.DC1/170363-guide-1-at-updated.pdf>.
7. Dunn KM, Saunders KW, Rutter CM, et al. Overdose and prescribed opioids: Associations among chronic non-cancer pain patients. *Ann Intern Med* 2010;152:85–92.
8. Bohnert AB, Valenstein M, Bair MJ, et al. Association between opioid prescribing patterns and opioid overdose-related deaths. *JAMA* 2011;305:1315–21.
9. Gomes T, Mamdani MM, Dhalla IA, Paterson JM, Juurlink DN. Opioid dose and drug-related mortality in patients with nonmalignant pain. *Arch Intern Med* 2011;171:686–91.
10. Chou R, Turner JA, Devine EB, et al. The effectiveness and risks of long-term opioid therapy for chronic pain: A systematic review for a National Institutes of Health Pathways to Prevention Workshop. *Ann Intern Med* 2015;162:276–86.
11. Volkow ND, McLellan TA, Cotto JH, Karithanom M, Weiss S. Characteristics of opioid prescriptions in 2009. *JAMA* 2011;305:1299–301.
12. Levy B, Paulozzi L, Mack KA, Jones CM. Trends in Opioid Analgesic-Prescribing Rates by Specialty, U.S., 2007–2012. *Am J Prev Med* 2015;49:409–13. Sep.
13. Health Quality Ontario. Starting on Opioids in Ontario [Internet]. 2017 [cited 2018 Jul 12]. Available from: <http://startingonopioids.hqontario.ca/>.
14. Kawamoto K, Houlihan CA, Balas EA, Lobach DF. Improving clinical practice using clinical decision support systems: A systematic review of trials to identify features critical to success. *BMJ* 2005;330:765.
15. Sim I, Gorman P, Greenes RA, et al. Clinical Decision support systems for the practice of evidence-based medicine. *J Am Med Inform Assoc JAMIA* 2001;8:527–34.
16. Haynes RB, Wilczynski NL. Effects of computerized clinical decision support systems on practitioner performance and patient outcomes: Methods of a decision-maker-researcher partnership systematic review. *Implement Sci* 2010;5:12.

17. Jia P, Zhang L, Chen J, Zhao P, Zhang M. The Effects of Clinical Decision Support Systems on Medication Safety: An Overview. *PLoS One* 2016;11:e0167683.
18. Lloyd R, Ramakrishna A, Mackay JA, et al. Computerized clinical decision support systems for acute care management: A decision-maker-researcher partnership systematic review of effects on process of care and patient outcomes. *Implement Sci IS* 2011;6:91.
19. Misra S, Gerstein HC, Garg AX, et al. Computerized clinical decision support systems for chronic disease management: A decision-maker-researcher partnership systematic review. *Implement Sci IS* 2011;6:92.
20. Moja L, Kwag KH, Lytras T, et al. Effectiveness of Computerized decision support systems linked to electronic health records: A systematic review and meta-analysis. *Am J Public Health* 2014;104:e12–22.
21. Jaspers MWM, Smeulders M, Vermeulen H, Peute LW. Effects of clinical decision-support systems on practitioner performance and patient outcomes: A synthesis of high-quality systematic review findings. *J Am Med Inform Assoc* 2011;18:327–34.
22. Nieuwlaat R, Connolly SJ, Mackay JA, et al. Computerized clinical decision support systems for therapeutic drug monitoring and dosing: A decision-maker-researcher partnership systematic review. *Implement Sci IS* 2011;6:90.
23. Sebaldt RJ, Mackay JA, Prorok JC, et al. Computerized clinical decision support systems for primary preventive care: A decision-maker-researcher partnership systematic review of effects on process of care and patient outcomes. *Implement Sci IS* 2011;6:87.
24. Hemens BJ, Holbrook A, Tonkin M, et al. Computerized clinical decision support systems for drug prescribing and management: A decision-maker-researcher partnership systematic review. *Implement Sci IS* 2011;6:89.
25. Bright TJ, Wong A, Dhurjati R, et al. Effect of clinical decision-support systems: A systematic review. *Ann Intern Med* 2012;157:29–43.
26. Roshanov PS, Fernandes N, Wilczynski JM, et al. Features of effective computerised clinical decision support systems: Meta-regression of 162 randomised trials. *BMJ* 2013;346:f657–f657.
27. Garg AX, Adhikari NKJ, McDonald H, et al. Effects of computerized clinical decision support systems on practitioner performance and patient outcomes: A systematic review. *JAMA* 2005;293:1223–38.
28. Holstiege J, Mathes T, Pieper D. Effects of computer-aided clinical decision support systems in improving antibiotic prescribing by primary care providers: A systematic review. *J Am Med Inform Assoc* 2015;22:236–42.
29. Stultz JS, Nahata MC. Computerized clinical decision support for medication prescribing and utilization in pediatrics. *J Am Med Inform Assoc* 2012;19:942–53.
30. Moxey A, Robertson J, Newby D, Hains I, Williamson M, Pearson SA. Computerized clinical decision support for prescribing: Provision does not guarantee uptake. *J Am Med Inform Assoc* 2010;17:25–33.
31. Patterson ES, Doebbeling BN, Fung CH, Militello L, Anders S, Asch SM. Identifying barriers to the effective use of clinical reminders: Bootstrapping multiple methods. *J Biomed Inform* 2005;38:189–99.
32. Arts DL, Medlock SK, van Weert H, Wyatt JC, Abu-Hanna A. Acceptance and barriers pertaining to a general practice decision support system for multiple clinical conditions: A mixed methods evaluation. *PLoS ONE* 2018;13:e0193187. [Internet]. 2018 Apr [cited 2018 19] Available from: <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5908177/>.
33. Dalleur O, Seger DL, Slight SP, et al. Inappropriate overrides of age-related alerts in prescriber order entry. *J Gen Intern Med*. 2015;S189–S190.
34. Slight SP, Beeler PE, Seger DL, et al. An evaluation of drug-allergy interaction alert overrides in inpatients. *J Gen Intern Med* 2015;S81.
35. Craig P, Dieppe P, Macintyre S, Michie S, Nazareth I, Petticrew M. Developing and evaluating complex interventions: the new Medical Research Council guidance. *BMJ*. 2008;337: a1655.
36. Craig P, Dieppe P, Macintyre S, Michie S, Nazareth I, Petticrew M. Developing and evaluating complex interventions: The new Medical Research Council guidance. *Int J Nurs Stud* 2013;50:587–92.
37. Catwell L, Sheikh A. Evaluating eHealth interventions: The need for continuous systemic evaluation. *PLoS Med* 2009;6:e1000126. [Internet] Aug 18 [cited 2018 Sep 24] Available from: <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2719100/>
38. Shah A. Characteristics of Initial prescription episodes and likelihood of long-term opioid use—United States, 2006–2015. *MMWR Morb Mortal Wkly Rep* [Internet]. 2017 [cited 2020 Feb 14];66. Available from: <https://www.facebook.com/CDCMMWR>.
39. Dasgupta N, Beletsky L, Ciccarone D. Opioid crisis: No Easy fix to its social and economic determinants. *Am J Public Health* 2018;108:182–6.
40. Brooks EA, Unruh A, Lynch ME. Exploring the lived experience of adults using prescription opioids to manage chronic noncancer pain. *Pain Res Manag J Can Pain Soc* 2015;20:15–22.
41. Anderson D, Zlateva I, Khatri K, Ciaburri N. Using health information technology to improve

- adherence to opioid prescribing guidelines in primary care. *Clin J Pain* 2015;31:573–9.
42. Patel S, Carmichael JM, Taylor JM, Bounthavong M, Higgins DT. Evaluating the Impact of a clinical decision support tool to reduce chronic opioid dose and decrease risk classification in a veteran population. *Ann Pharmacother* 2018;52:325–31.
43. Canada RE, DiRocco D, Day S. A better approach to opioid prescribing in primary care. *J Fam Pract* 2014;63:E1–E8.
44. Gugelmann H, Shofer FS, Meisel ZF, Perrone J. Multidisciplinary intervention decreases the use of opioid medication discharge packs from 2 urban EDs. *Am J Emerg Med* 2013;31:1343–8.
45. Sproule BA. Prescription Monitoring programs in Canada: best practice and program review [Internet]. Canadian Centre on Substance Abuse, Ottawa, ON 2015;[cited 2018 Jun 14]. Available from: <http://www.ccsa.ca/Resource%20Library/CCSA-Prescription-Monitoring-Programs-in-Canada-Report-2015-en.pdf>.
46. Rutkow L, Smith KC, Lai AY, Vernick JS, Davis CS, Alexander GC. Prescription drug monitoring program design and function: A qualitative analysis. *Drug Alcohol Depend* 2017;180:395–400.
47. Baehren DF, Marco CA, Droz DE, Sinha S, Callan EM, Akpunonu P. A statewide prescription monitoring program affects emergency department prescribing behaviors. *Ann Emerg Med* 2010;56:19–23.e1–3.
48. Lin HC, Wang Z, Boyd C, Simoni-Wastila L, Buu A. Associations between statewide prescription drug monitoring program (PDMP) requirement and physician patterns of prescribing opioid analgesics for patients with non-cancer chronic pain. *Addict Behav* 2018;76:348–54.
49. Kohlbeck S, Akert B, Pace C, Zosel A. A multistep approach to address clinician knowledge, attitudes, and behavior around opioid prescribing. *Wis Med J* 2018;117:38–41.
50. Click IA, Basden JA, Bohannon JM, Anderson H, Tudiver F. Opioid Prescribing in rural family practices: A qualitative study. *Subst Use Misuse* 2018;53:533–40.
51. Coleman CD. Evidence-Based strategies to minimize risk for opioid pain medication misuse among patients with chronic pain in a primary care setting. *Diss Abstr Int Sect B Sci Eng* 2016;64:
52. Chaudhary S, Compton P. Use of risk mitigation practices by family nurse practitioners prescribing opioids for the management of chronic nonmalignant pain. *Subst Abuse* 2017;38:95–104.
53. Arksey H, O'Malley L. Scoping studies: towards a methodological framework. *Int J Soc Res Methodol* 2005;8:19–32.
54. Levac D, Colquhoun H, O'Brien KK. Scoping studies: Advancing the methodology. *Implement Sci* 2010;5:69.
55. Colquhoun HL, Levac D, O'Brien KK, et al. Scoping reviews: time for clarity in definition, methods, and reporting. *J Clin Epidemiol* 2014;67:1291–4.
56. Peters M, Godfrey C, McInerney P, Soares C, Khalil H, Parker D. The Joanna Briggs Institute Reviewers' Manual 2015: Methodology for JBI scoping reviews. 2015 Jan 1 [cited 2018 Feb 7]; Available from: <https://espace.library.uq.edu.au/view/UQ:371443>.
57. Tricco AC, Lillie E, Zarin W, et al. PRISMA extension for scoping reviews (PRISMA-ScR): Checklist and explanation. *Ann Intern Med* 2018;169:467. [Internet] Sep 4 [cited 2018 Sep 21]; Available from: <http://annals.org/article.aspx?doi=10.7326/M18-0850>.
58. Li G, Abbade LPF, Nwosu I, et al. A scoping review of comparisons between abstracts and full reports in primary biomedical research. *BMC Med Res Methodol* 2017;17:181.
59. Main C, Moxham T, Wyatt JC, Kay J, Anderson R, Stein K. Computerised decision support systems in order communication for diagnostic, screening or monitoring test ordering: systematic reviews of the effects and cost-effectiveness of systems [Internet]. NIHR J Library 2010; [cited 2018 Jan 7]. Available from: <https://www.ncbi.nlm.nih.gov/books/NBK56829/>.
60. Grey matters: A practical tool for searching health-related grey literature | CADTH.ca [Internet]. [cited 2018 Sep 24]. Available from: <https://www.cadth.ca/resources/finding-evidence/grey-matters>.
61. Liebschutz JM, Xuan Z, Shanahan CW, et al. Improving adherence to long-term opioid therapy guidelines to reduce opioid misuse in primary care: A cluster-randomized clinical trial. *JAMA Intern Med* 2017;177:1265–72.
62. Downes JM, Klepser DG, Foster J, Nelson M. Development of a standardized approach for managing opioids in adults with chronic noncancer pain. *Am J Health-Syst Pharm* 2018;75:321–6.
63. Seal KH, Borsari B, Tighe J, et al. Optimizing pain treatment interventions (OPTI): A pilot randomized controlled trial of collaborative care to improve chronic pain management and opioid safety-rationale, methods, and lessons learned. *Contemp Clin Trials* 2019;77:76–85.
64. Binswanger IA, Joseph N, Hanratty R, et al. Novel opioid safety clinic initiative to deliver guideline-concordant chronic opioid therapy in primary care. *Mayo Clin Proc Innov Qual Outcomes* 2018;2:309–16.
65. Freeman PR, Curran GM, Drummond KL, et al. Utilization of prescription drug monitoring programs for prescribing and dispensing decisions: Results from a multi-site qualitative study. *Res Soc Adm Pharm RSAP* 2019;15:754–60.

66. Patchett D, Grover M, Kresin M, et al. The benefits of a standardized approach to opioid prescribing. *J Fam Pract* 2019;68:E1–E7.
67. Prescribing and Dispensing policies to address harms associated with prescription drug abuse | CADTH.ca [Internet]. [cited 2018 Aug 20]. Available from: <https://www.cadth.ca/prescribing-and-dispensing-policies-address-harms-associated-prescription-drug-abuse>.
68. McCarthy M. Containing the opioid overdose epidemic. *BMJ* 2012;345:e8340–e8340.
69. Trafton J, Martins S, Michel M, et al. Evaluation of the acceptability and usability of a decision support system to encourage safe and effective use of opioid therapy for chronic, noncancer pain by primary care providers. *Pain Med* 2010;11:575–85.
70. Furlan AD, Reardon R, Salach L. The opioid manager: a point-of-care tool to facilitate the use of the Canadian Opioid Guideline. *J Opioid Manag* 2012;8:57–61.
71. Midboe AM, Lewis ET, Cronkite RC, et al. Behavioral medicine perspectives on the design of health information technology to improve decision-making, guideline adherence, and care coordination in chronic pain management. *Behav Med Pract Policy Res* 2011;1:35–44.
72. Harle CA, Bauer SE, Hoang HQ, Cook RL, Hurley RW, Fillingim RB. Decision support for chronic pain care: how do primary care physicians decide when to prescribe opioids? a qualitative study. *BMC Fam Pract* 2015;16:48.
73. Finley EP, Garcia A, Rosen K, McGeary D, Pugh MJ, Potter JS. Evaluating the impact of prescription drug monitoring program implementation: a scoping review. *BMC Health Serv Res* 2017;17:420.
74. Fink DS, Schleimer JP, Sarvet A, et al. Association between prescription drug monitoring programs and nonfatal and fatal drug overdoses: A systematic review. *Ann Intern Med* 2018;168:783. Jun 5.
75. Grading quality of evidence and strength of recommendations. *BMJ* 2004;328:1490.
76. Health Quality Ontario. Recommendations for adoption: Opioid Prescribing for chronic pain [Internet]. 2018 [cited 2018 Jul 20]. Available from: <http://www.hqontario.ca/Portals/0/documents/evidence/quality-standards/qs-opioid-chronic-pain-recommendations-for-adoption-en.pdf>.
77. Ciani O, Buyse M, Drummond M, Rasi G, Saad ED, Taylor RS. Time to review the role of surrogate end points in health policy: State of the art and the way forward. *Value Health* 2017;20:487–95.
78. Wittes J, Lakatos E, Probstfield J. Surrogate end-points in clinical trials: Cardiovascular diseases. *Statist Med* 1989;8:415–25.
79. Gomes T, Khuu W, Martins D, et al. Contributions of prescribed and non-prescribed opioids to opioid related deaths: Population based cohort study in Ontario. Canada. *BMJ* 2018;362:k3207.
80. Mars SG, Bourgois P, Karandinos G, Montero F, Ciccarone D. “Every ‘never’ I ever said came true”: Transitions from opioid pills to heroin injecting. *Int J Drug Policy* 2014;25:257–66.
81. Compton WM, Jones CM, Baldwin GT. Relationship between nonmedical prescription-opioid use and heroin use. *N Engl J Med* 2016;374:154–63.
82. Baldwin N, Gray R, Goel A, Wood E, Buxton JA, Rieb LM. Fentanyl and heroin contained in seized illicit drugs and overdose-related deaths in British Columbia, Canada: An observational analysis. *Drug Alcohol Depend* 2018;185:322–7.
83. Rubin R. Illicit fentanyl driving opioid overdose deaths. *JAMA* 2017;318:2174.
84. O'Donnell J, Gladden RM, Seth P. Trends in deaths involving heroin and synthetic opioids excluding methadone, and law enforcement drug product reports, by census region—United States. [Internet] 2017;2006–2015.[cited 2018 Oct 1]. Available from: https://www-cdc-gov.myaccess.library.utoronto.ca/mmwr/volumes/66/wr/mm6634a2.htm?s_cid=mm6634a2_e.
85. Rhodes E, Wilson M, Robinson A, Hayden JA, Asbridge M. The effectiveness of prescription drug monitoring programs at reducing opioid-related harms and consequences: A systematic review. *BMC Health Serv Res* 2019;19:784.
86. Heres S, Davis J, Maino K, Jetzinger E, Kissling W, Leucht S. Why olanzapine beats risperidone, risperidone beats quetiapine, and quetiapine beats olanzapine: An exploratory analysis of head-to-head comparison studies of second-generation antipsychotics. *AJP* 2006;163:185–94.
87. Lexchin J, Bero LA, Djulbegovic B, Clark O. Pharmaceutical industry sponsorship and research outcome and quality: Systematic review. *BMJ* 2003;326:1167–1170.
88. Lesser LI, Ebbeling CB, Goozner M, Wypij D, Ludwig DS. Relationship between funding source and conclusion among nutrition-related scientific articles. *PLoS Med* 2007;4:e5.
89. Procyshyn RM, Chau A, Fortin P, Jenkins W. Prevalence and outcomes of pharmaceutical industry-sponsored clinical trials involving clozapine, risperidone, or olanzapine. *Can J Psychiatry* 2004;49:601–6.
90. Turner EH, Matthews AM, Linardatos E, Tell RA, Rosenthal R. Selective publication of antidepressant trials and its influence on apparent efficacy. *N Engl J Med* 2008;358:252–60.
91. Doshi P, Dickersin K, Healy D, Vedula SS, Jefferson T. Restoring invisible and abandoned trials: A call for people to publish the findings. *BMJ* 2013;346:f2865–f2865.

92. Chan AW, Hróbjartsson A, Haahr MT, Gotzsche PC, Altman DG. Empirical evidence for selective reporting of outcomes in randomized trials: Comparison of protocols to published articles. *JAMA* 2004;291:2457–65.
93. Electronic health records vendor to pay \$145 million to resolve criminal and civil investigations [Internet]. 2020 [cited 2020 Feb 14]. Available from: <https://www.justice.gov/opa/pr/electronic-health-records-vendor-pay-145-million-resolve-criminal-and-civil-investigations-0>.
94. Mair FS, May C, O'Donnell C, Finch T, Sullivan F, Murray E. Factors that promote or inhibit the implementation of e-health systems: an explanatory systematic review. *Bull World Health Organ* 2012;90:357–64.
95. Black AD, Car J, Pagliari C, et al. The impact of eHealth on the quality and safety of health care: A systematic overview. *PLoS Med* 2011;8:e1000387. [Internet] Jan 18 [cited Jul 18] Available from: <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3022523/>.
96. National Center for Injury Prevention and Control. Integrating & Expanding Prescription Drug Monitoring Program Data: Lessons from Nine States. Centre for Disease Control and Prevention 2017.
97. Moore GF, Audrey S, Barker M, et al. Process evaluations of complex interventions: UK Medical Research Council (MRC) guidance [Internet]. MRC Population Health Science Research Network 2014;[cited 2018 Jul 23]. Available from: <https://mrc.ukri.org/documents/pdf/mrc-phsrn-process-evaluation-guidance-final/>
98. Chan CL, Leyrat C, Eldridge SM. Quality of reporting of pilot and feasibility cluster randomised trials: A systematic review. *BMJ Open* 2017;7:e016970.
99. Liu H, Muhunthan J, Hayek A, et al. Examining the use of process evaluations of randomised controlled trials of complex interventions addressing chronic disease in primary health care—A systematic review protocol. *Syst Rev* 2016;5. [cited 2018 Sep 24] Available from: <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4986376/>.
100. Oakley A, Strange V, Bonell C, Allen E, Stephenson J. Process evaluation in randomised controlled trials of complex interventions. *BMJ* 2006;332:413–6.
101. Greenhalgh T, Russell J. Why do evaluations of eHealth programs fail? An alternative set of guiding principles. *PLoS Med* 2010;7:e1000360. Available from: <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2970573/>.
102. Basit MA, Baldwin KL, Kannan V, et al. Agile acceptance test-driven development of clinical decision support advisories: feasibility of using open source software. *JMIR Med Inform* 2018;6:e23. [Internet] Apr 13 [cited 2019 May] Available from: <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5924365/>.
103. Kannry J, McCullagh L, Kushniruk A, Mann D, Edonyabo D, McGinn T. A Framework for Usable and Effective Clinical Decision Support: Experience from the iCPR Randomized Clinical Trial.eGEMs 2015;3:10.
104. Kannan V, Fish J, Mutz J, et al. Rapid Development of specialty population registries and quality measures from electronic health record data: An agile framework. *Methods Inf Med* 2017;56:e74–83.

Appendix 1: Medline Search Strategy

#	Searches
1	narcotics/ or analgesics, opioid/ or alfentanil/ or alphaprodine/ or buprenorphine/ or buprenorphine, naloxone drug combination/ or butorphanol/ or codeine/ or dextromoramide/ or dextropropoxyphene/ or dihydromorphine/ or diphenoxylate/ or "enkephalin, ala ² -mephe ⁴ -gly ⁵ -"/ or "enkephalin, d-penicillamine ^{2,5} -"/ or ethylketocyclazocine/ or ethylmorphine/ or etorphine/ or fentanyl/ or heroin/ or hydrocodone/ or hydromorphone/ or levorphanol/ or meperidine/ or meptazinol/ or methadone/ or methadyl acetate/ or morphine/ or nalbuphine/ or opiate alkaloids/ or oxycodone/ or oxymorphone/ or pentazocine/ or phenazocine/ or phenoperidine/ or pirinitramide/ or promedol/ or sufentanil/ or tilidine/ or tramadol/
2	(analgesic* or opioid* or opiate* or narcotic* or alfentanil or alphaprodine or buprenorphine or (buprenorphine adj2 naloxone) or suboxone or subutex or butorphanol or codeine or dihydrocodeine or dextromoramide or dextropropoxyphene or dihydromorphine or diphenoxylate or ethylketocyclazocine or ethylmorphine or etorphine or fentanyl or duragesic or hydrocodone or hydromorphone or levorphanol or meperidine or meptazinol or methadone or methadyl acetate or morphine or nalbuphine or oxycodone or oxymorphone or pentazocine or phenazocine or phenoperidine or pirinitramide or promedol or sufentanil or tilidine or tramadol or dilaudid or OPANA or targin or tapendatol or nalbuphine or trama*).kf,tw.
3	1 or 2
4	hospital information systems/ or ambulatory care information systems/ or medical order entry systems/ or point-of-care systems/
5	Prescription Drug Monitoring Programs/
6	medical informatics/ or health information exchange/ or medical informatics applications/ or decision making, computer-assisted/ or diagnosis, computer-assisted/ or therapy, computer-assisted/ or drug therapy, computer-assisted/ or decision support techniques/ or "information storage and retrieval"/ or data mining/ or health information interoperability/ or information systems/ or community networks/ or decision support systems, clinical/ or health information systems/ or integrated advanced information management systems/ or management information systems/ or clinical pharmacy information systems/ or database management systems/ or decision support systems, management/ or medical order entry systems/ or reminder systems/ or medical informatics computing/ or nursing informatics/
7	microcomputers/ or computers, handheld/or smartphone/
8	prescription drug misuse/or prescription drug overuse/
9	"Drug and Narcotic Control"/mt, og, pc, sn, td [Methods, Organization & Administration, Prevention & Control, Statistics & Numerical Data, Trends]
10	Drug Monitoring/mt, sn [Methods, Statistics & Numerical Data]
11	4 or 5 or 6 or 7 or 8 or 9 or 10
12	((health* or clinical or medic*) adj3 informatic*).kw, tw.
13	((electronic* or computer*) adj3 order entr*).kw, tw.
14	((phone* or cellphone* or smartphone* or handheld) adj4 (app\$1 or application*)).kw, tw.
15	((decision* or reminder* or point of care or point-of-care or alert*) adj3 (tool* or system* or electronic* or computer-assisted)).kw, tw.
16	(PDMP or PDMPs).kw, tw.
17	(PMP or PMPs).kw, tw.
18	((prescri* or drug* or narcotic*) adj3 monitor*).tw, kw.
19	12 or 13 or 14 or 15 or 16 or 17 or 18
20	11 or 19
21	3 and 20

Database(s): Ovid MEDLINE: Epub Ahead of Print, In-Process & Other Non-Indexed Citations, Ovid MEDLINE® Daily and Ovid MEDLINE® 1946-Present Search Strategy.

Appendix 2: Gray Literature Search

Date	Database	URL	Search Strategy	No. Items Retrieved*	No. Screened (Uploaded to Citation Manager)
January 14, 2020	ClinicalTrials.gov	ClinicalTrials.gov	(Informatics OR order entry OR phone OR handheld OR application OR reminder OR alert OR electronic OR tool OR prescription drug monitoring database OR decision support) AND opioids	119	15
January 14, 2020	WHO ICTRP	https://www.who.int/ictcp/en/	Opioid AND decision support	7	2
			Opioid AND electronic	14	1
			Opioid AND informatics	0	0
			Opioid AND reminder	0	0
			Opioid AND prescription drug monitoring program	1	0
			Opioid AND alert	1	0
			Opioid AND application	5	0
			Opioid AND tool	6	0
			Opioid AND phone	29	0
			Opioid AND handheld	7	1

Databases Gray Literature

Date	Organization	URL	Search Mechanism	Search Strategy	No. Items Retrieved	No. Screened (uploaded to Citation Manager)
January 14, 2020	Open Gray or Open Sige	www.open grey.eu	Site search	(Informatics OR order entry OR phone OR handheld OR application OR reminder OR alert OR electronic OR tool OR prescription drug monitoring database OR decision support) AND opioid	13	0
January 14, 2020	Gray literature Report	https://www.greylit.org/ (no longer updated as of January 2017)	Site search	Opioid	42	0
January 14, 2020	Department of Veterans Affairs	www.va.gov	Site search using Google	(Informatics OR order entry OR phone OR handheld OR application OR reminder OR alert OR electronic OR tool OR prescription drug monitoring program OR decision support) AND opioid	331	0
January 14, 2020	Regenstrief Institute	Regenstrief.org	Site search using Google	(Informatics OR order entry OR phone OR handheld OR application OR reminder OR alert OR electronic OR tool OR prescription drug monitoring program OR decision support) AND opioid	11	4

Continued

Continued

Date	Organization	URL	Search Mechanism	Search Strategy	No. Items Retrieved	No. Screened (uploaded to Citation Manager)
January 14, 2020	Canadian Agency for Drugs and Technologies in Health (CADTH)	https://www.cadth.ca/	Site search	(Informatics OR order entry OR phone OR handheld OR application OR reminder OR alert OR electronic OR tool OR prescription drug monitoring program OR decision support) AND opioid	228	1
January 14, 2020	Institute of Health Economics	http://www.ihe.ca/index.php?publications	Site search	Opioid	2	0
January 14, 2020	Pan-Canadian HTA collaborative	http://www.crd.york.ac.uk/PanHTA/	Site search	Opioid	9	0
January 14, 2020	Programs for Assessment of Technology in Health (Canada)	https://www.cadth.ca/	Site search using Google	Opioid	0	0
January 14, 2020	The International Network of Agencies for Health Technology Assessment (INATHA)	http://www.inahta.org/publications/	Site search	Opioid	11	0
January 14, 2020	Agency for Healthcare Research and Quality (AHRQ)	https://search.ahrq.gov/search?q=opioid+&search_icon.x=0&search_icon.y=0	Site search using Google	Informatics “order entry” phone handheld application reminder alert electronic tool “prescription drug monitoring program” “decision support”AND opioid	103	14
January 14, 2020	Centre for Disease Control (CDC)	CDC.gov	Site search	Opioid “decision support” Opioid “electronic tool” Opioid “reminder system” Opioid “informatics” Opioid “prescription drug monitoring program” Opioid “alert” STOPPED search as no relevant results Opioid “application” Opioid “tool” Opioid “phone” Opioid “handheld”	151 9 17 243 385 945	0 0 0 0 0 0
January 14, 2020	Health IT	https://www.healthit.gov/	Site search using Google	(Informatics OR order entry OR phone OR handheld OR application OR reminder OR alert OR electronic OR tool OR prescription drug monitoring program OR decision support) AND opioid	497	1

Continued

Continued

Date	Organization	URL	Search Mechanism	Search Strategy	No. Items Retrieved	No. Screened (uploaded to Citation Manager)
January 14, 2020	Healthcare Information and Management Systems Society (HIMSS)	HIMSS.org	Site search	Opioid	231	5
January 14, 2020	Ontario MD	OntarioMD.ca	Site search	Opioid	18	0
January 14, 2020	Health ITanalytics	https://www.healthitanalytics.com/	Site search using Google	(Informatics OR order entry OR phone OR handheld OR application OR reminder OR alert OR electronic OR tool OR prescription drug monitoring program OR decision support) AND opioid	15	0
January 14, 2020	National Technical Information Service (NTIS)	https://www.ntis.gov/	Site search	Opioid	0	0

Search Engine Searching

Date	Search Engine	Search Strategy	# screened (Uploaded to citation manager)
January 14, 2020	Google advanced search	opioid decision support OR electronic OR reminder OR informatics OR alert OR application OR tool OR phone OR handheld	First 100 results 37
January 14, 2020	Google advanced search	Opioid AND "prescription drug monitoring program" AND "chronic pain"	First 100 results 5

*Screening first 100 results.