The effectiveness of toolkits as knowledge translation strategies for integrating evidence into clinical care: a systematic review

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ABSTRACT

Objectives: The aim of this systematic review was to evaluate the effectiveness of toolkits as a knowledge translation (KT) strategy for facilitating the implementation of evidence into clinical care. Toolkits include multiple resources for educating and/or facilitating behaviour change.

Design: Systematic review of the literature on toolkits.

Methods: A search was conducted on MEDLINE, EMBASE, PsycINFO and CINAHL. Studies were included if they evaluated the effectiveness of a toolkit to support the integration of evidence into clinical care, and if the KT goal(s) of the study were to inform, share knowledge, build awareness, change practice, change behaviour, and/or clinical outcomes in healthcare settings, inform policy, or to commercialise an innovation. Screening of studies, assessment of methodological quality and data extraction for the included studies were conducted by at least two reviewers.

Results: 39 relevant studies were included for full review; 8 were rated as moderate to strong methodologically with clinical outcomes that could be somewhat attributed to the toolkit. Three of the eight studies evaluated the toolkit as a single KT intervention, while five embedded the toolkit into a multistrategy intervention. Six of the eight toolkits were partially or mostly effective in changing clinical outcomes and six studies reported on implementation outcomes. The types of resources embedded within toolkits varied but included predominantly educational materials.

Conclusions: Future toolkits should be informed by high-quality evidence and theory, and should be evaluated using rigorous study designs to explain the factors underlying their effectiveness and successful implementation.

INTRODUCTION

Knowledge translation (KT) is a complex process occurring between researchers and knowledge users that includes the “synthesis, dissemination, exchange and ethically sound application of knowledge to improve health...provide more effective health services and products, and strengthen the health care system.”¹ The degree of engagement in the KT process may be influenced by factors such as the research results and needs of the knowledge user.¹ Clinical practice audits have demonstrated that health professionals do not consistently or effectively use current and high-quality research evidence as a basis for clinical care.² Despite strategies to facilitate the process of implementing research into practice, such as the development and evaluation of clinical practice guidelines, a major disconnect remains between evidence-based practice and actual clinical practice.³

Evidence-based KT strategies for linking research evidence and clinical practice include but are not limited to printed educational materials, educational meetings, educational outreach, the use of local opinion leaders, audit and feedback, and reminders.² These strategies have been used alone as single KT intervention or as multifaceted KT interventions, which consist of two or more strategies or variations of the same strategies (eg, educational materials) delivered in combination to change practice.¹⁻⁶ The benefit of multifaceted versus single KT interventions to change clinical and practice outcomes

Strengths and limitations of this study

- This systematic review on toolkits critically appraises research on strategies to facilitate practice change among health professionals.
- Results highlight the importance of evaluating implementation outcomes in addition to behavioural and clinical outcomes.
- This review was limited by a lack of an accepted definition for the term toolkit.
remains unclear, with some investigators reporting they are no more effective. 4,7,8

A variation on multifaceted KT interventions is the toolkit. Toolkits offer greater flexibility of use, and for the purposes of this review, are defined as a packaged grouping of multiple KT tools and strategies that codify explicit knowledge (eg, templates, pocket card guidelines, algorithms), and are used to educate and/or facilitate behaviour change. 9 Use of KT strategies housed within a toolkit are not necessarily prescribed in any combination or temporality (eg, Strategy A+/or Strategy B+/or Strategy C, etc). The goal is for the user to select KT strategies in the toolkit that are supported by evidence of effectiveness and for use at their own discretion, according to their aims, resources and context. Toolkits differ from multifaceted interventions in which the coupling of more than one KT strategy must be implemented together to comprise the ‘KT intervention’; for example, Strategy A +Strategy B=multifaceted KT strategy.

Evidence-based toolkits can be used to facilitate practice change, and can include strategies for guideline implementation, informing policy, practitioner training, and provide quality audit materials. 10,11 Currently, a wide range of toolkits address various clinical disease entities, such as diabetes and cancer care. For instance, the Registered Nurses Association of Ontario offers a toolkit on Best Practice Guidelines for patient care. 12 Despite the uncertainty surrounding the effectiveness of multifaceted KT interventions, organisations are investing resources in the development of KT toolkits because they provide a simple, more flexible and expedient method for promoting and utilising best healthcare practices. Whether these toolkits or their components are effectively implemented and positively associated with clinical outcomes remains unknown.

Toolkits comprise KT strategies that can be effective in supporting a range of KT aims if they are based on a clear rationale, quality evidence of their effectiveness, supported by a conceptual framework and built on a careful assessment of contextual barriers. 5 To be effective, toolkits should also provide high-quality evidence to guide their use or implementation. Currently, little is known about the effectiveness, feasibility and acceptability of toolkits. The aim of this systematic review was to identify and evaluate the effectiveness of toolkits for facilitating the implementation of evidence into clinical care and to inform future development, implementation and evaluation of toolkits.

METHODS

The methods for this review were based on the PRISMA checklist (http://www.prisma-statement.org/2.1.2%20-%20PRISMA%202009%20Checklist.pdf).

Search strategy

A systematic literature search of four electronic databases, MEDLINE (1946–November 2013), EMBASE (1947–November 2013), PsycINFO (1806–November 2013) and CINAHL (1981–November 2013), was conducted by a library information specialist. Search terms included database subject headings and text words for the following concepts: toolkits or toolboxes; evaluation, adherence or outcome assessment; and hospitals and hospitalised patients. The evaluation search terms used in MEDLINE, EMBASE and PsycINFO were based on published optimised search strategies 13–15 CINAHL evaluation terms were based on the optimised MEDLINE strategy. No date, age or language limits were applied (see online supplementary appendix).

Study selection

Study selection was conducted in two stages. First, all titles and abstracts were screened independently by two reviewers (Winnie Lam and Tissari Hewaranasinghage). To establish inter-rater reliability of study selection, each reviewer pilot tested 10 studies using the inclusion criteria. There was 95% agreement on the selected review articles. If necessary, a third reviewer (AS) who was not involved in the selection process resolved any disagreements. In the second stage, the full texts of all selected studies were screened to assess study eligibility and determine the final list of included studies.

Studies were included if: (1) they evaluated the effectiveness of a toolkit to support the integration of evidence into clinical care, either alone or embedded within a larger multistrategy intervention (toolkit +); (2) the KT goals(s) were to inform, share knowledge, build awareness, change practice, change behaviour (in the public), and/or clinical outcomes in healthcare settings, inform policy, or to commercialise an innovation; and (3) they included a comparison group. Studies published in languages other than English, thesis dissertations and studies published in non-peer-reviewed journals or in abstract form only were excluded. All study designs were included. Reference lists from included papers were screened for additional studies.

Methodological quality ratings

The methodological quality of included studies was assessed using the Effective Public Health Practice Project’s (EPHPP) Quality Assessment Tool for Quantitative Studies. 16 The EPHPP assesses methodological quality in systematic reviews of effectiveness. 17 Reliability and content and construct validity of the tool have been established. 18

The EPHPP tool can be used to evaluate multiple study designs that include comparison groups. Six categories, each consisting of a series of questions, are used to rate each study: (1) selection bias (two questions); (2) study design (four questions); (3) confounders (two questions); (4) blinding (two questions); (5) data collection methods (two questions) and (6) withdrawals and drop-outs (two questions). Each category is then assigned a rating (strong, moderate or weak), and based on these individual category ratings, a global rating is
assigned for the study (strong, moderate or weak). Additionally, the integrity of the study intervention and analyses is also examined; however, they do not contribute to the overall global rating.16

All studies were rated independently by two reviewers (AS and JY) using the EPHP tool. Prior to rating the studies, the tool was pilot tested on 10 studies. Overall per cent agreement was 88.5% (% = 0.84, 95% CI 0.72 to 0.96). When necessary, consensus meetings were held between reviewers to compare results and reach agreement on all studies. A third reviewer (KW) who was not involved in the quality assessment process resolved any disagreements.

**Data extraction and analysis**

Utilising a standardised data extraction chart, three reviewers (AS, KW and JY) independently extracted the following data from the studies that received a strong or moderate methodological global rating: study type, type of study participants, toolkit content, KT strategy and clinical outcome measures, including implementation outcomes as defined by Proctor et al88 (ie, acceptability, adoption, appropriateness, feasibility, fidelity, implementation cost, penetration and sustainability) and study results. Because many studies embedded the toolkit into a multistrategy intervention (ie, toolkit plus an additional KT strategy(ies)) and did not evaluate the toolkit alone, information regarding all of the components of the KT intervention was extracted. As well, the type of evidence, if any, underpinning the toolkits’ contents (KT strategies, tools) was extracted.

To determine toolkit effectiveness, Lugtenberg et al’s29 method was adopted to assign outcomes from each toolkit to one of three categories: (1) not effective (if no significant effects were demonstrated); (2) partially effective (if half or less of the outcome measures showed significant effects) or (3) mostly effective (if more than half the outcome measures showed significant effects). When study outcomes could not be at least partially attributed to the toolkit (eg, the toolkit was used in the multistrategy intervention and the control group), the study was excluded from detailed reporting.

If similar data from studies were available (eg, means, SDs, proportions), meta-analyses would be conducted. A weighted mean difference, or a standardised mean difference, relative risk, risk difference all with 95% CIs would be conducted using a fixed effects model. If pooling of results would not be possible, a narrative descriptive review of study results would be presented.

**RESULTS**

The search strategy yielded 39 unique studies for inclusion in this review (figure 1). Given the diversity of studies in terms of participants and outcomes, a meta-analysis was not possible; therefore, we chose to report on all studies with a strong or moderate global ratings rather than focusing only on randomised controlled trials (RCTs) of potentially weak quality.

The majority of the studies were RCTs (n=11)26 30 33 36 44 45 49 51 53 57 or one-group cohort studies (n=13).21 24 25 27 32 34 45 47 51 54 55 58 Eighteen of the included studies had toolkits embedded within a larger multistrategy KT intervention.22 23 26 28 31 32 38 42 45 49 53 55 57 58 and 21 studies evaluated toolkits as standalone KT interventions.11 21 24 25 29 30 33–37 39–41 44–46 48 50–52 56

Among all of the toolkits, 20 were developed for a specific disease context,21 22 26–28 31–34 40–45 47 51 54 55 57 most commonly for cancer (n=8),27 28 31 40 43 54 57 and diabetes (n=3).29 26 51 The remaining toolkits were developed for disease prevention (n=5),23 28 36 52 58 infection prevention (n=2),41 53 postoperative pain (n=1),48 smoking cessation (n=1),49 care in the geriatric population (n=8)24 25 29 30 35 36 39 56 patient safety (n=1)50 and general hospital quality improvement (n=1).57

Toolkits were targeted to health professionals (n=29),11 21 23–27 29–32 35 37–39 42 44 47 48 52 53 58–60 and caregivers (n=10)21 22 28 33 34 40 41 43 48 54. In one study, the intervention included separate toolkits for primary care physicians and patients.18

Only 2611 21 24–28 32 34 55 57 59–61 43 44 46–51 54 of the included studies specifically indicated the clinical evidence, rationale or theoretical basis underlying the toolkit strategies.

**Methodological quality of the studies**

The majority of studies (n=26)21–25 27–29 31 32 34 35 37–41 43–48 50 52 55 56 58 were rated as methodologically weak on the EPHP tool (ie, in terms of study design, selection bias, confounders, blinding, data collection methods and withdrawals and drop-outs); with 8 studies81 26 30 42 44 49 51 53 rated as moderate; and 53 34 43 54 57 as strong. The 13 moderate and strongly rated studies still had some general weaknesses. In 7 of the 13 studies21 26 30 33 42 44 57 blinding of outcome assessors and/or blinding of study participants to the research question were not explicitly stated. In the selection bias category, only 4 of the 13 studies26 36 44 57 reported the proportion of eligible participants who agreed to participate in the study. As well, in 6 of the 13 studies,11 26 30 36 42 44 raters agreed that the study participants were only somewhat likely, as opposed to very likely to represent the study population, introducing the potential for selection bias.

**Evaluation of the effectiveness of the toolkits**

In 5 of the 13 moderate to strongly rated studies,11 43 49 53 54 it was not possible to determine if clinical outcomes were attributable to the toolkit because all study participants received the toolkit in some variation. These five studies explored the effectiveness of the toolkit, either alone or paired with minimal additional interventions (multistrategy). A summary of the remaining eight studies is provided in table 1.26 30 33 36 42 44 51 57

Among the remaining eight studies, three\textsuperscript{33, 36, 51} evaluated the toolkit as a single KT intervention against a no KT intervention group, while five\textsuperscript{26, 30, 42, 44, 57} evaluated a multistrategy KT intervention against a no KT intervention group. Only four of five multistategy intervention studies\textsuperscript{26, 30, 42, 44} demonstrated partial to mostly effective results. Of the three single KT intervention studies, two\textsuperscript{33, 36} were mostly effective at changing clinical outcomes. Additionally, no studies evaluated the relative effectiveness of each KT strategy (eg, use of audit and feedback); therefore, it was not possible to determine which components contributed to the change in outcomes.

The majority of the studies\textsuperscript{26, 30, 33, 36, 44, 51} aimed to evaluate the toolkit’s effectiveness for a variety of KT goals. One study focused on changing patient clinical outcomes (eg, myocardial infarction, number of falls); two studies also evaluated change in patient behaviour;\textsuperscript{26, 33} and one evaluated behavioural change in family caregivers.\textsuperscript{36} Two studies\textsuperscript{44, 51} focused on toolkit effectiveness for changing clinician behaviour in addition to improving patient clinical outcomes, and two studies\textsuperscript{42, 57} were solely focused on improving clinician behaviour.

Implementation outcomes were mentioned in six studies.\textsuperscript{26, 30, 33, 36, 42, 44} Dykes \textit{et al}\textsuperscript{30} included a process for assessing fidelity of the KT intervention; Goeppinger \textit{et al}\textsuperscript{44} examined the adoption, appropriateness and sustainability of the toolkit; Horvath \textit{et al}\textsuperscript{26} provided information about the fidelity of the KT intervention and cost of the toolkit but did not conduct a cost/benefit analysis; and Cavanaugh \textit{et al}, Majumdar \textit{et al} and Menchetti \textit{et al}\textsuperscript{44} examined the sustainability of improved clinical outcomes over time.

Toolkit content varied across studies. Two studies included self-management toolkits for patients and caregivers with a focus on arthritis\textsuperscript{35} and Alzheimer’s.\textsuperscript{36} Six studies evaluated toolkits for health professionals on fall prevention,\textsuperscript{30} gastro-oesophageal reflux,\textsuperscript{12} depression,\textsuperscript{44} diabetes\textsuperscript{26, 51} and cancer.\textsuperscript{57} Toolkit resources included information/handout sheets, posters, pocket guides and educational modules. Wright \textit{et al}\textsuperscript{57} included reminder packages for participants comprised of a cover letter from an expert opinion leader, a peer-reviewed article
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<th>Study design, participants</th>
<th>Toolkit and intervention</th>
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<tr>
<td>Cavanaugh et al&lt;sup&gt;26&lt;/sup&gt; Two RCTs at 2 academic medical centres N=198 adult patients with diabetes n=99 (control) n=99 (intervention) Country: USA</td>
<td>Intervention components: enhanced diabetes care programme; training sessions; DLNET Control components: enhanced diabetes care programme Toolkit target: health professionals Toolkit contents: customisable 24 instructive modules about diabetes self-management activities, including blood glucose monitoring, nutrition management, foot care, administration of medications</td>
<td>Incorporated communication principles</td>
<td>1. Glycaemic control (A1c) 2. Patient-reported self-efficacy of diabetes self-management 3. Self-management behaviours 4. Treatment satisfaction 5. Sustainability</td>
<td>Significant improvements in A1c levels in intervention and control groups at 3 months (adjusted analyses showed greater improvement in the intervention group (p=0.03)) Significant improvement in self-efficacy from baseline in both groups (p&lt;0.01, 0.02) (but NS differences between groups in adjusted analyses) NS differences between intervention and control for self-management behaviour or treatment satisfaction NS differences between intervention and control groups at 6 months Implementation outcomes: sustainability of outcomes measured at 3 and 6 months Toolkit effectiveness: partially effective</td>
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<td>Dykes et al&lt;sup&gt;30&lt;/sup&gt; Cluster RCT N=8 units in 4 urban US hospitals, N=10,264 patients n=5160 (intervention) n=5104 (control) Country: USA</td>
<td>Intervention components: FPTK; local champions Control components: usual care Toolkit target: health professionals Toolkit contents: Morse Falls Scale to assess fall risk; interventions tailored to patient-specific areas of risk; bed poster, patient/family education handout, fall prevention plan (tailored for each patient) Literature review, focus groups with nurses+nursing assistants; assessment of barriers and facilitators to optimal practice</td>
<td>1. Patient falls per 1000 patient-days 2. Fall-related injuries 3. Fidelity</td>
<td>Significantly fewer patients with falls in intervention versus control units (p&lt;0.02) Significantly lower adjusted fall rates in intervention versus control units per 1000 patient-days (p=0.04) NS difference in fall-related injuries Implementation outcomes: protocol adherence &gt;89% Toolkit effectiveness: mostly effective</td>
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<td>Majumdar et al&lt;sup&gt;2&lt;/sup&gt; Controlled clinical trial N=14 managed care practices Country: USA</td>
<td>Low-intensity intervention components: evidence-based guideline on Helicobacter pylori; Toolkit High-intensity intervention components: evidence-based guideline on H. pylori; Toolkit; academic detailing of guideline dissemination by a PCP champion using persuasive educational session; 1 month reinforcement of guideline message; and reminder about eligible patients by a pharmacist Toolkit target: health professionals Toolkit contents: customised list of eligible patients from participating practice; educational materials for patients; patient letters used to arrange for test or follow-up appointment; pre-printed materials including: (A) H. pylori serology test requisitions, (B) preapproved prescriptions, (C) progress notes for patient charts Not specified</td>
<td>1. Rate of testing 2. Rate of continued use of acid suppressing medications 3. Sustainability</td>
<td>Significant increase in H. pylori test-ordering in high-intensity intervention versus usual care (p=0.02) Significant decrease in proton pump inhibitor use by 9% per year in high-intensity intervention versus usual care (p=0.028) Implementation outcomes: sustainability of outcomes measured at 12 months Toolkit effectiveness: mostly effective</td>
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MNS differences between groups in remission of depression at 3, 6, 12 months; however in patients with minor/major depression, intervention was more effective than usual care at 3 months (p=0.015). 1. Clinical remission of Intervention group showed significantly higher treatment response rates at 3(p=0.016) and 6 months (p=0.049). PCP increased use of appropriate antidepressants and decreased use of sedatives, hypnotics at 3 months. Implementation outcomes: sustainability of outcomes measured at 3, 6, 12 months Toolkit effectiveness: partially effective NS difference between groups in death or non-fatal MI (p=0.07) and use of a statin (p=0.26). Decreased use of ECG (p=0.02) and cardiac stress tests (p=0.04) in intervention group Implementation outcomes: not specified Toolkit effectiveness: not effective

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<th>Table 1 Continued</th>
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<tr>
<td>Menchetti et al</td>
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<td>Horvath et al 2015 RCT</td>
<td>N=108 dyads of patients with progressive dementia of Alzheimer’s type/caregiver n=48 (control) n=60 (intervention) Country: USA</td>
<td>Self-management program (eg, decision-making); Arthritis Help Book; audio relaxation and exercise CDs; audio CD of all material from information sheets</td>
<td>Principles of health literacy, patient-centred care and self-efficacy</td>
<td>1. Caregiver self-efficacy 2. Caregiver strain 3. Home safety 4. Risky behaviours and accidents 5. Fidelity 6. Costs</td>
<td>97% of participants reported use of the toolkit and found it useful. The Book was rated the most useful part</td>
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**Evidence informing toolkit development**
- Principles of health literacy, patient-centred care and self-efficacy

**Outcomes measured**
- 1. Caregiver self-efficacy
- 2. Caregiver strain
- 3. Home safety
- 4. Risky behaviours and accidents
- 5. Fidelity
- 6. Costs

**Results**
- 97% of participants reported use of the toolkit and found it useful. The Book was rated the most useful part

**Quality**
- S: Strong

**DLNET, Diabetes Literacy Numeracy Education Toolkit; FPTK, Fall Prevention Toolkit; LN, lymph node; M, moderate; NS, non significant; PCP, primary care physician; PHQ-9, Patient Health Questionnaire-9; RCT, randomised controlled trial; S, strong.**
practice; rationale for their inclusion in the toolkit, given the toolkit aims; and guidance on the implementation process—how they are to be used. Although the eight studies in this review mentioned some form of evidence underlying each component, descriptions were vague and non-descriptive, and few mentioned high-quality evidence, such as systematic reviews. Often, evidence was provided for only one component of the toolkit. Cavanaugh et al used communication theory to design the ‘Diabetes Literacy and Numeracy Education Toolkit’, and did not specify any underlying evidence for their content. Shah et al, however, provided evidence for using educational materials as a resource within their educational toolkit, which focused on cardiovascular disease screening and risk reduction in patients with diabetes. Nevertheless, their content was not based on a barriers assessment, quality improvement or educational theory.

Multiple barriers have been identified to account for the knowledge to practice gap, and many are intrinsic to health professionals and their practice environment or context. For example, organisational constraints, such as lack of time or an inability to access resources, are common barriers to KT. LaRocca et al suggested that the more successful KT intervention strategies were those that were accessible and could be tailored to the needs and preferences of the users. Components of the fall prevention toolkit by Dykes et al included patient/family education handouts that were tailored by the nurse based on the knowledge of the patient, thereby capitalising on high tension for change; adaptability, strength and quality of the intervention; and low complexity. The effects of tailoring strategies to address identified barriers to change require more clarity, but may improve care and patient outcomes, particularly when KT approaches can capitalise on what we know works in implementation. Only one of the eight reviewed studies assessed barriers and facilitators to inform the toolkit’s components. Furthermore, determining the influence of modifiable components of context (eg, leadership support, culture, evaluation) would further allow for customisation of KT strategies to facilitate practice change and clinical outcomes. Further research is needed on how the toolkit was developed, and the influence of the practice context as these factors may influence study outcomes.

Consideration should also be given to factors implicated in successful implementation. Proctor’s taxonomy for implementation outcomes was extracted from studies where possible, as these outcomes could be used to indicate successful implementation of the toolkit within the healthcare system. Developing toolkits supported by implementation guidance would go a long way in demonstrating how toolkits contribute to good clinical and implementation outcomes. Descriptions of most toolkits lacked details about the implementation process and outcomes. Evidence Based Practice for Improving Quality (EPIQ) is an example of a KT intervention that combines evidence, continuous quality improvement, an implementation process and assessment of implementation outcomes. In phase 1 (Preparation), the hospital unit identifies an implementation team, who are trained to review existing unit pain practices, guidelines and research evidence to inform targeted practice changes. In phase 2 (Implementation), the team identifies specific pain practice aims and KT strategies (eg, educational outreach, reminders and audit and feedback) to implement using quality improvement cycles. EPIQ was effective in improving pain process outcomes (ie, pain assessment and management) and reducing the odds of having severe pain by 51%.

Only two studies reported on fidelity of toolkit implementation. To be clinically effective, healthcare interventions need to be effectively implemented. Yet, implementation outcomes are often overlooked in research and KT practice, creating high potential for type III errors; lack of clarity about whether the intervention or its implementation have been unsuccessful. This type of error can reduce the power to detect significant effects of an intervention. Assessing the fidelity of implementing complex interventions addresses type III error and provides evidence of variability in implementation of interventions, which could also contribute to limited effectiveness.

All eight studies in this review used RCT designs to evaluate toolkit effectiveness. There is a common methodological challenge to RCT studies of KT effectiveness, in that this design could block important contextual factors that now have burgeoning evidence of their importance in successful implementation. Caution is required in interpreting which KT strategies are evidence-based, and new studies need to utilise more appropriate mixed methodologies or other types of randomised designs, such as wait listed or stepped-wedged designs, to address what works in implementation of practice changes.

Several limitations to this systematic review warrant discussion. The term ‘toolkit’ was used in the studies included in this systematic review. However, there is currently no accepted definition for toolkits in existing taxonomies related to quality improvement and behavioural change strategies (eg, Cochrane Effective Practice and Organisation of Care Group). Although we chose a term that had some consistency in the literature, based on the evidence reported in this review, there is no consensus on key content, implementation strategies to promote behavioural change or theoretical approaches that should be included in implementation toolkits. These findings could explain the heterogeneity of the toolkits included in this review. Therefore, capturing all relevant literature was challenging because of the lack of standard terminology used for toolkits. As a result, relevant studies might have been missed by the search. The majority of studies had significant methodological shortcomings and were rated as weak, mostly due to the study
designs. One of the limitations was that we focused on studies that evaluated the effectiveness of a toolkit to support the facilitation of evidence into clinical care; therefore, the studies included in this review reported quantitative results. The literature search was limited to toolkits used in hospital and other clinical settings. Broadening the search to community or public health settings may have yielded additional studies for the review.9

In summary, toolkits have potential as a promising KT strategy for facilitating practice change in healthcare. To fully understand their effectiveness, a systematic approach to planning and reporting their development, the evidence underlying each component, and any direction regarding appropriate implementation is required. Toolkits should have (1) a clearly described purpose, rationale for each component; (2) components that are rigorously developed and informed by high-quality evidence, such as systematic reviews; (3) delivery methods that are guided by a comprehensive implementation process (eg, self-directed, facilitation, reminders) with consideration for fidelity of implementation where appropriate; and (4) a rigorous evaluation plan and study design that can help explain the factors underlying their effectiveness and successful implementation (ie, combining outcome and process measures including context).9

Only a few of the toolkits in this review met all of these criteria.33 34 Ideally future studies of toolkit effectiveness should also be informed by a theoretical approach. In conclusion, this study provides some evidence for the utility of the toolkit.

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Contributors JY led the writing of the manuscript, organised all aspects of the systematic review, participated in the screening of abstracts, rating of methodological quality, data extraction and analysis. She also drafted the initial manuscript, made revisions and approved the final manuscript as submitted. As participated in the rating of methodological quality, data extraction, and analysis of articles included in the review. She also participated in drafting the initial manuscript and revisions. MB provided guidance and expertise in the overall conceptualisation of the review, revised and critically reviewed the manuscript, and approved the final manuscript as submitted. KW participated in reviewing the methodological quality, data extraction and analysis of all articles included in the report. She also participated in drafting the initial manuscript and revisions. BS provided guidance and expertise in the overall conceptualisation of the review, critically reviewed the manuscript and approved the final manuscript as submitted.

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