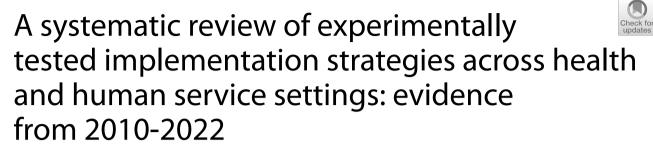
SYSTEMATIC REVIEW

Implementation Science

Open Access



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Abstract

Background Studies of implementation strategies range in rigor, design, and evaluated outcomes, presenting interpretation challenges for practitioners and researchers. This systematic review aimed to describe the body of research evidence testing implementation strategies across diverse settings and domains, using the Expert Recommendations for Implementing Change (ERIC) taxonomy to classify strategies and the Reach Effectiveness Adoption Implementation and Maintenance (RE-AIM) framework to classify outcomes.

Methods We conducted a systematic review of studies examining implementation strategies from 2010-2022 and registered with PROSPERO (CRD42021235592). We searched databases using terms "implementation strategy", "intervention", "bundle", "support", and their variants. We also solicited study recommendations from implementation science experts and mined existing systematic reviews. We included studies that quantitatively assessed the impact of at least one implementation strategy to improve health or health care using an outcome that could be mapped to the five evaluation dimensions of RE-AIM. Only studies meeting prespecified methodologic standards were included. We described the characteristics of studies and frequency of implementation strategy use across study arms. We also examined common strategy pairings and cooccurrence with significant outcomes.

Findings Our search resulted in 16,605 studies; 129 met inclusion criteria. Studies tested an average of 6.73 strategies (0-20 range). The most assessed outcomes were Effectiveness (n=82; 64%) and Implementation (n=73; 56%). The implementation strategies most frequently occurring in the experimental arm were Distribute Educational Materials (n=99), Conduct Educational Meetings (n=96), Audit and Provide Feedback (n=76), and External Facilitation (n=59). These strategies were often used in combination. Nineteen implementation strategies were frequently tested and associated with significantly improved outcomes. However, many strategies were not tested sufficiently to draw conclusions.

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Conclusion This review of 129 methodologically rigorous studies built upon prior implementation science data syntheses to identify implementation strategies that had been experimentally tested and summarized their impact on outcomes across diverse outcomes and clinical settings. We present recommendations for improving future similar efforts.

Keywords Implementation strategy, Systematic review, Health, Health-related outcomes

Contributions to the literature

- While many implementation strategies exist, it has been challenging to compare their effectiveness across a wide range of trial designs and practice settings
- This systematic review provides a transdisciplinary evaluation of implementation strategies across population, practice setting, and evidence-based interventions using a standardized taxonomy of strategies and outcomes.
- Educational strategies were employed ubiquitously; nineteen other commonly used implementation strategies, including External Facilitation and Audit and Provide Feedback, were associated with positive outcomes in these experimental trials.
- This review offers guidance for scholars and practitioners alike in selecting implementation strategies and suggests a roadmap for future evidence generation.

Background

Implementation strategies are "methods or techniques used to enhance the adoption, implementation, and sustainment of evidence-based practices or programs" (EBPs) [1]. In 2015, the Expert Recommendations for Implementing Change (ERIC) study organized a panel of implementation scientists to compile a standardized set of implementation strategy terms and definitions [2-4]. These 73 strategies were then organized into nine "clusters" [5]. The ERIC taxonomy has been widely adopted and further refined [6-13]. However, much of the evidence for individual or groups of ERIC strategies remains narrowly focused. Prior systematic reviews and metaanalyses have assessed strategy effectiveness, but have generally focused on a specific strategy, (e.g., Audit and Provide Feedback) [14–16], subpopulation, disease (e.g., individuals living with dementia) [16], outcome [15], service setting (e.g., primary care clinics) [17–19] or geography [20]. Given that these strategies are intended to have broad applicability, there remains a need to understand how well implementation strategies work across EBPs and settings and the extent to which implementation knowledge is generalizable.

There are challenges in assessing the evidence of implementation strategies across many EBPs, populations, and settings. Heterogeneity in population characteristics, study designs, methods, and outcomes have made it difficult to quantitatively compare which strategies work and under which conditions [21]. Moreover, there remains significant variability in how researchers operationalize, apply, and report strategies (individually or in combination) and outcomes [21, 22]. Still, synthesizing data related to using individual strategies would help researchers replicate findings and better understand possible mediating factors including the cost, timing, and delivery by specific types of health providers or key partners [23–25]. Such an evidence base would also aid practitioners with implementation planning such as when and how to deploy a strategy for optimal impact.

Building upon previous efforts, we therefore conducted a systematic review to evaluate the level of evidence supporting the ERIC implementation strategies across a broad array of health and human service settings and outcomes, as organized by the evaluation framework, RE-AIM (Reach, Effectiveness, Adoption, Implementation, Maintenance) [26–28]. A secondary aim of this work was to identify patterns in scientific reporting of strategy use that could not only inform reporting standards for strategies but also the methods employed in future. The current study was guided by the following research questions¹:

- 1. What implementation strategies have been most commonly and rigorously tested in health and human service settings?
- 2. Which implementation strategies were commonly paired?
- 3. What is the evidence supporting commonly tested implementation strategies?

Methods

We used the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA-P) model [29–31] to develop and report on the methods for this systematic

¹ We modestly revised the following research questions from our PROS-PERO registration after reading the articles and better understanding the nature of the literature: 1) What is the available evidence regarding the effectiveness of implementation strategies in supporting the uptake and sustainment of evidence intended to improve health and healthcare outcomes? 2) What are the current gaps in the literature (i.e., implementation strategies that do not have sufficient evidence of effectiveness) that require further exploration?

review (Additional File 1). This study was considered to be non-human subjects research by the RAND institutional review board.

Registration

The protocol was registered with PROSPERO (PROS-PERO 2021 CRD42021235592).

Eligibility criteria

This review sought to synthesize evidence for implementation strategies from research studies conducted across a wide range of health-related settings and populations. Inclusion criteria required studies to: 1) available in English; 2) published between January 1, 2010 and September 20, 2022; 3) based on experimental research (excluded protocols, commentaries, conference abstracts, or proposed frameworks); 4) set in a health or human service context (described below); 5) tested at least one quantitative outcome that could be mapped to the RE-AIM evaluation framework [26-28]; and 6) evaluated the impact of an implementation strategy that could be classified using the ERIC taxonomy [2, 32]. We defined health and human service setting broadly, including inpatient and outpatient healthcare settings, specialty clinics, mental health treatment centers, long-term care facilities, group homes, correctional facilities, child welfare or youth services, aging services, and schools, and required that the focus be on a health outcome. We excluded hybrid type I trials that primarily focused on establishing EBP effectiveness, qualitative studies, studies that described implementation barriers and facilitators without assessing implementation strategy impact on an outcome, and studies not meeting standardized rigor criteria defined below.

Information sources

Our three-pronged search strategy included searching academic databases (i.e., CINAHL, PubMed, and Web of Science for replicability and transparency), seeking recommendations from expert implementation scientists, and assessing existing, relevant systematic reviews and meta-analyses.

Search strategy

Search terms included "implementation strateg*" OR "implementation intervention*" OR "implementation bundl*" OR "implementation support*." The search, conducted on September 20, 2022, was limited to English language and publication between 2010 and 2022, similar to other recent implementation science reviews [22]. This timeframe was selected to coincide with the advent of

Title and abstract screening process

Each study's title and abstract were read by two reviewers, who dichotomously scored studies on each of the six eligibility criteria described above as yes=1 or no=0, resulting in a score ranging from 1 to 6. Abstracts receiving a six from both reviewers were included in the full text review. Those with only one score of six were adjudicated by a senior member of the team (MJC, SSR, DEG). The study team held weekly meetings to trouble-shoot and resolve any ongoing issues noted through the abstract screening process.

Full text screening

During the full text screening process, we reviewed, in pairs, each article that had progressed through abstract screening. Conflicts between reviewers were adjudicated by a senior member of the team for a final inclusion decision (MJC, SSR, DEG).

Review of study rigor

After reviewing published rigor screening tools [34–36], we developed an assessment of study rigor that was appropriate for the broad range of reviewed implementation studies. Reviewers evaluated studies on the following: 1) presence of a concurrent comparison or control group (=2 for traditional randomized controlled trial or stepped wedge cluster randomized trial and =1 for pseudo-randomized and other studies with concurrent control); 2) EBP standardization by protocol or manual (=1 if present); 3) EBP fidelity tracking (=1 if present); 4) implementation strategy standardization by operational description, standard training, or manual (=1 if present); 5) length of follow-up from full implementation of intervention (=2 for twelve months or longer, =1 for six to eleven months, or =0 for less than six months); and 6) number of sites (=1 for more than one site). Rigor scores ranged from 0 to 8, with 8 indicating the most rigorous. Articles were included if they 1) included a concurrent control group, 2) had an experimental design, and 3) received a score of 7 or 8 from two independent reviewers.

Outside expert consultation

We contacted 37 global implementation science experts who were recognized by our study team as leaders in the field or who were commonly represented among first or senior authors in the included abstracts. We asked each expert for recommendations of publications meeting study inclusion criteria (i.e., quantitatively evaluating the effectiveness of an implementation strategy). Recommendations were recorded and compared to the full abstract list.

Systematic reviews

Eighty-four systematic reviews were identified through the initial search strategy (See Additional File 3). Systematic reviews that examined the effectiveness of implementation strategies were reviewed in pairs for studies that were not found through our initial literature search.

Data abstraction and coding

Data from the full text review were abstracted in pairs, with conflicts resolved by senior team members (DEG, MJC) using a standard Qualtrics abstraction form. The form captured the setting, number of sites and participants studied, evidence-based practice/program of focus, outcomes assessed (based on RE-AIM), strategies used in each study arm, whether the study took place in the U.S. or outside of the U.S., and the findings (i.e., was there significant improvement in the outcome(s)?). We coded implementation strategies used in the Control and Experimental Arms. We defined the Control Arm as receiving the lowest number of strategies (which could mean zero strategies or care as usual) and the Experimental Arm as the most intensive arm (i.e., receiving the highest number of strategies). When studies included multiple Experimental Arms, the Experimental Arm with the least intensive implementation strategy(ies) was classified as "Control" and the Experimental Arm with the most intensive implementation strategy(ies) was classified as the "Experimental" Arm.

Implementation strategies were classified using standard definitions (MJC, SSR, DEG), based on minor modifications to the ERIC taxonomy [2-4]. Modifications resulted in 70 named strategies and were made to decrease redundancy and improve clarity. These modifications were based on input from experts, cognitive interview data, and team consensus [37] (See Additional File 4). Outcomes were then coded into RE-AIM outcome domains following best practices as recommended by framework experts [26-28]. We coded the RE-AIM domain of Effectiveness as either an assessment of the effectiveness of the EBP or the implementation strategy. We did not assess implementation strategy fidelity or effects on health disparities as these are recently adopted reporting standards [27, 28] and not yet widely implemented in current publications. Further, we did not include implementation costs as an outcome because reporting guidelines have not been standardized [38, 39].

Assessment and minimization of bias

Assessment and minimization of bias is an important component of high-quality systematic reviews. The Cochrane Collaboration guidance for conducting highquality systematic reviews recommends including a specific assessment of bias for individual studies by assessing the domains of randomization, deviations of intended intervention, missing data, measurement of the outcome, and selection of the reported results (e.g., following a prespecified analysis plan) [40, 41]. One way we addressed bias was by consolidating multiple publications from the same study into a single finding (i.e., N=1), so-as to avoid inflating estimates due to multiple publications on different aspects of a single trial. We also included high-quality studies only, as described above. However, it was not feasible to consistently apply an assessment of bias tool due to implementation science's broad scope and the heterogeneity of study design, context, outcomes, and variable measurement, etc. For example, most implementation studies reviewed had many outcomes across the RE-AIM framework, with no one outcome designated as primary, precluding assignment of a single score across studies.

Analysis

We used descriptive statistics to present the distribution of health or healthcare area, settings, outcomes, and the median number of included patients and sites per study, overall and by country (classified as U.S. vs. non-U.S.). Implementation strategies were described individually, using descriptive statistics to summarize the frequency of strategy use "overall" (in any study arm), and the mean number of strategies reported in the Control and Experimental Arms. We additionally described the strategies that were *only* in the experimental (and not control) arm, defining these as strategies that were "tested" and may have accounted for differences in outcomes between arms.

We described frequencies of pair-wise combinations of implementation strategies in the Experimental Arm. To assess the strength of the evidence supporting implementation strategies that were used in the Experimental Arm, study outcomes were categorized by RE-AIM and coded based on whether the association between use of the strategies resulted in a significantly positive effect (yes=1; no=0). We then created an indicator variable if at least one RE-AIM outcome in the study was significantly positive (yes=1; no=0). We plotted strategies on a graph with quadrants based on the combination of median number of studies in which a strategy appears and the median percent of studies in which a strategy was associated with at least one positive RE-AIM outcome. The upper right quadrant—higher number of studies overall and higher percent of studies with a significant RE-AIM outcome represents a superior level of evidence. For implementation strategies in the upper right quadrant, we describe each RE-AIM outcome and the proportion of studies which have a significant outcome.

Results

Search results

We identified 14,646 articles through the initial literature search, 17 articles through expert recommendation (three of which were not included in the initial search), and 1,942 articles through reviewing prior systematic reviews (Fig. 1). After removing duplicates, 9,399 articles were included in the initial abstract screening. Of those, 48% (n=4,075) abstracts were reviewed in pairs for inclusion. Articles with a score of five or six were reviewed a second time (n=2,859). One quarter of abstracts that scored lower than five were reviewed for a second time at random. We screened the full text of 1,426 articles in pairs. Common reasons for exclusion were 1) study rigor, including no clear delineation between the EBP and implementation strategy, 2) not testing an implementation strategy, and 3) article type that did not meet inclusion criteria (e.g., commentary, protocol, etc.). Six hundred seventeen articles were reviewed for study rigor with 385 excluded for reasons related to study design and rigor, and 86 removed for other reasons (e.g., not a research article). Among the three additional expertrecommended articles, one met inclusion criteria and was added to the analysis. The final number of studies abstracted was 129 representing 143 publications.

Descriptive results

Of 129 included studies (Table 1; see also Additional File 5 for Summary of Included Studies), 103 (79%) were conducted in a healthcare setting. EBP health care setting varied and included primary care (n=46; 36%), specialty care (n=27; 21%), mental health (n=11; 9%), and public health (n=30; 23%), with 64 studies (50%) occurring in an outpatient health care setting. Studies included a median of 29 sites and 1,419 target population (e.g., patients or students). The number of strategies varied widely across studies, with Control Arms averaging approximately two strategies (Range = 0-20, including studies with no strategy in the comparison group) and Experimental Arms averaging eight strategies (Range = 1-21). Non-US studies (n=73) included more sites and target population on average, with an overall median of 32 sites and 1,531 patients assessed in each study.

Organized by RE-AIM, the most evaluated outcomes were Effectiveness (n = 82, 64%) and Implementation (n = 73, 56%); followed by Maintenance (n=40; 31%), Adoption (n=33; 26%), and Reach (n=31; 24%). Most

studies (n = 98, 76%) reported at least one significantly positive outcome. Adoption and Implementation outcomes showed positive change in three-quarters of studies (n=78), while Reach (n=18; 58%), Effectiveness (n=44; 54%), and Maintenance (n=23; 58%) outcomes evidenced positive change in approximately half of studies.

The following describes the results for each research question.

1. What implementation strategies have been most commonly and rigorously tested in health and human service settings?

Table 2 shows the frequency of studies within which an implementation strategy was used in the Control Arm, Experimental Arm(s), and tested strategies (those used exclusively in the Experimental Arm) grouped by strategy type, as specified by previous ERIC reports [2, 6].

Control arm

In about half the studies (53%; n=69), the Control Arms were "active controls" that included at least one strategy, with an average of 1.64 (and up to 20) strategies reported in control arms. The two most common strategies used in Control Arms were: Distribute Educational Materials (n=52) and Conduct Educational Meetings (n=30).

Experimental arm

Experimental conditions included an average of 8.33 implementation strategies per study (Range = 1-21). Figure 2 shows a heat map of the strategies that were used in the Experimental Arms in each study. The most common strategies in the Experimental Arm were Distribute Educational Materials (n=99), Conduct Educational Meetings (n=96), Audit and Provide Feedback (n=76), and External Facilitation (n=59).

Tested strategies

The average number of implementation strategies that were included in the Experimental Arm only (and not in the Control Arm) was 6.73 (Range = 0-20).² Overall, the top 10% of tested strategies included Conduct Educational Meetings (n=68), Audit and Provide Feedback (n=63), External Facilitation (n=54), Distribute Educational Materials (n=49), Tailor Strategies (n=41), Assess for Readiness and Identify Barriers and Facilitators (n=38) and Organize Clinician Implementation Team Meetings (n=37). Few studies tested a single strategy

² Tested strategies are those which exist in the Experimental Arm but not in the Control Arm. Comparative effectiveness or time staggered trials may not have any unique strategies in the Experimental Arm and therefore in our analysis would have no Tested Strategies.

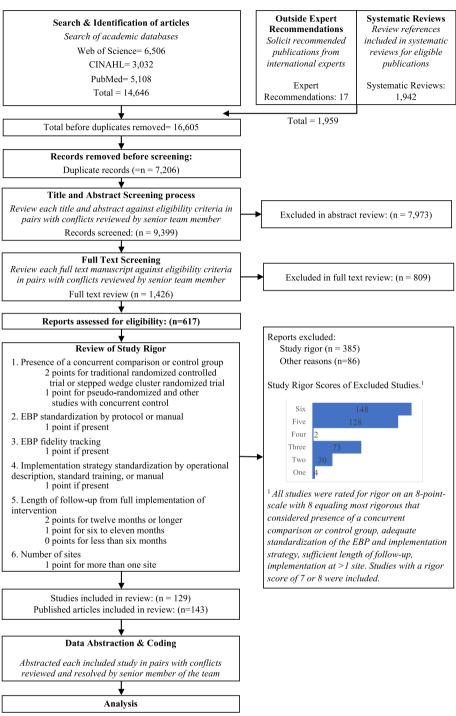


Fig. 1 Expanded PRISMA Flow Diagram

The expanded PRISMA flow diagram provides a description of each step in the review and abstraction process for the systematic review

(n=9). These strategies included, Audit and Provide Feedback, Conduct Educational Meetings, Conduct Ongoing Training, Create a Learning Collaborative, External Facilitation (n=2), Facilitate Relay of Clinical Data To

Providers, Prepare Patients/Consumers to be Active Participants, and Use Other Payment Schemes. Three implementation strategies were included in the Control or Experimental Arms but were not Tested including, Use

Table 1 Study characteristics

| | Total (n=129) | United States (<i>n</i> =56) | Non-U.S. (<i>n</i> =73) |
|---|-------------------|-------------------------------|--------------------------|
| Country (n, %) | | | |
| United States (U.S.) | 56 (43%) | | |
| Other Country ^a | 28 (22%) | | |
| Australia | 18 (14%) | | |
| The Netherlands | 14 (11%) | | |
| United Kingdom | 7 (5%) | | |
| Canada ^b | 6 (5%) | | |
| Area of health or healthcare (n,%) ^c | | | |
| Primary care | 46 (36%) | 21 (36%) | 25 (34%) |
| Public health | 30 (23%) | 12 (21%) | 18 (25%) |
| Specialty care | 27 (21%) | 10 (18%) | 17 (23%) |
| Mental health | 11 (9%) | 8 (14%) | 3 (4%) |
| Other | 18 (14%) | 5 (9%) | 13 (18%) |
| Setting (n, %) ^c | | | |
| Outpatient/clinic | 64 (50%) | 32 (57%) | 32(44%) |
| Inpatient/hospital—floors | 20 (15%) | 4 (7%) | 16 (22%) |
| School | 13 (10%) | 1 (2%) | 12 (16%) |
| Community center | 7 (5%) | 5 (9%) | 2 (3%) |
| Intensive Care Unit (ICU) | 5 (5%) | 1 (2%) | 4 (5%) |
| Emergency/urgent care | 3 (2%) | 1 (2%) | 2 (3%) |
| Home-based | 1 (1%) | 0 (0%) | 1 (1%) |
| Other | 29 (23%) | 15 (58%) | 14 (19%) |
| Number of sites, Median (IQR) | 29 (12-49) | 24 (12-47) | 32 (12-55) |
| Number of participants, Median (IQR) | 1,419 (306-5,957) | 1,152 (239-4,040) | 1,531 (365-7,279) |
| Implementation Strategies | | | |
| Control Arm [mean (SD) range] | 1.64 (2.86) 0-20 | 1.39 (2.30) 0-9 | 1.82 (3.23) 0-20 |
| Experimental Arm [mean (SD) range] | 8.33 (4.71) 1-21 | 8.66 (4.49) 1-20 | 8.07 (4.88) 1-21 |
| Tested [mean (SD) range] | 6.73 (4.45) 0-20 | 7.30 (4.46) 1-20 | 6.29 (4.43) 0-20 |
| Outcome Assessed ^d (n, %) | | | |
| Reach | 31 (24%) | 12 (21%) | 19 (26%) |
| Effectiveness | 82 (64%) | 31 (55%) | 51 (70%) |
| Adoption | 33 (26%) | 13 (23%) | 20 (27%) |
| Implementation | 73 (56%) | 29 (52%) | 44 (60%) |
| Maintenance | 40 (31%) | 18 (32%) | 22 (30%) |
| Positive outcome ^{e,f} (n, %) | | | |
| Reach | 18 (58%) | 6 (50%) | 12 (63%) |
| Effectiveness | 44 (54%) | 14 (45%) | 30 (59%) |
| Adoption | 24 (73%) | 10 (77%) | 14 (70%) |
| Implementation | 54 (74%) | 19 (66%) | 35 (80%) |
| Maintenance | 23 (58%) | 9 (50%) | 14 (64%) |

^a Other Countries included: Sweden (*n*=4), Germany (*n*=4), France (*n*=3), Spain (*n*=2), Zimbabwe, Uganda, Switzerland, Portugal, Norway, Nigeria, Kenya, Japan, Finland, Denmark, The Bahamas, New Zealand and three multi-country studies: Senegal and Mali; Australia and New Zealand; and England, Sweden, the Netherlands, and Republic of Ireland.

^b One study was conducted in both the U.S. and Canada with 7 sites in Canada and 1 U.S. site. This study was coded as being conducted in Canada

^c Studies could occur in more than one area of health/healthcare and setting. The denominator for reported percentages is the total number studies reviewed by column (Total *N* = 129; U.S. *N* = 56; Non-U.S. *N* = 73)).

^d The denominator for reported percentages is the total number studies reviewed by column (Total N = 129; U.S. N = 56; Non-U.S. N = 73)); studies could assess more than one outcome

^e The denominator for reported percentages is the total number studies reviewed by column (Total N = 129; U.S. N = 56; Non-U.S. N = 73));

^f Studies may have tested several measures of a RE-AIM outcome. The designation of "positive outcome" indicates studies with at least one measure of the RE-AIM outcome that reached statistical significance when comparing the Experimental to Control Arm. For example, if a study used several measures of Reach, the outcomes would be considered a positive outcome if at least one of those measures was statistically significant.

Table 2 Frequency of ERIC implementation strategy use

| Implementation Strategy | Overall | Control Arm (Least intensive arm) | Experimental Arm (Most intensive arm) | Tested |
|---|---------|---|--|----------|
| Use evaluative and iterative strategies | | | | |
| Assess for readiness and identify barriers and facilitators | 43 | 5 | 43 | 38 |
| Audit and provide feedback | 76 | 13 | 76 | 63 |
| Conduct cyclical small tests of change | 22 | 1 | 22 | 21 |
| Conduct local need assessment | 20 | 4 | 20 | 16 |
| Develop a formal implementation blueprint | 41 | 5 | 42 | 36 |
| Develop and implement tools for quality monitoring | 29 | 4 | 29 | 25 |
| Obtain and use patients/consumers and family feedback | 5 | 0 | 5 | 5 |
| Purposefully reexamine the implementation | 33 | 3 | 33 | 30 |
| Stage implementation scale up | 1 | 1 | 1 | 0 |
| Assess and redesign workflows ^a | 19 | 2 | 19 | 17 |
| Provide interactive assistance | | | | |
| Centralize technical assistance | 16 | 8 | 13 | 8 |
| Internal Facilitation ^a | 10 | 1 | 9 | 9 |
| External Facilitation ^a | 59 | 5 | 59 | 54 |
| Provide clinical supervision | 4 | 0 | 4 | 4 |
| Provide local technical assistance | 9 | 2 | 9 | 7 |
| Create an online learning community ^a | 8 | 1 | 8 | 7 |
| Adapt and tailor to context | | | | |
| Promote adaptability | 38 | 2 | 37 | 36 |
| Tailor strategies | 43 | 2 | 43 | 41 |
| Use data experts | 0 | 0 | 0 | 0 |
| Use data warehousing techniques | 2 | 0 | 2 | 2 |
| Develop stakeholder interrelationships | | | | |
| Build a coalition | 4 | 1 | 4 | 3 |
| Capture and share local knowledge | 13 | 2 | 13 | 11 |
| Conduct local consensus discussions | 19 | 2 | 19 | 17 |
| Develop academic partnerships | 1 | 0 | 1 | 1 |
| Identify and prepare champions | 22 | 1 | 22 | 21 |
| Identify early adopters | 0 | 0 | 0 | 0 |
| Inform local opinion leaders | 11 | 1 | 11 | 10 |
| Involve executive boards | 6 | 0 | 6 | 6 |
| Model and simulate change | 2 | 0 | 2 | 2 |
| Obtain formal commitments | 12 | 4 | 12 | 8 |
| Organize clinician implementation team meetings | 43 | 6 | 42 | 37 |
| Promote network weaving | 1 | 0 | 1 | 1 |
| Recruit, designate, and train for leadership | 19 | 2 | 19 | 17 |
| Use advisory boards and workgroups | 11 | 1 | 10 | 10 |
| Visit other sites | 1 | 0 | 1 | 10 |
| Engage community resources outside the practice* | 4 | 0 | 4 | 4 |
| Train and educate stakeholders | , | 0 | 1 | |
| Conduct educational meetings | 97 | 29 | 96 | 68 |
| Conduct educational meetings | 20 | 29 | 19 | 18 |
| Conduct educational outreach visits | 34 | 5 | 34 | 29 |
| Create a learning collaborative | 15 | 4 | 15 | 29 11 |
| Distribute educational materials | 100 | 4 51 | 99 | 49 |
| Make training dynamic | 20 | 3 | 20 | 49 17 |
| Provide ongoing consultation | 19 | 3 | 19 | 16 |

Table 2 (continued)

| Implementation Strategy | Overall | Control Arm (Least intensive arm) | Experimental Arm (Most intensive arm) | Tested ^b |
|--|---------|---|--|---------------------|
| Shadow other experts | 1 | 0 | 1 | 1 |
| Use train-the-trainer strategies | 10 | 2 | 9 | 8 |
| Support clinicians | | | | |
| Create new clinical teams | 1 | 0 | 1 | 1 |
| Develop resource sharing agreements | 0 | 0 | 0 | 0 |
| Facilitate relay of clinical data to providers | 10 | 2 | 10 | 8 |
| Remind clinicians | 22 | 5 | 22 | 17 |
| Revise professional roles | 2 | 0 | 2 | 2 |
| Engage consumers | | | | |
| Increase demand | 6 | 1 | 6 | 5 |
| Intervene with patients/consumers to enhance uptake and adher- ence | 18 | 8 | 16 | 10 |
| Involve patients/consumers and family members | 8 | 1 | 8 | 7 |
| Prepare patients/consumers to be active participants | 20 | 5 | 20 | 15 |
| Use mass media | 2 | 2 | 2 | 0 |
| Utilize financial strategies | | | | |
| Access new funding | 4 | 0 | 4 | 4 |
| Alter incentive/allowance structures | 9 | 4 | 8 | 5 |
| Alter patient/consumer fees | 0 | 0 | 0 | 0 |
| Develop disincentives | 0 | 0 | 0 | 0 |
| Fund and contract for the clinical innovation | 1 | 1 | 1 | 0 |
| Make billing easier | 0 | 0 | 0 | 0 |
| Place innovation on fee for service lists/formularies | 1 | 0 | 1 | 1 |
| Use capitated payments | 0 | 0 | 0 | 0 |
| Use other payment schemes | 1 | 0 | 1 | 1 |
| Change infrastructure | | | | |
| Change accreditation or membership requirements | 2 | 0 | 2 | 2 |
| Change liability laws | 0 | 0 | 0 | 0 |
| Change record systems | 12 | 3 | 12 | 9 |
| Change service sites | 0 | 0 | 0 | 0 |
| Mandate change | 5 | 1 | 5 | 4 |
| Start a dissemination organization | 0 | 0 | 0 | 0 |

^a Indicates implementation strategies new to ERIC

^b The Tested column includes only those strategies used exclusively in the Experimental Arm. Tested strategies may not be the difference between Experimental Arm and Control Arm as strategies may occur in the Control Arm but not the Experimental Arm

The following strategies were determined to be duplicative or subsumed in other strategies: Develop and Organize Quality Monitoring Systems, Facilitation, Develop an Implementation Glossary, Use an Implementation Advisor, Develop Educational Materials, Work with Educational Institutions, Change Physical Structure and equipment, and Create or Change Credentialing and/or Licensure Standards. See Additional File 3 for full details.

Ten implementation strategies were not used in any studies reviewed, including: Use Data Experts, Change Liability Laws, Change Service Sites, Start a Dissemination Organization, Identify Early Adopters, Develop Resource Sharing Agreements, Alter Patient/Consumer Fees, Make Billing Easier, Use Capitated Payments, and Develop Disincentives

We acknowledge that the term stakeholder can be problematic in that it connotes the violent power differential for indigenous populations

Mass Media, Stage Implementation Scale Up, and Fund and Contract for the Clinical Innovation.

2. Which implementation strategies were commonly paired?

Table 3 shows the five most used strategies in Experimental Arms with their top ten most frequent pairings, excluding Distribute Educational Materials and Conduct Educational Meetings, as these strategies were included in almost all Experimental and half of Control Arms. The five most used strategies in the Experimental

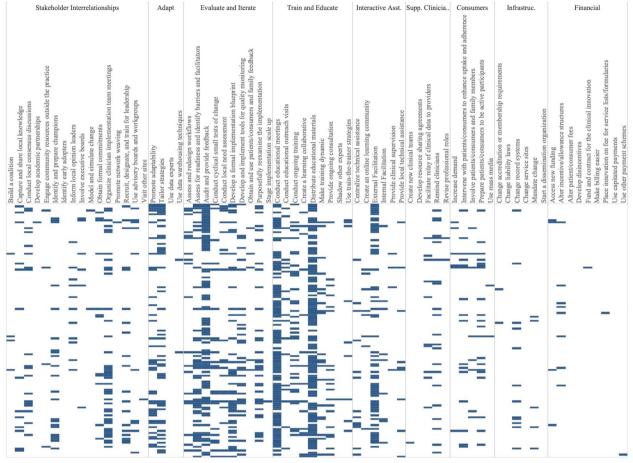


Fig. 2 Implementation strategies used in the Experimental Arm of included studies. Explore more here: https://public.tableau.com/views/Figure2_ 16947070561090/Figure2?:language=en-US&display_count=n&:origin=viz_share_link

Arm included Audit and Provide Feedback (n=76), External Facilitation (n=59), Tailor Strategies (n=43), Assess for Readiness and Identify Barriers and Facilitators (n=43), and Organize Implementation Teams (n=42).

Strategies frequently paired with these five strategies included two educational strategies: Distribute Educational Materials and Conduct Educational Meetings. Other commonly paired strategies included Develop a Formal Implementation Blueprint, Promote Adaptability, Conduct Ongoing Training, Purposefully Reexamine the Implementation, and Develop and Implement Tools for Quality Monitoring.

3. What is the evidence supporting commonly tested implementation strategies?

We classified the strength of evidence for each strategy by evaluating both the number of studies in which each strategy appeared in the Experimental Arm and the percentage of times there was at least one significantly positive RE-AIM outcome. Using these factors, Fig. 3 shows the number of studies in which individual strategies were evaluated (on the y axis) compared to the percentage of times that studies including those strategies had at least one positive outcome (on the x axis). Due to the non-normal distribution of both factors, we used the median (rather than the mean) to create four quadrants. Strategies in the lower left quadrant were tested in fewer than the median number of studies (8.5) and were less frequently associated with a significant RE-AIM outcome (75%). The upper right quadrant included strategies that occurred in more than the median number of studies (8.5) and had more than the median percent of studies with a significant RE-AIM outcome (75%); thus those 19 strategies were viewed as having stronger evidence. Of those 19 implementation strategies, Conduct Educational Meetings, Distribute Educational Materials, External Facilitation, and Audit and Provide Feedback continued to occur frequently, appearing in 59-99 studies.

| Implementation Strategy | Audit and provide feedback^ [76 studies] | External Facilitation^ [59 studies] | Tailor Strategies [43 studies] | Assess for readiness and identify barriers and facilitators^ [43 studies] | Organize clinician implementation team meetings [42 studies] |
|--|--|-------------------------------------|--------------------------------|--|---|
| Distribute educational materials^ | 57 | 48 | 36 | 36 | 35 |
| Conduct educational meetings^ | 55 | 52 | 33 | 39 | 36 |
| Audit and provide feedback^ | 0 | 38 | 27 | 28 | 22 |
| External Facilitation^ | 38 | 0 | 27 | 30 | 31 |
| Tailor strategies | 27 | 27 | 0 | 25 | 19 |
| Assess for readiness and identify barriers and facilitators^ | 28 | 30 | 25 | 0 | 22 |
| Organize clinician implementation team meetings | 22 | 31 | 19 | 22 | 0 |
| Develop a formal implementation blueprint | 24 | 27 | 20 | 25 | 26 |
| Promote adaptability | 22 | 22 | 19 | 22 | 19 |
| Conduct ongoing training | 15 | 20 | 15 | 15 | 15 |
| Purposefully reexamine the implementation^ | 27 | 24 | 15 | 24 | 20 |
| Develop and implement tools for quality monitoring [^] | 22 | 13 | 10 | 14 | 13 |

Table 3 Top 5 commonly used strategies in the Experimental Arm and their 10 most common pairings, organized by cluster +

Darker colors indicate a high frequency of pairings between implementation strategies.

[†] These top 5 strategies exclude Distribute Educational Materials and Conduct Educational Meetings, as these strategies were included in almost all Experimental and half of Control arms.

^denotes an implementation strategy in the top right quadrant represented in Fig.3. Explore more here: https://public.tableau.com/views/Table3_16947882124200/ Table3?:language=en-US&:display_count=n&:origin=viz_share_link

Figure 4 graphically illustrates the proportion of significant outcomes for each RE-AIM outcome for the 19 commonly used and evidence-based implementation strategies in the upper right quadrant. These findings again show the widespread use of Conduct Educational Meetings and Distribute Educational Materials. Implementation and Effectiveness outcomes were assessed most frequently, with Implementation being the mostly commonly reported significantly positive outcome.

Discussion

This systematic review identified 129 experimental studies examining the effectiveness of implementation strategies across a broad range of health and human service studies. Overall, we found that evidence is lacking for most ERIC implementation strategies, that most studies employed combinations of strategies, and that implementation outcomes, categorized by RE-AIM dimensions, have not been universally defined or applied. Accordingly, other researchers have described the need for universal outcomes definitions and descriptions across implementation research studies [28, 42]. Our findings have important implications not only for the current state of the field but also for creating guidance to help investigators determine which strategies and in what context to examine.

The four most evaluated strategies were Distribute Educational Materials, Conduct Educational Meetings, External Facilitation, and Audit and Provide Feedback. Conducting Educational Meetings and Distributing Educational Materials were surprisingly the most common. This may reflect the fact that education strategies are generally considered to be "necessary but not sufficient" for successful implementation [43, 44]. Because education is often embedded in interventions, it is critical to define the boundary between the innovation and the implementation strategies used to support the innovation. Further specification as to when these strategies are EBP core components or implementation strategies (e.g., booster trainings or remediation) is needed [45, 46].

We identified 19 implementation strategies that were tested in at least 8 studies (more than the median) and were associated with positive results at least 75% of the time. These strategies can be further categorized as being used in early or pre-implementation versus later in implementation. Preparatory activities or pre-implementation, strategies that had strong evidence included educational activities (Meetings, Materials, Outreach visits, Train for Leadership, Use Train the Trainer Strategies) and site diagnostic activities (Assess for Readiness, Identify Barriers and Facilitators, Conduct Local Needs Assessment, Identify and Prepare Champions, and Assess and Redesign Workflows). Strategies that target the implementation phase include those that provide coaching and support (External and Internal Facilitation), involve additional key partners (Intervene with Patients to Enhance

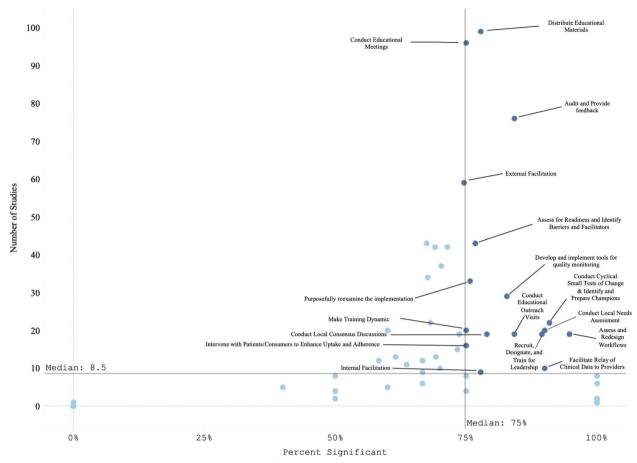


Fig. 3 Experimental Arm Implementation Strategies with significant RE-AIM outcome. Explore more here: https://public.tableau.com/views/Figur e3_16947017936500/Figure3?:language=en-US&publish=yes&:display_count=n&:origin=viz_share_link

Uptake and Adherence), and engage in quality improvement activities (Audit and Provide Feedback, Facilitate the Relay of Clinical Data to Providers, Purposefully Reexamine the Implementation, Conduct Cyclical Small Tests of Change, Develop and Implement Tools for Quality Monitoring).

There were many ERIC strategies that were not represented in the reviewed studies, specifically the financial and policy strategies. Ten strategies were not used in any studies, including: Alter Patient/Consumer Fees, Change Liability Laws, Change Service Sites, Develop Disincentives, Develop Resource Sharing Agreements, Identify Early Adopters, Make Billing Easier, Start a Dissemination Organization, Use Capitated Payments, and Use Data Experts. One of the limitations of this investigation was that not all individual strategies or combinations were investigated. Reasons for the absence of these strategies in our review may include challenges with testing certain strategies experimentally (e.g., changing liability laws), limitations in our search terms, and the relative paucity of implementation strategy trials compared to clinical trials. Many "untested" strategies require large-scale structural changes with leadership support (see [47] for policy experiment example). Recent preliminary work has assessed the feasibility of applying policy strategies and described the challenges with doing so [48–50]. While not impossible in large systems like VA (for example: the randomized evaluation of the VA Stratification Tool for Opioid Risk Management) the large size, structure, and organizational imperative makes these initiatives challenging to experimentally evaluate. Likewise, the absence of these ten strategies may have been the result of our inclusion criteria, which required an experimental design. Thus, creative study designs may be needed to test high-level policy or financial strategies experimentally.

Some strategies that were likely under-represented in our search strategy included electronic medical record reminders and clinical decision support tools and systems. These are often considered "interventions" when used by clinical trialists and may not be indexed as studies involving 'implementation strategies' (these tools have

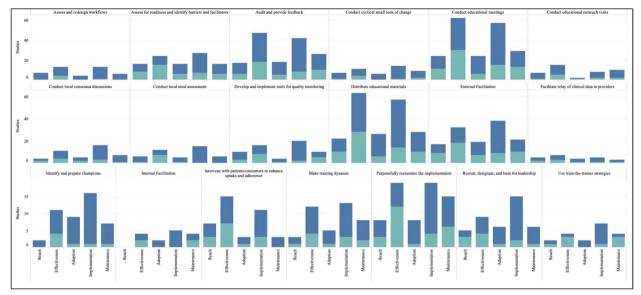


Fig. 4 RE-AIM outcomes for the 19 Top-Right Quadrant Implementation Strategies. The y-axis is the number of studies and the x-axis is a stacked bar chart for each RE-AIM outcome with R=Reach, E=Effectiveness, A=Adoption, I=Implementation, M=Maintenance. Blue denotes at least one significant RE-AIM outcome; Light blue denotes studies which used the given implementation strategy and did not have a significant RE-AIM. Explore more here: https://public.tableau.com/views/Figure4_16947017112150/Figure4?:language=en-US&publish=yes&:display_count=n&:origin=viz_share_link

been reviewed elsewhere [51-53]). Thus, strategies that are also considered interventions in the literature (e.g., education interventions) were not sought or captured. Our findings do not imply that these strategies are ineffective, rather that more study is needed. Consistent with prior investigations [54], few studies meeting inclusion criteria tested financial strategies. Accordingly, there are increasing calls to track and monitor the effects of financial strategies within implementation science to understand their effectiveness in practice [55, 56]. However, experts have noted that the study of financial strategies can be a challenge given that they are typically implemented at the system-level and necessitate research designs for studying policy-effects (e.g., quasi-experimental methods, systems-science modeling methods) [57]. Yet, there have been some recent efforts to use financial strategies to support EBPs that appear promising [58] and could be a model for the field moving forward.

The relationship between the number of strategies used and improved outcomes has been described inconsistently in the literature. While some studies have found improved outcomes with a bundle of strategies that were uniquely combined or a standardized package of strategies (e.g., Replicating Effective Programs [59, 60] and Getting To Outcomes [61, 62]), others have found that "more is not always better" [63–65]. For example, Rogal and colleagues documented that VA hospitals implementing a new evidence-based hepatitis C treatment chose >20 strategies, when multiple years of data linking strategies to outcomes showed that 1-3 specific strategies would have yielded the same outcome [39]. Considering that most studies employed multiple or multifaceted strategies, it seems that there is a benefit of using a targeted bundle of strategies that are purposefully aligns with site/clinic/population norms, rather than simply adding more strategies [66].

It is difficult to assess the effectiveness of any one implementation strategy in bundles where multiple strategies are used simultaneously. Even a 'single' strategy like External Facilitation is, in actuality, a bundle of narrowly constructed strategies (e.g., Conduct Educational Meetings, Identify and Prepare Champions, and Develop a Formal Implementation Blueprint). Thus, studying External Facilitation does not allow for a test of the individual strategies that comprise it, potentially masking the effectiveness of any individual strategy. While we cannot easily disaggregate the effects of multifaceted strategies, doing so may not yield meaningful results. Because strategies often synergize, disaggregated results could either underestimate the true impact of individual strategies or conversely, actually undermine their effectiveness (i.e., when their effectiveness comes from their combination with other strategies). The complexity of health and human service settings, imperative to improve public health outcomes, and engagement with community partners often requires the use of multiple strategies

simultaneously. Therefore, the need to improve realworld implementation may outweigh the theoretical need to identify individual strategy effectiveness. In situations where it would be useful to isolate the impact of single strategies, we suggest that the same methods for documenting and analyzing the critical components (or core functions) of complex interventions [67–70] may help to identify core components of multifaceted implementation strategies [71–74].

In addition, to truly assess the impacts of strategies on outcomes, it may be necessary to track fidelity to implementation strategies (not just the EBPs they support). While this can be challenging, without some degree of tracking and fidelity checks, one cannot determine whether a strategy's apparent failure to work was because it 1) was ineffective or 2) was not applied well. To facilitate this tracking there are pragmatic tools to support researchers. For example, the Longitudinal Implementation Strategy Tracking System (LISTS) offers a pragmatic and feasible means to assess fidelity to and adaptations of strategies [75].

Implications for implementation science: four recommendations

Based on our findings, we offer four recommended "best practices" for implementation studies.

- 1. Prespecify strategies using standard nomenclature. This study reaffirmed the need to apply not only a standard naming convention (e.g., ERIC) but also a standard *reporting* of for implementation strategies. While reporting systems like those by Proctor [1] or Pinnock [75] would optimize learning across studies, few manuscripts specify strategies as recommended [76, 77]. Pre-specification allows planners and evaluators to assess the feasibility and acceptability of strategies with partners and community members [24, 78, 79] and allows evaluators and implementers to monitor and measure the fidelity, dose, and adaptations to strategies delivered over the course of implementation [27]. In turn, these data can be used to assess the costs, analyze their effectiveness [38, 80, 81], and ensure more accurate reporting [82–85]. This specification should include, among other data, the intensity, stage of implementation, and justification for the selection. Information regarding why strategies were selected for specific settings would further the field and be of great use to practitioners. [63, 65, 69, 79, 86].
- 2. Ensure that standards for measuring and reporting implementation outcomes are consistently applied and account for the complexity of implementation studies. Part of improving standardized reporting

must include clearly defining outcomes and linking each outcome to particular implementation strategies. It was challenging in the present review to disentangle the impact of the intervention(s) (i.e., the EBP) versus the impact of the implementation strategy(ies) for each RE-AIM dimension. For example, often fidelity to the EBP was reported but not for the implementation strategies. Similarly, Reach and Adoption of the intervention would be reported for the Experimental Arm but not for the Control Arm, prohibiting statistical comparisons of strategies on the relative impact of the EBP between study arms. Moreover, there were many studies evaluating numerous outcomes, risking data dredging. Further, the significant heterogeneity in the ways in which implementation outcomes are operationalized and reported is a substantial barrier to conducting large-scale meta-analytic approaches to synthesizing evidence for implementation strategies [67]. The field could look to others in the social and health sciences for examples in how to test, validate, and promote a common set of outcome measures to aid in bringing consistency across studies and real-world practice (e.g., the NIH-funded Patient-Reported Outcomes Measurement Information System [PROMIS], https://www.healthmeasures.net/explore-measu rement-systems/promis).

- 3. Develop infrastructure to learn cross-study lessons in implementation science. Data repositories, like those developed by NCI for rare diseases, U.S. HIV Implementation Science Coordination Initiative [87], and the Behavior Change Technique Ontology [88], could allow implementation scientists to report their findings in a more standardized manner, which would promote ease of communication and contextualization of findings across studies. For example, the HIV Implementation Science Coordination Initiative requested all implementation projects use common frameworks, developed user friendly databases to enable practitioners to match strategies to determinants, and developed a dashboard of studies that assessed implementation determinants [89–94].
- 4. Develop and apply methods to rigorously study common strategies and bundles. These findings support prior recommendations for improved empirical rigor in implementation studies [46, 95]. Many studies were excluded from our review based on not meeting methodological rigor standards. Understanding the effectiveness of discrete strategies deployed alone or in combination requires reliable and low burden tracking methods to collect information about strategy use and outcomes. For example, frameworks like the Implementation Replication Framework [96]

could help interpret findings across studies using the same strategy bundle. Other tracking approaches may leverage technology (e.g., cell phones, tablets, EMR templates) [78, 97] or find novel, pragmatic approaches to collect recommended strategy specifications over time (e.g., dose, deliverer, and mechanism) [1, 9, 27, 98, 99]. Rigorous reporting standards could inform more robust analyses and conclusions (e.g., moving toward the goal of understanding causality, microcosting efforts) [24, 38, 100, 101]. Such detailed tracking is also required to understand how site-level factors moderate implementation strategy effects [102]. In some cases, adaptive trial designs like sequential multiple assignment randomized trials (SMARTs) and just-in-time adaptive interventions (JITAIs) can be helpful for planning strategy escalation.

Limitations

Despite the strengths of this review, there were certain notable limitations. For one, we only included experimental studies, omitting many informative observational investigations that cover the range of implementation strategies. Second, our study period was centered on the creation of the journal Implementation Science and not on the standardization and operationalization of implementation strategies in the publication of the ERIC taxonomy (which came later). This, in conjunction with latency in reporting study results and funding cycles, means that the employed taxonomy was not applied in earlier studies. To address this limitation, we retroactively mapped strategies to ERIC, but it is possible that some studies were missed. Additionally, indexing approaches used by academic databases may have missed relevant studies. We addressed this particular concern by reviewing other systematic reviews of implementation strategies and soliciting recommendations from global implementation science experts.

Another potential limitation comes from the ERIC taxonomy itself—i.e., strategy listings like ERIC are only useful when they are widely adopted and used in conjunction with guidelines for specifying and reporting strategies [1] in protocol and outcome papers. Although the ERIC paper has been widely cited (over three thousand times, accessed about 186 thousand times), it is still not universally applied, making tracking the impact of specific strategies more difficult. However, our experience with this review seemed to suggest that ERIC's use was increasing over time. Also, some have commented that ERIC strategies can be unclear and are missing key domains. Thus, researchers are making definitions clearer for lay users [37, 103], increasing the number of discrete strategies for specific domains

like HIV treatment, acknowledging strategies for new functions (e.g., de-implementation [104], local capacity building), accounting for phases of implementation (dissemination, sustainment [13], scale-up), addressing settings [12, 20], actors roles in the process, and making mechanisms of change to select strategies more user-friendly through searchable databases [9, 10, 54, 73, 104–106]. In sum, we found the utility of the ERIC taxonomy to outweigh any of the taxonomy's current limitations.

As with all reviews, the search terms influenced our findings. As such, the broad terms for implementation strategies (e.g., "evidence-based interventions"[7] or "behavior change techniques" [107]) may have led to inadvertent omissions of studies of specific strategies. For example, the search terms may not have captured tests of policies, financial strategies, community health promotion initiatives, or electronic medical record reminders, due to differences in terminology used in corresponding subfields of research (e.g., health economics, business, health information technology, and health policy). To manage this, we asked experts to inform us about any studies that they would include and cross-checked their lists with what was identified through our search terms, which yielded very few additional studies. We included standard coding using the ERIC taxonomy, which was a strength, but future work should consider including the additional strategies that have been recommended to augment ERIC, around sustainment [13, 79, 106, 108], community and public health research [12, 109–111], consumer or service user engagement [112], de-implementation [104, 113-117] and related terms [118].

We were unable to assess the bias of studies due to non-standard reporting across the papers and the heterogeneity of study designs, measurement of implementation strategies and outcomes, and analytic approaches. This could have resulted in over- or underestimating the results of our synthesis. We addressed this limitation by being cautious in our reporting of findings, specifically in identifying "effective" implementation strategies. Further, we were not able to gather primary data to evaluate effect sizes across studies in order to systematically evaluate bias, which would be fruitful for future study.

Conclusions

This novel review of 129 studies summarized the body of evidence supporting the use of ERIC-defined implementation strategies to improve health or healthcare. We identified commonly occurring implementation strategies, frequently used bundles, and the strategies with the highest degree of supportive evidence, while simultaneously identifying gaps in the literature. Additionally, we identified several key areas for future growth and operationalization across the field of implementation science with the goal of improved reporting and assessment of implementation strategies and related outcomes.

Abbreviations

| CDC | Centers for Disease Control | | |
|-----------|---|--|--|
| CINAHL | Cumulated Index to Nursing and Allied Health Literature | | |
| D&I | Dissemination and Implementation | | |
| EBP | Evidence-based practices or programs | | |
| ERIC | Expert Recommendations for Implementing Change | | |
| MOST | Multiphase Optimization Strategy | | |
| NCI | National Cancer Institute | | |
| NIH | National Institutes of Health | | |
| Pitt DISC | The Pittsburgh Dissemination and Implementation Science | | |
| | Collaborative | | |
| SMART | Sequential Multiple Assignment Randomized Trial | | |
| U.S. | United States | | |
| VA | Department of Veterans Affairs | | |
| | | | |

Supplementary Information

The online version contains supplementary material available at https://doi. org/10.1186/s13012-024-01369-5.

Supplementary Material 1. Supplementary Material 2. Supplementary Material 3.

Supplementary Material 4.

Supplementary Material 5.

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Author's contributions

LEA, SSR, and MJC conceptualized the study. LEA, SSR, MJC, and JOH developed the study design. LEA and JOH acquired the data. LEA, DEG, AP, RLB, DAQ, LGL, LPS, SSR, NQ, and MJC conducted the abstract, full text review, and rigor assessment. LEA, DEG, JOH, AP, RLB, DAQ, NQ, NCE, SSR, and MJC conducted the data abstraction. DEG, SSR, and MJC adjudicated conflicts. LEA and SSR analyzed the data. LEA, SSR, JOH, and MJC interpreted the data. LEA, SSR, and MJC drafted the work. All authors substantially revised the work. All authors approved the submitted version and agreed to be personally accountable for their contributions and the integrity of the work.

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Availability and materials

All data for this study are included in this published article and its supplementary information files.

Declarations

Ethics approval and consent to participate Not applicable.

Consent for publication

The manuscript does not contain any individual person's data.

Competing interests

None.

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