

BMJ Open Characteristics of knowledge translation platforms and methods for evaluating them: a scoping review protocol

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ABSTRACT

Introduction Knowledge translation platforms (KTPs) are intermediary organisations, initiatives or networks whose intent is to bridge the evidence into action divide. Strategies and tools include collaborative knowledge production, capacity building, information exchange and dialogue to facilitate relevant and timely engagement between researchers and decision-makers and other relevant stakeholders. With the wide range of definitions and descriptions of KTPs, there is a need to (1) provide a nuanced understanding of characteristics of KTPs and (2) assess and consolidate research methods used in mapping and evaluating KTPs to inform standardised process and impact evaluation.

Methods and analysis This scoping review will follow the recommended and accepted methods for scoping reviews and reporting guidelines. Eligibility for inclusion is any conceptual or empirical health-related qualitative, quantitative and/or mixed method studies including (1) definitions, descriptions and models or frameworks of KTPs (including those that do not self-identify as KTPs, eg, university research centres) and (2) research methods for mapping and/or evaluating KTPs. Searches will be carried out in PubMed, Scopus, CINAHL, Embase, Global Health and Web of Science using a predetermined search strategy, without any date, language or geographical restrictions. Two reviewers will independently screen titles and abstracts. One reviewer will complete data extraction for all included studies, and another will check a sample of 50% of the included studies. The analysis and synthesis will provide (1) an understanding of the various characteristics of KTPs; (2) insight into characteristics or factors that make them resilient and/or adaptive to facilitate impact (ie, influence policy and practice); and (3) an overview of the various methods for mapping and evaluating KTPs. We will explore enhancing an existing framework for classifying KTPs, or perhaps even developing a new framework for identifying and monitoring KTPs if necessary and relevant.

Ethics and dissemination This scoping review does not require ethics approval, as we will only include information from previously conducted studies and we will not involve human participants. The results will be submitted to a peer-reviewed scientific journal for publication and as conference presentations.

BACKGROUND

The global focus on achieving the sustainable development goals (SDGs) and implementing

STRENGTHS AND LIMITATIONS OF THIS STUDY

- ⇒ This scoping review will identify different qualitative, quantitative and mixed methods that can be used or combined to develop robust evaluations of knowledge translation platforms (KTPs).
- ⇒ The analysis and synthesis will guide the review authors in enhancing an existing framework for classifying KTPs or developing a new framework for identifying and monitoring KTPs if necessary and relevant.
- ⇒ This scoping review will include a consultation of authors of included studies and/or experts in the field, as well as other knowledge translation researchers and practitioners, to ensure its relevance and applicability.

universal health coverage (UHC) has resulted in an increased demand for research evidence to inform policy and practice.^{1 2} SDG 3 is about ensuring healthy lives and promoting well-being at all ages, which is closely linked with the aims of UHC.³ UHC aims to ensure that all people have access to needed and effective health services (including prevention, promotion, treatment, rehabilitation and palliation) of sufficient quality, without exposing users to financial hardship.⁴ However, despite the widely agreed SDGs and the UHC model, there is limited use of high-quality research on the effectiveness, acceptability and cost implications of health system, healthcare or public health interventions to inform policy, practice and implementation.⁵ Although research evidence on health-related interventions and systems is increasingly available, there are challenges around translating research evidence into policy and practice.⁵⁻⁸

Translating research evidence into policy and practice, or knowledge translation (KT), entails a 'dynamic and iterative process that includes synthesis, dissemination, exchange and ethically sound application of knowledge to improve health, provide more effective health services and products and strengthen the healthcare system'.⁹ Limited institutional



support or buy-in from leadership and relevant stakeholders for KT has been linked to several challenges,^{10–14} for example, poor infrastructure and inadequate financial and technical resources specifically for KT; inadequate soft skills, relationships and networks among evidence producers; evidence users' negative attitudes and poor knowledge about what KT is and how to do it; and scarce local research evidence relevant to microlevel policy and practice, among others.^{10–14}

Knowledge translation platforms (KTPs) are intermediary organisations, initiatives or networks whose intent is to overcome a range of inter-relationship and contextual challenges (see previous examples) using a multitude of strategies and tools.^{15 16} These include collaborative knowledge production, capacity building, information exchange and dialogue to facilitate relevant and timely engagement between researchers and different health decision-makers (eg, patients, health practitioners, healthcare managers, policy-makers and funders). There are currently a wide range of definitions and descriptions^{17–27} as well as models and frameworks^{28 29} of KTPs in the literature. However, there is only one published systematic review on KTPs by Partridge *et al*, which primarily synthesised the lessons learnt about activities, outputs, outcomes and impacts from KTPs specifically in low-income and middle-income countries.³⁰

First, there is a need to synthesise the different definitions and descriptions of KTPs in the literature so as to provide a common and nuanced understanding of what KTPs are. This is necessary for planning and carrying out comparisons and evaluations of KTPs as one mechanism for strengthening their overall usefulness. Second, there is a need to synthesise a wider range of characteristics of KTPs beyond those characteristics explored in the Partridge review; for example, strengths and limitations of different KTP models, funding and sustainability of KTPs, current operational status (determined using evidence from the scoping review and institutional websites), and the relationship between design and implementation factors or characteristics and successful functioning of KTPs and their influence to policy and practice. Having a better understanding of the different characteristics and types of KTPs can inform funders and governments about their implementation, sustainability and overall support for evidence-informed policy and practice. Additionally, identifying the different types of KTPs that can support evidence-informed policy and practice in local settings is critical, especially in the context of many public health and health system interventions for achieving UHC and the SDGs. For example, in the case that a KTP exists at the local level, decision-makers need to understand what KTPs are (including what KT is in general) and their role in supporting UHC decision-making processes. Third, there is a need to synthesise the research methods used in the literature for mapping and evaluating KTPs. Identifying the different qualitative, quantitative and mixed methods used for mapping and evaluating KTPs is an important step in exploring how different methods can

be used or combined to address the gap for more robust evaluations of KTPs. At a practical level, mapping KTPs can help like-minded organisations to identify opportunities that avoid duplication and amplify collaboration, particularly in settings where there are limited resources and expertise.

This scoping review therefore aims (1) to provide a more nuanced understanding of the characteristics of KTPs and (2) to assess and consolidate research methods used in mapping and evaluating KTPs to inform standardised process and impact evaluation. The review authors intend to use the review findings to inform a future research study on mapping, evaluating and strengthening KTPs that can support UHC policy, practice and implementation in South Africa.

IDENTIFYING THE RESEARCH QUESTIONS

This scoping review is on the characteristics and methods for mapping and evaluating KTPs. The specific objectives of the scoping review are

- ▶ To identify and synthesise definitions as well as characteristics of KTPs (eg, activities, processes, outputs, purpose, functions, models, stakeholders, positioning, funding, strengths, limitations, monitoring, impact and status).
- ▶ To identify and synthesise the design and implementation factors or characteristics that facilitate and/or hinder the successful functioning of KTPs and their influence to policy and practice.
- ▶ To identify and summarise the various research methods for mapping and evaluating KTPs (ie, qualitative, quantitative and mixed) and where evaluations were conducted, to synthesise their outcomes (eg, impact, success or failure).
- ▶ To explore the potential for developing a new or enhancing an existing framework for classifying KTPs.

IDENTIFYING RELEVANT STUDIES

We will use the scoping review methods outlined by Arksey and O'Malley.³⁰ The proposed steps are identifying the research question; identifying relevant studies; study selection; charting the data; collating, summarising and reporting the results; and consultation (ie, seeking insights beyond those in the literature from content experts). To report our review findings, we will follow the Preferred Reporting Items for Systematic Reviews and Meta-Analyses Extension for Scoping Reviews: Checklist and Explanation (see online supplemental file 1).³¹ We aim to conduct the proposed steps by February 2023.

Any published, empirical and conceptual, health research studies from anywhere on KTPs will be eligible. Qualitative, quantitative and/or mixed method studies are eligible for inclusion as long as they include a definition or description of what a KTP is and/or include one or more methods for mapping and/or evaluating KTPs. Eligible participants are KTP staff and users, for example, researchers, knowledge brokers, policy-makers, health

practitioners and managers, patients and community representatives, and journalists. KTPs can exist within and between different settings, for example, universities, research councils, professional bodies, civil society organisations, community organisations and government.^{15 16}

The search will identify all relevant studies without data, language or geographical restrictions. We will search the following electronic databases: PubMed, Scopus, CINAHL, Embase, Global Health and Web of Science. Search strings will include keywords and Medical Subject Headings (MeSH) terms related to KTP (concept A) (eg, policy brief, deliberate dialogue and knowledge exchange) and evidence-informed decision-making (concept B) (eg, health policy and policymaking). We have developed a preliminary search strategy in the PubMed database (see online supplemental file 2). To finalise our search strategy, we will apply an iterative approach to check whether known articles that meet our eligibility criteria were found by the search. We will also identify missing keywords and MeSH terms to add to our search strategy from the iterative process. In addition to the electronic searches, review authors will search the reference lists of all included studies and key references (eg, relevant systematic reviews) and contact authors of included studies and/or experts in the field for additional references.

STUDY SELECTION

The search across databases will identify titles and abstracts of relevant studies. The search results will be merged in the EndNote reference management programme where duplicates will be removed. The titles and abstracts will then be uploaded to an electronic programme, such as Covidence or Rayyan, for screening and data extraction. An eligibility form will be developed before screening starts. The following inclusion criteria will be used.

Focus of studies

- ▶ Definitions and characteristics of KTPs (eg, activities, processes, outputs, purpose, functions, models, stakeholders, positioning, funding, strengths, limitations, monitoring, impact and status).
- ▶ Factors or characteristics that facilitate and/or hinder the successful functioning of KTPs and their influence to policy and practice.
- ▶ Research methods for mapping and evaluating KTPs (ie, qualitative, quantitative and mixed).
- ▶ Evaluation outcomes of KTPs (eg, impact, success or failure).
- ▶ Existing frameworks for classifying KTPs.

The outcomes of KTPs (success or failure) will be based on what the KTPs themselves aim to achieve through their activities and processes. Where a KTP has been evaluated, the review team will use the evaluation results to determine which factors or characteristics contributed to its success or challenges. Where a KTP has not been evaluated, the review team will explore what the primary

authors deem as important factors or characteristics that facilitate and/or hinder its successful functioning and influence on policy and practice

Participants

KTP staff and users (eg, researchers, knowledge brokers, policy-makers, health practitioners and managers, patients and community representatives and journalists).

Setting

KTPs can exist within and between different settings (eg, research centres or departments, universities, research councils, professional bodies, civil society organisations, community organisations and government).

Two review authors will independently screen the titles and abstracts to determine their eligibility for full-text screening. We will retrieve full texts for titles and abstracts deemed relevant. One review author (B-MS) will screen all full texts and make a final decision about inclusion. Another review author will check the eligibility of a random sample of 50% of the full texts. Conflicts will be resolved with a third reviewer. The study selection process will be summarised using a Preferred Reporting Items for Systematic Reviews and Meta-Analyses flow diagram.

CHARTING THE DATA

Data extraction or 'charting of data' will be carried out once we have a final list of all the studies to be included in the review. Data extraction will be conducted by one review author (B-MS) who will collect, sift and sort data according to the objectives. The review author will extract information on the study and author details; research type and study design; research setting and participants; definitions, descriptions and characteristics of KTPs; and methods used for mapping and/or evaluating KTPs. A second review author will check data extraction of all included studies. Data extraction will be done in Excel to allow for comparison of key items across studies and to allow for synthesis within and across data items. Once all the data have been extracted and checked, studies will be categorised or 'charted' according to the following criteria: (1) what is a KTP? and/or (2) what methods are used to map and/or evaluate a KTP? Additional categories may be identified during the data extraction process, in consultation with the review team. We will not assess the methodological quality of the included studies, as that is the convention for such scoping reviews.^{29 30}

COLLATING, SUMMARISING AND REPORTING THE RESULTS

One review author (B-MS) will conduct data analysis using manual coding and data synthesis methods on the extracted and charted data. A second review author will check the data analysis work on an ongoing basis to ensure quality of the process. We will synthesise the data according to variation (breadth) and key components (depth) across definitions, characteristics and methods of

KTPs. The analysis will combine quantitative and qualitative syntheses to provide an overview of our findings. First, we will conduct a numerical analysis of all the included studies according to different categories, for example, study design (qualitative, quantitative and mixed), participants (KTP staff vs users), KTP characteristics (model, function and institution), and income level of country (low, middle and high). Second, we will conduct a qualitative narrative synthesis of the definitions, characteristics and methods of KTPs by looking for the key components across the data. The numerical and narrative syntheses will provide (1) an overview of the key aspects of KTP studies, (2) definitions and conceptualisations of KTPs, and (3) research methods for mapping and/or evaluating KTPs.

CONSULTATIONS

As mentioned earlier, to identify additional relevant studies, we will contact authors of included studies and/or experts in the field. We will engage with other KT researchers (ie, those undertaking scientific research in the KT field) and KT practitioners (ie, those designing, implementing, monitoring and evaluating KT interventions) as we carry out different stages of this scoping review to ensure its relevance and applicability. It is an advantage that our review team is made up of both KT researchers and practitioners who have the appropriate content and method expertise to ensure scientific rigour of the review. We will shape the review process and findings according to what we know is most useful from experience and also draw on colleagues working in the field to validate the findings and extract key messages or implications for research and practice.

PATIENT AND PUBLIC INVOLVEMENT

There was no patient or public involvement in the design of this protocol.

DISCUSSION

To our knowledge, this is the first scoping review of the evidence on KTPs globally. Our synthesis will be on the wide range of definitions, descriptions and characteristics of KTPs and aims to provide a better understanding of the relationship between some of the characteristics of KTPs (eg, we will explore the relationship between KTP design, success factors and effectiveness or impact). A key methodological strength of the scoping review is that we will scope for both conceptual and empirical studies, using any study design, and without applying any date, language or geographical restrictions.

Mapping KTPs in this scoping review can help researchers and other stakeholders leading KTPs identify opportunities for collaborating with other KTPs in their local setting or globally. Collaborations among KTPs can potentially reduce duplication of efforts and optimise the

use of already-limited resources and expertise to effectively engage decision-makers. Additionally, a synthesis of the characteristics of KTPs can help researchers and other stakeholders leading KTPs compare and evaluate the usefulness and effectiveness of different KTPs in relation to theirs. Evidence on the usefulness and effectiveness of KTPs, areas of improvement and the gaps in research can guide government and funding agencies' agendas. Further, the review findings can help decision-makers identify KTPs to collaborate with and provide them with an understanding of how KTPs can support decision-making processes. Lastly, the synthesis will guide review authors in developing a matrix or framework for classifying KTPs and inform the design of a future study to map and evaluate KTPs that can support UHC policy, practice and implementation in South Africa.

ETHICS AND DISSEMINATION

This is a scoping review of completed studies. As such, our research does not require ethics approval, as we do not involve human participants. The results will be submitted to a peer-reviewed scientific journal for publication and as conference presentations.

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Contributors B-MS conceptualised the protocol together with TY and NSJ. B-MS drafted the protocol with SC and TY providing methodological expertise and NSJ providing content expertise. All authors reviewed and approved the final manuscript before final submission for peer review.

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Competing interests None declared.

Patient and public involvement Patients and/or the public were not involved in the design, conduct, reporting, or dissemination plans of this research.

Patient consent for publication Not applicable.

Provenance and peer review Not commissioned; externally peer reviewed.

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Preferred Reporting Items for Systematic reviews and Meta-Analyses extension for Scoping Reviews (PRISMA-ScR) Checklist

SECTION	ITEM	PRISMA-ScR CHECKLIST ITEM	REPORTED ON PAGE #
TITLE			
Title	1	Identify the report as a scoping review.	1
ABSTRACT			
Structured summary	2	Provide a structured summary that includes (as applicable): background, objectives, eligibility criteria, sources of evidence, charting methods, results, and conclusions that relate to the review questions and objectives.	2-3
INTRODUCTION			
Rationale	3	Describe the rationale for the review in the context of what is already known. Explain why the review questions/objectives lend themselves to a scoping review approach.	5-6
Objectives	4	Provide an explicit statement of the questions and objectives being addressed with reference to their key elements (e.g., population or participants, concepts, and context) or other relevant key elements used to conceptualize the review questions and/or objectives.	7
METHODS			
Protocol and registration	5	Indicate whether a review protocol exists; state if and where it can be accessed (e.g., a Web address); and if available, provide registration information, including the registration number.	n/a
Eligibility criteria	6	Specify characteristics of the sources of evidence used as eligibility criteria (e.g., years considered, language, and publication status), and provide a rationale.	8
Information sources*	7	Describe all information sources in the search (e.g., databases with dates of coverage and contact with authors to identify additional sources), as well as the date the most recent search was executed.	8
Search	8	Present the full electronic search strategy for at least 1 database, including any limits used, such that it could be repeated.	8 & Additional file 2
Selection of sources of evidence†	9	State the process for selecting sources of evidence (i.e., screening and eligibility) included in the scoping review.	9
Data charting process‡	10	Describe the methods of charting data from the included sources of evidence (e.g., calibrated forms or forms that have been tested by the team before their use, and whether data charting was done independently or in duplicate) and any processes for obtaining and confirming data from investigators.	9
Data items	11	List and define all variables for which data were sought and any assumptions and simplifications made.	9
Critical appraisal of individual sources of evidence§	12	If done, provide a rationale for conducting a critical appraisal of included sources of evidence; describe the methods used and how this information was used in any data synthesis (if appropriate).	n/a



SECTION	ITEM	PRISMA-ScR CHECKLIST ITEM	REPORTED ON PAGE #
Synthesis of results	13	Describe the methods of handling and summarizing the data that were charted.	10
RESULTS			
Selection of sources of evidence	14	Give numbers of sources of evidence screened, assessed for eligibility, and included in the review, with reasons for exclusions at each stage, ideally using a flow diagram.	n/a
Characteristics of sources of evidence	15	For each source of evidence, present characteristics for which data were charted and provide the citations.	n/a
Critical appraisal within sources of evidence	16	If done, present data on critical appraisal of included sources of evidence (see item 12).	n/a
Results of individual sources of evidence	17	For each included source of evidence, present the relevant data that were charted that relate to the review questions and objectives.	n/a
Synthesis of results	18	Summarize and/or present the charting results as they relate to the review questions and objectives.	n/a
DISCUSSION			
Summary of evidence	19	Summarize the main results (including an overview of concepts, themes, and types of evidence available), link to the review questions and objectives, and consider the relevance to key groups.	n/a
Limitations	20	Discuss the limitations of the scoping review process.	n/a
Conclusions	21	Provide a general interpretation of the results with respect to the review questions and objectives, as well as potential implications and/or next steps.	11
FUNDING			
Funding	22	Describe sources of funding for the included sources of evidence, as well as sources of funding for the scoping review. Describe the role of the funders of the scoping review.	18

JBI = Joanna Briggs Institute; PRISMA-ScR = Preferred Reporting Items for Systematic reviews and Meta-Analyses extension for Scoping Reviews.

* Where *sources of evidence* (see second footnote) are compiled from, such as bibliographic databases, social media platforms, and Web sites.

† A more inclusive/heterogeneous term used to account for the different types of evidence or data sources (e.g., quantitative and/or qualitative research, expert opinion, and policy documents) that may be eligible in a scoping review as opposed to only studies. This is not to be confused with *information sources* (see first footnote).

‡ The frameworks by Arksey and O'Malley (6) and Levac and colleagues (7) and the JBI guidance (4, 5) refer to the process of data extraction in a scoping review as data charting.

§ The process of systematically examining research evidence to assess its validity, results, and relevance before using it to inform a decision. This term is used for items 12 and 19 instead of "risk of bias" (which is more applicable to systematic reviews of interventions) to include and acknowledge the various sources of evidence that may be used in a scoping review (e.g., quantitative and/or qualitative research, expert opinion, and policy document).

From: Tricco AC, Lillie E, Zarin W, O'Brien KK, Colquhoun H, Levac D, et al. PRISMA Extension for Scoping Reviews (PRISMA-ScR): Checklist and Explanation. *Ann Intern Med.* 2018;169:467–473. doi: 10.7326/M18-0850.



Database: PubMed

Search	Query
#3	Search: #1 AND #2
#2	Search: health policy[mh] OR policy maker[tiab] OR policy makers[tiab] OR policymaker[tiab] OR policymakers[tiab] OR policy making[tiab] OR policymaking[tiab] OR decision making[mh] OR decision making[tiab] OR decisionmaking[tiab] OR decision maker[tiab] OR decision makers[tiab] OR stakeholder[tiab] OR stakeholders[tiab]
#1	Search: translational medical research[mh] OR translational medical research[tiab] OR translational medical science[tiab] OR medical translational research[tiab] OR translational medicine[tiab] OR knowledge translation[tiab] OR translational research[tiab] OR evidence-informed[tiab] OR evidence brief[tiab] OR evidence briefs[tiab] OR policy brief[tiab] OR policy briefs[tiab] OR deliberative dialogue[tiab] OR deliberative dialogues[tiab] OR knowledge synthesis[tiab] OR knowledge syntheses[tiab] OR knowledge transfer[tiab] OR research uptake[tiab] OR evidence summary[tiab] OR evidence summaries[tiab] OR knowledge uptake[tiab] OR knowledge exchange[tiab] OR research to action[tiab] OR knowledge to action[tiab]