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

SECTION	ITEM	DDICMA CAD CHECKLICT ITEM	REPORTED
SECTION	ITEM	PRISMA-ScR CHECKLIST ITEM	ON PAGE #
TITLE			
Title	1	Identify the report as a scoping review.	P1
ABSTRACT	I		
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.	P3
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.	P7-9
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.	P9

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			ON PAGE #
METHODS			
		Indicate whether a review protocol exists;	
Protocol and		state if and where it can be accessed (e.g., a	
	5	Web address); and if available, provide	P9
registration		registration information, including the	
		registration number.	
		Specify characteristics of the sources of	
Eligibility criteria	6	evidence used as eligibility criteria (e.g.,	P11-12
Englothity Citiena	0	years considered, language, and publication	P11-12
		status), and provide a rationale.	
		Describe all information sources in the	
Information		search (e.g., databases with dates of	
	7	coverage and contact with authors to	P10-11
sources*		identify additional sources), as well as the	
		date the most recent search was executed.	
		Present the full electronic search strategy	
Search	8	for at least 1 database, including any limits	P 10
		used, such that it could be repeated.	
Selection of		State the process for selecting sources of	
sources of	9	evidence (i.e., screening and eligibility)	P13
evidence†		included in the scoping review.	
Data charting	10	Describe the methods of charting data from	D12
process‡		the included sources of evidence (e.g.,	P13

CECTION	TOTAL	DDICMA C-D CHECKLICE ITEM	REPORTED
SECTION	ITEM	PRISMA-ScR CHECKLIST ITEM	ON PAGE #
		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.	
Data items	11	List and define all variables for which data were sought and any assumptions and simplifications made.	P10
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
Synthesis of results	13	Describe the methods of handling and summarizing the data that were charted.	P13
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.	P14 and Figure

SECTION	ITEM	DDICMA C.D CHECKLICT ITEM	REPORTED
SECTION		PRISMA-ScR CHECKLIST ITEM	ON PAGE #
Characteristics of sources of evidence	15	For each source of evidence, present characteristics for which data were charted and provide the citations.	P12-14 Table 3
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.	P15-17
Synthesis of results	18	Summarize and/or present the charting results as they relate to the review questions and objectives.	P15-17
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.	P18-19
Limitations	20	Discuss the limitations of the scoping review process.	P20
Conclusions	21	Provide a general interpretation of the results with respect to the review questions	P21

SECTION	ITEM	PRISMA-ScR CHECKLIST ITEM	REPORTED ON PAGE #
		and objectives, as well as potential implications and/or next steps.	
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.	P22

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 (PRISMAScR): Checklist and Explanation. Ann Intern Med. 2018;169:467–473. doi: 10.7326/M18-0850.