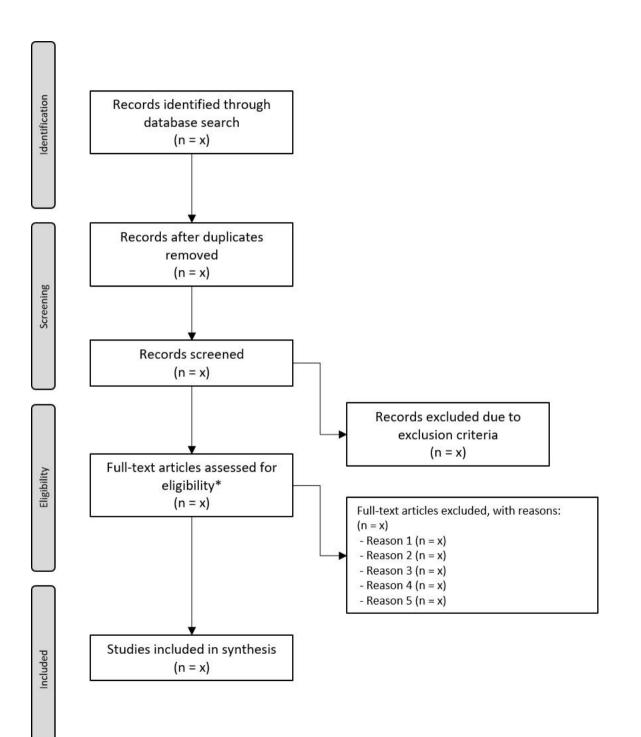
Supplementary 1: Preferred Reporting Items for Systematic reviews and Meta-Analyses extension for Scoping Reviews (PRISMA-ScR) Checklist

Section/topic	#	Checklist item	Reported on page #		
TITLE					
Title	1	Identify the report as a systematic review, meta-analysis, or both.			
ABSTRACT					
Structured summary	2	Provide a structured summary including, as applicable: background; objectives; data sources; study eligibility criteria, participants, and interventions; study appraisal and synthesis methods; results; limitations; conclusions and implications of key findings; systematic review registration number.	INSERT PAGE NUMBER		
INTRODUCTION					
Rationale	3	Describe the rationale for the review in the context of what is already known.	INSERT PAGE NUMBER		
Objectives	4	Provide an explicit statement of questions being addressed with reference to participants, interventions, comparisons, outcomes, and study design (PICOS).			
METHODS					
Protocol and registration	5	Indicate if a review protocol exists, if and where it can be accessed (e.g., Web address), and, if available, provide registration information including registration number.	INSERT PAGE NUMBER		
Eligibility criteria	6	Specify study characteristics (e.g., PICOS, length of follow-up) and report characteristics (e.g., years considered, language, publication status) used as criteria for eligibility, giving rationale.	INSERT PAGE NUMBER		
Information sources	7	Describe all information sources (e.g., databases with dates of coverage, contact with study authors to identify additional studies) in the search and date last searched.	INSERT PAGE NUMBER		
Search	8	Present full electronic search strategy for at least one database, including any limits used, such that it could be repeated.	INSERT PAGE NUMBER		
Study selection	9	State the process for selecting studies (i.e., screening, eligibility, included in systematic review, and, if applicable, included in the meta-analysis).	INSERT PAGE NUMBER		
Data collection process	10	Describe method of data extraction from reports (e.g., piloted forms, independently, in duplicate) and any processes for obtaining and confirming data from investigators.	INSERT PAGE NUMBER		

Data items	11	List and define all variables for which data were sought (e.g., PICOS, funding sources) and any assumptions and simplifications made.	INSERT PAGE NUMBER
Risk of bias in individual studies	12	Describe methods used for assessing risk of bias of individual studies (including specification of whether this was done at the study or outcome level), and how this information is to be used in any data synthesis.	INSERT PAGE NUMBER
Summary measures	13	State the principal summary measures (e.g., risk ratio, difference in means).	INSERT PAGE NUMBER
Synthesis of results	14	Describe the methods of handling data and combining results of studies, if done, including measures of consistency (e.g., I ²) for each meta-analysis.	INSERT PAGE NUMBER

Supplementary 2: PRISMA Flow Diagram template used within search methodology



Supplementary 3: Example of Data Extraction Tool

Author, Year	Title	Objective	Study type	Population (Size, Age, M/F)	Condition / Pathology	Theoretical Framework	Outcome Measures	Key Findings

Author, Year	Title	Health Knowledge	Self- management skills	Active information seeking and use	Actively communicating with health professionals	Seeking and negotiating treatment options	Decision making	Influences on health literacy	Health literacy outcomes

Item	Item	Where lo	cated **
number		Primary paper (page or appendix number)	Other † (details)
	BRIEF NAME		
1.	Provide the name or a phrase that describes the intervention.		
	WHY		_
2.	Describe any rationale, theory, or goal of the elements essential to the intervention. WHAT		
3.	Materials: Describe any physical or informational materials used in the intervention, including those provided to participants or used in intervention delivery or in training of intervention providers. Provide information on where the materials can be accessed (e.g. online appendix, URL).		
4.	Procedures: Describe each of the procedures, activities, and/or processes used in the intervention, including any enabling or support activities. WHO PROVIDED		
5.	For each category of intervention provider (e.g. psychologist, nursing assistant), describe their expertise, background and any specific training given.		
6.	HOW Describe the modes of delivery (e.g. face-to-face or by some other mechanism, such as internet or telephone) of the intervention and whether it was provided individually or in a group. WHERE		
7.	Describe the type(s) of location(s) where the intervention occurred, including any necessary infrastructure or relevant features.		

	WHEN and HOW MUCH
8.	Describe the number of times the intervention was delivered and over what period of time
	including the number of sessions, their schedule, and their duration, intensity or dose.
	TAILORING
9.	If the intervention was planned to be personalised, titrated or adapted, then describe what, why,
	when, and how.
	MODIFICATIONS
10. ‡	If the intervention was modified during the course of the study, describe the changes (what,
	why, when, and how).
	HOW WELL
11.	Planned: If intervention adherence or fidelity was assessed, describe how and by whom, and if
	any strategies were used to maintain or improve fidelity, describe them.
12. ‡	Actual: If intervention adherence or fidelity was assessed, describe the extent to which the
	intervention was delivered as planned.