Section/topic	Item No	Checklist item	Reported on Page Number/Line Number	Reported on Section/Paragraph
TITLE				
Title	1	Identify the report as a systematic review, meta-analysis, or both.	Page1	Title
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.	Page2	Abstract
INTRODUCTION				
Rationale	3	Describe the rationale for the review in the context of what is already known.	Page3-4	Background
Objectives	4	Provide an explicit statement of questions being addressed with reference to participants, interventions, comparisons, outcomes, and study design (PICOS).	Page4, Line14-18	Background, Paragraph3
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.	N/A *	N/A *
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.	Page5-6	Methods, Paragraph3-4
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.	Page5, Line1-16	Methods, Paragraph1-2
Search	8	Present full electronic search strategy for at least one database, including any limits used, such that it could be repeated.	Page23-35	Supplementary Material
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).	Page6-7	Methods, Paragraph5-6
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.	Page7, Line9-13	Methods, Paragraph7
Data items	11	List and define all variables for which data were sought (e.g., PICOS, funding sources) and any assumptions and simplifications made.	Page7, Line14-23	Methods, Paragraph8-9

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.	Page8, Line2-11	Methods, Paragraph10-11
13	State the principal summary measures (e.g., risk ratio, difference in means).	Page9	Methods, Paragraph12-13
14	Describe the methods of handling data and combining results of studies, if done, including measures of consistency (e.g., l²) for each meta-analysis.	Page8-9	Methods, Paragraph12-13
15	Specify any assessment of risk of bias that may affect the cumulative evidence (e.g., publication bias, selective reporting within studies).	N/A †	N/A †
16	Describe methods of additional analyses (e.g., sensitivity or subgroup analyses, meta-regression), if done, indicating which were pre-specified.	N/A‡	N/A ‡
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17	Give numbers of studies screened, assessed for eligibility, and included in the review, with reasons for exclusions at each stage, ideally with a flow diagram.	Page10, Line2-7	Results, Paragraph1
18	For each study, present characteristics for which data were extracted (e.g., study size, PICOS, follow-up period) and provide the citations.	Page 10, Line9-14	Results, Paragraph2
19	Present data on risk of bias of each study and, if available, any outcome level assessment (see item 12).	Page10, Line15-19	Results, Paragraph3
20	For all outcomes considered (benefits or harms), present, for each study: (a) simple summary data for each intervention group (b) effect estimates and confidence intervals, ideally with a forest plot.	Page10-12	Results, Paragraph4-7,9
21	Present results of each meta-analysis done, including confidence intervals and measures of consistency.	N/A §	N/A §
22	Present results of any assessment of risk of bias across studies (see Item 15).	N/A †	N/A †
23	Give results of additional analyses, if done (e.g., sensitivity or subgroup analyses, meta-regression [see Item 16]).	N/A‡	N/A‡
•			
24	Summarize the main findings including the strength of evidence for each main outcome; consider their relevance to key groups (e.g., healthcare providers, users, and policy makers).	Page12-13	Discussion, Paragraph1
25	Discuss limitations at study and outcome level (e.g., risk of bias), and at review-level (e.g., incomplete retrieval of identified research, reporting bias).	Page14, Line1-8	Discussion, Paragraph3
26	Provide a general interpretation of the results in the context of other evidence, and implications for future research.	Page14, Line13-19	Conclusions
	13 14 15 16 17 18 19 20 21 22 23 24 25	done at the study or outcome levelly, and how this information is to be used in any data synthesis. State the principal summary measures (e.g., risk ratio, difference in means). Describe the methods of handling data and combining results of studies, if done, including measures of consistency (e.g., if) for each meta-analysis. Specify any assessment of risk of bias that may affect the cumulative evidence (e.g., publication bias, selective reporting within studies). Describe methods of additional analyses (e.g., sensitivity or subgroup analyses, meta-regression), if done, indicating which were pre-specified. Give numbers of studies screened, assessed for eligibility, and included in the review, with reasons for exclusions at each stage, ideally with a flow diagram. For each study, present characteristics for which data were extracted (e.g., study size, PICOS, follow-up period) and provide the citations. Present data on risk of bias of each study and, if available, any outcome level assessment (see item 12). For all outcomes considered (benefits or harms), present, for each study: (a) simple summary data for each intervention group (b) effect estimates and confidence intervals, ideally with a forest plot. Present results of each meta-analysis done, including confidence intervals and measures of consistency. Present results of any assessment of risk of bias across studies (see Item 15). Give results of additional analyses, if done (e.g., sensitivity or subgroup analyses, meta-regression [see Item 16]). Summarize the main findings including the strength of evidence for each main outcome; consider their relevance to key groups (e.g., healthcare providers, users, and policy makers). Discuss limitations at study and outcome level (e.g., risk of bias), and at review-level (e.g., incomplete retrieval of identified research, reporting bias).	done at the study or outcome level), and how this information is to be used in any data synthesis. Page9 State the principal summary measures (e.g., risk ratio, difference in means). Page9 Page9 Page9 Page9 Page9 Page8-9 Page8-9 (e.g., F) for each meta-analysis. Describe the methods of handling data and combining results of studies, if done, including measures of consistency (e.g., F) for each meta-analysis. Describe methods of additional analyses (e.g., sensitivity or subgroup analyses, meta-regression), if done, indicating which were pre-specified. N/A ‡ Give numbers of studies screened, assessed for eligibility, and included in the review, with reasons for exclusions at each stage, ideally with a flow diagram. Page10, Line2-7 each study, present characteristics for which data were extracted (e.g., study size, PICOS, follow-up period) and provide the citations. Page10, Line9 14 Present data on risk of bias of each study and, if available, any outcome level assessment (see item 12). Page10, Line15-19 Present results of each meta-analysis done, including confidence intervals, ideally with a forest plot. Present results of each meta-analysis done, including confidence intervals and measures of consistency. N/A ‡ Present results of any assessment of risk of bias across studies (see Item 15). N/A † Summarize the main findings including the strength of evidence for each main outcome; consider their relevance to key groups (e.g., healthcare providers, users, and policy makers). Discuss limitations at study and outcome level (e.g., risk of bias), and at review-level (e.g., incomplete retrieval of identified research, reporting bias).

FUNDING							
Funding 2	27	Describe sources of funding for the systematic review and other support (e.g., supply of data); role of funders for the systematic review.	Page15	Funding			

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*As the checklist was provided upon initial submission, the page number/line number reported may be changed due to copyediting and may not be referable in the published version. In this case, the section/paragraph may be used as an alternative reference.

^{*} Since the outbreak of COVID-19 is a public health emergency, in order to complete the rapid review as fast as possible, this review was not registered.

 $[\]dagger$ Since meta-analysis is not available, and the number of studies included is small, this study didn't assess risk of bias across studies.

[‡] Since meta-analysis is not available in this study, there is no additional analyses.

[§] Due to the significant heterogeneity, this study only conducted qualitative synthesis instead of a meta-analysis.