

Section/topic (page no)	Item	PRISMA checklist item	Location / line in original manuscript
<b>Title</b>			
Title (3)	1	Identify the report as a systematic review, meta-analysis, or both.	title
<b>Abstract</b>			
Structured summary (4)	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.	abstract 5-33
<b>Introduction</b>			
Rationale (5)	3	Describe the rationale for the review in the context of what is already known.	48-52, 61-68, 74-78
Objectives (5)	4	Provide an explicit statement of questions being addressed with reference to participants, interventions, comparisons, outcomes, and study design (PICOS).	69-73
<b>Methods</b>			
Protocol and registration (6)	5	Indicate if a review protocol exists, if and where it can be accessed (eg, web address), and, if available, provide registration information including registration number.	156
Eligibility criteria (6)	6	Specify study characteristics (eg, PICOS, length of follow-up) and report characteristics (eg, years considered, language, publication status) used as criteria for eligibility, giving rationale.	81-92, Table 2
Information sources (7)	7	Describe all information sources (eg, databases with dates of coverage, contact with study authors to identify additional studies) in the search and date last searched.	77-79, 102-103
Search (7)	8	Present full electronic search strategy for at least one database, including any limits used, such that it could be repeated.	Table 1
Study selection (8)	9	State the process for selecting studies (ie, screening, eligibility, included in systematic review, and, if applicable, included in the meta-analysis).	84-92
Data collection process (9)	10	Describe method of data extraction from reports (eg, piloted forms, independently, in duplicate) and any processes for obtaining and confirming data from investigators.	94-103
Data items (9)	11	List and define all variables for which data were sought (eg, PICOS, funding sources) and any assumptions and simplifications made.	116-118
Risk of bias in individual studies (10)	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.	153-158
Summary measures (11)	13	State the principal summary measures (eg, risk ratio, difference in means).	116-119
Synthesis of results (11)	14	Describe the methods of handling data and combining results of studies, if done, including measures of consistency (eg, $I^2$ ) for each meta-analysis.	121-151
Risk of bias across studies (11)	15	Specify any assessment of risk of bias that may affect the cumulative evidence (eg, publication bias, selective reporting within studies).	159-160

Additional analyses (12)	16	Describe methods of additional analyses (eg, sensitivity or subgroup analyses, meta-regression), if done, indicating which were prespecified.	130-132
<b>Results</b>			
Study selection (13)	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	167-177 Fig.1
Study characteristics (14)	18	For each study, present characteristics for which data were extracted (eg, study size, PICOS, follow-up period) and provide the citations.	186-206, Table 5, table6, table 8 (supplement ary), table 10 (supplement ary)
Risk of bias within studies (15)	19	Present data on risk of bias of each study and, if available, any outcome level assessment (see item 12).	242-248 Table 9
Results of individual studies (16)	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.	224-237, Table 6, table 5, table 8(suppl), table 10 (suppl) figures 2-4,
Synthesis of results (17)	21	Present results of each meta-analysis done, including confidence intervals and measures of consistency.	Table 6, fig.2, fig 3, fig 4
Risk of bias across studies (18)	22	Present results of any assessment of risk of bias across studies (see item 15).	249-252 Table 6
Additional analysis (18)	23	Give results of additional analyses, if done (eg, sensitivity or subgroup analyses, meta-regression (see item 16)).	214-217  Table 6
<b>Discussion</b>			
Summary of evidence (18)	24	Summarise the main findings including the strength of evidence for each main outcome; consider their relevance to key groups (eg, healthcare providers, users, and policy makers).	255-302
Limitations (18)	25	Discuss limitations at study and outcome level (eg, risk of bias), and at review level (eg, incomplete retrieval of identified research, reporting bias).	304-362
Conclusions (19)	26	Provide a general interpretation of the results in the context of other evidence, and implications for future research.	365-370
<b>Funding</b>			
Funding (19)	27	Describe sources of funding for the systematic review and other support (eg, supply of data); role of funders for the systematic review.	On title page (only available to GSJ)