


Table S1. PRISMA Checklist.

Section and Topic	Item #	Checklist item	Location where item is reported
TITLE			
Title	1	Identify the report as a systematic review.	1
ABSTRACT			
Abstract	2	See the PRISMA 2020 for Abstracts checklist.	1
INTRODUCTION			
Rationale	3	Describe the rationale for the review in the context of existing knowledge.	2
Objectives	4	Provide an explicit statement of the objective(s) or question(s) the review addresses.	2
METHODS			
Eligibility criteria	5	Specify the inclusion and exclusion criteria for the review and how studies were grouped for the syntheses.	2
Information sources	6	Specify all databases, registers, websites, organisations, reference lists and other sources searched or consulted to identify studies. Specify the date when each source was last searched or consulted.	2
Search strategy	7	Present the full search strategies for all databases, registers and websites, including any filters and limits used.	2
Selection process	8	Specify the methods used to decide whether a study met the inclusion criteria of the review, including how many reviewers screened each record and each report retrieved, whether they worked independently, and if applicable, details of automation tools used in the process.	2
Data collection process	9	Specify the methods used to collect data from reports, including how many reviewers collected data from each report, whether they worked independently, any processes for obtaining or confirming data from study investigators, and if applicable, details of automation tools used in the process.	3
Data items	10a	List and define all outcomes for which data were sought. Specify whether all results that were compatible with each outcome domain in each study were sought (e.g. for all measures, time points, analyses), and if not, the methods used to decide which results to collect.	3
	10b	List and define all other variables for which data were sought (e.g. participant and intervention characteristics, funding sources). Describe any assumptions made about any missing or unclear information.	3
Study risk of bias assessment	11	Specify the methods used to assess risk of bias in the included studies, including details of the tool(s) used, how many reviewers assessed each study and whether they worked independently, and if applicable, details of automation tools used in the process.	3
Effect measures	12	Specify for each outcome the effect measure(s) (e.g. risk ratio, mean difference) used in the synthesis or presentation of results.	3
Synthesis methods	13a	Describe the processes used to decide which studies were eligible for each synthesis (e.g. tabulating the study intervention characteristics and comparing against the planned groups for each synthesis (item #5)).	Table 1
	13b	Describe any methods required to prepare the data for presentation or synthesis, such as handling of missing summary statistics, or data conversions.	3
	13c	Describe any methods used to tabulate or visually display results of individual studies and syntheses.	3
	13d	Describe any methods used to synthesize results and provide a rationale for the choice(s). If meta-analysis was performed, describe the model(s), method(s) to identify the presence and extent of statistical heterogeneity, and software package(s) used.	-
	13e	Describe any methods used to explore possible causes of heterogeneity among study results (e.g. subgroup analysis, meta-regression).	-
	13f	Describe any sensitivity analyses conducted to assess robustness of the synthesized results.	-
Reporting bias assessment	14	Describe any methods used to assess risk of bias due to missing results in a synthesis (arising from reporting biases).	3
Certainty assessment	15	Describe any methods used to assess certainty (or confidence) in the body of evidence for an outcome.	-

Section and Topic	Item #	Checklist item	Location where item is reported
RESULTS			
Study selection	16a	Describe the results of the search and selection process, from the number of records identified in the search to the number of studies included in the review, ideally using a flow diagram.	4 (Fig. 1)
	16b	Cite studies that might appear to meet the inclusion criteria, but which were excluded, and explain why they were excluded.	3
Study characteristics	17	Cite each included study and present its characteristics.	5
Risk of bias in studies	18	Present assessments of risk of bias for each included study.	Table 2
Results of individual studies	19	For all outcomes, present, for each study: (a) summary statistics for each group (where appropriate) and (b) an effect estimate and its precision (e.g. confidence/credible interval), ideally using structured tables or plots.	Table 3-6, Fig. 2
Results of syntheses	20a	For each synthesis, briefly summarise the characteristics and risk of bias among contributing studies.	Table 2
	20b	Present results of all statistical syntheses conducted. If meta-analysis was done, present for each the summary estimate and its precision (e.g. confidence/credible interval) and measures of statistical heterogeneity. If comparing groups, describe the direction of the effect.	-
	20c	Present results of all investigations of possible causes of heterogeneity among study results.	-
	20d	Present results of all sensitivity analyses conducted to assess the robustness of the synthesized results.	-
Reporting biases	21	Present assessments of risk of bias due to missing results (arising from reporting biases) for each synthesis assessed.	-
Certainty of evidence	22	Present assessments of certainty (or confidence) in the body of evidence for each outcome assessed.	-
DISCUSSION			
Discussion	23a	Provide a general interpretation of the results in the context of other evidence.	8
	23b	Discuss any limitations of the evidence included in the review.	9
	23c	Discuss any limitations of the review processes used.	9
	23d	Discuss implications of the results for practice, policy, and future research.	9
OTHER INFORMATION			
Registration and protocol	24a	Provide registration information for the review, including register name and registration number, or state that the review was not registered.	2
	24b	Indicate where the review protocol can be accessed, or state that a protocol was not prepared.	2
	24c	Describe and explain any amendments to information provided at registration or in the protocol.	-
Support	25	Describe sources of financial or non-financial support for the review, and the role of the funders or sponsors in the review.	9
Competing interests	26	Declare any competing interests of review authors.	9
Availability of data, code and other materials	27	Report which of the following are publicly available and where they can be found: template data collection forms; data extracted from included studies; data used for all analyses; analytic code; any other materials used in the review.	9

Table S2. Risk of bias checklist.

		Methodology Checklist 2: Controlled Trials	
Author: Camidge et al.			
Reviewer: Kang and Woo		Date: 2022/03/02	No: 1
<p>Before completing this checklist, consider:</p> <ol style="list-style-type: none"> Is the paper a randomised controlled trial or a controlled clinical trial? If in doubt, check the study design algorithm available from SIGN and make sure you have the correct checklist. If it is a controlled clinical trial questions 1.2, 1.3, and 1.4 are not relevant, and the study cannot be rated higher than 1+ Is the paper relevant to key question? Analyse using PICO (Patient or Population Intervention Comparison Outcome). IF NO REJECT (give reason below). IF YES complete the checklist. 			
Reason for rejection: 1. Paper not relevant to key question <input type="checkbox"/> 2. Other reason <input type="checkbox"/> (please specify):			
SECTION 1: INTERNAL VALIDITY			
<i>In a well conducted RCT study...</i>		<i>Does this study do it?</i>	
1.1	The study addresses an appropriate and clearly focused question.	Yes <input checked="" type="checkbox"/>	No <input type="checkbox"/> Can't say <input type="checkbox"/>
1.2	The assignment of subjects to treatment groups is randomised.	Yes <input checked="" type="checkbox"/>	No <input type="checkbox"/> Can't say <input type="checkbox"/>
1.3	An adequate concealment method is used.	Yes <input checked="" type="checkbox"/>	No <input type="checkbox"/> Can't say <input type="checkbox"/>
1.4	The design keeps subjects and investigators 'blind' about treatment allocation.	Yes <input checked="" type="checkbox"/>	No <input type="checkbox"/> Can't say <input type="checkbox"/>
1.5	The treatment and control groups are similar at the start of the trial.	Yes <input checked="" type="checkbox"/>	No <input type="checkbox"/> Can't say <input type="checkbox"/>
1.6	The only difference between groups is the treatment under investigation.	Yes <input checked="" type="checkbox"/>	No <input type="checkbox"/> Can't say <input type="checkbox"/>
1.7	All relevant outcomes are measured in a standard, valid and reliable way.	Yes <input type="checkbox"/> Can't say <input type="checkbox"/>	No <input checked="" type="checkbox"/>
1.8	What percentage of the individuals or clusters recruited into each treatment arm of the study dropped out before the study was completed?	0%	
1.9	All the subjects are analysed in the groups to which they were randomly allocated (often referred to as intention to treat analysis).	Yes <input checked="" type="checkbox"/> Can't say <input type="checkbox"/>	No <input type="checkbox"/> Does not apply <input type="checkbox"/>

1.10	Where the study is carried out at more than one site, results are comparable for all sites.	Yes <input type="checkbox"/> No <input type="checkbox"/> Can't say <input checked="" type="checkbox"/> Does not apply <input type="checkbox"/>
SECTION 2: OVERALL ASSESSMENT OF THE STUDY		
2.1	How well was the study done to minimise bias? <i>Code as follows:</i>	High quality (++) <input type="checkbox"/> Acceptable (+) <input checked="" type="checkbox"/> Low quality (-) <input type="checkbox"/> Unacceptable – reject 0 <input type="checkbox"/>
2.2	Taking into account clinical considerations, your evaluation of the methodology used, and the statistical power of the study, are you certain that the overall effect is due to the study intervention?	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/>
2.3	Are the results of this study directly applicable to the patient group targeted by this guideline?	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/>
2.4	Notes. Summarise the authors' conclusions. Add any comments on your own assessment of the study, and the extent to which it answers your question and mention any areas of uncertainty raised above.	



Methodology Checklist 2: Controlled Trials

Author: Hida et al.

Reviewer: Kang and Woo

Date: 2022/03/02

No: 2

Before completing this checklist, consider:

1. Is the paper a **randomised controlled trial** or a **controlled clinical trial**? If in doubt, check the study design algorithm available from SIGN and make sure you have the correct checklist. If it is a **controlled clinical trial** questions 1.2, 1.3, and 1.4 are not relevant, and the study cannot be rated higher than 1+
2. Is the paper relevant to key question? Analyse using PICO (Patient or Population Intervention Comparison Outcome). IF NO REJECT (give reason below). IF YES complete the checklist.

Reason for rejection: 1. Paper not relevant to key question ☐ 2. Other reason ☐ (please specify):

SECTION 1: INTERNAL VALIDITY

<i>In a well conducted RCT study...</i>		<i>Does this study do it?</i>
1.1	The study addresses an appropriate and clearly focused question.	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/>
1.2	The assignment of subjects to treatment groups is randomised.	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/>
1.3	An adequate concealment method is used.	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/>
1.4	The design keeps subjects and investigators 'blind' about treatment allocation.	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/>
1.5	The treatment and control groups are similar at the start of the trial.	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/>
1.6	The only difference between groups is the treatment under investigation.	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/>
1.7	All relevant outcomes are measured in a standard, valid and reliable way.	Yes <input type="checkbox"/> No <input checked="" type="checkbox"/> Can't say <input type="checkbox"/>
1.8	What percentage of the individuals or clusters recruited into each treatment arm of the study dropped out before the study was completed?	0%
1.9	All the subjects are analysed in the groups to which they were randomly allocated (often referred to as intention to treat analysis).	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/> Does not apply <input type="checkbox"/>

1.10	Where the study is carried out at more than one site, results are comparable for all sites.	Yes <input type="checkbox"/> No <input type="checkbox"/> Can't say <input checked="" type="checkbox"/> Does not apply <input type="checkbox"/>
SECTION 2: OVERALL ASSESSMENT OF THE STUDY		
2.1	How well was the study done to minimise bias? <i>Code as follows:</i>	High quality (++) <input type="checkbox"/> Acceptable (+) <input checked="" type="checkbox"/> Low quality (-) <input type="checkbox"/> Unacceptable – reject 0 <input type="checkbox"/>
2.2	Taking into account clinical considerations, your evaluation of the methodology used, and the statistical power of the study, are you certain that the overall effect is due to the study intervention?	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/>
2.3	Are the results of this study directly applicable to the patient group targeted by this guideline?	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/>
2.4	Notes. Summarise the authors' conclusions. Add any comments on your own assessment of the study, and the extent to which it answers your question and mention any areas of uncertainty raised above.	



Methodology Checklist 2: Controlled Trials

Author: Mok et al.

Reviewer: Kang and Woo

Date: 2022/03/02

No: 3

Before completing this checklist, consider:

1. Is the paper a **randomised controlled trial** or a **controlled clinical trial**? If in doubt, check the study design algorithm available from SIGN and make sure you have the correct checklist. If it is a **controlled clinical trial** questions 1.2, 1.3, and 1.4 are not relevant, and the study cannot be rated higher than 1+
2. Is the paper relevant to key question? Analyse using PICO (Patient or Population Intervention Comparison Outcome). IF NO REJECT (give reason below). IF YES complete the checklist.

Reason for rejection: 1. Paper not relevant to key question ☐ 2. Other reason ☐ (please specify):

SECTION 1: INTERNAL VALIDITY

<i>In a well conducted RCT study...</i>		<i>Does this study do it?</i>
1.1	The study addresses an appropriate and clearly focused question.	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/>
1.2	The assignment of subjects to treatment groups is randomised.	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/>
1.3	An adequate concealment method is used.	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/>
1.4	The design keeps subjects and investigators 'blind' about treatment allocation.	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/>
1.5	The treatment and control groups are similar at the start of the trial.	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/>
1.6	The only difference between groups is the treatment under investigation.	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/>
1.7	All relevant outcomes are measured in a standard, valid and reliable way.	Yes <input type="checkbox"/> No <input checked="" type="checkbox"/> Can't say <input type="checkbox"/>
1.8	What percentage of the individuals or clusters recruited into each treatment arm of the study dropped out before the study was completed?	0%
1.9	All the subjects are analysed in the groups to which they were randomly allocated (often referred to as intention to treat analysis).	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/> Does not apply <input type="checkbox"/>

1.10	Where the study is carried out at more than one site, results are comparable for all sites.	Yes <input type="checkbox"/> Can't say <input checked="" type="checkbox"/>	No <input type="checkbox"/> Does not apply <input type="checkbox"/>
SECTION 2: OVERALL ASSESSMENT OF THE STUDY			
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2.4	Notes. Summarise the authors' conclusions. Add any comments on your own assessment of the study, and the extent to which it answers your question and mention any areas of uncertainty raised above.		



Methodology Checklist 2: Controlled Trials

Author: Novello et al.

Reviewer: Kang and Woo

Date: 2022/03/02

No: 4

Before completing this checklist, consider:

1. Is the paper a **randomised controlled trial** or a **controlled clinical trial**? If in doubt, check the study design algorithm available from SIGN and make sure you have the correct checklist. If it is a **controlled clinical trial** questions 1.2, 1.3, and 1.4 are not relevant, and the study cannot be rated higher than 1+
2. Is the paper relevant to key question? Analyse using PICO (Patient or Population Intervention Comparison Outcome). IF NO REJECT (give reason below). IF YES complete the checklist.

Reason for rejection: 1. Paper not relevant to key question ☐ 2. Other reason ☐ (please specify):

SECTION 1: INTERNAL VALIDITY

<i>In a well conducted RCT study...</i>		<i>Does this study do it?</i>
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1.2	The assignment of subjects to treatment groups is randomised.	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/>
1.3	An adequate concealment method is used.	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/>
1.4	The design keeps subjects and investigators 'blind' about treatment allocation.	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/>
1.5	The treatment and control groups are similar at the start of the trial.	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/>
1.6	The only difference between groups is the treatment under investigation.	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/>
1.7	All relevant outcomes are measured in a standard, valid and reliable way.	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/>
1.8	What percentage of the individuals or clusters recruited into each treatment arm of the study dropped out before the study was completed?	0%
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1.10	Where the study is carried out at more than one site, results are comparable for all sites.	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/> Does not apply <input type="checkbox"/>
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Methodology Checklist 2: Controlled Trials

Author: Peters et al.

Reviewer: Kang and Woo

Date: 2022/03/02

No: 5

Before completing this checklist, consider:

1. Is the paper a **randomised controlled trial** or a **controlled clinical trial**? If in doubt, check the study design algorithm available from SIGN and make sure you have the correct checklist. If it is a **controlled clinical trial** questions 1.2, 1.3, and 1.4 are not relevant, and the study cannot be rated higher than 1+
2. Is the paper relevant to key question? Analyse using PICO (Patient or Population Intervention Comparison Outcome). IF NO REJECT (give reason below). IF YES complete the checklist.

Reason for rejection: 1. Paper not relevant to key question ☐ 2. Other reason ☐ (please specify):

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1.3	An adequate concealment method is used.	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/>
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2.4	Notes. Summarise the authors' conclusions. Add any comments on your own assessment of the study, and the extent to which it answers your question and mention any areas of uncertainty raised above.	



Methodology Checklist 2: Controlled Trials

Author: Zhou et al.

Reviewer: Kang and Woo

Date: 2022/03/02

No: 6

Before completing this checklist, consider:

1. Is the paper a **randomised controlled trial** or a **controlled clinical trial**? If in doubt, check the study design algorithm available from SIGN and make sure you have the correct checklist. If it is a **controlled clinical trial** questions 1.2, 1.3, and 1.4 are not relevant, and the study cannot be rated higher than 1+
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1.3	An adequate concealment method is used.	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Can't say <input type="checkbox"/>
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