

Methodological Expectations of  
Cochrane Intervention Reviews (MECIR)  
Standards for the conduct and reporting of  
new Cochrane Intervention Reviews,  
reporting of protocols and the planning,  
conduct and reporting of updates

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Version February 2021



## Cochrane Methods ‘What are MECIR Standards?’

URL: <https://methods.cochrane.org/methodological-expectations-cochrane-intervention-reviews>

### Methodological Expectations of Cochrane Intervention Reviews

The Methodological Expectations of Cochrane Intervention Reviews (known as MECIR Standards) are available [online](#). The online version includes links to the *Cochrane Handbook for Systematic Reviews of Interventions*, Cochrane Training and other Cochrane resources to provide additional explanation of how to implement the standard. The online version is kept up-to-date with amendments listed [here](#).

### Standards for Cochrane Reviews of interventions

The MECIR Standards are methodological standards to which all Cochrane Protocols, Reviews, and Updates are expected to adhere. They are divided into four sections:

1. Standards for the conduct of new Cochrane Intervention Reviews (C1-C75).
2. Standards for reporting of protocols of new Cochrane Intervention Reviews (PR1-PR44).
3. Standards for reporting of new Cochrane Intervention Reviews (R1-R109).
4. Standards for planning, conducting and reporting of updates of Cochrane Intervention Reviews (U1-U11, UR1-UR7).

These expectations are intended for both internal and external audiences. They provide authors and users of the Cochrane Library with clear and transparent expectations of review conduct and reporting.

### Implementation

The MECIR Standards have been integrated into the following Cochrane systems:

- The RevMan guidance panel.
- Editorial checklists.
- *The Cochrane Handbook for Systematic Reviews of Interventions*.

### Other key resources

- Introducing new MECIR Standards for trainers ([introductory videos via Cochrane Training](#)).
- [Version and changes to MECIR](#) - details on changes and developments to the MECIR Standards since 2016.
- *Cochrane Handbook for Systematic Reviews of Interventions*.
- [Cochrane Interaction Learning e-learning modules: Conducting an Intervention Review](#)

### Standards for the reporting of plain language summaries of new Reviews of interventions

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Plain language summaries fulfil an important function in Cochrane Reviews. Aimed at a broad readership they convey the review question and the findings in terms that are accessible to consumers and non-expert readers. [The standards](#) presented here were developed by a committee led by the Consumer Coordinator. They were finalized based on comments received from a consultation process that involved stakeholders who were internal and external to Cochrane.

URL: <https://community.cochrane.org/mecir-manual>

## Methodological Expectations of Cochrane Intervention Reviews (MECIR)

[Version February 2021](#)

Standards for the conduct and reporting of new Cochrane Intervention Reviews, reporting of protocols and the planning, conduct and reporting of updates

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These MECIR Standards present a guide to the conduct and reporting of Cochrane Intervention Reviews. Each set of Standards includes links to Cochrane Training resources, the *Cochrane Handbook for Systematic Reviews of Interventions* (the *Handbook*) and other available resources.

This online version will be kept up to date. A PDF of each section can be generated. All substantive changes will be noted [here](#).

- If available, MECIR Standards link to the most up-to-date version of the *Handbook* chapters.
- Where links to external resources are included [Cochrane Interactive Learning](#) is referred to as 'CIL'.
- We welcome your feedback on MECIR, or if you have any general queries related to the MECIR Standards, please contact [Cochrane Methods](#).

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## Key points and introduction

Key points:

- The MECIR Standards represent a true collaborative effort across our community.
- **They are an essential part of Cochrane's quality assurance strategy.**
- The MECIR Standards represent a living programme of work, and will be adapted over time as methods, and expectations change.

Ensuring that Cochrane Reviews represent the highest possible quality is critical if they are to inform decision making in clinical practice and health policy. Methodological Expectations of Cochrane Intervention Reviews (MECIR) are Standards that should guide the conduct and reporting of Cochrane Intervention Reviews. They are drawn from the *Cochrane Handbook for Systematic Reviews of Interventions* (the '*Handbook*'). The development of the Standards has been a collaborative effort over several years, involving review authors, editors and methodologists from all corners of our community. In this document we present a complete set of Standards for intervention reviews.



## Development and consultation

We established working groups in 2011 to develop minimum standards based on early proposals and groundwork by many groups and individuals within Cochrane. We agreed the need to identify methodological expectations for Cochrane protocols, reviews and updates of reviews on the effects of interventions that could be implemented across Cochrane. Six Working Groups covered six core methodological aspects of Cochrane Intervention Reviews:

- developing a question and deciding the scope of the review,
- searching for studies,
- selecting studies and collecting data,
- assessing risk of bias in studies,
- analysing data and undertaking meta-analyses,
- interpretation and presenting results.

For each of these areas, we set out to identify the following in respect of intervention reviews:

- A. essential minimum standards (*must do*);
- B. desirable standards (*should do*);
- C. common errors (*should not do*);
- D. fatal flaws (*must not do*) and identification of any important methodological uncertainties.

The existing Standards address A and B. At least one methodologist and one Co-ordinating Editor (clinical specialist) jointly led each working group. We sought to ensure that groups reflected divergent views and had access to appropriate expertise. We co-opted other people from across Cochrane as necessary to ensure co-ordination and consistency of approach (training and knowledge translation). From an initial draft set of Standards based primarily on the 2011 version of the *Handbook*, we consulted widely throughout Cochrane, after which the MECIR co-ordinating author team collated responses to produce the full original set of Standards.

We have updated the standards regularly since their first publication. They now reflect the guidance available in the most up-to-date publicly available version of the *Handbook*.

## Implementation of the Standards

The Methodological Expectations for Cochrane Intervention Review (MECIR) are the Standards that each Cochrane Intervention Review should meet. Review authors and Cochrane Review Groups are expected to adhere or oversee adherence to these Standards across different stages of the review process: protocols, reviews and updates.

All Standards are qualified with the status of ‘mandatory’ or ‘highly desirable’. Mandatory Standards should always be met unless an appropriate justification for not doing so can be provided. Highly desirable Standards should generally be implemented but justification for not implementing them is unnecessary. We introduce each set of Standards with key points and where necessary additional explanatory notes. The MECIR conduct Standards ([C1-C75](#)) are included in the [Cochrane Handbook for Systematic Reviews of Interventions](#).

Since the MECIR Standards were launched in 2011, technology has developed and changed how reviews are being produced. The development of web-based platforms such as Covidence, EPPI-Reviewer, and GRADEpro GDT, as well as tools supporting semi-automation, have changed the way that systematic reviews are produced. Whilst we can expect technology to develop and help improve efficiency in production of Cochrane Reviews, these Standards remain a fundamental element of the preparation and quality assurance of individual Cochrane Intervention Reviews.

The MECIR Standards represent a considerable amount of work from many people within the Cochrane community. The core team of Julian Higgins, Rachel Churchill, Toby Lasserson, my predecessor, David Tovey, and Jackie Chandler have made substantial contributions to the process. I am delighted to welcome James Thomas and Ella Flemyng to an expanded team of authors to coincide with the launch of version 6 of the *Handbook*.

We continue to welcome feedback from all of you who are responsible for delivering the Standards, and hope that they are useful to you in producing and maintaining high quality, relevant reviews that can guide decision makers throughout the world, in pursuit of better health.

Karla Soares-Weiser  
Editor in Chief  
The Cochrane Library

## Acknowledgements

We thank the following working group leads and contributors for their early development of the Standards: Doug Altman, Mohammed Ansari (Methods lead), Sally Bell-Syer, Patrick Bossuyt, Deborah Caldwell, Christopher Cates, Rachel Churchill (Co-ordinating Editors (Co-Eds) lead, Co-ordinating team), Mike Clarke (Co-Eds co-lead), Jan Clarkson (Co-Eds co-lead), Philippa Davies, Marina Davoli (Co-Eds lead), Ruth Foxlee, Chantelle Garritty, Davina Ghera (Co-Eds co-lead), Julie Glanville (Methods co-lead), Peter Herbison, Julian Higgins (Co-ordinating team), Sophie Hill (Co-Eds lead), Toby Lasserson (Co-ordinating team), Edith Leclercq, Carol Lefebvre (Methods co-lead), Jessie McGowan, Rachel Marshall, **Ruth Mitchell, Donal O'Mathuna, Anna Noel-Storr**, Georgia Salanti (Methods lead), Doug Salzwedel, Margaret Sampson, Jelena Savovic, Holger Schünemann (Methods lead), Ian Shemilt, Nandi Siegfried Jonathan Sterne (Methods lead), Britta Tendal (Methods lead), David Tovey (Co-ordinating team), Peter Tugwell, Lucy Turner, Claire Vale, Julia Walters, Helen Worthington (Co-Eds lead), and Janelle Yorke. We also thank all those Cochrane members of Review Groups, Methods Groups, Fields, Centres and Training who responded in some detail to MECIR Standards consultations, allowing us to improve these Standards to ensure relevance and comprehension.

URL: <https://community.cochrane.org/mecir-manual/introduction-key-points/versions-and-changes-mecir>

## Versions and changes to MECIR

### Process for updating MECIR

- For details on when and how updates to MECIR are made, please see [here](#).

### Updates pending for the next version (2022)

No updates pending.

### Version February 2021 ([click here for the PDF version](#))

- [Version March 2020](#) -changed to- Version February 2021
- [C56: Highly desirable](#) -changed to- [C56: Mandatory](#)
- [R106: 'Declarations of interest'](#), updated to reflect Cochrane's new Conflict of interest policy.
- See *Handbook* (Version 6) Section xxx – changed to- See *Handbook* Section xxx

### Version March 2020 ([click here for the PDF version](#))

- [Version October 2019](#) – changed to- [Version March 2020](#)
- During February and March 2020 edits were made to the PR, R, U and UR Standards in MECIR to update referencing to the new *Handbook* (version 6). All changes are reflected at the bottom of each page.
- PR14: Define in advance which outcomes are primary outcomes and which are secondary outcomes. -changed to- Define in advance outcomes that are critical to the review, and any additional important outcomes.
- PR27: Assess the risk of bias for each included study. For randomized trials, the Cochrane 'Risk of bias' tool should be used, involving judgements and supports for those judgements across a series of domains of bias, as described in Chapter 8 of the *Handbook* (version 5 or later). -changed to-  
Assess the risk of bias in at least one specific result for each included study. For randomized trials, the RoB 2 tool should be used, involving judgements and support for those judgements across a series of domains of bias, as described in *Handbook* (version 6).
- PR28: If the Risk of Bias 2 tool (see *Handbook* Chapter 8) is to be used, state whether interest will be in the effect of assignment to intervention or the effect of adhering to intervention, and explain how results will be selected to be assessed for risk of bias (i.e. for which outcome domains, outcome measures, time points and analyses). ADDED
- PR35: according to summary risk of bias, or restricted to studies at low risk of bias. -changed to- according to summary risk of bias, restricted to studies at low risk of bias or restricted to low- and-some-concerns of risk of bias.
- R32: Define in advance which outcomes are primary outcomes and which are secondary outcomes. -changed to- Define in advance outcomes that are critical to the review, and any additional important outcomes.
- R45: Assess the risk of bias for each included study. For randomized trials, the Cochrane 'Risk of bias' tool should be used, involving judgements and supports for those judgements across a series of domains of bias, as described in Chapter 8 of the *Handbook* (version 5 or later). - changed to- Assess the risk of bias in at least one specific result for each included study. For

randomized trials, the RoB 2 tool should be used, involving judgements and support for those judgements across a series of domains of bias, as described in Handbook version 6.

- R53: according to summary risk of bias, or restricted to studies at low risk of bias. -changed to- according to summary risk of bias, restricted to studies at low risk of bias or restricted to low-and-some-concerns of risk of bias.
- **R55: (Include a ‘Summary of Findings’ table according to recommendations described in Chapter 10 of the Cochrane Handbook (version 5 or later). Specifically:**  
include results for one population group (with few exceptions);  
indicate the intervention and the comparison intervention;  
include seven or fewer patient-important outcomes;  
describe the outcomes (e.g. scale, scores, follow-up);  
indicate the number of participants and studies for each outcome;  
present at least one baseline risk for each dichotomous outcome (e.g. study population or median/medium risk) and baseline scores for continuous outcomes (if appropriate);  
summarize the intervention effect (if appropriate); and  
include a measure of the certainty of the body of evidence)  
-changed to-  
Justify and document all assessments of the certainty of the body of evidence (for example downgrading or upgrading if using GRADE).
- R55: MECIR conduct standard 76 (Use the five GRADE considerations (study limitations, consistency of effect, imprecision, indirectness and publication bias) to assess the certainty of the body of evidence for each outcome, and to draw conclusions about the certainty of evidence within the text of the review.) [PRISMA item 12]  
- changed to-  
MECIR conduct standard 74: Use the five GRADE considerations (risk of bias, consistency of effect, imprecision, indirectness and publication bias) to assess the certainty of the body of evidence for each outcome, and to draw conclusions about the certainty of evidence within the text of the review.
- R56: to complete a PRISMA type flow chart -changed to- to be able to complete a flow diagram
- **R73: Present a ‘Risk of bias’ table for each included study -changed to- Present at least one ‘Risk of bias’ table for each study that is included in a synthesis**
- **R73: The ‘Risk of bias’ table in RevMan should be used, this is an extension of the table of ‘Characteristics of included studies’.** -changed to- **‘Risk of bias’ presentation tools in RevMan should be used wherever possible.**
- R73: Assess the risk of bias for each included study. For randomized trials, the Cochrane 'Risk of bias' tool should be used, involving judgements and supports for those judgements across a series of domains of bias, as described in Chapter 8 of the *Handbook* (version 5 or later) -changed to- Assess the risk of bias in at least one specific result for each included study. For randomized trials, the RoB 2 tool should be used, involving judgements and support for those judgements across a series of domains of bias, as described in *Handbook* (version 6).
- R74: Summarize the risk of bias -changed to- Present an overall risk of bias assessment
- R76: the heading hierarchy -changed to- any heading hierarchy
- R76: in RevMan5 ADDED
- R76: This standard will not be required when using the study-centric data structure of RevMan Web. ADDED
- R101: Consider the potential impact of reporting biases -changed to- Consider the potential impact of non-reporting biases
- U9: For randomized trials, they must be assessed using a currently accepted version of the **Cochrane ‘Risk of bias’ tool. The separation of performance bias and detection bias in the evaluation of blinding is highly desirable.** -changed to- If the previous version used the original risk of bias tool to assess randomised trials, consider whether or not to switch to the Risk of Bias

2 tool (see *Handbook* Chapter 8), including how many randomised trials were assessed in the previous version, how many new studies are expected for inclusion in the update, how well it was implemented in the previous version and whether it is feasible to switch.

#### Version October 2019

- Version July 2019 - changed to- Version October 2019
- Updates made to MECIR authors' affiliations
- Links to version 6 of the *Cochrane Handbook for Systematic Reviews of Interventions* added to all relevant standards (Conduct Standards C1-C75)
- Links to the Cochrane Editorial and Publishing Policy Resource updated
- James Thomas and Ella Flemyng added as co-authors
- Edits made to the MECIR Standards '[Key points and introduction](#)' page (see 'Section info' on the page for details).
- Edits made to the '[Development and consultation](#)' page (see 'Section info' on the page for details)
- New '[Implementation of the standards](#)' section written by Karla Soares-Weiser (see 'Section info' on the page for details)
- Edits made to the 'Key points and introduction' pages for each of the four sections (see 'Section info' on the [conduct](#), [reporting of protocols](#), [reporting](#) and [updates](#) pages for details)
- Added a new '[Translations of the MECIR Standards](#)' section
- Citation to the MECIR Manual as a whole and each section updated to reflect version October 2019
- [U11](#), column 2: quality -changed to- certainty (x2)
- [UR5](#), column 3: quality -changed to- certainty
- [UR7](#), column 3: quality -changed to- certainty
- [PR39](#) column 2 and 3: quality -changed to- certainty (x4)
- [PR40](#) column 3: quality -changed to- certainty
- [R12](#), column 3: quality -changed to- certainty
- [R55](#): column 2 and 3: quality -changed to- certainty (x4)
- [R96](#): column 3: quality -changed to- certainty
- [R98](#): column 3: quality -changed to- certainty (x2)
- [R99](#) column 2 and 3: quality -changed to- certainty (x5)
- [R100](#), column 3: Quality -changed to- Certainty

#### Version July 2019

- v1.06 2018 - changed to- Version July 2019
- Previous pages titled 'Latest substantive changes' and 'Versions' have been merged into one page titled 'Versions and changes to MECIR'
- Citation to the MECIR Manual as a whole and each section updated to reflect version July 2019
- [C1](#): See Handbook 2.3.2, 2.3.4, 17.2, 20.2.2 -changed to- See Handbook (version 6), Section 2.1
- [C2](#): See Handbook 5.1.1 -changed to- See Handbook (version 6), Section 2.3
- [C3](#): See Handbook 5.4.3, 14.1.1, 14.3 -changed to- See Handbook Section 2.1
- [C4](#): added: See Handbook (version 6), Section 2.4
- [C5](#): Handbook 5.2 -changed to- Handbook (version 6), Section 3.2.1
- [C6](#): Handbook 5.2 -changed to- Handbook (version 6), Section 3.2.1
- [C7](#): Handbook 5.3 -changed to- Handbook (version 6), Section 3.2.2
- [C8](#): Handbook 5.1.2 -changed to- Handbook (version 6), Section 3.2.4.1
- [C9](#): Handbook 5.5, 13.2.2 -changed to- Handbook (version 6), Section 3.3
- [C10](#): Handbook 5.5, 13.1.3 -changed to- Handbook (version 6), Section 3.3.1
- [C11](#): Handbook 13.1.2 -changed to- Handbook (version 6), Section 3.3

- [C12](#): Handbook 10.3.2 -changed to- Handbook (version 6), Section 3.4
- [C13](#): Handbook 5.2, 5.7 -changed to- Handbook (version 6), Section 3.2.1
- [C14](#), column 2: Define in advance which outcomes are primary outcomes and which are secondary outcomes. -changed to- Define in advance outcomes that are critical to the review, and any additional important outcomes.
- [C14](#), column 3: The primary outcomes -changed to- The critical outcomes
- [C14](#), column 3: It is important to identify up to seven outcomes from the primary and secondary outcomes that will form the basis of the GRADE assessment. -changed to- Additional important outcomes may also be specified. Up to seven critical and important outcomes will form the basis of the GRADE assessment and summarized in the review's abstract and other summary formats, although the review may measure more than seven outcomes.
- [C14](#), column 4: Handbook 5.4.2 -changed to- Handbook (version 6), Section 3.2.4.1
- [C15](#), column 2: that are important -changed to- that are critical or important
- [C15](#), column 3: that are important -changed to- that are critical or important
- [C15](#), column 3 new text: Any outcomes that would not be described as critical or important can be left out of the review.
- [C15](#), column 4: Handbook 5.4.2 -changed to- Handbook (version 6), Section 3.2.4.1
- [C16](#), column 4: Handbook 5.4.1 -changed to- Handbook (version 6), Section 3.2.4.1
- [C19](#), column 4: Handbook 6.3, 6.4 -changed to- Handbook (version 6), Section 1.5; 4.3.1.1
- [C20](#), column 3: ' Risk of bias' -changed to- 'risk of bias'
- [C20](#), column 4: Handbook 8.3 -changed to- Handbook (version 6), Section 1.5
- [C21](#), column 4: Handbook 9.1.2 -changed to- Handbook (version 6), Section 1.5
- [C22](#), column 4: Handbook 9.6.5 -changed to- Handbook (version 6), Section 1.5
- [C23](#), column 4: Handbook 11.5 -changed to- Handbook (version 6), Section 1.5
- [C24](#) column 3: Supplementary searches should be performed as described in sections 6.3.2 and 6.3.3 of the Handbook. -changed to- DELETED
- [C24](#): BLANK -changed to- See Handbook Section 4.3.1.1
- [C25](#): Handbook 6.2.1.4, 6.2.1.5 -changed to- Handbook (version 6), Section 4.3.1.4
- [C26](#): Handbook 13.3; 14.5; 15.3; 20.3.2.1 -changed to- Handbook (version 6), Section 4.4.1
- [C27](#): Handbook 6.2.3.1, 6.2.3.2, 6.2.3.3 -changed to- Handbook (version 6), Section 4.4.3
- [C28](#): Handbook 6.2.1.7, 6.2.1.8, 6.2.2 -changed to- Handbook (version 6), Section 4.3.5
- [C29](#): Handbook 6.2.2.5 -changed to- Handbook (version 6), Section 4.3.5
- [C30](#): Handbook 6.2.2.5 -changed to- Handbook (version 6), Section 4.3.5
- [C31](#): Handbook 6.2.3 -changed to- Handbook (version 6), Section 4.3.2
- [C32](#): Handbook 6.4.2, 6.4.4, 6.4.7 -changed to- Handbook (version 6), Section 4.4.2
- [C33](#): Handbook 6.4.5, 6.4.6, 6.4.8 -changed to- Handbook (version 6), Section 4.4.4
- [C34](#): Handbook 6.4.11, 6.4.2; 13.3.1.2; 14.5.2; 15.3.1; 17.5; 20.3.2.1 -changed to- Handbook (version 6), Section 4.4.7
- [C35](#): Handbook 6.4.9 -changed to- Handbook (version 6), Section 4.4.5
- [C36](#): Handbook 6.6.1 -changed to- Handbook (version 6), Section 4.5
- [C37](#): BLANK -changed to- Handbook (version 6), Section 4.4.10
- [C38](#): BLANK -changed to- Handbook (version 6), Section 4.4.10
- [C39](#) column 4: Handbook 7.2.4 -changed to- Handbook (version 6), Section 4.6.4
- [C40](#) column 4: Handbook 5.4.1 -changed to- Handbook (version 6), Section 4.6.3
- [C41](#) column 3: A PRISMA type flow diagram and a **table of 'Characteristics of excluded studies'** will need to be completed in the final review. -changed to- DELETED
- [C41](#) column 4: Handbook 6.6.1; 11.2.1 -changed to- Handbook (version 6), Section 4.6.4
- [C42](#) column 4: Handbook 7.2.1, 7.2.2, 7.6.4 -changed to- Handbook (version 6), Section 4.6.2; 5.2.1
- [C43](#) column 2: that has been -changed to- which has been



- [C43](#) column 3: Piloting the form within the review team using a sample of included studies is highly desirable -changed to- Piloting the form within the review team is highly desirable.
- [C43](#) column 4: Handbook 7.5 -changed to- Handbook (version 6), Section 5.4.1
- [C44](#) column 3: Details of funding source for each study and the declarations of interest for the primary investigators should also be collected during this process. TiDieR (Hoffman 2014) will assist selection of which characteristics of interventions should be sought. -changed to- DELETED
- [C44](#) column 4: Handbook 7.3; 11.2 -changed to- Handbook (version 6), Section 5.3.1
- [C45](#) column 3: not a mandatory standard for study characteristics. -changed to- not a mandatory standard for the former.
- [C45](#) column 4: Handbook 7.6.2, 7.6.5 -changed to- Handbook (version 6), Section 5.5.2
- [C46](#) column 4: Handbook 7.6.2 -changed to- Handbook (version 6), Section 5.5.2
- [C47](#) column 4: Handbook 7.7 -changed to- Handbook (version 6), Section 5.3.6
- [C48](#) column 4: Handbook 6.4.10 -changed to- Handbook (version 6), Section 4.4.6; 5.2
- [C49](#) column 3: Risk of bias -changed to- risk of bias
- [C49](#) column 4: Handbook 7.4.2 -changed to- Handbook (version 6), Section 5.2.3
- [C50](#) title: Choosing intervention groups in multi-arm studies -changed to- Choosing interventions in multi-arm studies
- [C50](#) column 2: include in the review only the intervention and control groups that meet -changed to- include in the review only the interventions that meet
- [C50](#) column 3: intervention groups (x2) -changed to- interventions (x2)
- [C50](#) column 4: Handbook 16.5.2 -changed to- Handbook (version 6), Section 5.3.6
- [C52](#) column 3: Recommendations for assessing bias in randomized studies included in Cochrane Reviews are now well established. -changed to- DELETED
- [C52](#) column 3: as described in this Handbook -changed to- as described in Handbook version 6
- [C52](#) column 4: See Handbook version 6 (Chapter 8) -changed to- See Handbook (version 6), Section 7.1.2; Chapter 8
- [C53](#) column 2: risk of bias tool -changed to- risk-of-bias-tool
- [C53](#) column 3: the risk of bias assessment -changed to- the risk-of-bias assessment
- [C53](#) column 4: See Handbook 8.3.4 -changed to- See Handbook (version 6), Section 7.3.2; Chapter 8
- [C54](#) column 2: risk of bias tables -changed to- risk-of-bias tables
- [C54](#) column 3: Items that are judged to be at an unclear risk of bias but are without accompanying information supporting the judgment appear as empty cells in the graphical plots based on the 'Risk of bias' tool in the published review. -changed to- DELETED
- [C54](#) column 4: Handbook 8.5.2 -changed to- Handbook (version 6), Section 7.3.2; Chapter 8
- [C55](#) column 2: risk of bias judgement -changed to- risk-of-bias judgement
- [C54](#) column 3: judgments -changed to- judgements
- [C55](#) column 4: Handbook 8.5.2 -changed to- Handbook (version 6), Section 7.3.2; Chapter 8
- [C56](#) column 4: Handbook 8.5.1, 8.11.2, 8.12.2 -changed to- Handbook (version 6), Section 7.3.2; Chapter 8
- [C57](#) title: Summarizing risk of bias assessments changed to- Summarizing risk-of-bias assessments
- [C57](#) column 4: Handbook 8.5.1, 8.13.2 -changed to- Handbook (version 6), Section 7.5; Chapter 8
- [C58](#) column 4: Handbook 8.7 -changed to- Handbook (version 6), Section 7.6.1; Chapter 8
- [C59](#) column 4: Handbook version 6 (Chapter 8) -changed to- Handbook (version 6), Section 7.6.1; Chapter 8
- [C60](#) column 3: “notable concern of conflicts of interest” -changed to- “notable concern about conflicts of interest”
- [C60](#) column 4: Handbook 8.8.1 -changed to- Handbook (version 6), Section 7.8.6; Chapter 8
- [C61](#) column 4: Handbook 9.2.3.2 -changed to- BLANK
- [C62](#) column 4: See Handbook 9.1.4 -changed to- BLANK



- [C63](#) column 4: See Handbook 9.5.2 -changed to- See Handbook (version 6), Section 10.10.2
- [C64](#) column 3: Risk of bias tool -changed to- 'risk-of-bias' tool
- [C64](#) column 4: See Handbook 16.2 -changed to- See Handbook (version 6), Section 10.12.1
- [C65](#) column 4 See Handbook 9.4.5.3 -changed to- See Handbook (version 6), Section 10.5.3
- [C66](#) column 3: and using multiple treatments meta-analysis. -changed to- and using network meta-analysis.
- [C66](#) column 4: See Handbook 7.7.3.8, 16.5.4 -changed to- See Handbook (version 6), Section 6.2.9 and Chapter 11.
- [C67](#) column 4: See Handbook 9.6.3.1 -changed to- See Handbook (version 6), Section 10.11.3.1
- [C68](#) column 4: See Handbook 9.6.5.2 -changed to- See Handbook (version 6), Section 10.11.5.2
- [C69](#) column 4: See Handbook 9.5.4 -changed to- See Handbook (version 6), Section 10.10.3
- [C70](#) column 3: of the study, i.e., to give it (x2) -changed to- of the study, that is, to give it (x2)
- [C70](#) column 4: see Handbook 9.3, 16.3, 16.4 -changed to- See Handbook (version 6), Section 6.2.1
- [C71](#) column 4: see Handbook 9.7 -changed to- See Handbook (version 6), Section 10.14
- [C72](#) column 2: Interpret a statistically non-significant P value (e.g. larger than 0.05) as a finding of uncertainty unless confidence intervals are sufficiently narrow to rule out an important magnitude of effect. -changed to- (Do not describe results as statistically significant or non-significant. Interpret the confidence intervals and their width.) Focus interpretation of results on estimates of effect and their confidence intervals, avoiding use of a distinction between **“statistically significant”** and **“statistically non-significant”**.
- [C72](#) column 4: See Handbook 12.4.2, 12.7.4 -changed to- See Handbook (version 6), Section 15.3.1
- [C73](#) column 4: See Handbook 10.1, 10.2 -changed to- See Handbook (version 6), Section 13.4
- [C74](#) column 2 title: Assessing the quality -changed to- Assessing the certainty
- [C74](#) column 2: quality of the body of evidence -changed to- certainty of the body of evidence
- [C74](#) column 2: quality of evidence -changed to- certainty of evidence
- [C74](#) column 3: quality of the body of evidence -changed to- certainty of the body of evidence
- [C74](#) column 4: See Handbook 12.2 -changed to- See Handbook Section 14.2.1
- [C75](#) column 2 title: quality of the body of evidence -changed to- certainty of the body of evidence
- [C75](#) column 2: quality of the body of evidence -changed to- certainty of the body of evidence
- [C75](#) column 4: See Handbook 12.2.1 -changed to- See Handbook Section 14.2.1

## Version 1.07

- [C56](#): "assess RoB due to lack of blinding....." replaced with NEW standard "Ensuring results of outcomes included in SoF are assessed for RoB....."
- [C57](#): "RoB due to incomplete outcome data...." replaced with "Summarizing RoB assessments...."
- [C58](#): "Summarizing RoB assessments...." replaced with "Addressing RoB in the synthesis...."
- [C59](#): "Addressing RoB in the synthesis...." replaced with "Incorporating assessments of RoB...."
- [C60](#): "Incorporating assessments of RoB...." replaced with NEW standard "Addressing Col in included trials....."

## Version 1.06

- [C73](#): Standard changed to: Consider the potential impact of non-reporting biases on the results of the review or the meta-analysis it contains. Rationale and elaboration changed to: There is overwhelming evidence of non-reporting biases of various types. These can be addressed at various points of the review. A thorough search, and attempts to obtain unpublished results, might minimize the risk. Analyses of the results of included studies, for example using funnel plots, can sometimes help determine the possible extent of the problem, as can attempts to identify study protocols, which should be a routine feature of Cochrane Reviews.

- [C24](#): Standard changed from “Planning the search” to “Searching general bibliographic databases and CENTRAL”
- [C41](#): Standard changed to: “Document the selection process in sufficient detail to be able to complete a flow diagram and a table of ‘Characteristics of excluded studies’. Change elaboration to read: “A PRISMA type flow diagram and a table of ‘Characteristics of excluded studies’ will need to be completed in the final review.....”
- [R56](#): Standard changed to: Provide information on the flow of studies....., ideally using a PRISMA type flow diagram.....individual studies”.
- [UR4](#): Elaboration changed to: “Provide information on the flow of studies into the updated review, ideally using a PRISMA type flow diagram.”
- [R98](#): Status changed to mandatory – Mandating SoF tables.
- [R102](#): **Changed elaboration to: “When formulating implications for practice base conclusions only on findings from the synthesis (quantitative or narrative) of studies included in the review. The conclusions of the review should convey the essence of the synthesis of included studies, without selective reporting of particular findings on the basis of the result, and without drawing on data that were not systematically compiled and evaluated as part of the review.”**

## Version 1.05

- [C48](#): Upgraded from 'highly desirable' to 'mandatory'.

## Version 1.04

- [R55](#): New Standard inserted. There is subsequent renumbering of all Standards in section up to R108.(23/01/2018)
- [C28](#): Changed from 'mandatory' to 'highly desirable'.(23/01/18)
- Links to Cochrane Interactive Learning modules have been added where needed.

## How to cite the MECIR Standards

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URL: <https://community.cochrane.org/mecir-manual/standards-conduct-new-cochrane-intervention-reviews-c1-75>

## STANDARDS FOR THE CONDUCT OF NEW COCHRANE INTERVENTION REVIEWS

Julian PT Higgins, Toby Lasserson, Jackie Chandler, David Tovey, James Thomas, Ella Flemyng and Rachel Churchill

URL: <https://community.cochrane.org/mecir-manual/standards-conduct-new-systematic-reviews/key-points-introduction>

## Key points and introduction

Key points:

- The conduct Standards should be consulted during preparation of the protocol for a Cochrane Intervention Review.
- They describe the methods that should be implemented throughout the review process.
- Few specific methods are mandatory, one notable exception being use of the Cochrane tool for assessing risk of bias when randomized trials are included in the review.

The MECIR Standards for conduct of a Cochrane Intervention Review provide expectations for the general methodological approach to be followed from designing the review up to interpreting the findings at the end. They should be consulted particularly when preparing the protocol for the review. The protocol describes the review question, the criteria for considering studies for the review, and the methods that will be followed to identify, appraise, summarize and synthesize the studies. Cochrane led the way in making protocols available to readers of the Cochrane Library. They ensure transparency in how reviews are prepared and allow the planned methods to be critiqued. Specification of the review **question (through setting the review's objectives) and the criteria for including studies are critical to the success of the review** and the first two sections of the standards address these tasks. The following section addresses selection of the outcomes of interest, an important aspect that should be prespecified carefully to avoid the need for post hoc decisions that could be influenced by the data.

The remaining Standards address the detailed methodology that will be followed during the review, covering the search for studies, selection of studies into the review, data collection, risk of bias assessment, synthesis (including any meta-analysis approaches), and overall assessment of the evidence. With few exceptions (such as use of the Cochrane Risk of Bias 2 tool for randomized trials), the precise methods to be used are not prescribed. For example, authors are free to use any meta-analysis method, although there is a potential convenience to both authors and readers if those implemented in Review Manager (RevMan) software are used.

Julian Higgins  
Professor of Evidence Synthesis  
University of Bristol

## Developing the protocol for the review

Cochrane Training resource: [writing a protocol](#) and [common errors and best practice: writing review protocols](#)

Cochrane Interactive Learning (CIL): [module 2 - writing the review protocol](#)

URL: <https://community.cochrane.org/mecir-manual/standards-conduct-new-cochrane-intervention-reviews-c1-75/developing-protocol-review-c1-23/setting-research-question-inform-scope-review-c1-4>

## 1.1 Setting the research question(s) to inform the scope of the review

Cochrane Training resource: [defining the review question](#)

Cochrane Interactive Learning (CIL): [module 1 - introduction to conducting systematic reviews](#)

Standard		Rationale and elaboration
C1	Formulating review questions	Mandatory
Ensure that the review question and particularly the outcomes of interest, address issues that are important to review users such as healthcare consumers, health professionals and policy makers.		Cochrane Reviews are intended to support clinical practice and policy, not just scientific curiosity. The needs of consumers play a central role in Cochrane Reviews and they can play an important role in defining the review question. Qualitative research, i.e. studies that explore the experience of those involved in providing and receiving interventions, and studies evaluating factors that shape the implementation of interventions, might be used in the same way. See <i>Handbook</i> Section 2.1
C2	Predefining objectives	Mandatory
Define in advance the objectives of the review, including participants, interventions, comparators and outcomes (PICO).		Objectives give the review focus and must be clear before appropriate eligibility criteria can be developed. If the review will address multiple interventions, clarity is required on how these will be addressed (e.g. summarized separately, combined or explicitly compared). See <i>Handbook</i> Section 2.3
C3	Considering potential adverse effects	Mandatory
Consider any important potential adverse effects of the intervention(s) and ensure that they are addressed.		It is important that adverse effects are addressed in order to avoid one-sided summaries of the evidence. At a minimum, the review will need to highlight the extent to which potential adverse effects have been evaluated in any included studies. Sometimes data on adverse effects are best obtained from non-randomized studies, or qualitative research studies. This does not mean however that all reviews must include non-randomized studies. See <i>Handbook</i> Section 2.1
C4	Considering equity and specific populations	Highly desirable

Consider in advance whether issues of equity and relevance of evidence to specific populations are important to the review, and plan for appropriate methods to address them if they are. Attention should be paid to the relevance of the review question to populations such as low-socioeconomic groups, low- or middle-income regions, women, children and older people.

Where possible reviews should include explicit descriptions of the effect of the interventions not only upon the whole population, but also on the disadvantaged, and/or the ability of the interventions to reduce socioeconomic inequalities in health, and to promote use of the interventions to the community.

See *Handbook* Section 2.4



URL: <https://community.cochrane.org/mecir-manual/standards-conduct-new-cochrane-intervention-reviews-c1-75/developing-protocol-review-c1-23/setting-eligibility-criteria-including-studies-review-c5-13>

## 1.2 Setting eligibility criteria for including studies in the review

Cochrane Training resource: [defining the review question](#)

Cochrane Interactive Learning (CIL): [module 2 - writing the review protocol](#)

	Standard	Rationale and elaboration	
C5	Predefining unambiguous criteria for participants		Mandatory
	Define in advance the eligibility criteria for participants in the studies.	<p>Predefined, unambiguous eligibility criteria are a fundamental prerequisite for a systematic review. The criteria for considering types of people included in studies in a review should be sufficiently broad to encompass the likely diversity of studies, but sufficiently narrow to ensure that a meaningful answer can be obtained when studies are considered in aggregate. Considerations when specifying participants include setting, diagnosis or definition of condition and demographic factors. Any restrictions to study populations must be based on a sound rationale, since it is important that Cochrane Reviews are widely relevant.</p> <p>See <i>Handbook</i> Section 3.2.1</p>	
C6	Predefining a strategy for studies with a subset of eligible participants		Highly desirable
	Define in advance how studies that include only a subset of relevant participants will be addressed.	<p><b>Sometimes a study includes some ‘eligible’ participants and some ‘ineligible’ participants, for example when an age cut-off is used in the review’s eligibility criteria. If data from the eligible participants cannot be retrieved, a mechanism for dealing with this situation should be prespecified.</b></p> <p>See <i>Handbook</i> Section 3, 3.2.1</p>	
C7	Predefining unambiguous criteria for interventions and comparators		Mandatory
	Define in advance the eligible interventions and the interventions against which these can be compared in the included studies.	<p>Predefined, unambiguous eligibility criteria are a fundamental prerequisite for a systematic review. Specification of comparator interventions requires particular clarity: are the experimental interventions to be compared with an inactive control intervention (e.g. placebo, no treatment, standard care, or a waiting list control), or with an active control intervention (e.g. a different variant of the same intervention, a different drug, a different kind of therapy)? Any restrictions on interventions and comparators, for example, regarding delivery, dose, duration, intensity, cointerventions and features of complex interventions should also be predefined and explained.</p> <p>See <i>Handbook</i> Section 3, 3.2.2</p>	

C8 Clarifying role of outcomes	Mandatory
<p>Clarify in advance whether outcomes listed under 'Criteria for considering studies for this review' are used as criteria for including studies (rather than as a list of the outcomes of interest within whichever studies are included).</p>	<p>Outcome measures should not always form part of the criteria for including studies in a review. However, some reviews do legitimately restrict eligibility to specific outcomes. For example, the same intervention may be studied in the same population for different purposes (e.g. hormone replacement therapy, or aspirin); or a review may address specifically the adverse effects of an intervention used for several conditions. If authors do exclude studies on the basis of outcomes, care should be taken to ascertain that relevant outcomes are not available because they have not been measured rather than simply not reported. See <i>Handbook</i> Section 3, 3.2.4.1</p>
C9 Predefining study designs	Mandatory
<p>Define in advance the eligibility criteria for study designs in a clear and unambiguous way, with a focus on features of a study's design rather than design labels.</p>	<p>Predefined, unambiguous eligibility criteria are a fundamental prerequisite for a systematic review. This is particularly important when non-randomized studies are considered. Some labels commonly used to define study designs can be <b>ambiguous</b>. For example a 'double blind' study may not make it clear who was blinded; a 'case control' study may be nested within a cohort, or be undertaken in a cross-sectional manner; or a 'prospective' study may have only some features defined or undertaken prospectively. See <i>Handbook</i> Section 3, 3.3</p>
C10 Including randomized trials	Mandatory
<p>Include randomized trials as eligible for inclusion in the review, <i>if it is feasible to conduct them to evaluate interventions and outcomes of interest</i>.</p>	<p>Randomized trials are the best study design for evaluating the efficacy of interventions. If it is feasible to conduct them to evaluate questions that are being addressed by the review, they must be considered eligible for the review. However, appropriate exclusion criteria may be put in place, for example regarding length of follow-up. See <i>Handbook</i> Section 3, 3.3.1</p>
C11 Justifying choice of study designs	Mandatory
<p>Justify the choice of eligible study designs.</p>	<p>It might be difficult to address some interventions or some outcomes in randomized trials. Authors should be able to justify why they have chosen either to restrict the review to randomized trials or to include non-randomized studies. The particular study designs included should be justified with regard to appropriateness to the review question and with regard to potential for bias. See <i>Handbook</i> Section 3, 3.3</p>
C12 Excluding studies based on publication status	Mandatory

Include studies irrespective of their publication status, unless exclusion is explicitly justified.

Obtaining and including data from unpublished studies (including grey literature) can reduce the effects of publication bias. However, the unpublished studies that can be located may be an unrepresentative sample of all unpublished studies.  
See *Handbook* Section 3, 3.4

### C13 Changing eligibility criteria

Mandatory

Justify any changes to eligibility criteria or outcomes studied. In particular, post hoc decisions about inclusion or exclusion of studies should keep faith with the objectives of the review rather than with arbitrary rules.

Following prespecified eligibility criteria is a fundamental attribute of a systematic review. However, unanticipated issues may arise. Review authors should make sensible post hoc decisions about exclusion of studies, and these should be documented in the review, possibly accompanied by sensitivity analyses. Changes to the protocol must not be made on the basis of the findings of the studies or the synthesis, as this can introduce bias.

See *Handbook* Section 3, 3.2.1

URL: <https://community.cochrane.org/mecir-manual/standards-conduct-new-cochrane-intervention-reviews-c1-75/developing-protocol-review-c1-23/selecting-outcomes-be-addressed-studies-included-review-c14-18>

### 1.3 Selecting outcomes to be addressed for studies included in the review

Cochrane Training resource: [defining the review question](#)

Cochrane Interactive Learning: [module 2 - writing the review protocol](#)

Standard		Rationale and elaboration
C14	Predefining outcome domains	Mandatory
Define in advance outcomes that are critical to the review, and any additional important outcomes.		Full specification of the outcomes includes consideration of outcome domains (e.g. quality of life) and outcome measures (e.g. SF-36). Predefinition of outcome reduces the risk of selective outcome reporting. The <i>critical outcomes</i> should be as few as possible and should normally reflect at least one potential benefit and at least one potential area of harm. It is expected that the review should be able to synthesize these outcomes if eligible studies are identified, and that the conclusions of the review will be based largely on the effects of the interventions on these outcomes. Additional important outcomes may also be specified. Up to seven critical and important outcomes will form the basis of the GRADE assessment and summarized in the <b>review's abstract and other summary</b> formats, although the review may measure more than seven outcomes See <i>Handbook</i> Section 3, 3.2.4.1
C15	Choosing outcomes	Mandatory
Choose only outcomes that are critical or important to users of the review such as healthcare consumers, health professionals and policy makers.		Cochrane Reviews are intended to support clinical practice and policy, and should address outcomes that are critical or important to consumers. These should be specified at protocol stage. Where available, established sets of core outcomes should be used. Patient-reported outcomes should be included where possible. It is also important to judge whether evidence of resource use and costs might be an important component of decisions to adopt the intervention or alternative management strategies around the world. Large numbers of outcomes, while sometimes necessary, can make reviews unfocussed, unmanageable for the user, and prone to selective outcome reporting bias. Biochemical, interim and process outcomes should be considered where they are important to decision makers. Any outcomes that would not be described as critical or important can be left out of the review. See <i>Handbook</i> Section 3, 3.2.4.1

C16	Predefining outcome measures	Highly desirable
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Define in advance details of what will constitute acceptable outcome measures (e.g. diagnostic criteria, scales, composite outcomes).

Having decided what outcomes are of interest to the review, authors should clarify acceptable ways in which these outcomes can be measured. It may be difficult, however, to predefine adverse effects.  
See *Handbook* Section 3, 3.2.4.1

C17	Predefining choices from multiple outcome measures	Highly desirable
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Define in advance how outcome measures will be selected when there are several possible measures (e.g. multiple definitions, assessors or scales).

Prespecification guards against selective outcome reporting, and allows users to confirm that choices were not overly influenced by the results. A predefined hierarchy of outcomes measures may be helpful. It may be difficult, however, to predefine adverse effects. A rationale should be provided for the choice of outcome measure.

C18	Predefining time points of interest	Highly desirable
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Define in advance the timing of outcome measurement.

Prespecification guards against selective outcome reporting, and allows users to confirm that choices were not overly influenced by the results. Authors may consider whether all time frames or only selected time points will be included in the review. These decisions should be based on outcomes important for making healthcare decisions. One strategy to make use of the available data could be to group time points **into prespecified intervals to represent ‘short-term’, ‘medium-term’ and ‘long-term’ outcomes and to take no more than one from each interval from each study for any particular outcome.**

URL: <https://community.cochrane.org/mecir-manual/standards-conduct-new-cochrane-intervention-reviews-c1-75/developing-protocol-review-c1-23/planning-review-methods-protocol-stage-c19-23>

## 1.4 Planning the review methods at protocol stage

Standard		Rationale and elaboration
C19	Planning the search	Mandatory
Plan in advance the methods to be used for identifying studies. Design searches to capture as many studies as possible that meet the eligibility criteria, ensuring that relevant time periods and sources are covered and not restricted by language or publication status.		Searches should be motivated directly by the eligibility criteria for the review, and it is important that all types of eligible studies are considered when planning the search. If searches are restricted by publication status or by language of publication, there is a possibility of publication bias, or language bias (whereby the language of publication is selected in a way that depends on the findings of the study), or both. Removing language restrictions in English language databases is not a good substitute for searching non-English language journals and databases. See <i>Handbook</i> Section 1, 1.5; Section 4, 4.3.1.1
C20	Planning the assessment of risk of bias in included studies	Mandatory
Plan in advance the methods to be used for assessing risk of bias in included studies, including the tool(s) to be used, how the tool(s) will be implemented, and the criteria used to assign studies, for example, to judgements of low risk, high risk and unclear risk of bias.		Predefining the methods and criteria for assessing risk of bias is important since analysis or interpretation of the review findings may be affected by the judgements made during this process. For randomized trials, use of the Cochrane ' <b>risk of bias</b> ' tool is Mandatory, so it is sufficient (and easiest) simply to refer to the definitions of low risk, unclear risk and high risk of bias provided in the <i>Handbook</i> . See <i>Handbook</i> Section 1, 1.5
C21	Planning the synthesis of results	Mandatory
Plan in advance the methods to be used to synthesize the results of the included studies, including whether a quantitative synthesis is planned, how heterogeneity will be assessed, choice of effect measure (e.g. odds ratio, risk ratio, risk difference or other for dichotomous outcomes), and methods for meta-analysis (e.g. inverse variance or Mantel Haenszel, fixed-effect or random-effects model).		Predefining the synthesis methods, particularly the statistical methods, is important, since analysis or interpretation of the review findings may be affected by the judgements made during this process. See <i>Handbook</i> Section 1, 1.5

C22	Planning subgroup analyses	Mandatory
	<p>Predefine potential effect modifiers (e.g. for subgroup analyses) at the protocol stage; restrict these in number, and provide rationale for each.</p>	<p>Pre-specification reduces the risk that large numbers of undirected subgroup analyses will lead to spurious explanations of heterogeneity. See <i>Handbook</i> Section 1, 1.5</p>
C23	Planning the GRADE assessment and ‘Summary of findings’ table	Mandatory
	<p>Plan in advance the methods to be used for assessing the certainty of the body of evidence, and summarizing the findings of the review.</p>	<p>Methods for assessing the certainty of evidence for the most important outcomes in the review need to be pre-specified. In <b>‘Summary of findings’ tables the most important feature is to</b> predefine the choice of outcomes in order to guard against selective presentation of results in the review. The table should include the essential outcomes for decision making (typically up to seven), which generally should not include surrogate or interim outcomes. The choice of outcomes should not be based on any anticipated or observed magnitude of effect, or because they are likely to have been addressed in the studies to be reviewed. See <i>Handbook</i> Section 1, 1.5</p>

## Performing the review



URL: <https://community.cochrane.org/mecir-manual/standards-conduct-new-cochrane-intervention-reviews-c1-c75/performing-review-c24-75/searching-studies-c24-38>

## 1.5 Searching for studies

Cochrane Training resource: [searching for studies](#)

Cochrane Interactive Learning (CIL): [module 3 - searching for studies](#)

Standard		Rationale and elaboration
C24	Searching general bibliographic databases and CENTRAL	Mandatory
Search the Cochrane Review Group's (CRG's) Specialized Register (internally, e.g. via the Cochrane Register of Studies, or externally via CENTRAL). Ensure that CENTRAL, MEDLINE (e.g. via PubMed) and Embase (if Embase is available to either the CRG or the review author), have been searched (either for the review or <b>for the Review Group's</b> Specialized Register).		Searches for studies should be as extensive as possible in order to reduce the risk of publication bias and to identify as much relevant evidence as possible. The minimum databases to be covered are the <b>CRG's</b> Specialized Register (if it exists and was designed to support reviews in this way), CENTRAL, MEDLINE and Embase (if Embase is available to either the CRG or the review author). Expertise may be required to avoid unnecessary duplication of effort. Some, but not all, reports of eligible studies from MEDLINE, Embase and the <b>CRGs'</b> Specialized Registers are already included in CENTRAL. See <i>Handbook</i> Section 4, 4.3.1.1
C25	Searching specialist bibliographic databases	Highly desirable
Search appropriate national, regional and subject-specific bibliographic databases.		Searches for studies should be as extensive as possible in order to reduce the risk of publication bias and to identify as much relevant evidence as possible. Databases relevant to the review topic should be covered (e.g. CINAHL for nursing-related topics, PsycINFO for psychological interventions), and regional databases (e.g. LILACS) should be considered. See <i>Handbook</i> Section 4, 4.3.1.4
C26	Searching for different types of evidence	Mandatory
<i>If the review has specific eligibility criteria around study design to address adverse effects, economic issues or qualitative research questions, undertake searches to address them.</i>		Sometimes different searches will be conducted for different types of evidence, such as for non-randomized studies for addressing adverse effects, or for economic evaluation studies. See <i>Handbook</i> Section 4, 4.4.1
C27	Searching trials registers	Mandatory

Search trials registers and repositories of results, where relevant to the topic, through ClinicalTrials.gov, the WHO International Clinical Trials Registry Platform (ICTRP) portal and other sources as appropriate.	Searches for studies should be as extensive as possible in order to reduce the risk of publication bias and to identify as much relevant evidence as possible. Although ClinicalTrials.gov is included as one of the registers within the WHO ICTRP portal, it is recommended that both ClinicalTrials.gov and the ICTRP portal are searched separately due to additional features in ClinicalTrials.gov. See <i>Handbook</i> Section 4, 4.3.3
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C28	Searching for grey literature	Highly desirable
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Search relevant grey literature sources such as reports, dissertations, theses, databases and databases of conference abstracts.	Searches for studies should be as extensive as possible in order to reduce the risk of publication bias and to identify as much relevant evidence as possible. See <i>Handbook</i> Section 4, 4.3.5
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C29	Searching within other reviews	Highly desirable
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Search within previous reviews on the same topic.	Searches for studies should be as extensive as possible in order to reduce the risk of publication bias and to identify as much relevant evidence as possible. See <i>Handbook</i> Section 4, 4.3.5
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C30	Searching reference lists	Mandatory
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Check reference lists in included studies and any relevant systematic reviews identified.	Searches for studies should be as extensive as possible in order to reduce the risk of publication bias and to identify as much relevant evidence as possible. See <i>Handbook</i> Section 4, 4.3.5
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C31	Searching by contacting relevant individuals and organizations	Highly desirable
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Contact relevant individuals and organizations for information about unpublished or ongoing studies.	Searches for studies should be as extensive as possible in order to reduce the risk of publication bias and to identify as much relevant evidence as possible. It is important to identify ongoing studies, so that these can be assessed for possible inclusion when a review is updated. See <i>Handbook</i> Section 4, 4.3.2
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C32	Structuring search strategies for bibliographic databases	Mandatory
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Inform the structure of search strategies in bibliographic databases around the main concepts of the review, using appropriate elements from PICO and study design. In structuring the search, maximize sensitivity whilst striving for reasonable precision.

**Ensure correct use of the 'AND' and 'OR' operators.**

Inappropriate or inadequate search strategies may fail to identify records that are included in bibliographic databases. **Expertise may need to be sought, in particular from the CRG's Information Specialist.** The structure of a search strategy should be based on the main concepts being examined in a review. In general databases, such as MEDLINE, a search strategy to identify studies for a Cochrane Review will typically have three sets of terms: 1) terms to search for the health condition of interest, i.e. the population; 2) terms to search for the intervention(s) evaluated; and 3) terms to search for the types of **study design to be included (typically a 'filter' for randomized trials)**. There are exceptions, however. For instance, for reviews of complex interventions, it may be necessary to search only for the population or the intervention. Within each concept, terms **are joined together with the Boolean 'OR' operator, and the concepts are combined with the Boolean 'AND' operator. The 'NOT' operator should be avoided where possible to avoid the danger of inadvertently removing records that are relevant from the search set.**

See *Handbook* Section 4, 4.4.2

### C33 Developing search strategies for bibliographic databases

Mandatory

Identify appropriate controlled vocabulary (e.g. MeSH, Emtree, including 'exploded' terms) and free-text terms (considering, for example, spelling variants, synonyms, acronyms, truncation and proximity operators).

Inappropriate or inadequate search strategies may fail to identify records that are included in bibliographic databases. Search strategies need to be customized for each database. It is **important that MeSH terms are 'exploded' wherever appropriate**, in order not to miss relevant articles. The same principle applies to Emtree when searching Embase and also to a number of other databases. The controlled vocabulary search terms for MEDLINE and Embase are not identical, and neither is the approach to indexing. In order to be as comprehensive as possible, it is necessary to include a wide range of free-text terms for each of the concepts selected. This might include the use of truncation and wildcards. Developing a search strategy is an iterative process in which the terms that are used are modified, based on what has already been retrieved.

See *Handbook* Section 4, 4.4.4

### C34 Using search filters

Highly desirable

Use specially designed and tested search filters where appropriate including the Cochrane Highly Sensitive Search Strategies for identifying randomized trials in MEDLINE, but do not use filters in pre-filtered databases e.g. do not use a randomized trial filter in

Inappropriate or inadequate search strategies may fail to identify records that are included in bibliographic databases. Search filters should be used with caution. They should be assessed not only for the reliability of their development and reported performance, but also for their current accuracy, relevance and effectiveness given the frequent interface and indexing changes affecting databases.

CENTRAL or a systematic review filter in DARE. See *Handbook* Section 4, 4.4.7

C35	Restricting database searches	Mandatory
	Justify the use of any restrictions in the search strategy on publication date and publication format.	Date restrictions in the search should only be used when there are date restrictions in the eligibility criteria for studies. They should be applied only if it is known that relevant studies could only have been reported during a specific time period, for example if the intervention was only available after a certain time point. Searches for updates to reviews might naturally be restricted by date of entry into the database (rather than date of publication) to avoid duplication of effort. Publication format restrictions (e.g. exclusion of letters) should generally not be used in Cochrane Reviews, since any information about an eligible study may be of value. See <i>Handbook</i> Section 4, 4.4.5
C36	Documenting the search process	Mandatory
	Document the search process in enough detail to ensure that it can be reported correctly in the review.	The search process (including the <i>sources</i> searched, when, by whom, and using which terms) needs to be documented in enough detail throughout the process to ensure that it can be reported correctly in the review, to the extent that all the searches of all the databases are reproducible. See <i>Handbook</i> Section 4, 4.4.5
C37	Rerunning searches	Mandatory
	Rerun or update searches for all relevant databases within 12 months before publication of the review or review update, and screen the results for potentially eligible studies.	The published review should be as up to date as possible. The search must be rerun close to publication, if the initial search date is more than 12 months (preferably six months) from the intended publication date, and the results screened for potentially eligible studies. Ideally the studies should be incorporated fully in the review. If not, then the potentially eligible studies will need to be <b>reported, at a minimum as a reference under ‘Studies awaiting</b>

**classification’ (or ‘Ongoing studies’ if they have not yet completed).**

See *Handbook* Section 4, 4.4.10

C38	Incorporating findings from rerun searches	Highly desirable
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Fully incorporate any studies identified in the rerun or update of the search within 12 months before publication of the review or review update.

The published review should be as up to date as possible. After the rerun of the search, the decision whether to incorporate any new studies fully into the review will need to be balanced against the delay in publication.

See *Handbook* Section 4, 4.4.10

URL: <https://community.cochrane.org/mecir-manual/standards-conduct-new-cochrane-intervention-reviews-c1-c75/performing-review-c24-75/selecting-studies-include-review-c39-42>

## 1.6 Selecting studies to include in the review

Cochrane Training resources: [selecting studies](#) and [Covidence webinar](#) (online tool for review production)

Cochrane Interactive Learning (CIL): [module 4 - selecting studies and collecting data](#)

Standard		Rationale and elaboration
C39	Making inclusion decisions	Mandatory
Use (at least) two people working independently to determine whether each study meets the eligibility criteria, and define in advance the process for resolving disagreements.		Duplicating the study selection process reduces both the risk of making mistakes and the possibility that selection is influenced by a <b>single person's biases. The inclusion decisions should be based on the full texts of potentially eligible studies when possible, usually after an initial screen of titles and abstracts.</b> It is desirable, but not mandatory, that two people undertake this initial screening, working independently. See <i>Handbook</i> Section 4, 4.6.4
C40	Excluding studies without useable data	Mandatory
Include studies in the review irrespective of whether measured outcome data are reported in a 'usable' way.		Systematic reviews typically should seek to include all relevant participants who have been included in eligible study designs of the relevant interventions and had the outcomes of interest measured. Reviews must not exclude studies solely on the basis of reporting of the outcome data, since this may introduce bias due to selective outcome reporting and risk undermining the systematic review process. While such studies cannot be included in meta-analyses, the implications of their omission should be considered. Note that studies may legitimately be excluded because outcomes were not measured. Furthermore, issues may be different for adverse effects outcomes, since the pool of studies may be much larger and it can be difficult to assess whether such outcomes were measured. See <i>Handbook</i> Section 4, 4.6.3
C41	Documenting decisions about records identified	Mandatory

Document the selection process in sufficient detail to complete a flow diagram and a table of **‘Characteristics of excluded studies’**.

Decisions should be documented for all records identified by the search. Numbers of records are sufficient for exclusions based on initial screening of titles and abstracts. Broad categorizations are sufficient for records classed as potentially eligible during an initial screen. **Studies listed in the table of ‘Characteristics of excluded studies’ should be those that a user might reasonably expect to find in the review.** At least one explicit reason for their exclusion must be documented. Authors will need to decide for each review when to map records to studies (if multiple records refer to one study). Lists of included and excluded studies must be based on studies rather than records.

See *Handbook* Section 4, 4.6.4

#### C42 Collating multiple reports

Mandatory

Collate multiple reports of the same study, so that each study, rather than each report, is the unit of interest in the review.

It is wrong to consider multiple reports of the same study as if they are multiple studies. Secondary reports of a study should not be discarded, however, since they may contain valuable information about the design and conduct. Review authors must choose and justify which report to use as a source for study results.

See *Handbook* Section 4, 4.6.2; Section 5, 5.2.1

## 1.7 Collecting data from included studies

Cochrane Training resources: [collecting data](#) and [Covidence webinar](#) (online tool for review production)

Cochrane Interactive Learning (CIL): [module 4 - selecting studies and collecting data](#)

Standard		Rationale and elaboration	
C43	Using data collection forms		Mandatory
Use a data collection form, which has been piloted.		Review authors often have different backgrounds and level of systematic review experience. Using a data collection form ensures some consistency in the process of data extraction, and is necessary for comparing data extracted in duplicate. The completed data collection forms should be available to the CRG on request. Piloting the form within the review team is highly desirable. At a minimum, the data collection form (or a very close variant of it) must have been assessed for usability. See Handbook Section 5, 5.4.1	
C44	Describing studies		Mandatory
Collect characteristics of the included studies in sufficient detail to populate a table of 'Characteristics of included studies'.		Basic characteristics of each study will need to be presented as part of the review, including details of participants, interventions and comparators, outcomes and study design. See Handbook Section 5, 5.3.1	
C45	Extracting study characteristics in duplicate		Highly desirable
Use (at least) two people working independently to extract study characteristics from reports of each study, and define in advance the process for resolving disagreements.		Duplicating the data extraction process reduces both the risk of making mistakes and the possibility that data selection is <b>influenced by a single person's biases</b> . Dual data extraction may be less important for study characteristics than it is for outcome data, so it is not a mandatory standard for the former. See Handbook Section 5, 5.5.2	
C46	Extracting outcome data in duplicate		Mandatory
Use (at least) two people working independently to extract outcome data from reports of each study, and define in advance the process for resolving disagreements.		Duplicating the data extraction process reduces both the risk of making mistakes and the possibility that data selection is <b>influenced by a single person's biases</b> . Dual data extraction is particularly important for outcome data, which feed directly into syntheses of the evidence, and hence to the conclusions of the review. See Handbook Section 5, 5.5.2	
C47	Making maximal use of data		Mandatory



Collect and utilize the most detailed numerical data that might facilitate similar analyses of included studies. Where 2×2 tables or means and standard deviations are not available, this might include effect estimates (e.g. odds ratios, regression coefficients), confidence intervals, test statistics (e.g. t, F, Z, Chi<sup>2</sup>) or P values, or even data for individual participants.

Data entry into RevMan is easiest when 2×2 tables are reported for dichotomous outcomes, and when means and standard deviations are presented for continuous outcomes. Sometimes these statistics are not reported but some manipulations of the reported data can be performed to obtain them. For instance, 2×2 tables can often be derived from sample sizes and percentages, while standard deviations can often be computed using confidence intervals or P values. Furthermore, the inverse-variance data entry format can be used even if the detailed data required for dichotomous or continuous data are not available, for instance if only odds ratios and their confidence intervals are presented. The RevMan calculator facilitates many of these manipulations.

See *Handbook* Section 5, 5.3.6

#### C48 Examining errata

Mandatory

Examine any relevant retraction statements and errata for information.

Some studies may have been found to be fraudulent or may have been retracted since publication for other reasons. Errata can reveal important limitations, or even fatal flaws, in included studies. All of these may lead to the potential exclusion of a study from a review or meta-analysis. Care should be taken to ensure that this information is retrieved in all database searches by downloading the appropriate fields, together with the citation data.

See *Handbook* Section 4, 4.4.6; Section 5, 5.2

#### C49 Obtaining unpublished data

Highly desirable

Seek key unpublished information that is missing from reports of included studies.

Contacting study authors to obtain or confirm data makes the review more complete, potentially enhances precision and reduces the impact of reporting biases. Missing information includes details to inform risk of bias assessments, details of interventions and outcomes, and study results (including breakdowns of results by important subgroups).

See *Handbook* Section 5, 5.2.3

#### C50 Choosing interventions in multi-arm studies

Mandatory

If a study is included with more than two intervention arms, include in the review only the interventions that meet the eligibility criteria.

There is no point including irrelevant interventions in the review. **Authors, however, should make it clear in the 'Table of characteristics of included studies' that these interventions were present in the study.**

See *Handbook* Section 5, 5.3.6

#### C51 Checking accuracy of numeric data in the review

Mandatory

Compare magnitude and direction of effects reported by studies with how they are presented in the review, taking account of legitimate differences.

This is a reasonably straightforward way for authors to check a number of potential problems, including typographical errors **in studies' reports, accuracy of data collection and** manipulation, and data entry into RevMan. For example, the direction of a standardized mean difference may accidentally be wrong in the review. A basic check is to ensure the same qualitative findings (e.g. direction of effect and statistical significance) between the data as presented in the review and the data as available from the original study. Results in forest plots should agree with data in the original report (point estimate and confidence interval) if the same effect measure and statistical model is used.

## 1.8 Assessing risk of bias in included studies

Cochrane Training resources: [assessing RoB](#) and [RoB 2.0 webinar](#)

Cochrane Interactive Learning (CIL): [module 5 - introduction to study quality and risk of bias](#)

Standard		Rationale and elaboration
C52	Assessing risk of bias	Mandatory
Assess the risk of bias in at least one specific result for each included study. For randomized trials, the RoB 2 tool should be used, involving judgements and support for those judgements across a series of domains of bias, as described in the Handbook.		The risk of bias in at least one specific result for every included study must be explicitly considered to determine the extent to which its findings can be believed, noting that risks of bias might vary by result. The RoB 2 tool – as described in the Handbook – must be used for all randomized trials in new reviews. This does not prevent other tools being used. See <i>Handbook</i> Section 7, 7.1.2
C53	Assessing risk of bias in duplicate	Mandatory
Use (at least) two people working independently to apply the risk-of-bias tool to each result in each included study, and define in advance the process for resolving disagreements.		Duplicating the risk-of-bias assessment reduces both the risk of making mistakes and the possibility that assessments are <b>influenced by a single person's biases</b> . See <i>Handbook</i> Section 7, 7.3.2
C54	Supporting judgements of risk of bias	Mandatory
Justify judgements of risk of bias (high, low and some concerns) and provide this information in the risk-of-bias <b>tables (as 'Support for judgement')</b> .		Providing support for the judgement makes the process transparent. See <i>Handbook</i> Section 7, 7.3.2
C55	Providing sources of information for risk-of-bias assessments	Mandatory
Collect the source of information for each risk-of-bias judgement (e.g. quotation, summary of information from a trial report, correspondence with investigator etc.). Where judgements are based on assumptions made on the basis of information provided outside publicly available documents, this should be stated.		Readers, editors and referees should have the opportunity to see for themselves from where supports for judgements have been obtained. See <i>Handbook</i> Section 7, 7.3.2

C56	Ensuring results of outcomes included in ‘Summary of findings’ tables are assessed for risk of bias	Mandatory
<p>Ensure that assessments of risk of bias cover the outcomes included in the ‘Summary of findings’ table.</p> <p>It may not be feasible to assess the risk of bias in every single result available across the included studies, particularly if a large number of studies and results are available. Review author should strive to assess risk of bias in the results of outcomes that are most important to patients. Such outcomes <b>will typically be included in ‘Summary of findings’ tables, which present the findings of seven or fewer patient-important outcomes.</b></p> <p>See <i>Handbook</i> Section 7, 7.3.2</p>		
C57	Summarizing the risk-of-bias assessments	Highly desirable
<p>Summarize the risk of bias for each key outcome for each study</p> <p>This reinforces the link between the characteristics of the study design and their possible impact on the results of the study, and is an important prerequisite for the GRADE approach to assessing the certainty of the body of evidence.</p> <p>See <i>Handbook</i> Section 7, 7.5</p>		
C58	Addressing risk of bias in the synthesis	Highly desirable
<p>Address risk of bias in the synthesis (whether quantitative or non-quantitative). For example, present analyses stratified according to summary risk of bias, restricted to studies at low risk of bias, or restricted to low-and-some-concerns of bias.</p> <p>Review authors should consider how study biases affect results. This is useful in determining the strength of conclusions and how future research should be designed and conducted.</p> <p>See <i>Handbook</i> Section 7, 7.6.1</p>		
C59	Incorporating assessments of risk of bias	Mandatory
<p><i>If randomized trials have been assessed using one or more tools in addition to the RoB 2 tool, use the RoB 2 tool as the primary assessment of bias for interpreting results, choosing the primary analysis, and drawing conclusions.</i></p> <p>For consistency of approach across Cochrane Intervention Reviews, the RoB 2 tool should take precedence when two or more tools are used for assessing risk of bias in randomized trials. The RoB 2 tool also feeds directly into the GRADE approach for assessing the certainty of the body of evidence.</p> <p>See <i>Handbook</i> Section 7, 7.6.1</p>		
C60	Addressing conflicts of interest in included trials	Highly desirable

Address conflict of interests in included trials, and reflect on possible impact on: a) differences in study design; b) risk of bias in trial result, and c) risk of bias in synthesis result

Review authors should consider assessing whether they judge a **trial to be of “notable concern about conflicts of interest”**. This assessment is useful for exploration of possible heterogeneity between trials (e.g. in a subgroup analysis), and for reflection on relevant mechanisms for how conflict of interest may have biased trial results and synthesis results. Concerns about conflicts of interest can be **reported in the ‘Characteristics of included studies’ table**.

See *Handbook* Section 7, 7.8.6

## 1.9 Synthesizing the results of included studies

Cochrane Interactive Learning (CIL): [module 6 - analysing the data](#)

Standard		Rationale and elaboration	
C61	Combining different scales		Mandatory
<p><i>If studies are combined with different scales, ensure that higher scores for continuous outcomes all have the same meaning for any particular outcome; explain the direction of interpretation; and report when directions are reversed.</i></p>		<p>Sometimes scales have higher scores that reflect a <b>‘better’ outcome and sometimes lower scores reflect ‘better’ outcome</b>. Meaningless (and misleading) results arise when effect estimates with opposite clinical meanings are combined.</p>	
C62	Ensuring meta-analyses are meaningful		Mandatory
<p>Undertake (or display) a meta-analysis only if participants, interventions, comparisons and outcomes are judged to be sufficiently similar to ensure an answer that is clinically meaningful.</p>		<p>Meta-analyses of very diverse studies can be misleading, for example where studies use different forms of control. Clinical diversity does not indicate necessarily that a meta-analysis should not be performed. However, authors must be clear about the underlying question that all studies are addressing.</p>	
C63	Assessing statistical heterogeneity		Mandatory
<p>Assess the presence and extent of between-study variation when undertaking a meta-analysis.</p>		<p>The presence of heterogeneity affects the extent to which generalizable conclusions can be formed. It is important to identify heterogeneity in case there is sufficient information to explain it and offer new insights. Authors should recognize that there is much uncertainty in measures such as <math>I^2</math> and <math>\tau^2</math> when there are few studies. Thus, use of simple thresholds to diagnose heterogeneity should be avoided. See <i>Handbook</i> Section 10, 10.10.2</p>	
C64	Addressing missing outcome data		Highly desirable
<p>Consider the implications of missing outcome data from individual participants (due to losses to follow-up or exclusions from analysis).</p>		<p>Incomplete outcome data can introduce bias. In most circumstances, authors should follow the principles of intention-to-treat analyses as far as possible (this may not be appropriate for adverse effects or if trying to demonstrate equivalence). Risk of bias due to incomplete outcome data is addressed in the Cochrane risk-of-bias tool. However, statistical analyses and careful interpretation of results are additional ways in which the issue can be addressed by review authors. Imputation methods</p>	

can be considered (accompanied by, or in the form of, sensitivity analyses).

See *Handbook* Section 10, 12.1

C65	Addressing skewed data	Highly desirable
	Consider the possibility and implications of skewed data when analysing continuous outcomes.	Skewed data are sometimes not summarized usefully by means and standard deviations. While statistical methods are approximately valid for large sample sizes, skewed outcome data can lead to misleading results when studies are small. See <i>Handbook</i> Section 10, 10.5.3
C66	Addressing studies with more than two groups	Mandatory
	<i>If multi-arm studies are included</i> , analyse multiple intervention groups in an appropriate way that avoids arbitrary omission of relevant groups and double-counting of participants.	Excluding relevant groups decreases precision and double-counting increases precision spuriously; both are inappropriate and unnecessary. Alternative strategies include combining intervention groups, separating comparisons into different forest plots and using network meta-analysis. See <i>Handbook</i> Section 6.2.9 and Section 11.
C67	Comparing subgroups	Mandatory
	If subgroup analyses are to be compared, and there are judged to be sufficient studies to do this meaningfully, use a formal statistical test to compare them.	Concluding that there is a difference in effect in different subgroups on the basis of differences in the level of statistical significance within subgroups can be very misleading. See <i>Handbook</i> Section 10, 10.11.3.1
C68	Interpreting subgroup analyses	Mandatory
	<i>If subgroup analyses are conducted</i> , follow the subgroup analysis plan specified in the protocol without undue emphasis on particular findings.	Selective reporting, or over-interpretation, of particular subgroups or particular subgroup analyses should be avoided. This is a problem especially when multiple subgroup analyses are performed. This does not preclude the use of sensible and honest post hoc subgroup analyses. See <i>Handbook</i> Section 10, 10.11.5.2
C69	Considering statistical heterogeneity when interpreting the results	Mandatory
	Take into account any statistical heterogeneity when interpreting the results, particularly when there is variation in the direction of effect.	The presence of heterogeneity affects the extent to which generalizable conclusions can be formed. If a fixed-effect analysis is used, the confidence intervals ignore the extent of heterogeneity. If a random-effects analysis is used, the result pertains to the mean effect across studies. In both cases, the implications of notable heterogeneity should be addressed. It may be possible to understand the reasons for the heterogeneity if there are sufficient studies. See <i>Handbook</i> Section 10, 10.10.3
C70	Addressing non-standard designs	Mandatory
	Consider the impact on the analysis of clustering, matching or	Cluster-randomized trials, cross-over trials, studies involving measurements on multiple body parts, and other designs need to be addressed specifically, since a naive analysis might

other non-standard design features of the included studies.

underestimate or overestimate the precision of the study. Failure to account for clustering is likely to overestimate the precision of the study, that is, to give it confidence intervals that are too narrow and a weight that is too large. Failure to account for correlation is likely to underestimate the precision of the study, that is, to give it confidence intervals that are too wide and a weight that is too small.

See *Handbook* Section 6, 6.2.1

## C71 Sensitivity analysis

Highly desirable

Use sensitivity analyses to assess the robustness of results, such as the impact of notable assumptions, imputed data, borderline decisions and studies at high risk of bias.

It is important to be aware when results are robust, since the strength of the conclusion may be strengthened or weakened.  
See *Handbook* Section 10, 10.14

## C72 Interpreting results

Mandatory

Focus interpretation of results on estimates of effect and their confidence intervals, avoiding use of a distinction between **“statistically significant”** and **“statistically non-significant”**.

Authors commonly mistake a lack of evidence of effect as evidence of a lack of effect.

See *Handbook* Section 15, 15.3.1

## C73 Investigating risk of bias due to missing results

Highly desirable

Consider the potential impact of non-reporting biases on the results of the review or the meta-analyses it contains.

There is overwhelming evidence of non-reporting biases of various types. These can be addressed at various points in the review. A thorough search, and attempts to obtain unpublished results, might minimize the risk. Analyses of the results of included studies, for example using funnel plots, can sometimes help determine the possible extent of the problem, as can attempts to identify study protocols, which should be a routine feature of Cochrane Reviews.

See *Handbook* Section 13, 13.4

URL: <https://community.cochrane.org/mecir-manual/standards-conduct-new-cochrane-intervention-reviews-c1-c75/performing-review-c24-75/assessing-quality-evidence-and-summarizing-findings-c74-75>

## 1.10 Assessing the certainty of evidence and summarizing the findings



Cochrane Training resource: [GRADE approach to evaluating evidence quality](#)

Cochrane Interactive Learning: [module 7 - interpreting the findings](#)

Standard		Rationale and Elaboration
C74	Assessing the certainty of the body of evidence	Mandatory
<p>Use the five GRADE considerations (risk of bias, consistency of effect, imprecision, indirectness and publication bias) to assess the certainty of the body of evidence for each outcome, and to draw conclusions about the certainty of evidence within the text of the review.</p>		<p>GRADE is the most widely used approach for summarizing confidence in effects of interventions by outcome across studies. It is preferable to use the online GRADEpro tool, and to use it as described in the help system of the software. This should help to ensure that author teams are accessing the same information to inform their judgments. Ideally, two people working independently should assess the certainty of the body of evidence and reach a consensus view on any downgrading decisions. The five GRADE considerations should be addressed irrespective of <b>whether the review includes a ‘Summary of findings’ table. It is helpful to draw on this information in the Discussion, in the Authors’ conclusions and to convey the certainty in the evidence in the Abstract and Plain language summary.</b></p> <p>See <i>Handbook</i> Section 14, 14.2.1</p>
C75	Justifying assessments of the certainty of the body of evidence	Mandatory
<p>Justify and document all assessments of the certainty of the body of evidence (for example downgrading or upgrading if using GRADE).</p>		<p>The adoption of a structured approach ensures transparency in formulating an interpretation of the evidence, and the result is more informative to the user.</p> <p>See <i>Handbook</i> Section 14, 14.2.1</p>

URL: <https://community.cochrane.org/mecir-manual/standards-conduct-new-systematic-reviews/performing-review/reference>

## Reference

Hoffmann TC, Glasziou PP, Boutron I, Milne R, Perera R, Moher D, et al. (2014) Better reporting of interventions: template for intervention description and replication (TIDieR) checklist and guide. *BMJ* 2014;348:g1687. doi: 10.1136/bmj.g1687

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URL: <https://community.cochrane.org/mecir-manual/standards-reporting-protocols-new-cochrane-intervention-reviews-pr1-44>

## STANDARDS FOR THE REPORTING OF PROTOCOLS FOR NEW COCHRANE INTERVENTION REVIEWS

Toby Lasserson, Rachel Churchill, Jackie Chandler, David Tovey, James Thomas, Ella Flemyng and Julian PT Higgins

URL: <https://community.cochrane.org/mecir-manual/standards-reporting-protocols-new-cochrane-intervention-reviews/key-points-introduction>

## Key points and introduction

Key points:

- Publishing a protocol for a Cochrane Review establishes a public record of the review question and planned methods.
- Reporting clear definitions will help authors to adhere to a well formulated approach.
- Readers need to determine how far the review will address their own questions of interest.
- Changes to the review question or methods will need to be clearly described and justified in the full review.

Publishing the protocol for a Cochrane Systematic Review is a key milestone in the review process. As with any other form of research, it finalizes the development of the research question and sets out the different methods that will be used to address it.

Preparing and publishing a clearly conceived and well-written protocol serves a number of purposes. Investment of effort in the development of the review question and methods and the definition of the different aspects of the eligibility criteria will provide review authors with a clear plan to guide implementation of methods and reporting the full review, reducing their reliance on post hoc decisions. Publishing the protocol gives readers access to the plan from which the review will develop. It also helps them to judge how the eligibility criteria of the review, stated outcomes and planned methods will address the intended question of interest.

The protocol is a public record of the question of interest and the intended methods before results of the studies are fully known. This helps anyone who evaluates the review to judge how far it fulfills the original objectives. One of the key parts of the Cochrane Review prepublication screening programme involves the comparison between the intended methods with those implemented during the preparation of the review. It is crucial that review authors acknowledge and justify important differences between methods stated in the protocol and those used to produce the review findings. This is key to supporting replication, and provides users of the review with a sense of how far the review preserves the research question. Particularly important changes concern eligibility criteria, the definition or status of outcome measurements and methods relating to effect measures, data analysis and exploration of heterogeneity. Any changes that are made to these aspects of the review could potentially impact on the overall objectives as well as the interpretation of the evidence summarized by the review.

On publication Cochrane systematic review protocols are automatically assigned a record on PROSPERO, the register of ongoing and completed systematic reviews. For more information see [www.crd.york.ac.uk/PROSPERO/](http://www.crd.york.ac.uk/PROSPERO/)

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URL: <https://community.cochrane.org/mecir-manual/standards-reporting-protocols-new-cochrane-intervention-reviews-pr1-44/reporting-review-plan-pr1-44>

## Reporting the review plan

Cochrane Training resources: [writing the protocol](#) and [common errors in protocols](#)

Cochrane Interactive Learning: [module 2 - writing the review protocol](#)

URL: <https://community.cochrane.org/mecir-manual/standards-reporting-protocols-new-cochrane-intervention-reviews-pr1-44/reporting-review-plan-pr1-44/title-and-authors-pr1-2>

## 1.11 Title and authors

Cochrane Interactive Learning: [module 2 - writing the review protocol](#)

Standard		Rationale and Elaboration
PR1	Format of title	Highly desirable
	Follow the standard template for a Cochrane Review title.	See <i>Handbook</i> <a href="#">Section II.1.3</a>
PR2	Authors	Mandatory
	List names and affiliations of all authors.	See <i>Handbook</i> <a href="#">Section II.2</a>

## 1.12 Background

Cochrane Training resource: [writing the protocol](#)

Cochrane Interactive Learning: [module 2 - writing the review protocol](#)

Standard		Rationale and elaboration
PR3	Background	Mandatory
Provide a concise description of the condition or problem addressed by the review question, a definition of the intervention and how it might work, and why it is important to do the review. Include the four standard RevMan headings when writing the Background.		<p>Systematic reviews should have a clearly defined and well-reasoned rationale that has been developed in the context of existing knowledge. Outlining the context of the review question is useful to readers and helps to establish key uncertainties that the review intends to address.</p> <p><b>Four standard headings are included in RevMan ('Description of the condition', 'Description of the intervention', 'How the intervention might work', and 'Why it is important to do this review').</b>  <i>See Handbook <a href="#">Section III.3.2</a></i></p>
PR4	Background references	Mandatory
Back up all key supporting statements with references.		Claims or statements regarding aspects such as disease burden, morbidity, prevalence and mechanisms of action should be substantiated and, where available, supported by evidence.



## 1.13 Objectives

Cochrane Training resource: [defining the review question](#) and [writing the protocol](#)

Cochrane Interactive Learning: [module 2 - writing the review protocol](#)

Standard		Rationale and elaboration
PR5	Main objective	Mandatory
State the main objective, where appropriate in a single concise sentence.		<p>The primary objective of a Cochrane Review should be to assess the effects of one or more healthcare interventions on user-important outcomes, both intended and unintended. The objective should be expressed in terms that relate to the population(s), intervention comparison(s) and, where appropriate to specify explicitly, the outcomes of interest (PICO). Review users may be patients, carers, policy makers, clinicians, practitioners or others.</p> <p><b>The format should be:</b> “To assess the effects of [intervention or comparison] for [health problem] for/in [types of people, disease or problem and setting if specified]”.</p> <p><i>MECIR conduct standard 2:</i> Define in advance the objectives of the review, including participants, interventions, comparators and outcomes (PICO). See <i>Handbook</i> <a href="#">Section III.3.2</a> and <a href="#">Section 2.3</a></p>
PR6	Secondary objectives	Highly desirable
State explicitly (as secondary objectives) any specific questions being addressed by the review, such as those relating to particular participant groups, intervention comparisons or outcome.		<p>The secondary objectives should be expressed in terms that relate to the population(s), intervention comparison(s) and, where appropriate, outcomes of interest.</p> <p><b>The format might be:</b> “To assess whether the effects of [intervention or comparison] differ according to [types of people, intervention or comparator characteristic, disease, problem, setting etc.]”.</p> <p>Secondary objectives should be kept succinct, since they will be published in the front sheet of the review protocol on the Cochrane Library.</p> <p><i>MECIR conduct standard 4:</i> Consider in advance whether issues of equity and relevance of evidence to specific populations are important to the review, and plan for appropriate methods to address them if they are. Attention should be paid to the relevance of the review question to</p>

populations such as low-socioeconomic groups, low- or middle-income regions, women, children and older people.  
See *Handbook* [Section III.3.2](#) and [Section 2.4](#)

## PR7 Economic evidence

Mandatory

*If health economics evidence is to be reviewed, state this explicitly in the Objectives (as a secondary objective).*

The primary aim of a Cochrane Review should be to assess the effects of one or more healthcare interventions on outcomes, both intended and unintended, that are important to review users. These outcomes may include economic outcomes. If health economics evidence is being reviewed as an integrated economics component, this should be stated as a secondary objective.

See *Handbook* [Section 20.2.2](#)

## PR8 Qualitative research evidence

Mandatory

*If qualitative research evidence is to be reviewed, state this explicitly in the Objectives (as a secondary objective).*

If qualitative research evidence is being **included to 'extend'** the review, this should be stated as a secondary objective.

See *Handbook* [Section 21.4](#)

## 1.14 Criteria for considering studies for this review

Cochrane Training resource: [defining the review question](#)

Cochrane Interactive Learning: [module 2 - writing the review protocol](#)

Standard		Rationale and elaboration
PR9	Eligibility criteria for types of study: study designs	Mandatory
State eligible study designs, using key study characteristics, and provide a justification for the choice.		<p>It is not necessary to explain why randomized trials are eligible (if that is the case), although it may be important to explain the eligibility or non-eligibility of other types of study.</p> <p>Particular care may be needed to explain whether cross-over trials and cluster-randomized trials are to be considered.</p> <p><b>Study characteristics might include details such as “with blind assessment of outcomes” or “with prospective identification of participants”, rather than ambiguous labels such as “double blind” or “prospective study”.</b></p> <p><b>If ‘conditional’ eligibility criteria are used that are based on absence of particular types of evidence (e.g. when no randomized trials are found), this must be stated unambiguously (and detailed methods for addressing all <i>potentially</i> eligible studies will need to be described).</b></p> <p><i>MECIR conduct standard 9:</i> Define in advance the eligibility criteria for study designs in a clear and unambiguous way, with a focus on features of a study's design rather than design labels.</p> <p><i>MECIR conduct standard 11:</i> Justify the choice of eligible study designs. See <i>Handbook Section III.3.3.1</i> and Section 3.3.3</p>
PR10	Eligibility criteria for types of study: study reports	Mandatory
<i>If studies will be excluded on the basis of publication status or language of publication, explain and justify this.</i>		<p>Studies should be included irrespective of their publication status and language of publication, unless exclusion is explicitly justified.</p> <p><i>MECIR conduct standard 12:</i> Include studies irrespective of their publication status, unless exclusion is explicitly justified. See <i>Handbook Section III.3.3.1</i> and Section 3.3.4</p>

PR11	Eligibility criteria for types of participants	Mandatory
	State eligibility criteria for participants, including any criteria around location, setting, diagnoses or definition of condition and demographic factors, and how studies including subsets of relevant participants will be addressed.	<p><i>MECIR conduct standard 5:</i> Define in advance the eligibility criteria for participants in the studies.</p> <p><i>MECIR conduct standard 6:</i> Define in advance how studies that include only a subset of relevant participants will be addressed.</p> <p>See <i>Handbook</i> <a href="#">Section III.3.3.1</a> and Section 3.2.1</p>
PR12	Eligibility criteria for types of interventions	Mandatory
	State eligibility criteria for interventions and comparators, including any criteria around delivery, dose, duration, intensity and cointerventions. Criteria for complex interventions should be made explicit, e.g. by stating mandatory components.	<p>Eligible interventions, and particularly the comparators, must address the stated objectives of the review. For example, inclusion of studies with an active comparator intervention is not consistent with an objective to look only at whether an experimental intervention is effective compared with an inactive control.</p> <p><i>MECIR conduct standard 7:</i> Define in advance the eligible interventions and the interventions against which these can be compared in the included studies.</p> <p>See <i>Handbook</i> <a href="#">Section III.3.3.1</a> and Section 3.2.2</p>
PR13	Role of outcomes	Mandatory
	Be explicit about the role of outcomes in determining eligibility of studies for the review.	<p>For most Cochrane Reviews of randomized trials of the intended effects of interventions, the aim should be to identify and include all relevant participants who have been randomized to the intervention comparisons of interest. The extent to which outcome data are available for these people can be affected by decisions made by the trialists – i.e. there is a risk of selective outcome reporting bias.</p> <p>An important distinction should be made between whether outcomes were measured, and whether the measured outcome data are available. Studies should not be excluded from a review solely because no outcome data are available. However, on occasion it will be appropriate to include only studies that <i>measured</i> particular outcomes. For example, a review of a multi-component public health intervention promoting healthy lifestyle choices, focussing on reduction in smoking prevalence, might legitimately exclude studies that do not measure smoking rates. Often it is difficult to know whether unreported outcomes were measured, so it is generally appropriate to include all studies irrespective of whether outcomes are reported.</p> <p><i>MECIR conduct standard 8:</i> Clarify in advance whether outcomes listed under 'Criteria for considering studies for this review' are used as criteria for including studies (rather</p>

than as a list of the outcomes of interest within whichever studies are included).

See *Handbook* [Section III.3.3.1](#) and [Section 3.2.4.1](#)

#### PR14 Outcome domains of interest

Mandatory

Define in advance outcomes that are critical to the review, and any additional important outcomes.

Up to seven outcomes should be prespecified for inclusion in a ‘**Summary of findings**’ table (see PR40); it may be convenient to highlight them here.

*MECIR conduct standard 14:* Define in advance outcomes that are critical to the review, and any additional important outcomes.

Also *MECIR conduct standards 15–18*

See *Handbook* [Section III.3.3.1](#) and [Section 3.2.4.1](#)

#### PR15 Outcome measures of interest

Mandatory

Define relevant outcome measures and time points for measurement, and any hierarchy for choosing among them.

Explain how multiple variants of outcome measures (e.g. definitions, assessors, scales, time points) will be addressed.

#### PR16 Minimally important difference

Highly desirable

Define minimally important differences for key outcome measures.

To facilitate interpretation of the size of effect of an intervention, it is important to understand the size of difference that is important to review users.

URL: <https://community.cochrane.org/mecir-manual/standards-reporting-protocols-new-cochrane-intervention-reviews-pr1-44/reporting-review-plan-pr1-44/search-methods-identification-studies-pr17-21>

## 1.15 Search methods for identification of studies

Cochrane Training resource: [searching for studies](#)

Cochrane Interactive Learning: [module 3 - searching for studies](#)

Standard		Rationale and elaboration
PR17	Search sources	Mandatory
List all sources that will be searched, including: CRG specialized register(s), CENTRAL, other databases, trials registers, websites and grey literature. State whether reference lists will be searched and whether individuals or organizations will be contacted.		<p><i>MECIR conduct standard 19:</i> Plan in advance the methods to be used for identifying studies. Design searches to capture as many studies as possible that meet the eligibility criteria, ensuring that relevant time periods and sources are covered and not restricted by language or publication status.</p> <p><i>MECIR conduct standard 36:</i> Document the search process in enough detail to ensure that it can be reported correctly in the review.</p> <p>Also <i>MECIR conduct standards 24–31</i> See <i>Handbook</i> <a href="#">Section III.3.3.2</a>, <a href="#">Section 1.5</a>; <a href="#">4.3.1.1</a> and <a href="#">Section 4.4.5</a></p>
PR18	Search restrictions	Mandatory
Specify and justify any restrictions to be placed on the search (e.g. time period or publication format).		<p><i>MECIR conduct standard 35:</i> Justify the use of any restrictions in the search strategy on publication date or publication format.</p> <p>See <i>Handbook</i> <a href="#">Section III.3.3.2</a> and <a href="#">Section 4.4.5</a></p>
PR19	Searches for different types of evidence	Mandatory
Some reviews extend beyond a focus on the effects of healthcare interventions and address specific additional types of evidence.		<p><i>MECIR conduct standard 26:</i> If the review has specific eligibility criteria around study design to address adverse effects, economic issues or qualitative research questions, undertake searches to address them.</p> <p>These are discussed in the <i>Handbook</i> Chapters <a href="#">19</a>, <a href="#">20</a> and <a href="#">21</a>.</p>
PR20	Search strategies for bibliographic databases	Mandatory

Present the complete search strategy (or strategies) to be implemented for at least one database in an Appendix, including any limits and filters to be used.

The line-by-line search string should be presented to facilitate peer review. Search strategies that are available elsewhere (e.g. standard methodological filters, or strategies used to populate a specialized register) may be referenced rather than reproduced. Note that when the full review is published, it is mandatory to report search strategies used for all databases.

*MECIR conduct standard 36:* Document the search process in enough detail to ensure that it can be reported correctly in the review.

Also *MECIR conduct standards 32–35*

See *Handbook* [Section III.3.3.2](#) and [Section 4.4.5](#)

PR21 Search strategies for other sources

Highly desirable

Report search terms that will be used to search any sources other than bibliographic databases (e.g. trials registers, the web).

Some of this information might be best placed in an Appendix.

*MECIR conduct standard 36:* Document the search process in enough detail to ensure that it can be reported correctly in the review.

See *Handbook* [Section III.3.3.2](#) and [Section 4.4.5](#)

## 1.16 Data collection and analysis

Standard		Rationale and elaboration
PR22	Inclusion decisions	Mandatory
State how inclusion decisions will be made (i.e. from search results to included studies), clarifying how many people will be involved and whether they will work independently.		<p><i>MECIR conduct standard 39:</i> Use (at least) two people working independently to determine whether each study meets the eligibility criteria, and define in advance the process for resolving disagreements.</p> <p>See Handbook <a href="#">Section III.3.3.3</a> and <a href="#">Section 4.4.10</a></p>
PR23	Data collection process	Mandatory
State how data will be extracted from reports of included studies, clarifying how many people will be involved (and whether they will work independently), and how disagreements will be resolved.		<p><i>MECIR conduct standard 43:</i> Use a data collection form that has been piloted.</p> <p><i>MECIR conduct standard 45:</i> Use (at least) two people working independently to extract study characteristics from reports of each study, and define in advance the process for resolving disagreements.</p> <p>See Handbook <a href="#">Section III.3.3.3</a>, <a href="#">Section 5.4.1</a> and <a href="#">Section 5.5.2</a></p>
PR24	Requests for data	Highly desirable
Describe what attempts will be made to obtain or clarify data from individuals or organizations.		<p><i>MECIR conduct standard 49:</i> Seek key unpublished information that is missing from reports of included studies.</p> <p>See Handbook <a href="#">Section III.3.3.3</a> and <a href="#">Section 5.2.3</a></p>
PR25	Data items	Mandatory
State the types of information that will be sought from reports of included studies.		<p>This information is a useful basis for the design of data collection forms and also indicates what sort of information about the included studies readers might anticipate seeing in the full text of the review. Detailed lists are not necessary. Instead, a broad outline of the summary information that authors might collect will suffice, for example:</p> <p><b>“We will collect information on study design and setting, participant characteristics (including disease severity and age), study eligibility criteria, details of the intervention(s) given, the outcomes assessed, the source of study funding and any conflicts of interest stated by the investigators.”</b></p> <p><i>MECIR conduct standard 44:</i> Collect characteristics of the included studies in sufficient detail to populate a table of ‘<b>Characteristics of included studies</b>’.</p> <p>See Handbook <a href="#">Section III.3.3.3</a> and <a href="#">Section 5.3.1</a></p>



PR26	Missing data	Highly desirable
<p>Comment on how missing data will be addressed.</p> <p>Briefly describe any planned strategies that will be used to address missing data. This might include imputation of missing outcome data for individuals within studies (such as worst-case or best-case scenarios), or imputations of missing standard deviations. Note that standard deviations can sometimes be computed from other reported statistics.</p> <p><i>MECIR conduct standard 47:</i> Collect and utilize the most detailed numerical data that might facilitate similar analyses of included studies. Where 2×2 tables or means and standard deviations are not available, this might include effect estimates (e.g. odds ratios, regression coefficients), confidence intervals, test statistics (e.g. t, F, Z, Chi<sup>2</sup>) or P values, or even data for individual participants.</p> <p><i>MECIR conduct standard 64:</i> Consider the implications of missing outcome data from individual participants (due to losses to follow-up or exclusions from analysis).</p> <p>See <i>Handbook</i> <a href="#">Section III.3.3.3</a>, <a href="#">Section 5.3.6</a> and <a href="#">Section 10.12.1</a></p>		
PR27	Tools to assess risk of bias in individual studies	Mandatory
<p>State and reference the tool(s) that will be used to assess risk of bias for included studies, how the tool(s) will be implemented, and the criteria that will be used to assign study results to judgements of low risk, high risk and unclear risk of bias.</p> <p>Different tools are likely to be appropriate for different types of studies (e.g. randomized trials and non-randomized studies). If the current <i>Handbook</i> guidance for undertaking ‘Risk of bias’ assessments will be followed in its entirety, then a reference to the <i>Handbook</i> is sufficient to provide the criteria used to assign judgements. Justify any intended deviations from the tool.</p> <p><i>MECIR conduct standard 20:</i> Plan in advance the methods to be used for assessing risk of bias in included studies, including the tool(s) to be used, how the tool(s) will be implemented, and the criteria used to assign study results to judgements of low risk, high risk and unclear risk of bias.</p> <p><i>MECIR conduct standard 52:</i> Assess the risk of bias in at least one specific result for each included study. For randomized trials, the RoB 2 tool should be used, involving judgements and support for those judgements across a series of domains of bias, as described in the <i>Handbook</i>.</p> <p>Also <i>MECIR conduct standards 53–60</i> See <i>Handbook</i> <a href="#">Section III.3.3.3</a>, <a href="#">Section 7.1.2</a> and <a href="#">Chapter 8</a></p>		
PR28	‘Risk of bias’ assessment process	Mandatory

State how risk of bias will be assessed, clarifying how many people will be involved (and whether they will work independently), and how disagreements will be resolved.	<p><i>MECIR conduct standard 53:</i> Use (at least) two people working independently to apply the ‘Risk of bias’ tool to each included study, and define in advance the process for resolving disagreements.</p> <p>If the Risk of Bias 2 tool (see <i>Handbook Chapter 8</i>) is to be used, state whether interest will be in the effect of assignment to intervention or the effect of adhering to intervention, and explain how results will be selected to be assessed for risk of bias (i.e. for which outcome domains, outcome measures, time points and analyses).</p> <p>See <i>Handbook Section III.3.3.3</i>, <a href="#">Section 7.3.2</a> and <a href="#">Chapter 8</a></p>
<b>PR29 Measures of effect</b>	<b>Mandatory</b>
State the effect measures that will be used to describe effect sizes in any included studies or meta-analyses, or both (e.g. risk ratio or odds ratio, mean difference or standardized mean difference).	See <i>Handbook Section III.3.3.3</i>
<b>PR30 Unit of analysis issues</b>	<b>Mandatory</b>
<i>If designs other than individually randomized, parallel-group randomized trials are likely to be included</i> , describe any methods that will be used to address clustering, matching or other design features of the included studies.	<p>In some circumstances, specific study designs are likely to be identified in which unit-of-analysis errors might arise. This includes cluster-randomized trials, cross-over trials, trials involving multiple body parts and non-randomized studies with clustered designs.</p> <p><i>MECIR conduct standard 70:</i> Consider the impact on the analysis of clustering, matching or other non-standard design features of the included studies.</p> <p>See <i>Handbook Section III.3.3.3</i> and <a href="#">Section 6.2.1</a></p>
<b>PR31 Studies with more than two groups</b>	<b>Highly desirable</b>
<i>If multi-arm studies are likely to be included</i> , explain how they will be addressed and incorporated into syntheses.	<p>Note that it is mandatory to describe these methods in the full version of the review if studies with more than one arm are identified and included.</p> <p><i>MECIR conduct standard 66:</i> <i>If multi-arm studies are included</i>, analyse multiple intervention groups in an appropriate way that avoids arbitrary omission of relevant groups and double-counting of participants.</p> <p>See <i>Handbook Section III.3.3.3</i>, <a href="#">Section 6.2.9</a> and <a href="#">Chapter 11</a>.</p>
<b>PR32 Quantitative synthesis</b>	<b>Mandatory</b>
Describe any intended statistical methods for combining results across studies (e.g. meta-analysis, subgroup analysis, meta-regression, sensitivity analysis), including methods for assessing heterogeneity (e.g. $I^2$ , $\text{Tau}^2$ , statistical test).	In the majority of reviews, most of this information is <b>located under the subheading ‘Data synthesis’</b> . Note, however, that additional subheadings should be used to provide details of subgroup analyses, assessment of heterogeneity and sensitivity analysis.

*MECIR conduct standard 21:* Plan in advance the methods to be used to synthesize the results of the included studies, including whether a quantitative synthesis is planned, how heterogeneity will be assessed, choice of effect measure (e.g. odds ratio, risk ratio, risk difference or other for dichotomous outcomes), and methods for meta-analysis (e.g. inverse variance or Mantel Haenszel, fixed-effect or random-effects model).

*MECIR conduct standard 62:* Undertake (or display) a meta-analysis only if participants, interventions, comparisons and outcomes are judged to be sufficiently similar to ensure an answer that is clinically meaningful.

*MECIR conduct standard 63:* Assess the presence and extent of between-study variation when undertaking a meta-analysis.

See Handbook [Section III.3.3.3](#), [Section 1.5](#) and [Section 10.10.2](#)

PR33	Non-quantitative synthesis	Mandatory
	Describe any intended non-statistical methods for synthesizing findings across studies (sometimes referred to as narrative or qualitative synthesis).	It may be apparent that a meta-analysis is unlikely, in which case methods should be prespecified for how the findings of the included studies will be compared and contrasted. See Handbook <a href="#">Chapter 12</a>
PR34	Risk of reporting bias across studies	Highly desirable
	Describe any methods that will be used for assessing the risk of reporting biases such as publication bias.	See Handbook <a href="#">Chapter 13</a>
PR35	Addressing risk of bias	Mandatory
	Describe how studies with high or variable risks of bias will be addressed in the synthesis.	Several options are available for addressing risk of bias in a synthesis, including reporting separate syntheses for studies at different risks of bias, restricting analysis to studies at low (or low and unclear) risk of bias only, and undertaking sensitivity analysis to examine the impact of risks of bias on the conclusions. An understanding of the impact of risks of bias is important to inform GRADE assessments.  <i>MECIR conduct standard 58:</i> Address risk of bias in the synthesis (whether quantitative or non-quantitative). For example, present analyses that are stratified according to summary risk of bias, restricted to studies at low risk of bias or restricted to low-and-some-concerns of risk of bias. See Handbook <a href="#">Section 7.6.1</a> and <a href="#">Chapter 8</a>
PR36	Subgroup analyses	Mandatory

<i>If subgroup analysis (or meta-regression) are planned, state the potential effect modifiers with rationale for each.</i>		<i>MECIR conduct standard 22: Predefine potential effect modifiers (e.g. for subgroup analyses) at the protocol stage, restrict these in number, and provide rationale for each.</i> <i>See Handbook <a href="#">Section III.3.3.3</a> and</i>
PR37	Methods for economic evidence	Mandatory
<i>If health economics evidence is to be reviewed, state the methods to be used to assess and synthesize this evidence.</i>		<i>See Handbook <a href="#">Chapter 20</a></i>
PR38	Methods for qualitative research evidence	Mandatory
<i>If qualitative research evidence is to be reviewed, state the methods to be used to assess and synthesize this evidence.</i>		<i>See Handbook <a href="#">Chapter 21</a></i>
PR39	Certainty of the evidence	Mandatory
State the methods to be used to assess the certainty of the body of evidence (using the five GRADE considerations).		If the current GRADE guidance for these assessments will be followed in its entirety (see <i>Handbook <a href="#">Chapter 14</a></i> ), then a reference to this is sufficient to provide the criteria used to make judgements.  <i>MECIR conduct standard 74: Use the five GRADE considerations (risk of bias, consistency of effect, imprecision, indirectness and publication bias) to assess the certainty of the body of evidence for each outcome, and to draw conclusions about the certainty of evidence within the text of the review.</i> <i>See Handbook <a href="#">Section III.3.3.3</a> and <a href="#">Section 14.2.1</a></i>
PR40	'Summary of findings' table	Mandatory
State which outcomes and comparisons it is planned will be included in a 'Summary of findings' table.		A maximum of seven important outcomes should be <b>prespecified for inclusion in a 'Summary of findings' table</b> (see <i>Handbook <a href="#">Chapter 14</a></i> ). If possible, sources of any <b>assumed risks to be presented in a 'Summary of findings' table</b> should be explained.  <i>MECIR conduct standard 23: Plan in advance the methods to be used for assessing the certainty of the body of evidence, and summarizing the findings of the review.</i> <i>See Handbook <a href="#">Section III.3.3.3</a> and <a href="#">Section 1.5</a></i>

# 1.17 Acknowledgements

PR41	Acknowledgements	Mandatory
	Acknowledge the contribution of people not listed as authors of the protocol, including any assistance from the Cochrane Review Group, non-author contributions and the role of any funders.	See <i>Handbook</i> <a href="#">Section III.3.7</a>

1.18 Contribution of authors

PR42	Contributions of authors	Mandatory
Describe the contributions of each author to the protocol.		See <i>Handbook</i> <a href="#">Section III.3.7</a>

## 1.19 Declarations of interest

PR43	Declarations of interests	Mandatory
	Report relevant present or recent (three years prior to declaration) affiliations or other involvement in any organization or entity with an interest <b>in the review's findings that might lead to a real or perceived conflict of interest.</b>	<p>The detailed policy for declaring relevant interests is available in the Cochrane Editorial and Publishing Policy Resource (EPPR). In brief, the nature and extent of the affiliation or involvement (whether financial or non-financial) should be described. Declarations of interest should be stated according to the relevant criteria from the International Committee of Medical Journal Editors (ICMJE), and must be consistent with interests declared on the Disclosure of Potential Conflicts of Interest form.</p> <p>See <i>Handbook</i> <a href="#">Section III.3.7</a></p>

## 1.20 Sources of support

PR44	Sources of support	Mandatory
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List sources of financial and non-financial support for the review and the role of the funder, if any.

See *Handbook* [Section III.3.7](#)



URL: <https://community.cochrane.org/mecir-manual/standards-reporting-protocols-new-cochrane-intervention-reviews-pr1-44/citation>

## Citation of the standards for reporting of protocols of new Cochrane Intervention Reviews

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## STANDARDS FOR THE REPORTING OF NEW COCHRANE INTERVENTION REVIEWS

Rachel Churchill, Toby Lasserson, Jackie Chandler, David Tovey, James Thomas, Ella Flemyng and Julian PT Higgins

URL: <https://community.cochrane.org/mecir-manual/standards-reporting-new-cochrane-intervention-reviews/key-points-introduction>

## Key points and introduction

Key points:

- Authors should consult the MECIR reporting standards before and during writing up of their review.
- The reporting standards are compatible with key reporting guidelines developed by different bodies, including PRISMA.
- Abstracts and Plain language summaries need to be consistent with each other, and with the main text of the review.
- Clear and consistent reporting supports replication of systematic reviews and should make updating easier.

Authors should consult these reporting standards before and during writing up of their review. Adherence to the standards will help authors to prepare an informative, readable review. It will also help to make editorial evaluation of their work efficient. It is especially important to declare and justify differences to the planned question or eligibility criteria, since these may indicate important changes to the scope of the review. Where any search, data collection and analysis methods used are different from those planned, this also needs to be reported and explained. The reporting standards are available from within Review Manager (RevMan) software according to the heading or subheading to which they relate.

Several reporting guidelines are already available for primary studies and systematic reviews, and have been compiled by the Equator Network<sup>[1]</sup>. MECIR Standards are compatible with the core items in two key sources of reporting guidance for systematic reviews: the Preferred Reporting Items for Systematic reviews and Meta-Analyses (PRISMA), and the US Institute of Medicine's standards for systematic reviews.

Accurately summarizing the key findings of a Cochrane Review in its Abstract and Plain language summary serves an important purpose in knowledge translation. These standalone summaries help to convey the results of the review to a broad audience. Authors should take particular care to ensure that **conclusions drawn in the main text of the review under 'Implications for practice' and 'Implications for research' take account of the strength of evidence presented in the review, and are appropriately distilled in the Abstract and Plain language summary.**

Authors and editors should ensure that all parts of the review are succinct and readable, so that someone who is not an expert in the area can understand it. The published review needs to signpost and structure information clearly to help orientate readers. Review methods should be reported in sufficient detail that others are in principle able to reproduce the findings. Clear reporting of the eligibility criteria and methods will also help future efforts to update and maintain the published version of the review.

Rachel Churchill  
Professor of Evidence Synthesis and Co-ordinating Editor  
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<sup>[1]</sup> The Equator Network is a Library for health research reporting that provides a searchable database and can be found at <http://www.equator-network.org/>

## Reporting review conduct

## 1.21 Title and Authors

Standard		Rationale and elaboration
R1	Format of title	Highly desirable
Follow the standard template for a Cochrane Review title.		See <i>Handbook</i> <a href="#">Section II.1.3</a>
R2	Authors	Mandatory
List names and affiliations of all authors.		See <i>Handbook</i> <a href="#">Section II.2</a>

## 1.22 Abstract

Cochrane Training resource: [common errors - summary versions of a review](#)

Cochrane Interactive Learning: [module 8 - reporting the review](#)

Standard		Rationale and elaboration
R3	Writing the Abstract	Mandatory
	Prepare a structured Abstract to provide a succinct summary of the review. In the interests of brevity it is highly desirable for authors to provide an Abstract of less than 700 words, and it should be no more than 1000 words in length.	Abstracts are a prominent, publically accessible summary of the review that need to stand alone. They should convey key information about the review question and its findings, and be informative to readers.
R4	Abstract, Background	Mandatory
	Summarize the rationale and context of the review.	See <i>Handbook</i> <a href="#">Section III.3.1</a>
R5	Abstract, Objectives	Mandatory
	State the main objective(s), preferably in a single concise sentence.	The objective(s) should be expressed in terms that relate to the population(s), intervention comparison(s) and, where appropriate, outcomes of interest. See <i>Handbook</i> <a href="#">Section III.3.1</a>
R6	Abstract, Search methods	Mandatory
	Provide the date of the last search from which records were evaluated and that any studies identified were incorporated into the review, and an indication of the databases and other sources searched.	Abstracts should aim to give readers brief, but key, information about the comprehensiveness of the search and the currency of the information summarized by the review.  The Abstract must include the month and year of the set of searches up to which the conclusions of the review are valid. This date should reflect the date of the most recent set of searches from which all records have been screened for relevance and any studies meeting the eligibility criteria have been fully incorporated into the review (studies may be awaiting classification if, for example, the review authors are awaiting translation or clarification from authors or sponsors).  <b>Abstracts do not need to report on recent repeat or ‘catch-up’ searches whose results have not been fully incorporated into the review. However, discretion should be applied if such searches identify a large body of evidence, the absence of which may affect the reliability of the conclusions.</b>

The amount of information regarding the search should be indicative of the process rather than provide specific details. In the interests of brevity certain details regarding the overall process may need to be moved to the full text of the review.

**Example:** “CENTRAL, MEDLINE, Embase, five other databases and three trials registers were searched on [date] together with reference checking, citation searching and contact with study authors to identify additional studies”.

See *Handbook* [Section III.3.1](#)

R7	Abstract, Selection criteria	Mandatory
	Summarize eligibility criteria of the review, including information on study design, population and comparison.	Any extensions to eligibility criteria to address adverse effects, economic issues or qualitative research should be mentioned. See <i>Handbook</i> <a href="#">Section III.3.1</a>
R8	Abstract, Data collection and analysis	Mandatory
	Summarize any noteworthy methods for selecting studies, collecting data, evaluating risk of bias and synthesizing findings. For many reviews it may be sufficient to state “We used standard methodological procedures expected by Cochrane.”	<p>This section of the Abstract should indicate the rigour of the methods that underpin the results reported subsequently in the Abstract. It does not need to replicate the detailed description of the methods given in the main text of the review.</p> <p>Details of how many people were involved in the screening process and collection of information about any included studies are not necessary in the Abstract. Key statistical methods may be given if not clear from the results that follow.</p> <p>The Abstract should prioritize the disclosure of non-standard approaches. For example, rather than disclosing all domains applied in the assessment of bias, notable variations on the standard approach should be given, such as use of non-standard tools.</p> <p>See <i>Handbook</i> <a href="#">Section III.3.1</a></p>
R9	Abstract, Main results: number of studies and participants	Mandatory
	Report the number of included studies and participants.	<p>The total number of included studies should be stated. It might be appropriate to provide numbers of studies and participants for specific comparisons and main outcomes if the amount of evidence differs substantially from the total. Numbers of participants <i>analysed</i> should generally be presented in preference to numbers <i>recruited</i> (e.g. randomized); it is important to be clear which numbers are being reported. For some types of data there may be preferable alternatives to the number of participants (e.g. person-years of follow-up, number of limbs).</p> <p>See <i>Handbook</i> <a href="#">Section III.3.1</a></p>
R10	Abstract, Main results: study characteristics	Highly desirable

Provide a brief description of key characteristics that will determine the applicability of the body of evidence (e.g. age, severity of condition, setting, study duration).

Summarizing the study characteristics will provide readers of the Abstract with important information about the applicability of the included studies. This is particularly important if the included studies reflect a subgroup of those eligible for inclusion in the review, for example, if the review intended to address the effects of interventions across all age groups, but included studies that only recruited adolescents.

See *Handbook* [Section III.3.1](#)

R11	Abstract, Main results: bias assessment	Mandatory
	Provide a comment on the findings of the bias assessment.	The ‘Risk of bias’ assessments are a key finding and form a fundamental part of the strength of the conclusions drawn in the review. If risks of bias differ substantially for different comparisons and outcomes, this should be mentioned. See <i>Handbook</i> <a href="#">Section III.3.1</a>
R12	Abstract, Main results: findings	Mandatory
	Report findings for all important outcomes, irrespective of the strength and direction of the result, and of the availability of data.	Findings should typically include concise information about the size of effect and certainty of evidence for the outcome (such as risk of bias, consistency of effect, imprecision, indirectness and publication bias), for example using GRADE.  Outcomes reported in the Abstract should not be selected solely on the basis of the findings. In general, the same outcomes in the Abstract should be presented in the Plain language summary and ‘Summary of findings’ tables. <b>If no studies measured the outcome</b> , then a comment should be made to that effect. See <i>Handbook</i> <a href="#">Section III.3.1</a>
R13	Abstract, Main results: adverse effects	Mandatory
	Ensure that any findings related to adverse effects are reported. If adverse effects data were sought, but availability of data was limited, this should be reported.	The Abstract of the review should aim to reflect a balanced summary of the benefits and harms of the intervention. See <i>Handbook</i> <a href="#">Section III.3.1</a>
R14	Abstract, Main results: format of numerical results	Mandatory
	Present summaries of statistical analyses in the same way as they are reported in the review and in a standard way, ensuring that readers will understand the direction of benefit and the measurement scale used, and that confidence intervals are included where appropriate.	The standard format for reporting the results of statistical analysis includes an indication of the summary measure, point estimate and confidence interval, e.g. odds ratio 0.75 (95% confidence interval 0.62 to 0.89).



R15	Abstract, Main results: interpretability of findings	Highly desirable
<p>Ensure that key findings are interpretable, or are re-expressed in an interpretable way. For instance, they might be re-expressed in absolute terms (e.g. assumed and corresponding risks, NNTBs, group means), and outcomes combined with a standardized scale (e.g. standardized mean difference) might be re-expressed in units that are more naturally understood.</p> <p>Absolute effects provide a useful illustration of the likely impact of intervention, and are usually easier to understand than relative effects. Units expressed on a standardized scale reflect the effect estimate as the number of standard deviations. This is not intuitive to many readers who may be more familiar with specific scales. Any re-expressed findings must have been presented in the same way in the main text of the review (see previous standard).</p>		
R16	Abstract, Authors' conclusions	Mandatory
<p>State key conclusions drawn.</p> <p><b>Authors' conclusions may include both</b> implications for practice and implications for research. Care must be taken to avoid interpreting lack of evidence of effect as evidence of lack of effect.</p> <p>Recommendations for practice should be avoided.</p> <p>See <i>Handbook</i> <a href="#">Section III.3.1</a> and <a href="#">Section 15.6.1</a></p>		
R17	Completeness of main review text	Mandatory
<p>Ensure that all findings reported in the Abstract and Plain language summary, including re-expressions of meta-analysis results, also appear in the main text of the review.</p> <p>See <i>Handbook</i> <a href="#">Section III.3.1</a> and <a href="#">Section III.4</a></p>		
R18	Consistency of summary versions of the review	Mandatory
<p>Ensure that reporting of objectives, important outcomes, results, caveats and conclusions is consistent across the main text, the Abstract, the Plain language summary and the 'Summary of findings' table (if included).</p> <p>Summary versions of the review should be written on the assumption that they are likely to be read in isolation from the rest of the review.</p>		

## 1.23 Background

Cochrane Training resource: [writing a protocol](#)

Cochrane Interactive Learning: [module 2 - writing the review protocol](#)

Standard		Rationale and elaboration
R19	Background	Mandatory
Provide a concise description of the condition or problem addressed by the review question, definition of the intervention and how it might work, and why it is important to do the review.		Systematic reviews should have a clearly defined and well-reasoned rationale that has been developed in the context of existing knowledge. Outlining the context of the review question is useful to readers and helps to establish key uncertainties that the review intends to address.
R20	Background headings	Highly desirable
Include the four standard RevMan headings when writing the Background.		Four standard headings are included in RevMan ('Description of the condition', 'Description of the intervention', 'How the intervention might work', and 'Why it is important to do this review'). See <i>Handbook</i> <a href="#">Section III.3.2</a>
R21	Background references	Mandatory
Back up all key supporting statements with references.		Claims or statements regarding aspects such as disease burden, morbidity, prevalence and mechanisms of action should be substantiated and, where available, supported by external evidence.
R22	Main objective	Mandatory
State the main objective, where appropriate in a single concise sentence.		The primary objective of a Cochrane Review should be to assess the effects of one or more healthcare interventions on user-important outcomes, both intended and unintended. The objective should be expressed in terms that relate to the population(s), intervention comparison(s) and, where appropriate, to specify the outcomes of interest explicitly. Review users may be patients, carers, policy makers, clinicians, practitioners or others.  <i>MECIR conduct standard 2:</i> Define in advance the objectives of the review, including participants, interventions, comparators and outcomes (PICO).  Where possible, the format should be of the form "To assess the effects of [intervention or comparison] for [health problem] for/in [types of people, disease or problem and setting if specified]". See <i>Handbook</i> <a href="#">Section III.3.2</a> and <a href="#">Section 2.3</a>
R23	Secondary objectives	Highly desirable

State explicitly (as secondary objectives) any specific questions being addressed by the review, such as those relating to particular participant groups, intervention comparisons or outcomes.

The objectives should be expressed in terms that relate to the population(s), intervention comparison(s) and, where appropriate, outcomes of interest.

*MECIR conduct standard 4:* Consider in advance whether issues of equity and relevance of evidence to specific populations are important to the review, and plan for appropriate methods to address them if they are. Attention should be paid to the relevance of the review question to populations such as low-socioeconomic groups, low- or middle-income regions, women, children and older people.

See *Handbook* [Section III.3.2](#) and [Section 2.4](#)

#### R24 Economic evidence

Mandatory

*If health economics evidence is being reviewed*, state this explicitly in the Objectives (as a secondary objective).

The primary aim of a Cochrane Review should be to assess the effects of one or more healthcare interventions on user-important outcomes, both intended and unintended. These outcomes may include economic outcomes. If health economics evidence is being reviewed as an integrated economics component, this should be stated as a secondary objective.

See *Handbook* [Section 20.2.2](#)

#### R25 Qualitative research evidence

Mandatory

*If qualitative research evidence is being reviewed*, state this explicitly in the Objectives (as a secondary objective).

The primary aim of a Cochrane Review should be to assess the effects of one or more healthcare interventions on user-important outcomes, both intended and unintended. If qualitative research **evidence is being included to 'extend' the review**, this should be stated as a secondary objective.

See *Handbook* [Section 21.4](#)

## 1.24 Methods

Standard		Rationale and elaboration
R26	Reference protocol	Highly desirable
Cite the protocol for the review.		<p>The reader should be made aware that the review is based on a published protocol. This is particularly important if the review has been split into multiple reviews since the protocol was published. The most convenient place to reference the protocol for the <b>review is under ‘Other published versions of this review’</b>. Since the protocol is usually no longer included in the CDSR once the review is published, it should be cited using the last publication citation for the protocol. Archived versions of protocols can be accessed via the current version of the review.</p>

## 1.25 Criteria for considering studies for this review

Cochrane Training resource: [defining the review question](#)

Cochrane Interactive Learning: [module 2 - writing the review protocol](#)

Standard		Rationale and elaboration
R27	Eligibility criteria for types of study: study designs	Mandatory
State eligible study designs, and provide a justification for the choice.		<p>It is not necessary to explain why randomized trials are eligible (if that is the case), although it may be important to explain why other types of study meet the eligibility criteria of the review.</p> <p><i>MECIR conduct standard 9:</i> Define in advance the eligibility criteria for study designs in a clear and unambiguous way, with a focus on features of a study's design rather than design labels.</p> <p><i>MECIR conduct standard 11:</i> Justify the choice of eligible study designs. See <i>Handbook</i> <a href="#">Section III.3.3.1</a> and <a href="#">Section 3.3</a>.</p>
R28	Eligibility criteria for types of study: study reports	Mandatory
If studies are excluded on the basis of publication status or language of publication, explain and justify this.		<p>Studies should be included irrespective of their publication status and language of publication, unless explicitly justified.</p> <p><i>MECIR conduct standard 12:</i> Include studies irrespective of their publication status, unless exclusion is explicitly justified. See <i>Handbook</i> <a href="#">Section III.3.3.1</a> and <a href="#">Section 3.4</a></p>
R29	Eligibility criteria for types of participants	Mandatory
State eligibility criteria for participants, including any criteria around location, setting, diagnosis or definition of condition and demographic factors, and how studies including subsets of relevant participants are addressed.		<p>Any notable restrictions on the eligibility criteria of the review should be given and explained (e.g. exclusion of people under or over a certain age, specific settings of intervention).</p> <p><i>MECIR conduct standard 5:</i> Define in advance the eligibility criteria for participants in the studies.</p> <p><i>MECIR conduct standard 6:</i> Define in advance how studies that include only a subset of relevant participants will be addressed. See <i>Handbook</i> <a href="#">Section III.3.3.1</a> and <a href="#">Section 3.2.1</a></p>
R30	Eligibility criteria for types of interventions	Mandatory

State eligibility criteria for interventions and comparators, including any criteria around delivery, dose, duration, intensity, co-interventions and characteristics of complex interventions.

*MECIR conduct standard 7:* Define in advance the eligible interventions and the interventions against which these can be compared in the included studies.  
See *Handbook* [Section III.3.3.1](#) and [Section 3.2.2](#)

### R31 Role of outcomes

Mandatory

*If measurement of particular outcomes is used as an eligibility criterion, state and justify this.*

Studies should never be excluded from a review solely because no outcomes of interest are *reported*. However, on occasion it will be appropriate to include only studies that *measured* particular outcomes. For example, a review of a multi-component public health intervention promoting healthy lifestyle choices, focussing on reduction in smoking prevalence, might legitimately exclude studies that do not measure smoking rates.

*MECIR conduct standard 8:* Clarify in advance whether outcomes listed under 'Criteria for considering studies for this review' are used as criteria for including studies (rather than as a list of the outcomes of interest within whichever studies are included).  
See *Handbook* [Section III.3.3.1](#) and [Section 3.2.4.1](#)

### R32 Outcomes of interest

Mandatory

Define in advance outcomes that are critical to the review, and any additional important outcomes and define acceptable ways of measuring them.

Explain how multiple variants of outcome measures (e.g. definitions, assessors, scales, time points) are addressed.

*MECIR conduct standard 14:* Define in advance outcomes that are critical to the review, and any additional important outcomes

Also *MECIR conduct standards 15–18*

See *Handbook* [Section III.3.3.1](#) and [Section 3.2.4.1](#)

URL: <https://community.cochrane.org/mecir-manual/standards-reporting-new-cochrane-intervention-reviews-r1-109/reporting-review-conduct-r1-55/search-methods-identification-studies-r33-38>

## 1.26 Search methods for identification of studies

Cochrane Training resource: [searching for studies](#)

Cochrane Interactive Learning: [module 3 - searching for studies](#)

Standard	Rationale and elaboration
R33 Search sources	Mandatory
List all sources searched, including: databases, trials registers, websites and grey literature. Database names should include platform or provider name (or both), and dates of coverage; websites should include full name and URL. State whether reference lists were searched and whether individuals or organizations were contacted.	<p><i>MECIR conduct standard 36:</i> Document the search process in enough detail to ensure that it can be reported correctly in the review.</p> <p>Also <i>MECIR conduct standards 24–31</i> See <i>Handbook</i> <a href="#">Section III.3.3.2</a>, <a href="#">Section 1.5</a>; <a href="#">4.3.1.1</a> and <a href="#">Section 4.4.5</a></p>
R34 Latest searches	Mandatory
Provide the date of the last search and the issue or version number (where relevant) for each database for which results were evaluated and incorporated into the review. If a search was rerun prior to publication, and its results were not incorporated, explain how the results were dealt with, and provide the date of the search.	<p>The review should provide the search date up to which studies have been retrieved and assessed for inclusion. This is the date to which the conclusions of the review are valid. It should reflect the date of the most recent set of searches from which all records have been screened for relevance and any studies meeting the eligibility criteria have been fully incorporated into the review (studies may be awaiting classification if, for example, the review authors are awaiting translation or clarification from authors or sponsors).</p> <p>Since the review is likely to have drawn on searches conducted across multiple databases, it is possible that searches were performed on more than one date. The earliest date of the most recent set of searches should be provided in the review text and as the hard-coded date of the last search. The remaining dates for other databases should be reported in an Appendix.</p> <p><b>If a ‘catch-up’ search was run subsequent to the review being written up, any relevant studies not yet assessed for inclusion should be listed in the section ‘Studies awaiting assessment’.</b></p> <p><i>MECIR conduct standard 37:</i> Rerun or update searches for all relevant databases within 12 months before publication of the review or review update, and screen the results for potentially eligible studies.</p>

*MECIR conduct standard 38:* Incorporate fully any studies identified in the rerun or update of the search within 12 months before publication of the review or review update.  
See *Handbook* [Section 4.4.10](#)

<b>R35</b>	<b>Search restrictions</b>	<b>Mandatory</b>
Specify and justify any restrictions placed on the time period covered by the search.	<i>MECIR conduct standard 35:</i> Justify the use of any restrictions in the search strategy on publication date or publication format. See <i>Handbook</i> <a href="#">Section III.3.3.2</a> and <a href="#">Section 4.4.5</a>	
<b>R36</b>	<b>Searches for different types of evidence</b>	<b>Mandatory</b>
<i>If the review has specific eligibility criteria concerning inclusion of additional studies such as studies of adverse effects, health economics evidence or qualitative research evidence, describe search methods for identifying such studies.</i>	Some reviews extend beyond a focus on the effects of healthcare interventions and address specific additional types of evidence. These are discussed in the <i>Handbook</i> .  <i>MECIR conduct standard 26:</i> <i>If the review has specific eligibility criteria around study design to address adverse effects, economic issues or qualitative research questions, undertake searches to address them.</i> These are discussed in the <i>Handbook</i> Chapters <a href="#">19</a> , <a href="#">20</a> and <a href="#">21</a> .	
<b>R37</b>	<b>Search strategies for bibliographic databases</b>	<b>Mandatory</b>
Present the exact search strategy (or strategies) used for each database in an Appendix, including any limits and filters used, so that it could be replicated.	Search strategies that are available elsewhere (e.g. standard methodological filters, or strategies used to populate a specialized register) may be referenced rather than reproduced. Including the number of hits for each line in the strategy is optional.  <i>MECIR conduct standard 36:</i> Document the search process in enough detail to ensure that it can be reported correctly in the review.  Also <i>MECIR conduct standards 32–35</i> See <i>Handbook</i> <a href="#">Section III.3.3.2</a> and <a href="#">Section 4.4.5</a>	
<b>R38</b>	<b>Search strategies for other sources</b>	<b>Highly desirable</b>
Report the search terms used to search any sources other than bibliographic databases (e.g. trials registers, the web), and the dates of the searches.	Some of this information might be better placed in an Appendix.  <i>MECIR conduct standard 36:</i> Document the search process in enough detail to ensure that it can be reported correctly in the review. See <i>Handbook</i> <a href="#">Section III.3.3.2</a> and <a href="#">Section 4.4.5</a>	





## 1.27 Data collection and analysis

Standard	Rationale and elaboration	
R39 Inclusion decisions		Mandatory
State how inclusion decisions were made (i.e. from search results to included studies), clarifying how many people were involved and whether they worked independently.	<p><i>MECIR conduct standard 39:</i> Use (at least) two people working independently to determine whether each study meets the eligibility criteria, and define in advance the process for resolving disagreements.</p> <p>See <i>Handbook</i> <a href="#">Section III.3.3.3</a> and <a href="#">Section 4.4.10</a></p>	
R40 Data collection process		Mandatory
State how data were extracted from reports of included studies, clarifying how many people were involved, whether they worked independently, and how disagreements were resolved. Describe data collection process for any reports requiring translation.	<p><i>MECIR conduct standard 43:</i> Use a data collection form that has been piloted.</p> <p><i>MECIR conduct standard 45:</i> Use (at least) two people working independently to extract study characteristics from reports of each study, and define in advance the process for resolving disagreements.</p> <p>See <i>Handbook</i> <a href="#">Section III.3.3.3</a>, <a href="#">Section 5.4.1</a> and <a href="#">Section 5.5.2</a></p>	
R41 Requests for data		Highly desirable
Describe attempts to obtain or clarify data from individuals or organizations.	<p><i>MECIR conduct standard 49:</i> Seek key unpublished information that is missing from reports of included studies.</p> <p>See <i>Handbook</i> <a href="#">Section III.3.3.3</a> and <a href="#">Section 5.2.3</a></p>	
R42 Data items		Mandatory
State the types of information that were sought from reports of included studies.	<p><i>MECIR conduct standard 44:</i> Collect characteristics of the included <b>studies in sufficient detail to populate a table of ‘Characteristics of included studies’</b>.</p> <p>See <i>Handbook</i> <a href="#">Section III.3.3.3</a> and <a href="#">Section 5.3.1</a></p>	
R43 Transformations of data		Mandatory
Explain any transformations of reported data prior to presentation in the review, along with any assumptions made. Explain any procedures for extracting numeric data from graphs.	<p><i>MECIR conduct standard 47:</i> Collect and utilize the most detailed numerical data that might facilitate similar analyses of included studies. Where 2×2 tables or means and standard deviations are not available, this might include effect estimates (e.g. odds ratios, regression coefficients), confidence intervals, test statistics (e.g. t, F, Z, Chi<sup>2</sup>) or P values, or even data for individual participants.</p> <p>See <i>Handbook</i> <a href="#">Section 5.3.6</a></p>	
R44 Missing outcome data		Highly desirable
Explain how missing outcome data were addressed.	Describe how assumptions are applied for missing data, e.g. last observation carried forward, or assumptions of particular values such as worst-case or best-case scenarios.	

See *Handbook* [Section III.3.3.3](#), [Section 5.3.6](#) and [Section 10.12.1](#)

R45	Tools to assess risk of bias in individual studies	Mandatory
	State and reference the tool(s) used to assess risk of bias for included studies, how the tool(s) was implemented, and the criteria used to assign studies to judgements of low risk, high risk and unclear risk of bias.	<p>If the <i>Handbook</i> <b>guidance for undertaking ‘Risk of bias’</b> assessments was followed in its entirety, then a reference to the <i>Handbook</i> is sufficient to provide the criteria used to assign judgements. Justify any deviations from the tool.</p> <p><i>MECIR conduct standard 52:</i> Assess the risk of bias in at least one specific result for each included study. For randomized trials, the RoB 2 tool should be used, involving judgements and support for those judgements across a series of domains of bias, as described in the <i>Handbook</i>.</p> <p><i>MECIR conduct standards 53–61</i> See <i>Handbook</i> <a href="#">Section III.3.3.3</a>, <a href="#">Section 7.1.2</a> and <a href="#">Chapter 8</a></p>
R46	Effect measures	Mandatory
	State the effect measures used by the review authors to describe effect sizes (e.g. risk ratio, mean difference) in any included studies or meta-analyses, or both.	See <i>Handbook</i> <a href="#">Section III.3.3.3</a>
R47	Non-standard designs	Mandatory
	<i>If designs other than individually randomized, parallel-group randomized trials are included,</i> describe any methods used to address clustering, matching or other design features of the included studies.	<p><i>MECIR conduct standard 70:</i> Consider the impact on the analysis of clustering, matching or other non-standard design features of the included studies.</p> <p>See <i>Handbook</i> <a href="#">Section 6.2.1</a></p>
R48	Studies with more than two groups	Mandatory
	<i>If multi-arm studies are included,</i> explain how they were addressed and incorporated into syntheses.	<p><i>MECIR conduct standard 66:</i> <i>If multi-arm studies are included,</i> analyse multiple intervention groups in an appropriate way that avoids arbitrary omission of relevant groups and double-counting of participants.</p> <p>See <i>Handbook</i> <a href="#">Section III.3.3.3</a>, <a href="#">Section 6.2.9</a> and <a href="#">Chapter 11</a>.</p>
R49	Assessing heterogeneity	Mandatory
	Describe the methods used to identify the presence of heterogeneity between the studies in the review (e.g. non-quantitative assessment, $I^2$ , $\text{Tau}^2$ or statistical test).	<p><i>MECIR conduct standard 69:</i> Take into account any statistical heterogeneity when interpreting the results, particularly when there is variation in the direction of effect.</p> <p><i>MECIR conduct standard 62:</i> Undertake (or display) a meta-analysis only if participants, interventions, comparisons and outcomes are judged to be sufficiently similar to ensure an answer that is clinically meaningful.</p>

*MECIR conduct standard 63:* Assess the presence and extent of between-study variation when undertaking a meta-analysis.  
See *Handbook* [Section 10.10.2](#) and [Section 10.10.3](#).

R50	Risk of reporting bias across studies	Highly desirable
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Describe any methods used for assessing the risk of reporting biases such as publication bias.

See *Handbook* [Chapter 13](#)

R51	Data synthesis	Mandatory
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Describe any methods used for combining results across studies. Where data have been combined in statistical software external to RevMan, reference the software, commands and settings used to run the analysis.

Decisions to depart from intended methods, for example an alternative statistical model, should be reported and justified.

*MECIR conduct standard 62:* Undertake (or display) a meta-analysis only if participants, interventions, comparisons and outcomes are judged to be sufficiently similar to ensure an answer that is clinically meaningful.

R52	Subgroup analyses	Mandatory
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*If subgroup analysis (or meta-regression) was performed*, state the potential effect modifiers with rationale for each, stating whether each was defined a priori or post hoc and how they were compared (e.g. statistical tests).

*MECIR conduct standard 22:* Predefine potential effect modifiers (e.g. for subgroup analyses) at the protocol stage, restrict these in number, and provide rationale for each.

*MECIR conduct standard 67:* If subgroup analyses are to be compared, and there are judged to be sufficient studies to do this meaningfully, use a formal statistical test to compare them.

See *Handbook* [Section III.3.3.3](#) and [Section 10.11.3.1](#)

R53	Addressing risk of bias	Mandatory
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Describe how studies with high or variable risks of bias are addressed in the synthesis.

*MECIR conduct standard 58:* Address risk of bias in the synthesis (whether quantitative or non-quantitative). For example, present analyses that are stratified according to summary risk of bias, restricted to studies at low risk of bias or restricted to low-and-some-concerns of risk of bias.

See *Handbook* [Section 7.6.1](#) and [Chapter 8](#)

R54	Sensitivity analysis	Mandatory
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State the basis for any sensitivity analyses performed.

*MECIR conduct standard 71:* Use sensitivity analyses to assess the robustness of results, such as the impact of notable assumptions, imputed data, borderline decisions and studies at high risk of bias.

See *Handbook* , [Section 10.14](#)

R55	Summary of findings	Highly desirable
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State any methods for summarizing the findings of the review, including the assessment of the certainty of the body of evidence for each outcome.

*MECIR conduct standard 75:* Justify and document all assessments of the certainty of the body of evidence (for example downgrading or upgrading if using GRADE).

*MECIR conduct standard 74:* Use the five GRADE considerations (risk of bias, consistency of effect, imprecision, indirectness and publication bias) to assess the certainty of the body of evidence for each outcome, and to draw conclusions about the certainty of evidence within the text of the review.

See *Handbook* [Section 14.2.1](#)

## Results

## 1.28 Description of studies

Standard		Rationale and elaboration
R56	Flow of studies	Mandatory
Provide information on the flow of studies from the number(s) of references identified in the search to the number of studies included in the review, ideally using a PRISMA type flow diagram. Clarify how multiple references for the same study relate to the individual studies.		<p><i>MECIR conduct standard 41:</i> Document the selection process in sufficient detail to be able to complete a flow diagram and a <b>table of ‘Characteristics of excluded studies’</b>.</p> <p><i>MECIR conduct standard 42:</i> Collate multiple reports of the same study, so that each study, rather than each report, is the unit of interest in the review.</p> <p>See <i>Handbook</i> <a href="#">Section III.3.4.1</a>, <a href="#">Section 4.6.4</a>, <a href="#">Section 4.6.2</a> and <a href="#">Section 5.2.1</a></p>
R57	Lack of included studies	Highly desirable
If a review identifies no eligible studies, restrict the Results section to a description of the flow of studies and any brief comments about reasons for exclusion of studies.		<p><b>Under ‘Risk of bias in included studies’ and ‘Effects of interventions’, state “No study met the eligibility criteria”. Any discussion of evidence not meeting the eligibility criteria of the review should be in the Discussion section.</b></p> <p>See <i>Handbook</i> <a href="#">Section III.3.4.1</a></p>
R58	Excluded studies	Mandatory
List key excluded studies and provide justification for each exclusion.		<p><b>The table of ‘Characteristics of excluded studies’ is intended as an aid to users rather than a comprehensive list of studies that were identified but not included. List here any studies that a user might reasonably expect to find in the review to explain why they are excluded.</b></p> <p>See <i>Handbook</i> <a href="#">Section III.3.4.1</a></p>
R59	Studies awaiting classification	Highly desirable
List the characteristics of any completed studies that have been identified as potentially eligible but have not been incorporated into the review.		<p>Users of the review will be interested to learn of any potentially relevant studies that have been conducted and are known to the review team, but have not yet been incorporated in to the review irrespective of their publication status. This will help them to assess the stability of the review findings. These should be listed <b>in the table of ‘Characteristics of studies awaiting classification’</b>, along with any details that are known. Authors should also consider the impact of not including these studies on the review findings as a potential limitation, and the extent to which they affect the implications for research.</p> <p>See <i>Handbook</i> <a href="#">Section III.3.4.1</a></p>
R60	Ongoing studies	Mandatory

Provide details of any identified studies that have not been completed.

Users of the review will be interested to learn of any potentially relevant studies that have not been completed. This will help them to assess the stability of the review findings. These should **be listed in the table of 'Characteristics of ongoing studies'**, along with any details that are known.

Cochrane Reviews should be mindful of research waste so it is useful to consider how ongoing studies might address the review question under **'Implications for research'**.

See *Handbook* [Section III.3.4.1](#)

R61	<b>Table of 'Characteristics of included studies'</b>	Mandatory
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**Present a table of 'Characteristics of included studies' using a uniform format across all studies.**

*MECIR conduct standard 44:* Collect characteristics of the included studies in sufficient detail to populate a table of **'Characteristics of included studies'**.

See *Handbook* [Section III.3.4.1](#) and [Section 5.3.1](#)

R62	<b>Included studies</b>	Mandatory
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Provide a brief narrative summary of any included studies. This should include the number of participants and a summary of the characteristics of the study populations and settings, interventions, comparators and funding sources.

See *Handbook* [Section III.3.4.1](#)

R63	<b>Table of 'Characteristics of included studies': methods</b>	Mandatory
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Provide the basic study design or design features (e.g. parallel group randomized trial, cluster-randomized trial, controlled before and after study).

Even if the review is restricted to one study design, these tables should provide a comprehensive summary of each study.

It is important that labels used to describe study designs are clearly defined in the review. See *Handbook* [Section III.3.4.1](#) and [Section 5.3](#)

R64	<b>Table of 'Characteristics of included studies': participants</b>	Mandatory
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Provide sufficient information about the study populations to enable a user of the review to assess the **applicability of the review's findings** to their own setting.

Information presented in this table should reflect the baseline demographics of the study sample. In addition, it is helpful to state the eligibility criteria of the study.

See *Handbook* [Section III.3.4.1](#) and [Section 5.3](#)

R65	<b>Table of 'Characteristics of included studies': sample sizes</b>	Mandatory
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Include the sample size for each included study in the table of **'Characteristics of included studies'**.

If sample sizes are available for each intervention group, these should be included. A convenient place is often within the box



for Interventions, e.g. inserting “(n = )” after each listed intervention group.

See *Handbook* [Section III.3.4.1](#) and [Section 5.3](#)

R66	Table of ‘Characteristics of included studies’: interventions	Mandatory
	Provide sufficient information to enable users of the review to assess the applicability of the intervention to their own setting, and if possible in a way that allows the intervention to be replicated.	The components of all interventions (drug, non-drug, simple or complex) should be reported. Reporting guidelines have been developed for describing interventions used in primary research and review authors may find it useful to structure their description of interventions around the core attributes highlighted by TIDieR (Hoffman 2014). Lengthy explanations of interventions should be avoided. Citations to sources of detailed descriptions can be included. See <i>Handbook</i> <a href="#">Section III.3.4.1</a> and <a href="#">Section 5.3</a>
R67	Table of ‘Characteristics of included studies’: outcomes	Mandatory
	Provide clear and consistent information about outcomes measured (or reported), how they were measured and the times at which they were measured.	It should be clear whether main outcomes of interest in the review were measured in the study. See <i>Handbook</i> <a href="#">Section III.3.4.1</a> and <a href="#">Section 5.3</a>
R68	Table of ‘Characteristics of included studies’: dates	Highly desirable
	Include the dates when the study was conducted in the table of ‘Characteristics of included studies’.	If dates are not available then this should be stated (e.g. “Study dates not reported”). See <i>Handbook</i> <a href="#">Section III.3.4.1</a> and <a href="#">Section 5.3</a>
R69	Table of ‘Characteristics of included studies’: funding source	Mandatory
	Include details of funding sources for the study, where available.	Details of funding sources should be placed in this table rather than as part of the ‘Risk of bias’ table. Including an extra row in the table of ‘Characteristics of included studies’ is encouraged. See <i>Handbook</i> <a href="#">Section III.3.4.1</a> and <a href="#">Section 5.3</a>
R70	Table of ‘Characteristics of included studies’: declarations of interest	Mandatory
	Include details of any declarations of interest among the primary researchers.	Declarations of interest should be placed in this table rather than as part of the ‘Risk of bias’ table. Including an extra row in the table of ‘Characteristics of included studies’ is encouraged. See <i>Handbook</i> <a href="#">Section III.3.4.1</a> and <a href="#">Section 5.3</a>

R71	Choice of intervention groups in multi-arm studies	Highly desirable
<p><i>If a study is included with more than two intervention arms, restrict comments on any irrelevant arms to a brief comment in the table of 'Characteristics of included studies'.</i></p> <p>Intervention arms that are not relevant to the review question should not be discussed in detail, although it is useful to clarify (in this table) that such arms were present.</p> <p><i>MECIR conduct standard 50:</i> If a study is included with more than two intervention arms, include in the review only intervention and control groups that meet the eligibility criteria.)</p> <p>See <i>Handbook</i> <a href="#">Section 5.3.6</a></p>		
R72	References to included studies	Mandatory
<p>List all reports of each included study under the relevant Study ID.</p> <p>It is important that all reports are listed, and are grouped by study. Marking one report as the primary reference is helpful where appropriate.</p>		

URL: <https://community.cochrane.org/mecir-manual/standards-reporting-new-cochrane-intervention-reviews-r1-109/results-r56-109/risk-bias-included-studies-r73-75>

## 1.29 Risk of bias in included studies

Cochrane Training resource: [assessing RoB included studies](#) and [RoB 2.0 webinar](#)

Cochrane Interactive Learning: [module 5 - introduction to study quality and risk of bias](#)

Standard		Rationale and elaboration
R73	<b>'Risk of bias' table</b>	Mandatory
Present at least one <b>'Risk of bias'</b> table for each study that is included in a synthesis, with judgements about risks of bias, and explicit support for these judgements.		<p><b>'Risk of bias'</b> presentation tools in RevMan should be used wherever possible.</p> <p><i>MECIR conduct standard 52:</i> Assess the risk of bias in at least one specific result for each included study. For randomized trials, the RoB 2 tool should be used, involving judgements and support for those judgements across a series of domains of bias, as described in the Handbook.</p> <p>Also <i>MECIR conduct standards 54–61</i> See Handbook <a href="#">Section 7.1.2</a> and <a href="#">Chapter 8</a></p>
R74	Summary assessments of risk of bias	Highly desirable
Present an overall risk of bias assessment across domains for each key outcome for each included study, and ensure that these are supported by the <b>information presented in the 'Risk of bias' tables</b> .		<p><i>MECIR conduct standard 57:</i> Summarize the risk of bias for each key outcome for each study.</p> <p>See Handbook <a href="#">Section 7.5</a> and <a href="#">Chapter 8</a></p>
R75	Risk of bias in included studies	Mandatory
Provide a brief narrative summary of the risks of bias among the included studies.		It may be helpful to identify any studies considered to be at low risk of bias for particular key outcomes.

## 1.30 Effects of interventions

Cochrane Interactive Learning: [module 8 - reporting the review](#)

Standard		Rationale and elaboration
R76	<b>Use of 'Data and analysis' headings</b>	Mandatory
Ensure appropriate use of any heading hierarchy of Comparisons, Outcomes, Subgroups and Study data in the 'Data and analysis' section.		Appropriate use of the hierarchy in RevMan 5 ensures consistency of structure across reviews. It is confusing for the user if outcomes <b>are listed against the heading 'Comparison' and interventions listed against the heading 'Outcome or subgroup'</b> . This standard will not be required when using the study-centric data structure of RevMan Web.
R77	<b>Presenting data</b>	Highly desirable
Ensure that simple summary data for each intervention group, as well as estimates of effect size (comparing the intervention groups), are available for each study for each outcome of interest to the review. These appear by default when dichotomous or continuous outcome data are analysed within RevMan.		Simple summaries such as numbers of events, means and standard deviations should be presented for each treatment group when available. This is achieved primarily by using the <b>'Data and analyses' section of the review</b> , for dichotomous and continuous outcomes. For other outcomes, these should typically be presented in tables labelled <b>'Other data'</b> . <b>When data for each separate intervention group are available for outcomes analysed as 'generic inverse-variance' data</b> , these might be presented in Additional tables. See <i>Handbook</i> <a href="#">Section III.3.4.4</a>
R78	<b>Number of studies and participants</b>	Mandatory
State how many studies and how many participants contributed data to results for each outcome, along with the proportion of the included studies and recruited participants potentially available for the relevant comparison.		It is unlikely that the same number of studies will contribute data to every outcome of interest. Specific studies may contribute different numbers of participants for different outcomes. Therefore, for each comparison, it is helpful to indicate to readers what proportion of the relevant included studies and recruited participants contribute data to each outcome. Failure to disclose this may be misleading.
R79	<b>Source of data</b>	Highly desirable
State the source of all data presented in the review, in particular, whether it was obtained from published literature, by correspondence, from a trials register, from a web-based data repository, etc.		Transparency of data source enables validation or verification of data by others, including editors or readers of the review.
R80	<b>Multiple outcome data</b>	Mandatory

Describe any post hoc decisions that might give rise to accusations of selective outcome reporting, for example when there were multiple outcome measures (e.g. different scales), multiple time points or multiple ways of presenting results.

Transparent disclosure of post hoc decisions will enable readers of the review to assess the credibility of the results of the review for themselves. Post hoc decisions to change the definition or priority of outcome measures must be reported and justified **under ‘Differences between the protocol and review’**.

*MECIR conduct standard 16:* Define in advance details of what are acceptable outcome measures (e.g. diagnostic criteria, scales, composite outcomes).

*MECIR conduct standard 17:* Define in advance how outcome measures will be selected when there are several possible measures (e.g. multiple definitions, assessors or scales).

*MECIR conduct standard 18:* Define in advance the timing of outcome measurement.

See *Handbook* [Section 3.2.4.1](#) and [Section 5.4.1](#)

#### R81 Ordering of results and ‘Data and analysis’ section

Highly desirable

Organize results to follow the order of comparisons and outcomes specified in the protocol, following in particular the distinction between primary and secondary outcomes.

Review authors must avoid selective reporting of analysis results in a way that depends on the findings. The best way to achieve this is to follow a well-structured protocol and present results as outlined in that protocol. However, sometimes a pragmatic decision needs to be made that an alternative arrangement is preferable, particularly with regard to comparisons. This choice should be explicitly justified.

#### R82 Prespecified outcomes

Mandatory

Report synthesis results for all prespecified outcomes, irrespective of the strength or direction of the result. Indicate when data were not available for outcomes of interest, and whether adverse effects data were identified.

To avoid selective outcome reporting (in truth or in perception), the review should address all outcomes specified in the protocol.

#### R83 Statistical uncertainty

Mandatory

Accompany all effect size estimates with a measure of statistical uncertainty (e.g. a confidence interval with a specified level of confidence such as 90%, 95% or 99%).

Confidence intervals are the preferred method for expressing statistical uncertainty.

#### R84 P values

Highly desirable

*If reporting P values*, provide exact P values (e.g.  $P = 0.08$  rather than  $P > 0.05$ ).

Effect estimates with confidence intervals are the preferred method of presenting numeric results. P values should not be used as an alternative to confidence intervals and should not be **used to divide results into ‘significant’ or ‘non-significant’**; exact P values portray the strength of evidence against the null hypothesis.

See *Handbook* [Section 15.3.2](#)

R85	Tables and Figures	Mandatory
Link to each Table and Figure.	All tables and figures should have a brief descriptive caption and must be referred to in numerical order in the review text.	
R86	Number of Tables and Figures	Highly desirable
Keep the number of Tables and Figures low to convey key findings without affecting the readability of the review text.	Tables (typically implemented as Additional tables) and Figures (including RevMan flow charts, RevMan forest plots and imported graphics) may be added to reviews and included in the body of the text. Reviews should try and avoid including more than six such Tables and Figures in total. Further Tables and Figures can <b>be included as supplementary material (e.g. as ‘Data and analysis’ forest plots or within Appendices).</b>	
R87	Consistency of results	Mandatory
Ensure that all statistical results presented in the main review text are consistent between the text and <b>the ‘Data and analysis’ tables.</b>	Errors can be introduced, particularly when analyses are rerun.	
R88	Direction of effect	Mandatory
State whether findings indicate a clear direction of benefit.	Where results indicate that an intervention is better or worse than another intervention, it is important to make it clear which intervention is favoured. This is the case particularly when different scales are combined using standardized mean differences.	
R89	Interpretability of results	Mandatory
Ensure that key findings are interpretable, or are re-expressed in an interpretable way. For instance, they might be re-expressed in absolute terms (e.g. assumed and corresponding risks, NNTBs, group means), and outcomes combined with a standardized scale (e.g. standardized mean difference) might be re-expressed in units that are more naturally understood. If minimally important differences were prespecified or are available, these should be provided to aid interpretation.	Absolute effects provide a useful illustration of the likely impact of an intervention, and are usually easier to understand than relative effects. They may need to be accompanied, however, with information about assumed baseline risks. Confidence intervals should be presented for NNTBs and similar summary measures. Re-expressing relative effects as absolute effects often requires the specification of assumed (e.g. untreated) risks, and the source of these should be provided. Results expressed as standardized mean differences reflect the number of standard <b>deviations’ difference between</b> mean responses. This is not intuitive to many readers who may be more familiar with specific scales. Ideally, minimally important effect sizes should be specified in the protocol.	
R90	Studies without usable data	Mandatory

Comment on the potential impact of studies that apparently measured outcomes, but did not contribute data that allowed the study to be included in syntheses.

There is good evidence of selective outcome reporting among clinical trials. Outcomes that are believed to have been measured but are not reported in a usable format may therefore be systematically different from those that are usable, and introduce **bias**. **‘Usable’ in this sense refers both to incorporation in a meta-analysis and to consideration in non-statistical syntheses of findings.** Authors might consider using a table to indicate which studies contributed data to the outcomes of interest in the review.

*MECIR conduct standard 40:* Include studies in the review irrespective of whether measured outcome data are reported in a **‘usable’ way**.

See Handbook [Section 4.6.3](#)

R91	Missing outcome data	Highly desirable
	Discuss the implications of missing outcome data from individual participants (due to losses to follow-up or exclusions from analysis).	<i>MECIR conduct standard 64:</i> Consider the implications of missing outcome data from individual participants (due to losses to follow-up or exclusions from analysis). See Handbook <a href="#">Section 10.12.1</a>
R92	Skewed data	Highly desirable
	Discuss the possibility and implications of skewed data when analysing continuous outcomes.	<i>MECIR conduct standard 65:</i> Consider the possibility and implications of skewed data when analysing continuous outcomes. See Handbook <a href="#">Section 10.5.3</a>
R93	Forest plots	Highly desirable
	Present data from multiple studies in forest plots (using the 'Data and analyses' structure in RevMan) wherever possible, providing it is reasonable to do so.	Presenting data in forest plots can be useful, even if the studies are not combined in a meta-analysis.
R94	Multiple subgroup analyses and sensitivity analyses	Highly desirable
	<i>If presenting multiple sensitivity analyses or different ways of subgrouping the same studies,</i> present these in summary form (e.g. a single Table or Figure) and not in multiple forest plots.	
R95	Labels on plots	Mandatory
	Label the directions of effect and the intervention groups in forest plots with the interventions being compared.	<b>By default, RevMan currently uses ‘experimental’ and ‘control’</b> within labels. It is helpful to replace these with more specific intervention names, and essential if the ordering is swapped (or for head-to-head comparisons). Directions of effect should be used as consistently as possible within a review.
R96	Risk of bias across studies	Highly desirable

Present results of the assessment of risk of bias across studies (and across domains) for each key outcome, and state whether this leads to concerns about the validity of the review's findings.

Considerations of risk of bias across studies are required for assessments of the certainty of the body of evidence (e.g. using GRADE).

#### R97 Reporting biases

Highly desirable

Present results of any assessment of the potential impact of reporting biases on the review's findings.

*MECIR conduct standard 73:* Consider the potential impact of reporting biases on the results of the review or the meta-analyses it contains.

See *Handbook* [Section 13.4](#)

#### R98 'Summary of findings' table

Mandatory

Present a 'Summary of findings' table according to recommendations described in the *Handbook* (version 5 or later).

Specifically: include results for one clearly defined population group (with few exceptions); indicate the intervention and the comparison intervention; include seven or fewer patient-important outcomes; describe the outcomes (e.g. scale, scores, follow-up); indicate the number of participants and studies for each outcome; present at least one baseline risk for each dichotomous outcome (e.g. study population or median/medium risk) and baseline scores for continuous outcomes (if appropriate); summarize the intervention effect (if appropriate); and include a measure of the certainty of the body of evidence for each outcome.

Efforts should be made to incorporate information presented in 'Summary of findings' tables (such as absolute effects, certainty ratings and downgrading decisions) in other parts of the review including the Abstract, Plain language summary, Effects of interventions, Discussion and Authors' conclusions.

#### R99 Assessments of the certainty of the body of evidence

Mandatory

Provide justification or rationale for any measures of the certainty of the body of evidence for each key outcome. If a 'Summary of findings' table is used, use footnotes to explain any downgrading or upgrading according to the GRADE approach.

*MECIR conduct standard 74:* Use the five GRADE considerations (risk of bias, consistency of effect, imprecision, indirectness and publication bias) to assess the certainty of the body of evidence for each outcome, and to draw conclusions about the certainty of evidence within the text of the review.

*MECIR conduct standard 75:* Justify and document all assessments of the certainty of the body of evidence (for example downgrading or upgrading if using GRADE).

See *Handbook* [Section 14.2.1](#)



## 1.31 Discussion

Cochrane Interactive Learning: [module 8 - reporting the review](#)

Standard		Rationale and elaboration
R100	Discussion headings	Highly desirable
Include the standard RevMan headings when writing the Discussion.		<p><b>Five standard headings</b> are included in RevMan ('Summary of main results', 'Overall completeness and applicability of evidence', 'Certainty of the evidence', 'Potential biases in the review process', 'Agreements and disagreements with other studies or reviews').</p> <p>See Handbook <a href="#">Section III.3.5</a></p>
R101	Limitations	Mandatory
Discuss limitations of the review at study and outcome level (e.g. regarding risk of bias), and at review level (e.g. incomplete identification of studies, reporting bias).		<p>Review authors must explicitly state the limitations of their review. One aspect that is easily overlooked is that of adverse effects. In particular, if the review methods do not allow for detection of serious or rare adverse events, or both, the review authors must explicitly state this as a limitation. Additional considerations here include currency and completeness of the search, completeness of data collection processes, assumptions made regarding classification of interventions, outcomes or subgroups, and methods used to account for missing data.</p> <p><i>MECIR conduct standard 73:</i> Consider the potential impact of non-reporting biases on the results of the review or the meta-analyses it contains.</p> <p>See Handbook <a href="#">Section 13.4</a></p>

## 1.32 Authors' conclusions

Cochrane Interactive Learning: [module 8 - reporting the review](#)

Standard		Rationale and elaboration
R102	Conclusions: implications for practice	Mandatory
Provide a general interpretation of the evidence so that it can inform healthcare or policy decisions. Avoid making recommendations for practice.		When formulating implications for practice base conclusions only on findings from the synthesis (quantitative or narrative) of studies included in the review. The conclusions of the review should convey the essence of the synthesis of included studies, without selective reporting of the particular findings on the basis of the result, and without drawing on data that were not systematically compiled and evaluated as part of the review.
		See <i>Handbook</i> <a href="#">Section III.3.6</a> and <a href="#">Section 15.6.1</a>
R103	Conclusions: implications for research	Mandatory
<i>If recommending further research</i> , structure the implications for research to address the nature of evidence required, including population, intervention comparison, outcome, and type of study.		Researchers and research funders are an important user group of Cochrane Reviews. Recommendations for future research should offer constructive guidance on addressing the remaining uncertainties identified by the review. This is particularly important for reviews that identify few or no studies. Include any information about completed or ongoing studies that are likely to address the review question.

## 1.33 Acknowledgements

Standard		Rationale and elaboration
R104	Acknowledgements	Mandatory
Acknowledge the contribution of people not listed as authors of the review, including any assistance from the Cochrane Review Group, non-author contributions to searching, data collection, study appraisal or statistical analysis, and the provision of funding.		See <i>Handbook</i> <a href="#">Section III.3.7</a>

## 1.34 Contributions of authors

Standard		Rationale and elaboration
R105	Contributions of authors	Mandatory
Describe the contributions of each author to the review.		See <i>Handbook</i> <a href="#">Section III.3.7</a>

## 1.35 Declarations of interest

Standard	Rationale and elaboration
R106 Declarations of interests	Mandatory
<p>Report any current or recent (within the 36 months prior to registration of the review) financial interests relevant to the topic of the review. This means payments from any commercial organization with an interest in the topic of the review. Include the dates of the involvement</p>	<p>Cochrane has two Conflict of Interest policies relating to Cochrane Library content. Which policy applies to a particular review depends on whether the review was registered before or after 14 October 2020.</p> <p>For reviews registered after 14 October 2020 the <a href="#">Col Policy for Cochrane Library Content (2020)</a> applies and for reviews registered before 14 October 2020 the <a href="#">Commercial Sponsorship Policy (2014)</a> applies.</p>
<p>Report any current or recent (within 36 months prior to registration of the review) non-financial relationships and activities that have a direct and obvious connection to the topic of the review. Include the dates of the involvement.</p>	<p>Declarations of interest should be stated according to the relevant Col policy and must be consistent with interests declared on the Disclosure of Potential Conflicts of Interest form.</p>
<p>Report involvement in any study that may be eligible for inclusion in the review.</p>	<p>See <i>Handbook</i> <a href="#">Section III.3.7</a> and <a href="#">EPPR Disclosure of potential conflict of interest by author policy</a></p> <p>Cochrane Training resource: <a href="#">writing a protocol</a></p>

## 1.36 Differences between protocol and review

Standard		Rationale and elaboration
R107	Changes from the protocol	Mandatory
Explain and justify any changes from the protocol (including any post hoc decisions about eligibility criteria or the addition of subgroup analyses).		<p><i>MECIR conduct standard 13:</i> Justify any changes to eligibility criteria or outcomes studied. In particular, post hoc decisions about inclusion or exclusion of studies should keep faith with the objectives of the review rather than with arbitrary rules.</p> <p>See <i>Handbook</i> <a href="#">Section 3.2.1</a></p>
R108	Methods not implemented	Mandatory
Document aspects of the protocol that were not implemented (e.g. because no studies, or few studies, were found) in the section <b>‘Differences between protocol and review’</b> , rather than in the <b>Methods</b> section.		<p>Including a record of methods that were not implemented helps to retain specific details of the protocol. By doing so, the next version of the review can be seen to be coherent with what was planned in the protocol.</p> <p>See <i>Handbook</i> <a href="#">Section III.3.7</a></p>

## 1.37 Sources of support

Standard		Rationale and elaboration
R109	Sources of support	Mandatory
List sources of financial and non-financial support for the review and the role of the funder, if any.		See <i>Handbook</i> <a href="#">Section III.3.7</a>

URL: <https://community.cochrane.org/mecir-manual/standards-reporting-new-cochrane-intervention-reviews-r1-109/reference>

## Reference

Hoffmann TC, Glasziou PP, Boutron I, Milne R, Perera R, Moher D, et al. (2014) Better reporting of interventions: template for intervention description and replication (TIDieR) checklist and guide. *BMJ* 2014;348:g1687. doi: 10.1136/bmj.g1687



URL: <https://community.cochrane.org/mecir-manual/standards-reporting-new-cochrane-intervention-reviews-r1-109/citation>

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## STANDARDS FOR THE PLANNING, CONDUCT AND REPORTING OF UPDATES OF COCHRANE INTERVENTION REVIEWS

Jackie Chandler, Toby Lasserson, Julian PT Higgins, David Tovey, James Thomas, Ella Flemyng and Rachel Churchill

URL: <https://community.cochrane.org/mecir-manual/standards-planning-conduct-and-reporting-updates-cochrane-intervention-reviews/key-points-introduction>

## Key points and introduction

Key points:

- Before undertaking an update, authors should consider the currency and relevance of the question, as well as the methodology used to address it.
- A new protocol will be required if important changes are made to the review question or the general methodology.
- An update should be conducted according to the standards required for any review, with the following additional requirements to ensure that any changes are managed appropriately and reported clearly to readers.

Since its inception, Cochrane has advocated for the routine updating of systematic reviews, in order to take account of new evidence. However, before undertaking an update, it is important to consider carefully whether an update is warranted. See [Handbook Chapter IV, section 2](#) for a framework and checklist on deciding whether or when to update a Cochrane Review. All CRGs are encouraged to classify their reviews by their update status, to denote whether the review is up to date, an update is pending or no update is planned (see the [Updating Classification System](#)).

Several important decisions are required at the beginning of the planning of an update. The first is whether the original review question is still relevant. The second is whether the general methodological approach is still appropriate to answer the review question: this will need a review of the original protocol. Third, authors need to address whether the scope of the review is appropriate, whether it should be split into two or more reviews, or whether it should be merged with other reviews. Important changes of this nature indicate a need for a new protocol.

The following updating standards reflect three key stages: planning, conducting and reporting the update. Expectations are that review authors will consider each of these sections before updating a review. Authors should examine and address any feedback on the original review before embarking on an update or a new derivative review. Planning an update should involve discussion with the Cochrane Review Group (CRG) over the adoption of new methods or changes to the review question proposed. The following standards for updates should be used in conjunction with the conduct and reporting standards for new Cochrane Reviews and these are cited where necessary.

Jackie Chandler  
Methods Co-ordinator (2011-2018)  
Editorial and Methods Department

## Deciding on and performing an update

URL: <https://community.cochrane.org/mecir-manual/standards-planning-conduct-and-reporting-updates-cochrane-intervention-reviews-u1-11-ur1-7/deciding-and-performing-update-u1-11-ur1-7/planning-update-u1-5>

## 1.38 Planning the update

Standard	Rationale and elaboration
U1 Reconsidering review questions	Mandatory
Confirm or amend review question (PICO) and objectives.	<p>Consider whether it is important to modify or add new objectives to make the review relevant to its users.</p> <p>Consider whether the review will be split, merged with another review or otherwise changed substantially. If so, a new protocol might be warranted and the <i>MECIR conduct standards</i> should be followed rather than these <i>update standards</i>. It will be necessary to agree the approach to updating the review with the CRG.</p> <p><i>MECIR conduct standards C1, C2</i>  See explanatory note 1  See Handbook <a href="#">Section IV.3.1</a>, <a href="#">Section 2.1</a> and <a href="#">Section 2.3</a></p>
U2 Reconsidering outcomes	Mandatory
Confirm or amend outcomes of interest.	<p>Consider whether it is necessary to modify or add outcomes to ensure all user-important outcomes, including adverse effects, are addressed. Define which outcomes are primary outcomes and which are secondary outcomes. Keep the total number of outcomes as small as possible. Consider core outcome sets where available. Prioritize outcomes that will be assessed with the GRADE considerations.</p> <p><i>MECIR conduct standards C3, C14-C18, C23</i>  See Handbook <a href="#">Section 1.5</a>, <a href="#">Section 2.1</a>, <a href="#">Section 3.2.4.1</a>, <a href="#">Section 5.4.1</a></p>
U3 Reconsidering eligibility criteria	Mandatory
Confirm or amend eligibility criteria.	<p>Changes to the review objectives (e.g. additional consideration of rare adverse effects, economic issues or qualitative issues) may require modification of the eligibility criteria, possibly extending the scope to additional types of studies.</p>
U4 Planning the search	Mandatory
Decide appropriate search methods.	<p>There are four considerations in planning search methods for updates:</p> <ol style="list-style-type: none"> <li>1. Changes to eligibility criteria may require the search methods to be modified, or additional search strategies to be developed.</li> <li>2. Additional sources might need to be searched (e.g. trials registers) if not searched for the last published version of the review. Consideration should also be given to the importance of</li> </ol>

- searching data repositories and information available from regulatory agencies.
3. The update search (for unchanged eligibility criteria) will normally be limited to material added or indexed after the date of the previous search. The yield of the previous searches may be useful to decide whether the full search is repeated or whether only a subset of sources should be searched for the update.
  4. The original database search strategies may need to be modified, for example by adding search terms, adding new database subject headings, or by removing unhelpful search terms that identified many irrelevant studies in the original search.

*MECIR update standards U6 and UR3*

See Handbook [Section IV.3.4](#)

U5    Reconsidering data collection and analysis methods	Mandatory
Consider whether methods for data collection and analysis (including a GRADE assessment) need to be amended in the light of recent methodological developments.	<p>Decide if changes are required to make better use of existing data or to incorporate new data by referring to the current version of the <i>Handbook</i>. Recent developments in 'Risk of bias' assessment, statistical methods or narrative synthesis approaches may lead to more inclusive or more robust synthesis of the evidence.</p> <p>The GRADE assessment will require evaluation of risk of bias, inconsistency, imprecision, indirectness and publication bias. See <i>MECIR update standard U11</i></p> <p><b>If a 'Summary of findings' table is not included in the current version</b>, decide on the main outcomes and comparisons to be included and ensure that the relevant data have been (or will be) collected. See <i>MECIR update standard UR5</i></p> <p><i>MECIR update standards U9-U10</i></p>

URL: <https://community.cochrane.org/mecir-manual/standards-planning-conduct-and-reporting-updates-cochrane-intervention-reviews-u1-11-ur1-7/deciding-and-performing-update-u1-11-ur1-7/conduct-standards-specific-updates-u6-11>

## 1.39 Conduct standards specific to updates

Standard	Rationale and elaboration
<b>U6 Searching</b>	<b>Mandatory</b>
Undertake a new search.	<p>An updated review must include an update search for new (or additional) studies. For issues to consider in planning the search, see <i>MECIR update standard U4</i>.</p> <p>The most recent search must be no more than 12 months (preferably six months) from the intended publication date, and the results screened for potentially eligible studies.</p> <p>See <i>MECIR conduct standard C37</i>: Rerun or update searches for all relevant databases within 12 months before publication of the review or review update, and screen the results for potentially eligible studies.</p> <p>See <i>Handbook Section IV.4</i> and <a href="#">Section 4.4.10</a></p>
<b>U7 Including new studies</b>	<b>Mandatory</b>
Implement conduct standards for study selection and data collection for any newly identified studies (with updated criteria or methods as determined above).	<p><i>MECIR conduct standards C39-C51</i></p> <p>See <i>Handbook</i> <a href="#">Section 4.4.6</a>, <a href="#">Section 5.3.6</a>, <a href="#">Section 4.6.3</a>, <a href="#">Section 4.6.4</a>, <a href="#">Section 4.6.2</a>, <a href="#">Section 5.2</a>, <a href="#">Section 5.2.1</a>, <a href="#">Section 5.2.3</a>, <a href="#">Section 5.3.1</a>, <a href="#">Section 5.3.6</a>, <a href="#">Section 5.4.1</a> and <a href="#">Section 5.5.2</a></p>
<b>U8 Reconsider previously identified studies</b>	<b>Mandatory</b>
Consider studies previously identified as included, awaiting classification, ongoing and excluded, and collect additional information from them if necessary.	<p>Ensure appropriate methodology is followed to select included studies and collect information from them.</p> <p>It will be necessary to establish whether any studies previously identified as ongoing have now been completed.</p> <p>Ensure that reasons for excluding studies are consistent with current eligibility criteria and methodological standards.</p> <p>A redesign of the data collection form may be required if review questions or objectives have been modified.</p>
<b>U9 Assessing risk of bias</b>	<b>Mandatory</b>

Ensure all studies are consistently assessed for risk of bias.

The updated review must include a ‘Risk of bias’ assessment of all new and previously included studies. If the previous version used the original risk of bias tool to assess randomised trials, consider whether or not to switch to the Risk of Bias 2 tool (see *Handbook Chapter 8*), including how many randomised trials were assessed in the previous version, how many new studies are expected for inclusion in the update, how well it was implemented in the previous version and whether it is feasible to switch.

*MECIR conduct standards C52-C60*

See *Handbook*, [Section 7.1.2](#), [Section 7.3.2](#), [Section 7.5](#), [Section 7.6.1](#), [Section 7.8.6](#) and [Chapter 8](#)

#### U10 Synthesizing results

Mandatory

Implement review synthesis methods (possibly revised for the update) according to conduct standards for synthesis, across all included studies.

*MECIR conduct standards C61-C73*

See *Handbook* [Section 6.2.1](#), [Section 6.2.9](#), [Section 10.5.3](#), [Section 10.10.2](#), [Section 10.10.3](#), [Section 10.11.3.1](#), [Section 10.11.5.2](#), [Section 10.12.1](#), [Section 10.14](#), [Chapter 11](#), [Section 13.4](#), [Section 15.3.1](#)

#### U11 Assessing certainty of the evidence

Mandatory

Assess certainty of evidence using GRADE considerations of risk of bias, inconsistency, imprecision, indirectness and publication bias.

This must be applied to the full body of evidence for the key outcomes included in the updated review. The most convenient way to present **GRADE assessments** is in a ‘Summary of findings’ table.

*MECIR conduct standards C74-C75 and MECIR reporting standard R97*  
See *Handbook* [Section 14.2.1](#)



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## 1.40 Reporting standards specific to updates

	Standard	Rationale and elaboration	
UR1	Background	Review and update background as necessary to reflect changes over time.	Mandatory
		Examples of changes that should be addressed include updated estimates of disease burden, new understanding of how people are affected by the disease or condition, new insights into mechanisms of action, or changes in policy or practice. Up-to-date references should be supplied to support this information. See <i>Handbook Section IV.5</i>	
UR2	Changes to scope	Explain any changes to questions, objectives or eligibility criteria.	Mandatory
		Motivations to amend review questions and objectives for the update (such as addition of new interventions, or concerns over adverse effects) should be explained in the Background, and changes to eligibility criteria should be explained, dated and <b>justified as ‘Differences between the protocol and the review’</b> .	
UR3	Search for studies	Describe the update search.	Mandatory
		Describe which sources of information were searched for the update, and how. If any of the sources originally searched were not searched for the update, this should be explained and justified. There are at least four possibilities for providing information about search methods in an updated review: <ol style="list-style-type: none"> <li>1. An <i>integrated</i> approach is to describe all searches together, which may be most feasible if the same search was repeated.</li> <li>2. An <i>incremental</i> approach is to add information at each update to describe explicitly which searches were done for the update, retaining all information about previous searches.</li> <li>3. A <i>replacement</i> approach is to describe only the searches done for the update, using the previous review as one source of studies.</li> <li>4. A <i>hybrid</i> approach is to describe only the searches done for the update in the main text, using Appendices to provide information about previous searches.</li> </ol> See <i>Handbook Section IV.5</i>	
UR4	Flow of studies	Record the flow of studies.	Mandatory
		Provide information on the flow of studies into the updated review, ideally using a PRISMA type flow diagram. There are two broad options for providing information about how studies were identified that are included in the updated version of the review:	

1. The results of previous searches can be retained in the review and supplemented with information about studies identified in the update.
2. Alternatively, only information about searches in the current update can be presented, with the previous version of the review serving as one particular source of studies.

Either approach is acceptable. If taking the latter approach, the flow diagram should show one box for the number of studies included in the original review or previous update and an additional box for the new studies retrieved for the current update. If multiple searches have been conducted for the current update, the results of all the searches should be added together.

See *Handbook* [Section IV.5](#)

UR5	<b>'Summary of findings' tables</b>	Highly desirable
<p><b>Present a 'Summary of findings' table</b> according to recommendations described in the <i>Handbook</i> (version 5 or later). Specifically, include results for one clearly defined population group (with few exceptions).</p> <p>Efforts should be made to incorporate information presented in <b>'Summary of findings' tables (such as absolute effects, GRADE certainty ratings and downgrading decisions)</b> in other parts of the review including the Abstract, Plain language summary, Effects of <b>interventions, Discussion and Authors' conclusions</b>. See <i>Handbook</i> <a href="#">Chapter 14</a></p>		
UR6	<b>Integrating findings</b>	Mandatory
<p>Present findings integrated across new and previously included studies and not just for the new studies (in the main text, <b>Abstract, 'Summary of findings' tables and Plain language summary</b>).</p> <p>The main findings should be presented for the totality of evidence: it is not helpful to a new reader to be told about incremental updates to the evidence base. However, the impact of new evidence on review findings may be useful to draw on when interpreting the results.</p>		
UR7	<b>What's new?</b>	Mandatory
<p>Explain <b>what's new</b>.</p> <p>It is important that changes are explained to inform returning <b>readers about what's new. This should be achieved in several ways.</b></p> <p>A comment should be inserted to explain that the review is an update of a previously published review. This might be placed at the beginning or end of the Background or the start of the section <b>'Search methods for identification of studies'</b>. <b>It can be helpful to explain also whether the article describes the first, second, third and so on update of the review.</b></p> <p>Changes in review questions, eligibility criteria and methods should <b>be reported in the section 'Differences between protocol and review'</b>, making it clear that they are changes since the previous version.</p>		

**Changes in findings must be reported and dated in the ‘What’s new’ section.** This should include the numbers of new studies and participants in those studies; and the nature of any changes in assessments of the certainty of the evidence (e.g. using GRADE) and in the clinical implications of the findings. For particularly notable changes it is useful to comment on these within the text of the review.

See *Handbook* [Section IV.5](#)

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## TRANSLATIONS OF THE MECIR STANDARDS

URL: <https://community.cochrane.org/mecir-manual/translations-mecir-standards/key-points-and-introduction>

## Key points and introduction

- Cochrane encourages translations of the MECIR Manual in order to support the engagement of people with different native languages in Cochrane Review production.
- Full details on the conditions and process for translating the MECIR Manual can be found in the [Cochrane MECIR translations guidance](#)
- If you are interested in translating the MECIR Manual, contact [methods@cochrane.org](mailto:methods@cochrane.org).

The Cochrane Editorial and Methods Department, Knowledge Translation Department and authors of the MECIR Standards encourage translations of the MECIR Standards in order to support the engagement of people with different native languages in Cochrane Review production.

The MECIR Standards are the ‘how-to’ guides for conducting, reporting and updating Cochrane Intervention Reviews, Protocols and Updates. The MECIR Standards for the conduct of new Cochrane Intervention Reviews are embedded throughout the [Cochrane Handbook for Systematic Reviews of Interventions](#).

Translation proposals will be assessed and approved by the Cochrane Methods team and the Translations Coordinator. Please see the [Cochrane MECIR translations guidance](#) for full details on the conditions that must be met for MECIR translations, how to initiate a MECIR translation and keeping it up-to-date.

If you are interested in translating the MECIR Manual, or have any questions about the process or other general queries, please contact [methods@cochrane.org](mailto:methods@cochrane.org).

*(Translations of MECIR will be hosted as PDFs within this section)*



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