

Review proposal form: intervention reviews

Version January 2021

Please complete this form to outline your proposal for a Cochrane Review. Complete all sections in full. Email the completed form to Tina Jakob, Managing Editor, Cochrane Metabolic and Endocrine Disorders: tina.jakob@uni-duesseldorf.de

**Data Protection**

The personal data included in this form will be used to complete your Cochrane author profiles if the title is accepted.

Both successful and unsuccessful submissions may be archived for the Review Group’s records.

This form will be anonymised before circulation to editors considering this title proposal, for reasons of equity and confidentiality.

Please see the [Cochrane Privacy Policy](https://community.cochrane.org/organizational-info/resources/policies/cochrane-privacy-policy) for further information. Please direct any queries about data protection to support@cochrane.org.

🞏 By submitting this form, we give Cochrane permission to process the data included here.

# IMPORTANT: Disclosure of Conflicts of interest

Please read Cochrane’s [Conflict of Interest Policy for Cochrane Library content](https://training.cochrane.org/online-learning/editorial-policies/coi-policy/coi-policy-cochrane-library) and confirm in Section 6 below whether any member of the author team has a potential Conflict of Interest.

If your title is accepted, the Review Group will request a full Declaration of Interest from each member of the author team. The title will not be registered until the Review Group has assessed any relevant Conflict of Interest.

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| Essential checks before title submission:* We have searched the [*Cochrane Database of Systematic Reviews*](https://www.cochranelibrary.com/advanced-search) in the Cochrane Library for published reviews and protocols and can confirm that this proposal has not been covered by another Cochrane Review.
* We have checked that this proposal falls within the scope of Cochrane Metabolic and Endocrine Disorders.
* We understand that all authors must follow the [*Cochrane Handbook for Systematic Reviews of Interventions*](https://training.cochrane.org/handbook/current).
* We have read Cochrane’s [Conflict of Interest Policy for Cochrane Library content](https://training.cochrane.org/online-learning/editorial-policies/coi-policy/coi-policy-cochrane-library) and have informed the Cochrane Metabolic and Endocrine Disorders' Managing Editor of any potential conflict of interest.
* We have read [Managing expectations: what does Cochrane expect of authors, and what can authors expect of Cochrane?](https://documentation.cochrane.org/pages/viewpage.action?pageId=117381161) and are aware that preparing a Cochrane Review requires a significant commitment from all authors.
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| Author registrationNOTE TO REVIEW GROUPS: PLEASE DELETE THIS SECTION BEFORE CIRCULATING THIS FORM. |
| All authors should create [Cochrane Accounts](https://account.cochrane.org/) before submitting this form.To enable editorial staff to identify you in our contributor management system, please list the email addresses used at account registration.  |
| Author 1 | Email used to register for Cochrane Account |
| Author 2 | Email used to register for Cochrane Account |
| Add other rows as required for other author team members.  |

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| Proposed title(see [Handbook sections II.1.3](https://training.cochrane.org/handbook/current/chapter-ii#section-ii-1-3) and [1.1.2](https://training.cochrane.org/handbook/current/chapter-01#section-1-2)). Your proposal should not overlap with an existing Cochrane Review.You must use one of the standard formats for Cochrane Review titles:* [Intervention] FOR [health problem/issue] e.g. Antibiotics for acute bronchitis
* [Intervention A] VERSUS [Intervention B] FOR [health problem/issue] e.g. Short-term versus long-term antibiotics for acute bronchitis
* [Intervention] FOR [health problem/issue] IN [participant group] e.g. Antibiotics for acute bronchitis in children
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| Title: |  |

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| Contact person Author who will take responsibility for the review, and communicate with the editorial base throughout review development; does not need to be the first listed author. |
| Name: |  |
| Email:  |  |

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| Review proposal and inclusion criteria(see [Handbook chapter 2](https://training.cochrane.org/handbook/current/chapter-02)) |
| Why is it important to do this review? | Why are you proposing to undertake this review? For example, is it particularly topical at the present time?Please use the ‘Review context’ section below to state if this review would form part of a Masters or Doctorate, or of a larger research project. |
| Description of the condition: |  |
| Description of the intervention: |  |
| How the intervention might work: |  |
| Review objectives: | Give a short statement of the primary aim of the review, e.g. to assess the effects of your intervention. |
| Types of study: ([section 3.3](https://training.cochrane.org/handbook/current/chapter-03#a-33-determining-which-study-designs-to-include)) | Outline the types of study that will be included in the review. Most Cochrane Reviews of interventions focus on randomised controlled trials (RCTs). If your review will include non-randomised studies, please provide specific reasons for this. |
| **P**articipants / population: ([section 3.2.1](https://training.cochrane.org/handbook/current/chapter-03#section-3-2-1)) | Outline the types of populations to be included and excluded. Consider demographic factors, the type/stage of disease/condition, and care setting. |
| **I**ntervention: ([section 3.2.2](https://training.cochrane.org/handbook/current/chapter-03#section-3-2-2)) | Outline the details of the intervention you wish to investigate. Consider the dose, intensity, mode of delivery, and combinations of interventions. Are there variations you wish to exclude?  |
| **C**omparison: ([section 3.2.3](https://training.cochrane.org/handbook/current/chapter-03#section-3-2-3)) | What will the intervention be compared to, e.g. placebo, no intervention, standard care? |
| Study length: | Minimum duration of intervention:  |
| Minimum duration of follow-up: |
| **O**utcomes and adverse effects: ([section 3.2.4](https://training.cochrane.org/handbook/current/chapter-03#section-3-2-4)) | List the primary and secondary outcomes you will measure. Include including outcomes important to people with the relevant disease/condition as well as those treating them. Specify how your outcomes may be measured, e.g. the type of scale or count likely to be used, and the timing of the measurement.Please note that all-cause mortality, morbidity/complications, adverse events, health-related quality of life and socioeconomic effects have to be investigated and reported in every CMED review. Adverse events should be a primary outcome parameter. Evaluation of surrogate outcomes only will not be possible. |
| Primary outcomes:(maximum of three) |
| Secondary outcomes: |
| Exclusion criteria:  |  |
| Subgroup analyses: ([section 10.11](https://training.cochrane.org/handbook/current/chapter-10#section-10-11)) | Outline any subgroups you plan to investigate for their influence on the size of the treatment effect, e.g. subgroups of the population, variations of the intervention |
| Potential included studies: | Please supply references for at least 2 RCTs relevant to this topic. Include ongoing studies listed in trial registries if relevant.1. …
2. …
 |
| Did a database search (at least PubMed) for systematic reviews / meta-analyses identify a publication relevant for your question? |
| Other information: | Outline any other factors you plan to consider in your review, or other information you would like to provide, e.g. relevance to consumers, how this review complements other published Cochrane Reviews.If there are no RCTs or ongoing studies, please explain why it is important to do this review. |
| Related Cochrane Reviews or protocols: |  |

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| Review context |
| Is the review subject to any specific funding? |  |
| Would the review form part of your postgraduate study, or of a larger research project? |  |
| Has the review already been submitted for publication or published elsewhere? |  |

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| Consumer/Patient involvement Enlisting a consumer author may not be possible or practical but we now require you to consider how you could involve consumers in the production and/or dissemination of your review. Consumers could, for example, help frame the research question, identify important outcomes and/or give feedback of their understanding of the background section of your protocol. Please follow the links for further [information](https://www.cochrane.org/news/statement-principles-consumer-involvement-cochrane) or [training](https://training.cochrane.org/involving-people).  |
| Please state if there has been any consumer involvement in identifying the topic of your review |  |
| Please state your plans for consumer involvement in your review |  |

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| Declarations of interest |
| All authors must read [Cochrane's Conflict of Interest Policy for Cochrane Library content](https://training.cochrane.org/online-learning/editorial-policies/coi-policy/coi-policy-cochrane-library). Before the title can be registered, each author must declare any relevant Conflicts of Interest (financial and non-financial) that exist or existed in the 36 months prior to this form being submitted.**Important information*** The following individuals are prohibited from being an author on a Cochrane Review:
	+ Anyone who is or has been employed in the 36 months prior to title registration by a commercial organization with a financial interest in the topic of the review.
	+ Anyone who owns a commercial organization with an interest in the topic of the review.
	+ Anyone who owns or has applied for a patent related to the topic of the review.
* Authors must declare all relevant financial interests within the 36 months prior to title registration. Such payments include (but are not limited to) speaker fees, honoraria, consultancies, membership of advisory boards and payment of travel, accommodation and conference registration expenses.
* Financial interests are considered relevant if a payment is made by a commercial organization that is developing, or manufactures, markets or distributes (anywhere in the world) an intervention or potential comparator related to the topic of the review. This applies regardless of the reported direction of effect and even if the payment was for work and advice that did not relate to the topic of the review.
* Overall, 67% (two thirds) of the author group must not have any relevant financial interests.
* The first and last author must not have any relevant financial interests and must not have been involved in industry-controlled studies (see [definitions](https://training.cochrane.org/online-learning/editorial-policies/coi-policy/coi-policy-cochrane-library#definitions) in the policy) that may be eligible for inclusion in the review.
* Anyone who has been involved in the conduct, analysis and publication of a study that could be included in the review cannot determine overall study inclusion and exclusion criteria or make study eligibility decision about, extract data from, carry out the risk of bias assessment for, or perform GRADE assessment of that study.
* Authors must remain in compliance with this policy through to the point that the review is published. If an author acquires any additional relevant financial interests while working on the review, they must inform the Review Group’s Managing Editor immediately.
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| **Have all members of the author team read** [Cochrane's Conflict of Interest Policy](https://training.cochrane.org/online-learning/editorial-policies/coi-policy/coi-policy-cochrane-library)**?** Yes 🞏 No 🞏**Do any members of the author team have a potential conflict of interest?** Yes 🞏 No 🞏 |
| If yes, you should discuss these potential conflicts with the Review Group’s Managing Editor before submitting this form. Failure to disclose relevant potential conflicts at this stage, or at any point during the writing of the review, may lead to it being rejected for publication or being removed from the Cochrane Library at a later date. Wilful failure to disclose relevant conflicts of interest will be considered a form of scientific misconduct. |

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| Authors' responsibilities |
| By completing this form, you accept responsibility for preparing, maintaining and updating the review in accordance with Cochrane policy. Cochrane Metabolic and Endocrine Disorders will provide support to assist with the preparation of the review.If drafts are not submitted by the agreed deadlines, or if the Review Group is unable to contact you for an extended period, Cochrane has the right to de‑register the title or transfer the title to alternative authors. Cochrane has the right to reject a Cochrane Review at any stage before publication (including unpublished protocols, unpublished Cochrane Reviews, and Cochrane Reviews that are being updated). Please see Cochrane’s [Rejection Policy](https://community.cochrane.org/editorial-and-publishing-policy-resource/cochrane-review-management/rejection-cochrane-reviews).You accept responsibility for maintaining the review in light of new evidence, comments and criticisms, and other developments, and updating the review based on need, or, if requested, transferring responsibility for maintaining the review to others. |
| Publication in the *Cochrane Database of Systematic Reviews (CDSR)* |
| Cochrane’s support in preparing your review is conditional upon your agreement to publish the protocol, finished review and subsequent updates in the *CDSR*. By completing this form you undertake to publish this review in the *CDSR* before publishing elsewhere (concurrent publication in other journals may be allowed in certain circumstances with prior permission).  |
| I understand the commitment required to undertake a Cochrane Review, and agree to publish first in the *CDSR*.**Signed on behalf of the authors:** |
| **Form completed by:** |
| **Date:** |

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| Proposed deadlines |
| Please be aware that in case you did not check a protocol draft into Archie (the Cochrane Collaboration’s information and management system) three months after the protocol template was sent to you, your title may be de-registered. |
| Date you plan to submit a draft protocol (within three months after the protocol template was sent to you): |  |
| Date you plan to submit a draft review (within 12 months after publication of protocol): |  |

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| Proposed referees for your review |
| Please suggest up to three referees for your review question and specify contact details. |
| Referee 1 |  |
| Referee 2 |  |

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| Review authors(see [Handbook sections II.2.1](https://training.cochrane.org/handbook/current/chapter-ii#section-ii-2-1) and [II.2.2](https://training.cochrane.org/handbook/current/chapter-ii#section-ii-2-2))In accordance with Cochrane’s [Publication Policy](https://community.cochrane.org/editorial-and-publishing-policy-resource/ethical-considerations/authorship-and-contributorship), each person named as an author must: * Make a substantial contribution to the conception and design, or analysis and interpretation of the data in the review
* Be involved in drafting the review
* Approve the final version of the review before publication
* Agree to be accountable for the accuracy and integrity of the review
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| Contact person / Author 1 Author who will take responsibility for the review, and communicate with the editorial base throughout review development; does not need to be the first listed author – please adjust numbering above. |
| Full name and qualifications: *e.g.* *Dr Xia Li, PhD* |  |
| Job title: *e.g. Registrar* |  |
| Department:  |  |
| Organisation: *e.g. West China Hospital, Sichuan University* |  |
| City:  |  |
| Country:  |  |
| Data protection and privacy If your title is accepted,as the review contact person, your affiliation and email address will be published with the completed protocol or review in the *Cochrane Database of Systematic Reviews*.Personal data collected and used for publication in the Cochrane Library are covered by the [Wiley Privacy policy](https://www.wiley.com/en-gb/privacy).Your Cochrane Account details will be visible to other groups and contributors in our contact database. If you are allocated a role as a Cochrane author, you will be able to update your profile and can choose to hide your email address and affiliation from contributors not in your primary group. |
| What expertise do you bring to the review (e.g. clinical, review methods, statistics)? |  |
| Have you prepared a systematic review before? | Yes 🞏 No 🞏 |
| If yes, have you prepared a Cochrane Review? | Yes 🞏 No 🞏 |
| If yes, please state most recent title: |  |
| Do you already have a role in another Cochrane Review Group? | Yes 🞏 No 🞏 |
| How many titles, protocols, unfinished reviews are you currently working on with Cochrane Review Groups? |  |
| If yes, which one(s)? |  |
| Level of spoken and written English:*First language / Basic / Average / Fluent* |  |
| Translating clinical trials published in languages other than English is a vital role in Cochrane. I would be willing to assist with translation of clinical trials published in these language(s): |  |

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| Author 2 You must have at least two authors to register a title. Copy this table for additional authors. |
| Full name and qualifications: *e.g.* *Dr Xia Li, PhD* |  |
| Job title: *e.g. Registrar* |  |
| Department:  |  |
| Organisation: *e.g. West China Hospital, Sichuan University* |  |
| City:  |  |
| Country:  |  |
| Data protection and privacy If your title is accepted,as the review contact person, your affiliation and email address will be published with the completed protocol or review in the *Cochrane Database of Systematic Reviews*.Personal data collected and used for publication in the Cochrane Library are covered by the [Wiley Privacy policy](https://www.wiley.com/en-gb/privacy).Your Cochrane Account details will be visible to other groups and contributors in our contact database. If you are allocated a role as a Cochrane author, you will be able to update your profile and can choose to hide your email address and affiliation from contributors not in your primary group. |
| What expertise do you bring to the review (e.g. clinical, review methods, statistics)? |  |
| Have you prepared a systematic review before? | Yes 🞏 No 🞏 |
| If yes, have you prepared a Cochrane Review? | Yes 🞏 No 🞏 |
| If yes, please state most recent title: |  |
| Do you already have a role in another Cochrane Review Group? | Yes 🞏 No 🞏 |
| How many titles, protocols, unfinished reviews are you currently working on with Cochrane Review Groups? |  |
| If yes, which one(s)? |  |
| Level of spoken and written English:*First language / Basic / Average / Fluent* |  |
| Translating clinical trials published in languages other than English is a vital role in Cochrane. I would be willing to assist with translation of clinical trials published in these language(s): |  |

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| Roles and responsibilitiesPlease advise who has agreed to undertake each of the following tasks. |
| Authors | Initials | Initials | Initials | Initials | Initials | Initials | Initials | Initials |
| Protocol |
| Draft the protocol: |  |  |  |  |  |  |  |  |
| **Review** |
| Develop and run the search strategy: \* |  |  |  |  |  |  |  |  |
| Obtain copies of studies: |  |  |  |  |  |  |  |  |
| Select which studies to include (2 people): |  |  |  |  |  |  |  |  |
| Extract data from studies (2 people): |  |  |  |  |  |  |  |  |
| Enter data into RevMan: |  |  |  |  |  |  |  |  |
| Carry out the analysis: |  |  |  |  |  |  |  |  |
| Interpret the analysis: |  |  |  |  |  |  |  |  |
| Draft the final review: |  |  |  |  |  |  |  |  |
| \*can be undertaken by our Group’s Information Specialist if needed |
| Preliminary order of review authors for citation (initials): |

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| Team resources |
| Have you read the [*Cochrane Handbook for Systematic Reviews of Interventions*](https://training.cochrane.org/handbook/current)?  | Yes 🞏 No 🞏 |
| Do you require training? | Yes 🞏 No 🞏 |
| If yes, on which topics? |  |
| Have you attended a Cochrane Review training workshop? | Yes 🞏 No 🞏 |
| If no, do you plan to register for a [future Cochrane training event](https://training.cochrane.org/search/site?f%5B0%5D=bundle%3Aworkshop&f%5B1%5D=bm_field_archived%3Afalse)? | Yes 🞏 No 🞏 |
| Which workshop did you/will you attend?  |  |
| Have you read the information for review authors on the Cochrane Metabolic and Endocrine Disorders website? (see <http://www.endoc.cochrane.org>)  | Yes 🞏 No 🞏 |
| Do you have access to the [*Cochrane Database of Systematic Reviews*](https://www.cochranelibrary.com/advanced-search)? | Yes 🞏 No 🞏 |
| Do you have access to MEDLINE (Ovid) and Embase (Elsevier.com)? | Yes 🞏 No 🞏 |
| Do you have access to a medical library? | Yes 🞏 No 🞏 |
| If yes, can you order journal articles not held in the library? | Yes 🞏 No 🞏 |
| Do you have access to advice from a medical librarian or information specialist? If yes, please provide information specialist name:  | Yes 🞏 No 🞏 |
| Do you have access to reference management software (e.g. Endnote)? | Yes 🞏 No 🞏 |
| If yes, which software, and what version? |  |
| Do you have access to a statistician (strongly recommended)? | Yes 🞏 No 🞏 |
| If yes, please provide statistician’s name: |  |
| Do you have contact with consumer groups relevant to this review? | Yes 🞏 No 🞏 |
| If yes, please list relevant consumer groups: |  |
| Have you identified appropriate time and resources to complete the review? | Yes 🞏 No 🞏 |