



Data collection form

Intervention review – RCTs and non-RCTs

This form can be used as a guide for developing your own data extraction form. Sections can be expanded and added, and irrelevant sections can be removed. It is difficult to design a single form that meets the needs of all reviews, so it is important to consider carefully the information you need to collect, and design your form accordingly. Information included on this form should be comprehensive, and may be used in the text of your review, 'Characteristics of included studies' table, risk of bias assessment, and statistical analysis.

Notes on using a data extraction form:

- Be consistent in the order and style you use to describe the information for each included study.
- Record any missing information as unclear or not described, to make it clear that the information was not found in the study report(s), not that you forgot to extract it.
- Include any instructions and decision rules on the data collection form, or in an accompanying document. It is important to practice using the form and give training to any other authors using the form.
- You will need to protect the document in order to use the form fields (Tools / Protect document)

Review title or ID

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Study ID (*surname of first author and year first full report of study was published e.g. Smith 2001*)

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Report IDs of other reports of this study (*e.g. duplicate publications, follow-up studies*)

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Notes:

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1... General Information

1. Date form completed (<i>dd/mm/yyyy</i>)	
2. Name/ID of person extracting data	
3. Report title (<i>title of paper/ abstract/ report that data are extracted from</i>)	
4. Report ID (<i>if there are multiple reports of this study</i>)	
5. Reference details	
6. Report author contact details	
7. Publication type (<i>e.g. full report, abstract, letter</i>)	

8. Study funding source <i>(including role of funders)</i>	
Possible conflicts of interest <i>(for study authors)</i>	
9. Notes:	

2... Eligibility

Study Characteristics	Review Inclusion Criteria <i>(Insert inclusion criteria for each characteristic as defined in the Protocol)</i>	Yes/ No / Unclear	Location in text <i>(pg & ¶/fig/table)</i>
10. Type of study	Randomised trial	.	
	Non-randomised trial	.	
	Controlled before-after study <ul style="list-style-type: none"> Contemporaneous data collection At least 2 intervention and 2 control clusters 	.	
	Interrupted time series OR Repeated measures study <ul style="list-style-type: none"> At least 3 timepoints before and 3 after the intervention Clearly defined intervention point 	.	
	Other design (specify):	.	
11. Participants		.	
12. Types of intervention		.	
13. Types of outcome measures		.	
14. Decision:			
15. Reason for exclusion			
16. Notes:			

DO NOT PROCEED IF STUDY EXCLUDED FROM REVIEW

3... Population and setting

	Description <i>Include comparative information for each group (i.e. intervention and controls) if available</i>	Location in text <i>(pg & ¶/fig/table)</i>
17. Population description <i>(from which study participants are drawn)</i>		
18. Setting <i>(including location and social context)</i>		
19. Inclusion criteria		
20. Exclusion criteria		
21. Method/s of recruitment of participants		
22. Notes:		

4... Methods

	Descriptions as stated in report/paper	Location in text <i>(pg & ¶/fig/table)</i>
23. Aim of study		
24. Design <i>(e.g. parallel, crossover, non-RCT)</i>		
25. Unit of allocation <i>(by individuals, cluster/groups or body parts)</i>		
26. Start date		
27. End date		
28. Duration of participation <i>(from recruitment to last follow-up)</i>		
29. Notes:		

5... Risk of Bias assessment

See [Chapter 8](#) of the Cochrane Handbook. Additional domains may be required for non-randomised studies.

Domain	Risk of bias <i>Low/ High/Unclear</i>	Support for judgement	Location in text <i>(pg & ¶/fig/table)</i>
30. Random sequence generation <i>(selection bias)</i>	.		
31. Allocation concealment <i>(selection bias)</i>	.		

Domain	Risk of bias <i>Low/ High/Unclear</i>	Support for judgement	Location in text <i>(pg & ¶/fig/table)</i>
32. Blinding of participants and personnel <i>(performance bias)</i>	.	Outcome group: All/	
<i>(if required)</i>	.	Outcome group:	
33. Blinding of outcome assessment <i>(detection bias)</i>	.	Outcome group: All/	
<i>(if required)</i>	.	Outcome group:	
34. Incomplete outcome data <i>(attrition bias)</i>	.		
35. Selective outcome reporting? <i>(reporting bias)</i>	.		
36. Other bias	.		
37. Notes:			

6... Participants

Provide overall data and, if available, comparative data for each intervention or comparison group.

	Description as stated in report/paper	Location in text <i>(pg & ¶/fig/table)</i>
38. Total no. randomised <i>(or total pop. at start of study for NRCTs)</i>		
39. Clusters <i>(if applicable, no., type, no. people per cluster)</i>		
40. Baseline imbalances		
41. Withdrawals and exclusions <i>(if not provided below by outcome)</i>		
42. Age		
43. Sex		
44. Race/Ethnicity		
45. Severity of illness		
46. Co-morbidities		
47. Other treatment received <i>(additional to study intervention)</i>		
48. Other relevant sociodemographics		
49. Subgroups measured		
50. Subgroups reported		
51. Notes:		

7... Intervention groups

Copy and paste table for each intervention and comparison group

Intervention Group 1

	Description as stated in report/paper	Location in text (pg & ¶/fig/table)
52. Group name		
53. No. randomised to group (specify whether no. people or clusters)		
54. Description (include sufficient detail for replication, e.g. content, dose, components; if it is a natural experiment, describe the pre-intervention)		
55. Duration of treatment period		
56. Timing (e.g. frequency, duration of each episode)		
57. Delivery (e.g. mechanism, medium, intensity, fidelity)		
58. Providers (e.g. no., profession, training, ethnicity etc. if relevant)		
59. Co-interventions		
60. Economic variables (i.e. intervention cost, changes in other costs as result of intervention)		
61. Resource requirements to replicate intervention (e.g. staff numbers, cold chain, equipment)		
62. Notes:		

8... Outcomes

Copy and paste table for each outcome.

Outcome 1

	Description as stated in report/paper	Location in text (pg & ¶/fig/table)
63. Outcome name		
64. Time points measured (specify whether from start or end of intervention)		
65. Time points reported		

	Description as stated in report/paper		Location in text (pg & ¶/fig/table)
66. Outcome definition <i>(with diagnostic criteria if relevant and note whether the outcome is desirable or undesirable if this is not obvious)</i>			
67. Person measuring/ reporting			
68. Unit of measurement <i>(if relevant)</i>			
69. Scales: upper and lower limits <i>(indicate whether high or low score is good)</i>			
70. Is outcome/tool validated?	Yes/No/Unclear		
71. Imputation of missing data <i>(e.g. assumptions made for ITT analysis)</i>			
72. Assumed risk estimate <i>(e.g. baseline or population risk noted in Background)</i>			
73. Notes:			

9... Results

Copy and paste the appropriate table for each outcome, including additional tables for each time point and subgroup as required.

For randomised or non-randomised trial - Dichotomous outcome

	Description as stated in report/paper				Location in text (pg & ¶/fig/table)
74. Comparison					
75. Outcome					
76. Subgroup					
77. Time point <i>(specify whether from start or end of intervention)</i>					
78. Results <i>Note whether:</i> <ul style="list-style-type: none"> • <i>post-intervention OR</i> • <i>change from baseline</i> <i>And whether</i> <ul style="list-style-type: none"> • <i>Adjusted OR</i> • <i>Unadjusted</i> 	Intervention		Comparison		
	No. events	No. participants	No. events	No. participants	
79. Baseline data	Intervention		Comparison		
	No. events	No. participants	No. events	No. participants	
80. No. missing participants and reasons					

	Description as stated in report/paper		Location in text (pg & ¶/fig/table)
81. No. participants moved from other group and reasons			
82. Any other results reported			
83. Unit of analysis (e.g. by individuals, health professional, practice, hospital, community)			
84. Statistical methods used and appropriateness of these methods (e.g. adjustment for correlation)			
85. Reanalysis required? (if yes, specify why, e.g. correlation adjustment)	Yes/No/Unclear		
86. Reanalysis possible?	Yes/No/Unclear		
87. Reanalysed results			
88. Notes:			

For randomised or non-randomised trial - Continuous outcome

	Description as stated in report/paper						Location in text (pg & ¶/fig/table)
89. Comparison							
90. Outcome							
91. Subgroup							
92. Time point (specify whether from start or end of intervention)							
93. Post-intervention or change from baseline?							
94. Results <i>Note whether:</i> <ul style="list-style-type: none"> • post-intervention OR • change from baseline <i>And whether</i> <ul style="list-style-type: none"> • Adjusted OR. • Unadjusted . 	Intervention			Comparison			
	Mean	SD (or other variance)	No. participants	Mean	SD (or other variance)	No. participants	
95. Baseline data	Intervention			Comparison			
	Mean	SD (or other variance)	No. participants	Mean	SD (or other variance)	No. participants	
96. No. missing participants and reasons							

	Description as stated in report/paper		Location in text (pg & ¶/fig/table)
97. No. participants moved from other group and reasons			
98. Any other results reported			
99. Unit of analysis (e.g. by individuals, health professional, practice, hospital, community)			
100. Statistical methods used and appropriateness of these methods (e.g. adjustment for correlation)			
101. Reanalysis required? (if yes, specify why)	Yes/No/Unclear		
102. Reanalysis possible?	Yes/No/Unclear		
103. Reanalysed results			
104. Notes:			

For randomised or non-randomised trial - Other outcome

	Description as stated in report/paper				Location in text (pg & ¶/fig/table)
105. Comparison					
106. Outcome					
107. Subgroup					
108. Time point (specify whether from start or end of intervention)					
109. Type of outcome					
110. Results	Intervention result	SD (or other variance)	Control result	SD (or other variance)	
	Overall results		SE (or other variance)		
111. No. participant	Intervention		Control		
112. No. missing participants and reasons					
113. No. participants moved from other group and reasons					
114. Any other results reported					

	Description as stated in report/paper		Location in text (pg & ¶/fig/table)
115. Unit of analysis (e.g. by individuals, health professional, practice, hospital, community)			
116. Statistical methods used and appropriateness of these methods			
117. Reanalysis required? (if yes, specify why)	.	.	
118. Reanalysis possible?	.	.	
119. Reanalysed results			
120. Notes:			

For controlled before-after study

	Description as stated in report/paper				Location in text (pg & ¶/fig/table)
121. Comparison					
122. Outcome					
123. Subgroup					
124. Timepoint (specify whether from start or end of intervention)					
125. Post-intervention or change from baseline?					
126. Results	Intervention result	SD (or other variance)	Control result	SD (or other variance)	
	Overall results		SE (or other variance)		
127. No. participants	Intervention		Control		
128. No. missing participants and reasons					
129. No. participants moved from other group and reasons					
130. Any other results reported					
131. Unit of analysis (individuals, cluster/groups or body parts)					
132. Statistical methods used and appropriateness of these methods					

	Description as stated in report/paper		Location in text (pg & ¶/fig/table)
133. Reanalysis required? <i>(specify)</i>	. . . <i>Yes/No/Unclear</i>		
134. Reanalysis possible?	. . . <i>Yes/No/Unclear</i>		
135. Reanalysed results			
136. Notes:			

For interrupted time series or repeated measures study

	Description as stated in report/paper		Location in text (pg & ¶/fig/table)	
137. Comparison				
138. Outcome				
139. Subgroup				
140. Length of timepoints measured <i>(e.g. days, months)</i>				
Total period measured				
141. No. participants measured				
142. No. missing participants and reasons				
143. No. timepoints measured	144. Pre-intervention	145. Post-intervention		
146. Mean value <i>(with variance measure)</i>				
147. Difference in means (post – pre)				
148. Percent relative change				
149. Result reported by authors <i>(with variance measure)</i>				
150. Unit of analysis <i>(individuals or cluster/groups)</i>				
151. Statistical methods used and appropriateness of these methods				
152. Reanalysis required? <i>(specify)</i>	. . . <i>Yes/No/Unclear</i>			
153. Reanalysis possible?	. . . <i>Yes/No/Unclear</i>			
154. Individual timepoint results				
155. Read from figure?	. . . <i>Yes/No/Unclear</i>			
156. Reanalysed results	Change in level	SE	Change in slope	SE
157. Notes:				

10. Applicability

158. Have important populations been excluded from the study? <i>(consider disadvantaged populations, and possible differences in the intervention effect)</i>	. <i>Yes/No/Unclear</i>	
159. Is the intervention likely to be aimed at disadvantaged groups? <i>(e.g. lower socioeconomic groups)</i>	. <i>Yes/No/Unclear</i>	
160. Does the study directly address the review question? <i>(any issues of partial or indirect applicability)</i>	. <i>Yes/No/Unclear</i>	
161. Notes:		

11. Other information

	Description as stated in report/paper	Location in text <i>(pg & ¶/fig/table)</i>
162. Key conclusions of study authors		
163. References to other relevant studies		
164. Correspondence required for further study information <i>(what and from whom)</i>		
165. Further study information requested <i>(from whom, what and when)</i>		
166. Correspondence received <i>(from whom, what and when)</i>		
167. Notes:		