



CONSORT 2010 checklist of information to include when reporting a randomised trial*

Section/Topic	Item No	Checklist item	Reported on page No
Title and abstract			
	1a	Identification as a randomised trial in the title	1, I. 3
	1b	Structured summary of trial design, methods, results, and conclusions (for specific guidance see CONSORT for abstracts)	2, II. 38 f.
Introduction			
Background and objectives	2a	Scientific background and explanation of rationale	3, II. 70 f.
	2b	Specific objectives or hypotheses	4, II. 111 f.
Methods			
Trial design	3a	Description of trial design (such as parallel, factorial) including allocation ratio	5, II. 125 f.
	3b	Important changes to methods after trial commencement (such as eligibility criteria), with reasons	not applicable
Participants	4a	Eligibility criteria for participants	5, II. 130 f.
	4b	Settings and locations where the data were collected	5, II. 130 f.
Interventions	5	The interventions for each group with sufficient details to allow replication, including how and when they were actually administered	6, II. 166 f. 7, II. 175 f.
Outcomes	6a	Completely defined pre-specified primary and secondary outcome measures, including how and when they were assessed	7, II. 184 f. 7, II. 194 f.
	6b	Any changes to trial outcomes after the trial commenced, with reasons	not applicable
Sample size	7a	How sample size was determined	8, II. 221 f.
	7b	When applicable, explanation of any interim analyses and stopping guidelines	not applicable
Randomisation:			
Sequence generation	8a	Method used to generate the random allocation sequence	5, II. 121 f.
	8b	Type of randomisation; details of any restriction (such as blocking and block size)	
Allocation concealment mechanism	9	Mechanism used to implement the random allocation sequence (such as sequentially numbered containers), describing any steps taken to conceal the sequence until interventions were assigned	5, II. 121 f.
Implementation	10	Who generated the random allocation sequence, who enrolled participants, and who assigned participants to interventions	5, II. 121 f.
Blinding	11a	If done, who was blinded after assignment to interventions (for example, participants, care providers, those	5, II. 147 f.

	assessing outcomes) and how	
	If relevant, description of the similarity of interventions	not applicable
11b	Statistical methods used to compare groups for primary and secondary outcomes	8, II. 207 f.
12a	Methods for additional analyses, such as subgroup analyses and adjusted analyses	8, II. 216 ff.
Results		
13a	For each group, the numbers of participants who were randomly assigned, received intended treatment, and were analysed for the primary outcome	5, II. 134 ff.
13b	For each group, losses and exclusions after randomisation, together with reasons	8, II. 228 - 229
14a	Dates defining the periods of recruitment and follow-up	5, II. 145 f.
14b	Why the trial ended or was stopped	not applicable
15	A table showing baseline demographic and clinical characteristics for each group	not applicable
16	For each group, number of participants (denominator) included in each analysis and whether the analysis was by original assigned groups	5, II. 134 ff.
17a	For each primary and secondary outcome, results for each group, and the estimated effect size and its precision (such as 95% confidence interval)	9, II. 232 f. 17, I. 467 17, I. 472
17b	For binary outcomes, presentation of both absolute and relative effect sizes is recommended	not applicable
18	Results of any other analyses performed, including subgroup analyses and adjusted analyses, distinguishing pre-specified from exploratory	not applicable
19	All important harms or unintended effects in each group (for specific guidance see CONSORT for harms)	not applicable
Discussion		
20	Trial limitations, addressing sources of potential bias, imprecision, and, if relevant, multiplicity of analyses	12, II. 321 f.
21	Generalisability (external validity, applicability) of the trial findings	not applicable
22	Interpretation consistent with results, balancing benefits and harms, and considering other relevant evidence	11, II. 300 f.
Other information		
23	Registration number and name of trial registry	5, II. 121 f.
24	Where the full trial protocol can be accessed, if available	not applicable
25	Sources of funding and other support (such as supply of drugs), role of funders	13, II. 375 f.

*We strongly recommend reading this statement in conjunction with the CONSORT 2010 Explanation and Elaboration for important clarifications on all the items. If relevant, we also recommend reading CONSORT extensions for cluster randomised trials, non-inferiority and equivalence trials, non-pharmacological treatments, herbal interventions, and pragmatic trials. Additional extensions are forthcoming: for those and for up to date references relevant to this checklist, see www.consort-statement.org.