



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

Section/Topic	Item No	Checklist item	Reported on Page Number/Line Number	Reported on Section/Paragraph
<b>Title and abstract</b>				
	1a	Identification as a randomised trial in the title	Page 1 Line 2	Section 1 Paragraph 1
	1b	Structured summary of trial design, methods, results, and conclusions (for specific guidance see Table 2)	Page 2 Line 2-24	Section 1 Paragraph 4
<b>Introduction</b>				
Background and objectives	2a	Scientific background and explanation of rationale	Page 3 Line 2-22	Section 2 Paragraph 1
	2b	Specific objectives or hypotheses	Page 4 Line 1-3	Section 2 Paragraph 2
<b>Methods</b>				
Trial design	3a	Description of trial design (such as parallel, factorial) including allocation ratio	Page 5 Line 9-13	Section 3 Paragraph 1
	3b	Important changes to methods after trial commencement (such as eligibility criteria), with reasons	Page 5 Line 15-23, Page 6 Line 1-5	Section 3 Paragraph 2
Participants	4a	Eligibility criteria for participants	Page 5 Line 3-7	Section 3 Paragraph 1
	4b	Settings and locations where the data were collected	Page 5 Line 5-7	Section 3 Paragraph 1
Interventions	5	The interventions for each group with sufficient details to allow replication, including how and when they were actually administered	Page 5 Line 9-13	Section 3 Paragraph 1
Outcomes	6a	Completely defined pre-specified primary and secondary outcome measures, including how and when they were assessed	Page 5 Line 15-23, Page 6 Line 1-5	Section 3 Paragraph 2
	6b	Any changes to trial outcomes after the trial commenced, with reasons	Page 5 Line 15-23, Page 6 Line 1-5	Section 3 Paragraph 2
Sample size	7a	How sample size was determined	Page 5 Line 10	Section 3 Paragraph 1
	7b	When applicable, explanation of any interim analyses and stopping guidelines	Page 5 Line 15-23, Page 6 Line 1-5	Section 3 Paragraph 2
Randomisation:				
Sequence generation	8a	Method used to generate the random allocation sequence	Page 5 Line 3-13	Section 3 Paragraph 1
	8b	Type of randomisation; details of any restriction (such as blocking and block size)	Page 5 Line 7-9	Section 3 Paragraph 1
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	Page 5 Line 3-13	Section 3 Paragraph 1

Implementation	10	Who generated the random allocation sequence, who enrolled participants, and who assigned participants to interventions	Page 5 Line 3-7	Section 3 Paragraph 1
Blinding	11a	If done, who was blinded after assignment to interventions (for example, participants, care providers, those assessing outcomes) and how	Page 5 Line 3-7	Section 3 Paragraph 1
	11b	If relevant, description of the similarity of interventions	Page 5 Line 5-7	Section 3 Paragraph 1
Statistical methods	12a	Statistical methods used to compare groups for primary and secondary outcomes	Page 6 Line 9-10	Section 3 Paragraph 3
	12b	Methods for additional analyses, such as subgroup analyses and adjusted analyses	Page 6 Line 7-9	Section 3 Paragraph 3
<b>Results</b>				
Participant flow (a diagram is strongly recommended)	13a	For each group, the numbers of participants who were randomly assigned, received intended treatment, and were analysed for the primary outcome	Page 5 Line 7-13	Section 3 Paragraph 1
	13b	For each group, losses and exclusions after randomisation, together with reasons	Page 5 Line 7-9	Section 3 Paragraph 1
Recruitment	14a	Dates defining the periods of recruitment and follow-up	Page 5 Line 5	Section 3 Paragraph 1
	14b	Why the trial ended or was stopped	Page 5 Line 5-7	Section 3 Paragraph 1
Baseline data	15	A table showing baseline demographic and clinical characteristics for each group	Page 5 Line 3-13	Section 3 Paragraph 1
Numbers analysed	16	For each group, number of participants (denominator) included in each analysis and whether the analysis was by original assigned groups	Page 5 Line 9-13	Section 3 Paragraph 1
Outcomes and estimation	17a	For each primary and secondary outcome, results for each group, and the estimated effect size and its precision (such as 95% confidence interval)	Page 7 Line 18-23, Page 8 Line 1-5	Section 4 Paragraph 2
	17b	For binary outcomes, presentation of both absolute and relative effect sizes is recommended	Page 7 Line 3-16	Section 4 Paragraph 1
Ancillary analyses	18	Results of any other analyses performed, including subgroup analyses and adjusted analyses, distinguishing pre-specified from exploratory	Page 7 Line 18-23, Page 8 Line 1-5	Section 4 Paragraph 2
Harms	19	All important harms or unintended effects in each group (for specific guidance see CONSORT for harms)	Page 7 Line 18-23, Page 8 Line 1-5	Section 4 Paragraph 2
<b>Discussion</b>				
Limitations	20	Trial limitations, addressing sources of potential bias, imprecision, and, if relevant, multiplicity of analyses	Page 10 Line 8-10	Section 5 Paragraph 3
Generalisability	21	Generalisability (external validity, applicability) of the trial findings	Page 9 Line 2-10	Section 5 Paragraph 1
Interpretation	22	Interpretation consistent with results, balancing benefits and harms, and considering other relevant evidence	Page 9 Line 11-23, Page 10 Line 1-7	Section 5 Paragraph 2
<b>Other information</b>				
Registration	23	Registration number and name of trial registry	Page 10 Line 18-20	Section 6 Paragraph 3

Protocol	24	Where the full trial protocol can be accessed, if available	Page 10 Line 22	Section 6 Paragraph 4
Funding	25	Sources of funding and other support (such as supply of drugs), role of funders	Page 11 Line 2-5	Section 1 Paragraph 5

\*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](http://www.consort-statement.org).

**Table 2 Items to include when reporting a randomized trial in a journal or conference abstract**

Item	Description	Reported on Page Number/Line Number	Reported on Section/Paragraph
Title	Identification of the study as randomized	Page 1 Line 2	Section 1 Paragraph 1
Authors *	Contact details for the corresponding author	Page 1 Line 19-22	Section 1 Paragraph 3
Trial design	Description of the trial design (e.g. parallel, cluster, non-inferiority)	Page 2 Line 6-14	Section 1 Paragraph 4
<b>Methods</b>			
Participants	Eligibility criteria for participants and the settings where the data were collected	Page 5 Line 4-7	Section 3 Paragraph 1
Interventions	Interventions intended for each group	Page 5 Line 10-13	Section 3 Paragraph 1
Objective	Specific objective or hypothesis	Page 5 Line 12-13	Section 3 Paragraph 1
Outcome	Clearly defined primary outcome for this report	Page 5 Line 15	Section 3 Paragraph 2
Randomization	How participants were allocated to interventions	Page 5 Line 5-9	Section 3 Paragraph 1
Blinding (masking)	Whether or not participants, care givers, and those assessing the outcomes were blinded to group assignment	Page 5 Line 3-13	Section 3 Paragraph 1
<b>Results</b>			
Numbers randomized	Number of participants randomized to each group	Page 5 Line 7-8	Section 3 Paragraph 1
Recruitment	Trial status	Page 5 Line 3-7	Section 3 Paragraph 1
Numbers analysed	Number of participants analysed in each group	Page 5 Line 7-8	Section 3 Paragraph 1
Outcome	For the primary outcome, a result for each group and the estimated effect size and its precision	Page 7 Line 3-13	Section 4 Paragraph 1
Harms	Important adverse events or side effects	Page 7 Line 15-23	Section 4 Paragraph 2

Conclusions	General interpretation of the results	Page 10 Line 5-7	Section 5 Paragraph 2
Trial registration	Registration number and name of trial register	Page 10 Line 18-20	Section 6 Paragraph 3
Funding	Source of funding	Page 11 Line 2-5	Section 6 Paragraph 5

\* this item is specific to conference abstracts

From: Hopewell S, Clarke M, Moher D, et al. CONSORT for reporting randomized controlled trials in journal and conference abstracts: explanation and elaboration. PLoS Med. 2008;5(1):e20

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\*As the checklist was provided upon initial submission, the page number/line number reported may be changed due to copyediting and may not be referable in the published version. In this case, the section/paragraph may be used as an alternative reference.