

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	N.A
	1b	Structured summary of trial design, methods, results, and conclusions (for specific guidance see CONSORT for abstracts)	p 4, 5
Introduction			
Background and objectives	2a	Scientific background and explanation of rationale	p 6
	2b	Specific objectives or hypotheses	p 6, 7
Methods			
Trial design			
	3a	Description of trial design (such as parallel, factorial) including allocation ratio	p 7
Participants			
	3b	Important changes to methods after trial commencement (such as eligibility criteria), with reasons	N.A
	4a	Eligibility criteria for participants	p 7
	4b	Settings and locations where the data were collected	p 7
Interventions			
	5	The interventions for each group with sufficient details to allow replication, including how and when they were actually administered	p 8
Outcomes			
	6a	Completely defined pre-specified primary and secondary outcome measures, including how and when they were assessed	p 10
Sample size			
	6b	Any changes to trial outcomes after the trial commenced, with reasons	N.A
	7a	How sample size was determined	N.A
	7b	When applicable, explanation of any interim analyses and stopping guidelines	N.A
Randomisation:			
Sequence generation			
	8a	Method used to generate the random allocation sequence	N.A
	8b	Type of randomisation; details of any restriction (such as blocking and block size)	N.A
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	N.A
Implementation			
	10	Who generated the random allocation sequence, who enrolled participants, and who assigned participants to interventions	N.A
Blinding			
	11a	If done, who was blinded after assignment to interventions (for example, participants, care providers, those	N.A

	assessing outcomes) and how	N.A
	11b If relevant, description of the similarity of interventions	N.A
Statistical methods	12a Statistical methods used to compare groups for primary and secondary outcomes	N.A
	12b Methods for additional analyses, such as subgroup analyses and adjusted analyses	N.A
Results		
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	P7
Recruitment	13b For each group, losses and exclusions after randomisation, together with reasons	N.A
	14a Dates defining the periods of recruitment and follow-up	P7
	14b Why the trial ended or was stopped	N.A
Baseline data	15 A table showing baseline demographic and clinical characteristics for each group	N.A
Numbers analysed	16 For each group, number of participants (denominator) included in each analysis and whether the analysis was by original assigned groups	N.A
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)	P10-13
	17b For binary outcomes, presentation of both absolute and relative effect sizes is recommended	N.A
Ancillary analyses	18 Results of any other analyses performed, including subgroup analyses and adjusted analyses, distinguishing pre-specified from exploratory	P10-13
Harms	19 All important harms or unintended effects in each group (for specific guidance see CONSORT for harms)	P11
Discussion		
Limitations	20 Trial limitations, addressing sources of potential bias, imprecision, and, if relevant, multiplicity of analyses	P17
Generalisability	21 Generalisability (external validity, applicability) of the trial findings	P13-16
Interpretation	22 Interpretation consistent with results, balancing benefits and harms, and considering other relevant evidence	P13-16
Other information		
Registration	23 Registration number and name of trial registry	N.A
Protocol	24 Where the full trial protocol can be accessed, if available	P7
Funding	25 Sources of funding and other support (such as supply of drugs), role of funders	P17

*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.