

STROBE Statement—checklist of items that should be included in reports of observational studies

	<b>Item No.</b>	<b>Recommendation</b>	<b>Page No.</b>
<b>Title and abstract</b>	1	(a) Indicate the study’s design with a commonly used term in the title or the abstract	Page 2, Line 36
		(b) Provide in the abstract an informative and balanced summary of what was done and what was found	Page 2, Line 36-44 Page 3, Line 45-51
<b>Introduction</b>			
Background/rationale	2	Explain the scientific background and rationale for the investigation being reported	Page 3-4 Lines 55-84
Objectives	3	State specific objectives, including any prespecified hypotheses	Page 4 Lines 84-87
<b>Methods</b>			
Study design	4	Present key elements of study design early in the paper	Page 5 Lines 88-94
Setting	5	Describe the setting, locations, and relevant dates, including periods of recruitment, exposure, follow-up, and data collection	Pages 5 Lines 88-100 Lines 113-116
Participants	6	(a) <i>Cohort study</i> —Give the eligibility criteria, and the sources and methods of selection of participants. Describe methods of follow-up	N/A
		<i>Case-control study</i> —Give the eligibility criteria, and the sources and methods of case ascertainment and control selection. Give the rationale for the choice of cases and controls	N/A
		<i>Cross-sectional study</i> —Give the eligibility criteria, and the sources and methods of selection of participants	Page 5-6 Line 104-116
		(b) <i>Cohort study</i> —For matched studies, give matching criteria and number of exposed and unexposed <i>Case-control study</i> —For matched studies, give matching criteria and the number of controls per case	N/A
Variables	7	Clearly define all outcomes, exposures, predictors, potential confounders, and effect modifiers. Give diagnostic criteria, if applicable	Page 6-7 Lines 125-135
Data sources/ measurement	8*	For each variable of interest, give sources of data and details of methods of assessment (measurement). Describe comparability of assessment methods if there is more than one group	Page 6 Lines 113-124

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Bias	9	Describe any efforts to address potential sources of bias	Page 5 Lines 94-100
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Study size	10	Explain how the study size was arrived at	Page 5 101-106
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Quantitative variables	11	Explain how quantitative variables were handled in the analyses. If applicable, describe which groupings were chosen and why	Page 7 Lines 145-146
Statistical methods	12	(a) Describe all statistical methods, including those used to control for confounding	Page 7 Line 146-152
		(b) Describe any methods used to examine subgroups and interactions	Page 7 Line 152-154
		(c) Explain how missing data were addressed	Page 7 Line 152-153
		(d) <i>Cohort study</i> —If applicable, explain how loss to follow-up was addressed <i>Case-control study</i> —If applicable, explain how matching of cases and controls was addressed <i>Cross-sectional study</i> —If applicable, describe analytical methods taking account of sampling strategy	N/A
		(e) Describe any sensitivity analyses	Page 7 Line 149-150
<b>Results</b>			
Participants	13*	(a) Report numbers of individuals at each stage of study—eg numbers potentially eligible, examined for eligibility, confirmed eligible, included in the study, completing follow-up, and analysed	Page 8 Line 157
		(b) Give reasons for non-participation at each stage	N/A
		(c) Consider use of a flow diagram	N/A
Descriptive data	14*	(a) Give characteristics of study participants (eg demographic, clinical, social) and information on exposures and potential confounders	Page 8-9 Lines 157-171 Page 12 Line 236
		(b) Indicate number of participants with missing data for each variable of interest	Page 8 Line 170 (Table 1) Page 12 Line 236 (Table 4)
		(c) <i>Cohort study</i> —Summarise follow-up time (eg, average and total amount)	N/A
Outcome data	15*	<i>Cohort study</i> —Report numbers of outcome events or summary measures over time	N/A
		<i>Case-control study</i> —Report numbers in each exposure category, or summary measures of exposure	N/A
		<i>Cross-sectional study</i> —Report numbers of outcome events or summary measures	Pages 9-10 Lines 177-194

Main results	16	(a) Give unadjusted estimates and, if applicable, confounder-adjusted estimates and their precision (eg, 95% confidence interval). Make clear which confounders were adjusted for and why they were included	Page 10 Line 195-199 (Figure 2)
		(b) Report category boundaries when continuous variables were categorized	Page 11 Line 213-219
		(c) If relevant, consider translating estimates of relative risk into absolute risk for a meaningful time period	Page 12 Line 227-234

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Other analyses	17	Report other analyses done—eg analyses of subgroups and interactions, and sensitivity analyses	Page 14 Line 262 -264
<b>Discussion</b>			
Key results	18	Summarise key results with reference to study objectives	Page 15 Line 266-272
Limitations	19	Discuss limitations of the study, taking into account sources of potential bias or imprecision. Discuss both direction and magnitude of any potential bias	Page 20-21 Line 373-395
Interpretation	20	Give a cautious overall interpretation of results considering objectives, limitations, multiplicity of analyses, results from similar studies, and other relevant evidence	Page 21 Line 397-400
Generalisability	21	Discuss the generalisability (external validity) of the study results	Page 21 Line 400-404
<b>Other information</b>			
Funding	22	Give the source of funding and the role of the funders for the present study and, if applicable, for the original study on which the present article is based	Page 22-23 Line 436-439

\*Give information separately for cases and controls in case-control studies and, if applicable, for exposed and unexposed groups in cohort and cross-sectional studies.

**Note:** An Explanation and Elaboration article discusses each checklist item and gives methodological background and published examples of transparent reporting. The STROBE checklist is best used in conjunction with this article (freely available on the Web sites of PLoS Medicine at <http://www.plosmedicine.org/>, Annals of Internal Medicine at <http://www.annals.org/>, and Epidemiology at <http://www.epidem.com/>). Information on the STROBE Initiative is available at [www.strobe-statement.org](http://www.strobe-statement.org).