

STROBE Statement—Checklist of items that should be included in reports of *cohort studies*

	Item No	Recommendation
Title and abstract	1	(a) Indicate the study's design with a commonly used term in the title or the abstract (First line, 1st page) (b) Provide in the abstract an informative and balanced summary of what was done and what was found (Abstract, 2nd page)
Introduction		
Background/rationale	2	Explain the scientific background and rationale for the investigation being reported (Paragraph 1-2, 4th page)
Objectives	3	State specific objectives, including any prespecified hypotheses (2nd Paragraph, 4th page)
Methods		
Study design	4	Present key elements of study design early in the paper hypotheses (2nd line, 5th page)
Setting	5	Describe the setting, locations, and relevant dates, including periods of recruitment, exposure, follow-up, and data collection (2nd paragraph, 5th page)
Participants	6	(a) Give the eligibility criteria, and the sources and methods of selection of participants. Describe methods of follow-up (2nd paragraph, 6th page) (b) For matched studies, give matching criteria and number of exposed and unexposed (N/A)
Variables	7	Clearly define all outcomes, exposures, predictors, potential confounders, and effect modifiers. Give diagnostic criteria, if applicable (2nd paragraph, 5th page ; 3rd paragraph, 6th page; and 1st paragraph, 7th page)
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 (2nd paragraph, 5th page and 3rd paragraph, 6th page, 1st paragraph, 7th page)
Bias	9	Describe any efforts to address potential sources of bias (2nd paragraph, 6th page)
Study size	10	Explain how the study size was arrived at (2nd paragraph, 6th page)
Quantitative variables	11	Explain how quantitative variables were handled in the analyses. If applicable, describe which groupings were chosen and why (Last sentence, 3rd paragraph, 8th page)
Statistical methods	12	(a) Describe all statistical methods, including those used to control for confounding (b) Describe any methods used to examine subgroups and interactions (c) Explain how missing data were addressed (d) If applicable, explain how loss to follow-up was addressed (e) Describe any sensitivity analyses (Last paragraph, 6th page to 2nd paragraph, 7th page; and 2nd paragraph, 9th page)
Results		
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 (b) Give reasons for non-participation at each stage (c) Consider use of a flow diagram (2nd paragraph, 6th page and last paragraph, 7th page)
Descriptive data	14*	(a) Give characteristics of study participants (eg demographic, clinical, social) and information on exposures and potential confounders

		(b) Indicate number of participants with missing data for each variable of interest
		(c) Summarise follow-up time (eg, average and total amount) (last paragraph, 7th page to 2nd paragraph, 8th page)
Outcome data	15*	Report numbers of outcome events or summary measures over time (2nd paragraph, 8th page)
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 (b) Report category boundaries when continuous variables were categorized (c) If relevant, consider translating estimates of relative risk into absolute risk for a meaningful time period (Page 8 to 11th)
Other analyses	17	Report other analyses done—eg analyses of subgroups and interactions, and sensitivity analyses (2nd paragraph, 9th page)
Discussion		
Key results	18	Summarise key results with reference to study objectives (1st paragraph, 12th page)
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 (1st paragraph, 14th page)
Interpretation	20	Give a cautious overall interpretation of results considering objectives, limitations, multiplicity of analyses, results from similar studies, and other relevant evidence (2nd paragraph, 12th page to 2nd paragraph, 13th page)
Generalisability	21	Discuss the generalisability (external validity) of the study results (Last sentence, 1st paragraph, 14th page)
Other information		
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 (Last sentence, page 1st)

*Give information separately for exposed and unexposed groups.

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 <http://www.strobe-statement.org>.