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

Section/item	Item No	Recommendation	Reported on Page Number/Line Number	Reported on Section/Paragraph
Title and abstract	1	(a) Indicate the study's design with a commonly used term in the title or the abstract	Line 32	Abstract
		(b) Provide in the abstract an informative and balanced summary of what was done and what was found	Page 2	Abstract
Introduction				
Background/ rationale	2	Explain the scientific background and rationale for the investigation being reported	Lines 57-69	Introduction Paragraph 2-3
Objectives	3	State specific objectives, including any prespecified hypotheses	Line 70-72 (see notes)	Introduction - last
Methods				
Study design	4	Present key elements of study design early in the paper	Lines 77-82; 98-107	Materials and methodes:
Setting	5	Describe the setting, locations, and relevant dates, including periods of recruitment, exposure, follow-up, and data collection	Lines 77-82, 98-107	Materials and Methodes: Paragraph 1 and 3
Participants	6	(a) ***Cohort study***—Give the eligibility criteria, and the sources and methods of selection of participants. Describe methods of follow-up ***Case-control study***—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 ***Cross-sectional study***—Give the eligibility criteria, and the sources and methods of selection of participants	Lines 77-82	Materials and Methodes: Paragraph 1
		(b) ***Cohort study***—For matched studies, give matching criteria and number of exposed and unexposed ***Case-control study***—For matched studies, give matching criteria and the number of controls per case	Lines 98-100 Lines 108-109 (see notes)	Materials and Methodes: Paragraph 3 and 4
Variables	7	Clearly define all outcomes, exposures, predictors, potential confounders, and effect modifiers. Give diagnostic criteria, if applicable	Lines 98-107	Materials and Methodes: Paragraph 3
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	Lines 83-92; (see notes)	Materials and Methodes: Paragraph 2
Bias	9	Describe any efforts to address potential sources of bias	108-109 (see notes)	Materials and Methodes:
Study size	10	Explain how the study size was arrived at	Lines 78-82	Materials and Methodes:
Quantitative variables	11	Explain how quantitative variables were handled in the analyses. If applicable, describe which groupings were chosen and why	Lines 108 - 117	Materials and Methodes: Paragraph 4

Statistical methods	12	(a) Describe all statistical methods, including those used to control for confounding	Lines 108 - 117	Materials and Methodes:
		(b) Describe any methods used to examine subgroups and interactions	Lines 108-117. (see notes)	Materials and Methodes:
		(c) Explain how missing data were addressed	Line 115	Materials and Methodes:
		(d) ***Cohort study***—If applicable, explain how loss to follow-up was addressed ***Case-control study***—If applicable, explain how matching of cases and controls was addressed ***Cross-sectional study***—If applicable, describe analytical methods taking account of sampling strategy	Lines 108-109 Table 2 (see notes)	Materials and Methodes: Paragraph 4
		(e) Describe any sensitivity analyses	Lines 108 - 117	Materials and Methodes:
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	Lines 120-139 (see notes)	Results: Paragraph 1
		(b) Give reasons for non-participation at each stage	Lines 120 - 139	Results: Paragraph 1
		(c) Consider use of a flow diagram	not used (see Notes)	
Descriptive data	14*	(a) Give characteristics of study participants (eg demographic, clinical, social) and information on exposures and potential confounders	Lines 120 - 139 Table 1 and 2.	Results: Paragraph 1
		(b) Indicate number of participants with missing data for each variable of interest	Lines 120 - 139	Results: Paragraph 1
		(c) ***Cohort study***—Summarise follow-up time (eg, average and total amount)	Line 120	Results: Paragraph 1
Outcome data	15*	***Cohort study***—Report numbers of outcome events or summary measures over time	no cohort study	
		Case-control study—Report numbers in each exposure category, or summary measures of exposure	Lines 140-141; Lines	Results: Paragraph 2; 4;
		Cross-sectional study—Report numbers of outcome events or summary measures	no cross sectional study	
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	Table 2	Table 2
		(b) Report category boundaries when continuous variables were categorized	no categorization of	
		(c) If relevant, consider translating estimates of relative risk into absolute risk for a meaningful time period	not relevant	
Other analyses	17	Report other analyses done—eg analyses of subgroups and interactions, and sensitivity analyses	no subgroups in study	
Discussion				
Key results	18	Summarise key results with reference to study objectives	Lines 189-204; Lines	Discussion: Paragraph 5-7;
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	Lines 219 - 222 (see notes)	Discussion: Paragraph 9

Interpretation	20	Give a cautious overall interpretation of results considering objectives, limitations, multiplicity of analyses, results from similar studies, and other relevant evidence	Lines 232-237	Discussion: last paragraph
Generalisability	21	Discuss the generalisability (external validity) of the study results	lines 235-237	Discussion: last paragraph
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	Line 243: no funding	

*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.

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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.