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	4/81-83, 88-89, 5/90-91	Abstract/1,2
		(b) Provide in the abstract an informative and balanced summary of what was done and what was found	4/88-89, 5/90-100	Abstract/2.3
Introduction				
Background/ rationale	2	Explain the scientific background and rationale for the investigation being reported	6/129-138, 7/139-140, 155-159	Background/3-6
Objectives	3	State specific objectives, including any prespecified hypotheses	7/155-159	Background/6
Methods				
Study design	4	Present key elements of study design early in the paper	8/164-166	Methods/1
Setting	5	Describe the setting, locations, and relevant dates, including periods of recruitment, exposure, follow-up, and data collection	8/164-166, 176-180, 9/189-190	Methods/1, 2, 4
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	7/147-153, 9/209-214	Background/5 Patient Selection/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	Our study is not matched	Our study is not matched
Variables	7	Clearly define all outcomes, exposures, predictors, potential confounders, and effect modifiers. Give diagnostic criteria, if applicable	8/176-189	Methods/2, 3,4
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	8/164-166, 176-180, 182-184, 186-1989	Methods/1,2,3,4
Bias	9	Describe any efforts to address potential sources of bias	7/147-153, 8/170-173	Backgr/ 5, Methods/1
Study size	10	Explain how the study size was arrived at	9/209-215	Patient selection/1
Quantitative variables	11	Explain how quantitative variables were handled in the analyses. If applicable, describe which groupings were chosen and why	9/93	Methods/5

Statistical methods	12	(a) Describe all statistical methods, including those used to control for confounding	9/192	Methods/5
		(b) Describe any methods used to examine subgroups and interactions	No sub-groups were	No sub-groups were
		(c) Explain how missing data were addressed	9/194-195	Methods/5
		(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	9/189-190	Methods/4
		(e) Describe any sensitivity analyses	No sensitivity analyses	No sensitivity analyses
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	12/256-257	Results/1
		(b) Give reasons for non-participation at each stage	All subjects participated	All subjects participated
		(c) Consider use of a flow diagram	No flow diagram was used	No flow diagram was used
Descriptive data	14*	(a) Give characteristics of study participants (eg demographic, clinical, social) and information on exposures and potential confounders	12/256- 262	Results/1
		(b) Indicate number of participants with missing data for each variable of interest	9/194-195	Methods/5
		(c) ***Cohort study***—Summarise follow-up time (eg, average and total amount)	9/189-190	Methods/4
Outcome data	15*	***Cohort study***—Report numbers of outcome events or summary measures over time	Table 2	Table 2
		Case-control study—Report numbers in each exposure category, or summary measures of exposure	Not a case-control study	Not case-control study
		Cross-sectional study—Report numbers of outcome events or summary measures	Not a cross-sectional study	Not 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	12/256-257, 269-294	Results/1,3-7
		(b) Report category boundaries when continuous variables were categorized	12/256-258, 12/ 264-265,	Results/1,2,3
		(c) If relevant, consider translating estimates of relative risk into absolute risk for a meaningful time period	Not relevant to the study	Not relevant to the study
Other analyses	17	Report other analyses done—eg analyses of subgroups and interactions, and sensitivity analyses	No other analysis was	No other analysis was
Discussion				
Key results	18	Summarise key results with reference to study objectives	14/315-319, 324-326,	Discussion/3,4,6
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	15/356-357	Conclusion/2

Interpretation	20	Give a cautious overall interpretation of results considering objectives, limitations, multiplicity of analyses, results from similar studies, and other relevant evidence	14/310-326, 15/333-343	Discussion/2-6
Generalisability	21	Discuss the generalisability (external validity) of the study results	15/347-352	Conclusion/1
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	No funding was used for this study	No funding was used for this study

*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.