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	Page 3 / Line 55	Methods section
		(b) Provide in the abstract an informative and balanced summary of what was done and what was found	Page 3 / Line 49	Abstract section
Introduction				
Background/ rationale	2	Explain the scientific background and rationale for the investigation being reported	Page 5 / Line 122-138	Introduction section
Objectives	3	State specific objectives, including any prespecified hypotheses	Page 5 / Line 139-145	Introduction section
Methods				
Study design	4	Present key elements of study design early in the paper	Page 6 / Line 148-162	Patients and Methods
Setting	5	Describe the setting, locations, and relevant dates, including periods of recruitment, exposure, follow-up, and data collection	Page 6 / Line 148-162	Patients and Methods section
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	Page 6 / Line 148-162 Retrospective, cohort study utilizing an institutional database	Patients and Methods section
		(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	Patients were not matched	
Variables	7	Clearly define all outcomes, exposures, predictors, potential confounders, and effect modifiers. Give diagnostic criteria, if applicable	Page 6 / Line 163-165	Patients and Methods section
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 / Line 148-162	Patients and Methods section
Bias	9	Describe any efforts to address potential sources of bias	Page 10 / Line 272-287	Comments section
Study size	10	Explain how the study size was arrived at	Page 6 / Line 148-162	Patients and Methods
Quantitative variables	11	Explain how quantitative variables were handled in the analyses. If applicable, describe which groupings were chosen and why	Page 6 / Line 163-165	Patients and Methods section

Statistical methods	12	(a) Describe all statistical methods, including those used to control for confounding	Page 6 / Line 166-164	Patients and Methods
		(b) Describe any methods used to examine subgroups and interactions	Page 6 / Line 166-164	Patients and Methods
		(c) Explain how missing data were addressed	Page 10 / Line 271-282	Comments section
		(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	Page 10 / Line 271-282	Comments section
		(e) Describe any sensitivity analyses	NA (none done)	NA
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	Page 7 / Line 177-179	Results section
		(b) Give reasons for non-participation at each stage	Page 7 / Line 181-184	Results section
		(c) Consider use of a flow diagram	NA (none done)	NA
Descriptive data	14*	(a) Give characteristics of study participants (eg demographic, clinical, social) and information on exposures and potential confounders	Page 7 / Line 186-207	Results section
		(b) Indicate number of participants with missing data for each variable of interest	Page 7 / Line 186-207	Results section
		(c) ***Cohort study***—Summarise follow-up time (eg, average and total amount)	Page 7 / Line 181-184	Results section
Outcome data	15*	***Cohort study***—Report numbers of outcome events or summary measures over time	Page 7 / Line 176-207	Results section
		Case-control study—Report numbers in each exposure category, or summary measures of exposure	NA (see above)	NA
		Cross-sectional study—Report numbers of outcome events or summary measures	NA (See above)	NA
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 7 / Line 176-207	Results section
		(b) Report category boundaries when continuous variables were categorized	Page 7 / Line 176-207	Results section
		(c) If relevant, consider translating estimates of relative risk into absolute risk for a meaningful time period	NA (none done)	NA
Other analyses	17	Report other analyses done—eg analyses of subgroups and interactions, and sensitivity analyses	NA (none done)	NA
Discussion				
Key results	18	Summarise key results with reference to study objectives	Page 8 / Line 209-270	Comments section
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 10 / Line 271-287	Comments section

Interpretation	20	Give a cautious overall interpretation of results considering objectives, limitations, multiplicity of analyses, results from similar studies, and other relevant evidence	Page 11 / Line 289-294	Conclusions section
Generalisability	21	Discuss the generalisability (external validity) of the study results	Page 11 / Line 289-294	Conclusions section
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	Page 1 / Line 12	Title page

*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.