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 61	Abstract, Background
		(b) Provide in the abstract an informative and balanced summary of what was done and what was found	page 3, line 77	Abstract, Summary
Introduction				
Background/ rationale	2	Explain the scientific background and rationale for the investigation being reported	page 5, line 93	Introduction, paragraph 1
Objectives	3	State specific objectives, including any prespecified hypotheses	page 5, line 125	Indroduction, paragraph 3
Methods				
Study design	4	Present key elements of study design early in the paper	page 6, line 135	methods, patient
Setting	5	Describe the setting, locations, and relevant dates, including periods of recruitment, exposure, follow-up, and data collection	page 6, line 136	methods, patient enrollment
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 135	methods, patient enrollment
		(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	NA, not a matched study	NA, not a matched study
Variables	7	Clearly define all outcomes, exposures, predictors, potential confounders, and effect modifiers. Give diagnostic criteria, if applicable	page 6, line 162	methods, Polymerase Chain Reaction
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 152	methods, data collection and calculation
Bias	9	Describe any efforts to address potential sources of bias	page 6, line 138	method, patient enrollment
Study size	10	Explain how the study size was arrived at	page 6, line 135	methods, patient
Quantitative variables	11	Explain how quantitative variables were handled in the analyses. If applicable, describe which groupings were chosen and why	page 6, line 153	methods, data collection and calculation

Statistical methods	12	(a) Describe all statistical methods, including those used to control for confounding	page 7, line 170	methods, stastics analysis
		(b) Describe any methods used to examine subgroups and interactions	page 7, line 171	methods, stastics analysis
		(c) Explain how missing data were addressed	NA, no missing data	NA, no missing data
		(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	NA, no loss of follow-up	NA, no loss of follow-up
		(e) Describe any sensitivity analyses	NA, not applicable	NA, not applicable
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	NA, not suitable for the study cohort and design	NA, not suitable for the study cohort and design
		(b) Give reasons for non-participation at each stage	NA, not suitable	NA, not suitable
		(c) Consider use of a flow diagram	NA, not suitable	NA, not suitable
Descriptive data	14*	(a) Give characteristics of study participants (eg demographic, clinical, social) and information on exposures and potential confounders	page 7, line 183	result, paragraph 1
		(b) Indicate number of participants with missing data for each variable of interest	NA, no missing data	NA, no missing data
		(c) ***Cohort study***—Summarise follow-up time (eg, average and total amount)	NA, not suitable	NA, not suitable
Outcome data	15*	***Cohort study***—Report numbers of outcome events or summary measures over time	page 7, line 187	result, complete blood
		Case-control study—Report numbers in each exposure category, or summary measures of exposure	NA, not suitable	NA, not suitable
		Cross-sectional study—Report numbers of outcome events or summary measures	NA, not suitable	NA, not suitable
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 187	result, Complete blood count profile
		(b) Report category boundaries when continuous variables were categorized	NA, not applicable	NA, not applicable
		(c) If relevant, consider translating estimates of relative risk into absolute risk for a meaningful time period	NA, not applicable	NA, not applicable
Other analyses	17	Report other analyses done—eg analyses of subgroups and interactions, and sensitivity analyses	NA, not applicable	NA, not applicable
Discussion				
Key results	18	Summarise key results with reference to study objectives	page 8, line 203	discussion, paragraph 1
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 8, line 229	discussion, paragraph 4

Interpretation	20	Give a cautious overall interpretation of results considering objectives, limitations, multiplicity of analyses, results from similar studies, and other relevant evidence	NA, no similar study	NA, no similar study
Generalisability	21	Discuss the generalisability (external validity) of the study results	NA, no similar study	NA, no similar study
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 2, line 34	acknowledgment, 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.