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 1, Line 3	Title
		(b) Provide in the abstract an informative and balanced summary of what was done and what was found	Page 2, Lines 13-22	Abstract, Paragraphs 2-3
Introduction				
Background/ rationale	2	Explain the scientific background and rationale for the investigation being reported	Page 3, Lines 31-57	Introduction, Paragraphs 1-2
Objectives	3	State specific objectives, including any prespecified hypotheses	Page 4, Lines 70-71; Page	Introduction, Paragraphs
Methods				
Study design	4	Present key elements of study design early in the paper	Pages 5-6, Lines 115-136	Methods, Section 2.2 and
Setting	5	Describe the setting, locations, and relevant dates, including periods of recruitment, exposure, follow-up, and data collection	Pages 4-5, Lines 84-89; Page 5, Lines 109-115	Methods, Section 2.1, Paragraph 1, and 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 5, Lines 90-99	Methods, Section 2.1, Paragraph 2
		(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	Page 5, Lines 139-140	Results, Section 3.1
Variables	7	Clearly define all outcomes, exposures, predictors, potential confounders, and effect modifiers. Give diagnostic criteria, if applicable	Page 5, Lines 100-107	Methods, Section 2.1, 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	Pages 5-6, Lines 109-136; Page 6, Lines 139-140	Methods, Section 2.2 and Section 2.3; Results,
Bias	9	Describe any efforts to address potential sources of bias	Page 4-5, Lines 84-107;	Methods, Section 2.1,
Study size	10	Explain how the study size was arrived at	Page 4, Lines 70-71; Page	Introduction, Paragraphs
Quantitative variables	11	Explain how quantitative variables were handled in the analyses. If applicable, describe which groupings were chosen and why	Page 6, Lines 125-136	Methods, Section 2.3

Statistical methods	12	(a) Describe all statistical methods, including those used to control for confounding	Page 6, Lines 125-136	Methods, Section 2.3
		(b) Describe any methods used to examine subgroups and interactions	N/A (The study did not	N/A (The study did not
		(c) Explain how missing data were addressed	Page 6, Lines 125-127	Methods, Section 2.3
		(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	N/A (This study employs no specific sampling strategy due to the comprehensive nature of	N/A (This study employs no specific sampling strategy due to the comprehensive nature of
		(e) Describe any sensitivity analyses	N/A (This study did not	N/A (This study did not
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 6, Lines 139-140	Results, Section 3.1
		(b) Give reasons for non-participation at each stage	Page 6-7. Lines 141-145	Results. Section 3.1
		(c) Consider use of a flow diagram	N/A (Based on the study's	N/A (Based on the study's
Descriptive data	14*	(a) Give characteristics of study participants (eg demographic, clinical, social) and information on exposures and potential confounders	Page 6-7. Lines 141-145	Results. Section 3.1
		(b) Indicate number of participants with missing data for each variable of interest	Page 6-7, Lines 141-145	Results. Section 3.1
		(c) ***Cohort study***—Summarise follow-up time (eg, average and total amount)	N/A (This study is not a	N/A (This study is not a
Outcome data	15*	***Cohort study***—Report numbers of outcome events or summary measures over time	N/A (This study is not a	N/A (This study is not a
		Case-control study—Report numbers in each exposure category, or summary measures of exposure	N/A (This study is not a	N/A (This study is not a
		Cross-sectional study—Report numbers of outcome events or summary measures	Page 7, Lines 146-167	Results, Section 3.2,
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 148, Lines 151-152, Lines 158-159,	Results, Section 3.2, Section 3.3, and Section
		(b) Report category boundaries when continuous variables were categorized	N/A (This study requires	N/A (This study requires
		(c) If relevant, consider translating estimates of relative risk into absolute risk for a meaningful time period	N/A (Not relavant to this	N/A (Not relavant to this
Other analyses	17	Report other analyses done—eg analyses of subgroups and interactions, and sensitivity analyses	N/A (This study involves	N/A (This study involves
Discussion				
Key results	18	Summarise key results with reference to study objectives	Page 7, Lines 169-182	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 11, Lines 256-267	Discussion, Section 4.4

Interpretation	20	Give a cautious overall interpretation of results considering objectives, limitations, multiplicity of analyses, results from similar studies, and other relevant evidence	Page 8, Lines 184-214	Discussion, Section 4.1
Generalisability	21	Discuss the generalisability (external validity) of the study results	N/A (This study involves	N/A (This study involves
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 13, Lines 291-294	Acknowledgements

*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.

Article information: https://dx.doi.org/10.21037/qims-24-753
*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.