STROBE Statement—Checklist of items that should be included in reports of *cohort studies*

	Item No	Recommendation
Title and abstract	1	(a) Indicate the study's design with a commonly used term in the title or the abstract
		PAGES 1 AND 2
		(b) Provide in the abstract an informative and balanced summary of what was done
		and what was found
		PAGE 2
Introduction		
Background/rationale	2	Explain the scientific background and rationale for the investigation being reported PAGES 3 AND 4
Objectives	3	State specific objectives, including any prespecified hypotheses PAGE 4
Methods		
Study design	4	Present key elements of study design early in the paper PAGE 4
Setting	5	Describe the setting, locations, and relevant dates, including periods of recruitment,
		exposure, follow-up, and data collection
		PAGES 4 AND 5
Participants	6	(a) Give the eligibility criteria, and the sources and methods of selection of
		participants. Describe methods of follow-up
		PAGES 4 AND 5. FIGURE 3
		(b) For matched studies, give matching criteria and number of exposed and
		unexposed
		NOT APPLICABLE
Variables	7	Clearly define all outcomes, exposures, predictors, potential confounders, and effect
		modifiers. Give diagnostic criteria, if applicable
		PAGE 6
Data sources/	8*	For each variable of interest, give sources of data and details of methods of
measurement		assessment (measurement). Describe comparability of assessment methods if there is
		more than one group
		PAGES 6 AND 7
Bias	9	Describe any efforts to address potential sources of bias
		PAGE 11
Study size	10	Explain how the study size was arrived at
		PAGE 6
Quantitative variables	11	Explain how quantitative variables were handled in the analyses. If applicable,
		describe which groupings were chosen and why
		PAGES 6 AND 7
Statistical methods	12	(a) Describe all statistical methods, including those used to control for confounding
		PAGES 6 AND 7
		(b) Describe any methods used to examine subgroups and interactions
		NOT APPLICABLE
		(c) Explain how missing data were addressed
		NOT APPLICABLE
		(d) If applicable, explain how loss to follow-up was addressed
		NOT APPLICABLE
		1

(\underline{e}) Describe any sensitivity analyses NOT APPLICABLE

		NOT AFFLICABLE
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, FIGURE 3
		(b) Give reasons for non-participation at each stage
		PAGE 7, FIGURE #
		(c) Consider use of a flow diagram
		FIGURE 3
Descriptive data	14*	(a) Give characteristics of study participants (eg demographic, clinical, social) and
		information on exposures and potential confounders TABLE 1
		(b) Indicate number of participants with missing data for each variable of interest NOT APPLICABLE
		(c) Summarise follow-up time (eg, average and total amount)
		FIGURE 4
Outcome data	15*	Report numbers of outcome events or summary measures over time
		TABLES 3 AND 4
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
		PAGES 7 AND 8, TABLES 2,3, AND 4
		(b) Report category boundaries when continuous variables were categorized
		TABLE 2
		(c) If relevant, consider translating estimates of relative risk into absolute risk for a
		meaningful time period
		NOT APPLICABLE
Other analyses	17	Report other analyses done—eg analyses of subgroups and interactions, and sensitivity analyses
		NOT APPLICABLE
Discussion		
Key results	18	Summarise key results with reference to study objectives PAGE 8
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
		PAGES 10 AND 11
Interpretation	20	Give a cautious overall interpretation of results considering objectives, limitations,
		multiplicity of analyses, results from similar studies, and other relevant evidence
		PAGES 8, 9 AND 10
Generalisability	21	Discuss the generalisability (external validity) of the study results
		PAGE 11
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 12