STROBE Statement—Checklist of items that should be included in reports of *cross-sectional studies* 

	Item		Reported
	No	Recommendation	
Title and abstract	1	(a) Indicate the study's design with a commonly used term in the title	Abstract
		or the abstract	
		(b) Provide in the abstract an informative and balanced summary of	Abstract
		what was done and what was found	
Introduction			
Background/rationale	2	Explain the scientific background and rationale for the investigation	Introduction
C		being reported	
Objectives	3	State specific objectives, including any prespecified hypotheses	Introduction
Methods			
Study design	4	Present key elements of study design early in the paper	Methods
Setting	5	Describe the setting, locations, and relevant dates, including periods of	Methods
C		recruitment, exposure, follow-up, and data collection	
Participants	6	(a) Give the eligibility criteria, and the sources and methods of	Methods
		selection of participants	
Variables	7	Clearly define all outcomes, exposures, predictors, potential	Methods
		confounders, and effect modifiers. Give diagnostic criteria, if applicable	
Data sources/	8*	For each variable of interest, give sources of data and details of	Methods
measurement		methods of assessment (measurement). Describe comparability of	
		assessment methods if there is more than one group	
Bias	9	Describe any efforts to address potential sources of bias	Not
2140		2 socios uni, criorio to accreso potentian socios er ente	applicable
Study size	10	Explain how the study size was arrived at	Methods
Quantitative	11	Explain how quantitative variables were handled in the analyses. If	Methods
variables		applicable, describe which groupings were chosen and why	
Statistical methods	12	(a) Describe all statistical methods, including those used to control for	Methods
		confounding	
		(b) Describe any methods used to examine subgroups and interactions	Not
			applicable
		(c) Explain how missing data were addressed	Methods
		(d) If applicable, describe analytical methods taking account of	Not
		sampling strategy	applicable
		(e) Describe any sensitivity analyses	Not
		(a) Describe any sensativity analyses	applicable
Results		<u>I</u>	··FF
Participants	13*	(a) Report numbers of individuals at each stage of study—eg numbers	Results
	13	potentially eligible, examined for eligibility, confirmed eligible,	Results
		included in the study, completing follow-up, and analysed	
		(b) Give reasons for non-participation at each stage	Not
		(b) Give reasons for non-participation at each stage	applicable
		(c) Consider use of a flow diagram	Not
		(c) Consider use of a now diagram	applicable
Descriptive data	14*	(a) Give characteristics of study participants (eg demographic, clinical,	Results
	14"	social) and information on exposures and potential confounders	Nesuits
			Not
		(b) Indicate number of participants with missing data for each variable	Not

		of interest	applicable
Outcome data	15*	Report numbers of outcome events or summary measures	Results
Main results	16	(a) Give unadjusted estimates and, if applicable, confounder-adjusted	Results
		estimates and their precision (eg, 95% confidence interval). Make clear	tables
		which confounders were adjusted for and why they were included	
		(b) Report category boundaries when continuous variables were	Not
		categorized	applicable
		(c) If relevant, consider translating estimates of relative risk into	Not
		absolute risk for a meaningful time period	applicable
Other analyses	17	Report other analyses done—eg analyses of subgroups and interactions,	Not
		and sensitivity analyses	applicable
Discussion			
Key results	18	Summarise key results with reference to study objectives	Interpretation
Limitations	19	Discuss limitations of the study, taking into account sources of	Limitations
		potential bias or imprecision. Discuss both direction and magnitude of	
		any potential bias	
Interpretation	20	Give a cautious overall interpretation of results considering objectives,	Interpretation
		limitations, multiplicity of analyses, results from similar studies, and	
		other relevant evidence	
Generalisability	21	Discuss the generalisability (external validity) of the study results	Limitations
Other information			
Funding	22	Give the source of funding and the role of the funders for the present	Funding
		study and, if applicable, for the original study on which the present	declaration
		article is based	statement
			and
			Introduction