

Evidence of SBA through People Centered Care - STROBE checklist

	Item No	Recommendation	Status
Title and abstract	1	(a) Indicate the study's design with a commonly used term in the title or the abstract	Completed
		(b) Provide in the abstract an informative and balanced summary of what was done and what was found	Completed
Introduction			
Background/rationale	2	Explain the scientific background and rationale for the investigation being reported	Completed
Objectives	3	State specific objectives, including any prespecified hypotheses	Completed
Methods			
Study design	4	Present key elements of study design early in the paper	Completed
Setting	5	Describe the setting, locations, and relevant dates, including periods of recruitment, exposure, follow-up, and data collection	Completed
Participants	6	<i>Cross-sectional study</i> —Give the eligibility criteria, and the sources and methods of selection of participants	Completed
Variables	7	Clearly define all outcomes, exposures, predictors, potential confounders, and effect modifiers. Give diagnostic criteria, if applicable	Completed
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	Completed
Bias	9	Describe any efforts to address potential sources of bias	Completed
Study size	10	Explain how the study size was arrived at	Completed
Quantitative variables	11	Explain how quantitative variables were handled in the analyses. If applicable, describe which groupings were chosen and why	Completed
Statistical methods	12	(a) Describe all statistical methods, including those used to control for confounding	Completed
		(b) Describe any methods used to examine subgroups and interactions	Completed
		(c) Explain how missing data were addressed	Completed
		<i>Cross-sectional study</i> —If applicable, describe analytical methods taking account of sampling strategy	Completed
		(e) Describe any sensitivity analyses	

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Results		Status
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
		(b) Give reasons for non-participation at each stage
		(c) Consider use of a flow diagram
Descriptive data	14*	(a) Give characteristics of study participants (eg demographic, clinical, social) and information on exposures and potential confounders
		<i>Cross-sectional study</i> —Report numbers of outcome events or summary measures
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
		(b) Report category boundaries when continuous variables were categorized
		(c) If relevant, consider translating estimates of relative risk into absolute risk for a meaningful time period
Other analyses	17	Report other analyses done—eg analyses of subgroups and interactions, and sensitivity analyses
Discussion		
Key results	18	Summarise key results with reference to study objectives
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
Interpretation	20	Give a cautious overall interpretation of results considering objectives, limitations, multiplicity of analyses, results from similar studies, and other relevant evidence
Generalisability	21	Discuss the generalisability (external validity) of the study results
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