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	PN: 1/LN: 8-9	Abstract/2nd paragraph
		(b) Provide in the abstract an informative and balanced summary of what was done and what was found	PN: 1/LN: 1-22	Abstract/1-4 paragraphs
Introduction				
Background/ rationale	2	Explain the scientific background and rationale for the investigation being reported	PN: 1-2/LN: 26-34	Introduction/1-3 paragraphs
Objectives	3	State specific objectives, including any prespecified hypotheses	PN: 2/LN: 35-44	Introduction/3-5
Methods				
Study design	4	Present key elements of study design early in the paper	PN: 3/LN: 58-61	Patients/3rd paragraph
Setting	5	Describe the setting, locations, and relevant dates, including periods of recruitment, exposure, follow-up, and data collection	PN: 2-3/LN: 46-51, 58-61	Patients/1,3 paragraphs
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	PN: 2-3/LN: 45-68	Patients/1-6 paragraphs
		(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	N/A, the study does not contain matched groups	N/A, the study does not contain matched groups
Variables	7	Clearly define all outcomes, exposures, predictors, potential confounders, and effect modifiers. Give diagnostic criteria, if applicable	PN: 3-6/LN: 76-165	Surgical technique /1-22 paragraphs
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	PN: 3-6/LN: 76-165	Surgical technique /1-22 paragraphs
Bias	9	Describe any efforts to address potential sources of bias	N/A, the article describes	N/A, the article describes
Study size	10	Explain how the study size was arrived at	PN: 2-3/LN: 46-61	Patients/1-3 paragraphs
Quantitative variables	11	Explain how quantitative variables were handled in the analyses. If applicable, describe which groupings were chosen and why	PN: 3/LN: 73-75	Patients/8th paragraph

Statistical methods	12	(a) Describe all statistical methods, including those used to control for confounding	PN: 3/LN: 58-61, 73-75	Patients/3, 8 paragraphs
		(b) Describe any methods used to examine subgroups and interactions	N/A, the study does not	N/A, the study does not
		(c) Explain how missing data were addressed	N/A, the study does not	N/A, the study does not
		(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, study contain only early postoperative follow-up	N/A, study contains only early postoperative follow-up
		(e) Describe any sensitivity analyses	N/A, the study does not	N/A, the study does 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	PN: 2-3/LN: 46-61 PN: 6/LN: 167-168	Patients/1-3 paragraphs Results/1st paragraph
		(b) Give reasons for non-participation at each stage	PN: 2-3/LN: 46-61	Patients/1-3 paragraphs
		(c) Consider use of a flow diagram	N/A, the article does not	N/A, the article does not
Descriptive data	14*	(a) Give characteristics of study participants (eg demographic, clinical, social) and information on exposures and potential confounders	PN: 3/LN: 62	Patients/4th paragraph
		(b) Indicate number of participants with missing data for each variable of interest	N/A, the study does not	N/A, the study does not
		(c) ***Cohort study***—Summarise follow-up time (eg, average and total amount)	PN: 3/LN: 60-61	Patients/3th paragraph
Outcome data	15*	***Cohort study***—Report numbers of outcome events or summary measures over time	PN: 6/LN: 167-168	Results/1st paragraph
		Case-control study—Report numbers in each exposure category, or summary measures of exposure	N/A, Cohort study	N/A, Cohort study
		Cross-sectional study—Report numbers of outcome events or summary measures	N/A, Cohort study	N/A, Cohort study
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	PN: 6-7/LN: 169-186	Results/2-4 paragraphs
		(b) Report category boundaries when continuous variables were categorized	N/A, the study does not	N/A, the study does not
		(c) If relevant, consider translating estimates of relative risk into absolute risk for a meaningful time period	N/A, the study does not	N/A, the study does not
Other analyses	17	Report other analyses done—eg analyses of subgroups and interactions, and sensitivity analyses	N/A, the study does not	N/A, the study does not
Discussion				
Key results	18	Summarise key results with reference to study objectives	PN: 9/LN: 249-257	Conclusions/1-2
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	PN: 3/LN: 63-68 PN: 9/LN: 245-248	Patients/5th paragraph Discussion /14th

Interpretation	20	Give a cautious overall interpretation of results considering objectives, limitations, multiplicity of analyses, results from similar studies, and other relevant evidence	PN: 7-9/LN: 187-248	Discussion /1-14 paragraphs
Generalisability	21	Discuss the generalisability (external validity) of the study results	PN: 8/LN: 205-214	Discussion /4-6
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	PN: 12/LN: 316	Conflicts of interest/1st paragraph

*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/jovs-22-2
*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.