| 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 3/Line 32-35                  | Abstract/Paragraph 2         |
|                  |         | (b) Provide in the abstract an informative and balanced summary of what was done and what was found | Page 3-4/Line 36-50                | Abstract/Paragraph 3-4       |
| Introduction     |         |                                                                                  |                                     |                              |
| Background/rationale | 2       | Explain the scientific background and rationale for the investigation being reported | Page 5/Line 55-73                  | Introduction/Paragraph 1     |
| Objectives       | 3       | State specific objectives, including any prespecified hypotheses                  | Page 5-6/Line 74-76                | Introduction/Paragraph 2     |
| Methods          |         |                                                                                  |                                     |                              |
| Study design     | 4       | Present key elements of study design early in the paper                           | Page 7/Line 79-86                  | Methods/Paragraph 1          |
| Setting          | 5       | Describe the setting, locations, and relevant dates, including periods of recruitment, exposure, follow-up, and data collection | Page 7-8/Line 79-99                | Methods/Paragraph 1-2        |
| 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 7-8/Line 79-99                | Methods/Paragraph 1-2        |
|                  |         | (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 |                              |                              |
| Variables        | 7       | Clearly define all outcomes, exposures, predictors, potential confounders, and effect modifiers. Give diagnostic criteria, if applicable | Page 7/Line 79-86                  | Methods/Paragraph 1          |
| 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 | Page 7/Line 79-86                  | Methods/Paragraph 1          |
| Bias             | 9       | Describe any efforts to address potential sources of bias                          | Page 7/Line 79-86                  | Methods/Paragraph 1          |
| Study size       | 10      | Explain how the study size was arrived at                                         | Page 7/Line 79-86                  | Methods/Paragraph 1          |
| Quantitative variables | 11     | Explain how quantitative variables were handled in the analyses. If applicable, describe which groupings were chosen and why | Page 7/Line 79-86                  | Methods/Paragraph 1          |
| Statistical methods | 12 | (a) Describe all statistical methods, including those used to control for confounding | Page 8/Line 101-105 | Methods/Paragraph 3 |
|---------------------|----|---------------------------------------------------------------------------------|---------------------|---------------------|
|                     |    | (b) Describe any methods used to examine subgroups and interactions               | Page 8/Line 101-105 | Methods/Paragraph 3 |
|                     |    | (c) Explain how missing data were addressed                                       | NA                  | Not performed       |
|                     |    | (d) **Cohort study**—If applicable, explain how loss to follow-up was addressed   | Page 8/Line 101-105 | Methods/Paragraph 3 |
|                     |    | **Case-control study**—If applicable, explain how matching of cases and controls was addressed | NA                  | Not performed       |
|                     |    | **Cross-sectional study**—If applicable, describe analytical methods taking account of sampling strategy | NA                  | Not performed       |
|                     |    | (e) Describe any sensitivity analyses                                             | NA                  | Not performed       |
| 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 9/Line 108-111 | Results/Paragraph 1 |
|                     |    | (b) Give reasons for non-participation at each stage                               | NA                  | No non-participation |
|                     |    | (c) Consider use of a flow diagram                                                | NA                  | Not used            |
| Descriptive data    | 14*| (a) Give characteristics of study participants (eg demographic, clinical, social) and information on exposures and potential confounders | Page 9/Line 108-111 | Results/Paragraph 1 |
|                     |    | (b) Indicate number of participants with missing data for each variable of interest | NA                  | No missing data     |
|                     |    | (c) **Cohort study**—Summarise follow-up time (eg, average and total amount)     | Page 11/Line 150-151 | Results/Paragraph 3 |
| Outcome data        | 15*| **Cohort study**—Report numbers of outcome events or summary measures over time    | Page 9-11/Line 112-151 | Results/Paragraph 2-3 |
|                     |    | **Case-control study**—Report numbers in each exposure category, or summary measures of exposure | NA                  | Not applicable to this |
|                     |    | **Cross-sectional study**—Report numbers of outcome events or summary measures     | NA                  | Not applicable to this |
| 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 9-11/Line 108-151 | Results/Paragraph 1-3 |
|                     |    | (b) Report category boundaries when continuous variables were categorized         | Table 1-3           | Table 1-3           |
|                     |    | (c) If relevant, consider translating estimates of relative risk into absolute risk for a meaningful time period | NA                  | Not applicable to this |
| Other analyses      | 17 | Report other analyses done—eg analyses of subgroups and interactions, and sensitivity analyses | Page 10-11/Line 161-199 | Results/Paragraph 3-5 |
| Discussion          |    |                                                                                  |                     |                     |
| Key results         | 18 | Summarise key results with reference to study objectives                          | Page 12-13/Line 154-178 | Discussion/Paragraph 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 | NA                  | Not available       |
| Interpretation                        | 20 | Give a cautious overall interpretation of results considering objectives, limitations, multiplicity of analyses, results from similar studies, and other relevant evidence | Page 13-14/Line 179-200 | Discussion/Paragraph 3 |
|--------------------------------------|----|---------------------------------------------------------------------------------------------------------------------------------|--------------------------|------------------------|
| Generalisability                    | 21 | Discuss the generalisability (external validity) of the study results                                                        | Page 13-14/Line 179-211 | Discussion/Paragraph 3-4 |
| **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 | NA                      | No source of funding   |

*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.

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*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.*