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Appendix

Authors: These minimum items of information are needed by our referees and Editors to evaluate your manuscript. Additional information may be appropriate, depending on your study design and objectives.

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http://www.consort-statement.org (for clinical trials; there are elaborations for abstracts, cluster designs, reporting of harms, herbal interventions, non-inferiority and equivalence studies, trials of non-pharmacologic interventions, and pragmatic trials)


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   b. Descriptions of how blindness was accomplished for all subjective evaluations

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   e. Distinction between assumptions, input data, calculations from intermediate steps in the modeling process, and model predictions
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   g. Sensitivity analyses of key assumptions and of the input variables that had the greatest uncertainty
   h. Descriptions of the variability in the "outputs" from stochastic models

5. For statistical-hypothesis tests:
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   b. Alpha and tails, and any methods used to adjust for multiple comparisons (to protect experiment-wise alpha from the problem of multiplicity)
   c. Methods used to adjust for clustering within the data
   d. Methods used to determine that the statistical assumptions were met (e.g., that the data were Gaussian or that the odds ratio or hazards ratio was constant across the observed range of the risk factor)
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