

Section/Topic	Peters et al. [17]	Brunoni et al. [18]	Aldoghachi et al. [19]	Froud et al. [20]	Ai et al. [21]	Kitzkerová et al. [22]	Bassi et al. [24]	Youssef et al. [15]	Caldieraro et al. [25]	Wang et al. [26]	Han et al. [27]	Su et al. [10]	Tatham et al. [28]	Tatham et al. [16]	Jaworska et al. [29]	Kostic et al. [30]	Cao et al. [31]
Results																	
Participants	14) Report the numbers of individuals at each stage of the study. Give reasons for nonparticipation at each stage. Report the number of participants not genotyped, and reasons why they were not genotyped.																
Descriptives: Population	15) Report demographic and clinical characteristics of the study population, including risk factors used in the risk modeling.																
Descriptives: Model estimates	16) Report unadjusted associations between the variables in the risk model(s) and the outcome. Report adjusted estimates and their precision from the full risk model(s) for each variable.																
Risk distributions	17) Descriptives: Model estimates																
Assessment	18) Report measures of model fit and predictive ability, and any other performance measures, if pertinent.																
Validation	19) Report any validation of the risk model(s).																
Other analyses	20) Present results of any subgroup, interaction, or exploratory analyses, whenever pertinent.																
Discussion																	
Limitations	21) Discuss limitations and assumptions of the study, particularly those concerning study design, selection of participants, and measurements and analyses, and discuss their impact on the results of the study.																
Interpretation	22) Give an overall interpretation of results considering objectives, limitations, multiplicity of analyses, results from similar studies, and other relevant evidence.																
Generalizability	23) Discuss the generalizability and, if pertinent, the health care relevance of the study results.																

X = Present.