

TRIPOD Checklist: Prediction Model Development

Section	Item	Checklist description	Reported on Page Number/Line Number	Reported on Section/Paragraph
Title and abstract				
Title	1	Identify the study as developing and/or validating a multivariable prediction model, the target population, and the outcome to be predicted.	Page1 /Line3-4	title/Paragraph2
Abstract	2	Provide a summary of objectives, study design, setting, participants, sample size, predictors, outcome, statistical analysis, results, and conclusions.	Page2 /Line35-62	Abstract/p13-p17
Introduction				
Background and objectives	3a	Explain the medical context (including whether diagnostic or prognostic) and rationale for developing or validating the multivariable prediction model, including references to existing models.	Page3-4 /Line70-109	INtroduction /P19-P23
	3b	Specify the objectives, including whether the study describes the development or validation of the model or both.	Page4 /Line110-130	INtroduction /P24
Methods				
Source of data	4a	Describe the study design or source of data (e.g., randomized trial, cohort, or registry data), separately for the development and validation data sets, if applicable.	Page4-5 /Line131-145	method/P25-P27
	4b	Specify the key study dates, including start of accrual; end of accrual; and, if applicable, end of follow-up.	Page5 /Line147-172	method/P28-P31
Participants	5a	Specify key elements of the study setting (e.g., primary care, secondary care, general population) including number and location of centres.	page5 /Line136-139	method/P27
	5b	Describe eligibility criteria for participants.	page5 /Line139-145	method/P25
	5c	Give details of treatments received, if relevant.	na	na
Outcome	6a	Clearly define the outcome that is predicted by the prediction model, including how and when assessed.	page6 /Line147-202	method/P33-p45
	6b	Report any actions to blind assessment of the outcome to be predicted.	na	na
Predictors	7a	Clearly define all predictors used in developing or validating the multivariable prediction model, including how and when they were measured.	page6 /Line176-202	method/P33-p53
	7b	Report any actions to blind assessment of predictors for the outcome and other predictors.	na	na
Sample size	8	Explain how the study size was arrived at.	page5 /Line139-145	method/P25

Missing data	9	Describe how missing data were handled (e.g., complete-case analysis, single imputation, multiple imputation) with details of any imputation method.	page5 /Line140-141	method/P27
Statistical analysis methods	10a	Describe how predictors were handled in the analyses.	page5 /Line176-187	method/P33
	10b	Specify type of model, all model-building procedures (including any predictor selection), and method for internal validation.	page5 /Line176-187	method/P33
	10d	Specify all measures used to assess model performance and, if relevant, to compare multiple models.	page5 /Line176-187	method/P33
Risk groups	11	Provide details on how risk groups were created, if done.	NA	NA
Results				
Participants	13a	Describe the flow of participants through the study, including the number of participants with and without the outcome and, if applicable, a summary of the follow-up time. A diagram may be helpful.	page5 /Line207-216	result/P38
	13b	Describe the characteristics of the participants (basic demographics, clinical features, available predictors), including the number of participants with missing data for predictors and outcome.	page5 /Line207-216	result/P38
Model development	14a	Specify the number of participants and outcome events in each analysis.	page8-9 /Line207-216	result/P46-47
	14b	If done, report the unadjusted association between each candidate predictor and outcome.	NA	NA
Model specification	15a	Present the full prediction model to allow predictions for individuals (i.e., all regression coefficients, and model intercept or baseline survival at a given time point).	page8 /Line246-253	result/P45
	15b	Explain how to use the prediction model.	page8-9 /Line286-292	result/P49
Model performance	16	Report performance measures (with CIs) for the prediction model.	page9 /Line279-281	result/P49
Discussion				
Limitations	18	Discuss any limitations of the study (such as nonrepresentative sample, few events per predictor, missing data).	page10-11 /Line367-374	Discussion/P54
Interpretation	19b	Give an overall interpretation of the results, considering objectives, limitations, and results from similar studies, and other relevant evidence.	page10-11 /Line367-374	Discussion/P54
Implications	20	Discuss the potential clinical use of the model and implications for future research.	page11 /Line372-374	Discussion/P54
Other information				
Supplementary information	21	Provide information about the availability of supplementary resources, such as study protocol, Web calculator, and data sets.	table s1/line232	Result/P40
Funding	22	Give the source of funding and the role of the funders for the present study.	NA	NA

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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.