TRIPOD Checklist: Prediction Model Validation

	1	Checklist Item	Page
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.	1
Abstract	2	Provide a summary of objectives, study design, setting, participants, sample size, predictors, outcome, statistical analysis, results, and conclusions.	2
Introduction		outcome, statistical analysis, results, and conclusions.	
Introduction		Explain the medical context (including whether diagnostic or prognostic) and rationale for	
Background and objectives	3a	developing or validating the multivariable prediction model, including references to existing models.	3, 4
	3b	Specify the objectives, including whether the study describes the development or validation of the model or both.	4
Methods		the model of both.	
Methous		Describe the study design or source of data (e.g., randomized trial, cohort, or registry data),	1
Source of data	4a	separately for the development and validation data sets, if applicable.	5
	4b	Specify the key study dates, including start of accrual; end of accrual; and, if applicable, end of follow-up.	5
Participants	5a	Specify key elements of the study setting (e.g., primary care, secondary care, general population) including number and location of centres.	5
	5b	Describe eligibility criteria for participants.	5
	5c	Give details of treatments received, if relevant.	NA
Outcome	6a	Clearly define the outcome that is predicted by the prediction model, including how and when assessed.	6, 7
	6b	Report any actions to blind assessment of the outcome to be predicted.	NA
	00	Clearly define all predictors used in developing or validating the multivariable prediction	IVA
	7a	model, including how and when they were measured.	5, 6
	7b	Report any actions to blind assessment of predictors for the outcome and other predictors.	NA
Sample size	8	Explain how the study size was arrived at.	5
Sample Size	0	Describe how missing data were handled (e.g., complete-case analysis, single imputation,	3
Missing data	9	multiple imputation) with details of any imputation method.	6, 7
Statistical analysis methods	10c	For validation, describe how the predictions were calculated.	10
	10d	Specify all measures used to assess model performance and, if relevant, to compare multiple models.	10
	10e	Describe any model updating (e.g., recalibration) arising from the validation, if done.	NA
Risk groups	11	Provide details on how risk groups were created, if done.	NA
Development vs. validation	12	For validation, identify any differences from the development data in setting, eligibility criteria, outcome, and predictors.	19, 22
Results		1 / / 1	
Participants		Describe the flow of participants through the study, including the number of participants with	
	13a	and without the outcome and, if applicable, a summary of the follow-up time. A diagram may be helpful.	8
		Describe the characteristics of the participants (basic demographics, clinical features, available	
	13b	predictors), including the number of participants with missing data for predictors and outcome.	19
	13c	For validation, show a comparison with the development data of the distribution of important	22
Model		variables (demographics, predictors and outcome).	_
performance	16	Report performance measures (with CIs) for the prediction model. If done, report the results from any model updating (i.e., model specification, model	8, 9
Model-updating	17	performance).	NA
Discussion			
Limitations	18	Discuss any limitations of the study (such as nonrepresentative sample, few events per predictor, missing data).	12
Interpretation	19a	For validation, discuss the results with reference to performance in the development data, and any other validation data.	11
	19b	Give an overall interpretation of the results, considering objectives, limitations, results from similar studies, and other relevant evidence.	10 - 12
Implications	20	Discuss the potential clinical use of the model and implications for future research.	12
Other information	120	Disease the potential entire as of the model and implications for future research.	12
Supplementary	21	Provide information about the availability of supplementary resources, such as study protocol,	13
information		Web calculator, and data sets.	10
Funding	22	Give the source of funding and the role of the funders for the present study.	13