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.	Page.1/Line.1-2	Title
Abstract	2	Provide a summary of objectives, study design, setting, participants, sample size, predictors, outcome, statistical analysis, results, and conclusions.	Page.1-2/Line.24-43	Abstract/Paragraph.1-4
Introduction				
Background and objectives	За	Explain the medical context (including whether diagnostic or prognostic) and rationale for developing or validating the multivariable prediction model, including references to existing models.	Page.2-3/Line.47-78	Introduction/Paragraph.1-3
	3b	Specify the objectives, including whether the study describes the development or validation of the model or both.	Page.3/Line.78-80	Introduction/Paragraph.3
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, ifapplicable.	Page.3-4/Line.82-117	Methods/Paragraph.1-5
	4b	Specify the key study dates, including start of accrual; end of accrual; and, if applicable, end of follow-up.	Page.3-4/Line.90-94	Methods/Paragraph.2
Participants	5a	Specify key elements of the study setting (e.g., primary care, secondary care, general population) including number and location of centres.	Page.4-5/Line.118-144	Methods/Paragraph.6-10
	5b	Describe eligibility criteria for participants.	Page.3-4/L.90-97	Methods/Paragraph.2
	5c	Give details of treatments received, if relevant.	N/A	N/A
Outcome	6a	Clearly define the outcome that is predicted by the prediction model, including how and when assessed.	Page.5-6/L.145-174	Methods/Paragraph.11-15
	6b	Report any actions to blind assessment of the outcome to be predicted.	N/A	N/A
Predictors	7a	Clearly define all predictors used in developing or validating the multivariable prediction model, including how and when they were measured.	Page.5-6/L.145-174	Methods/Paragraph.11-15
	7b	Report any actions to blind assessment of predictors for the outcome and other predictors.	N/A	N/A
Sample size	8	Explain how the study size was arrived at.	Page.3-4/L.90-97	Methods/Paragraph.2

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9	Describe how missing data were handled (e.g., complete-case analysis, single imputation, multiple imputation) with details of any imputation method.	N/A	N/A
10a	Describe how predictors were handled in the analyses.	N/A	N/A
10b	Specify type of model, all model-building procedures (including any predictor selection), and method for internal validation.	N/A	N/A
10d	Specify all measures used to assess model performance and, if relevant, to compare multiple models.	Page.5-6/L.145-174	Methods/Paragraph.11-15
11	Provide details on how risk groups were created, if done.	N/A	N/A
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.	N/A	N/A
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.	N/A	N/A
14a	Specify the number of participants and outcome events in each analysis.	Page.6-7/Line.177-198	Results/Paragraph.1
14b	If done, report the unadjusted association between each candidate predictor and outcome.	N/A	N/A
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).	N/A	N/A
15b	Explain how to the use the prediction model.	Page.6-7/Line.177-198	Results/Paragraph.1
16	Report performance measures (with CIs) for the prediction model.	N/A	N/A
18	Discuss any limitations of the study (such as nonrepresentative sample, few events per predictor, missing data).	Page.9/Line.265-269	Discussion/Paragraph.5
19b	Give an overall interpretation of the results, considering objectives, limitations, and results from similar studies, and other relevant evidence.	Page.9-10/Line.271-277	Conclusion
20	Discuss the potential clinical use of the model and implications for future research.	Page.9-10/Line.271-277	Conclusion
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21	Provide information about the availability of supplementary resources, such as study protocol, Web calculator, and data sets.	N/A	N/A
	10a 10b 10d 11 13a 13a 13b 14a 14b 15a 15b 16 16 18	imputation) with details of any imputation method. 10a Describe how predictors were handled in the analyses. 10b Specify type of model, all model-building procedures (including any predictor selection), and method for internal validation. 10d Specify all measures used to assess model performance and, if relevant, to compare multiple models. 11 Provide details on how risk groups were created, if done. 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. 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. 14a Specify the number of participants and outcome events in each analysis. 14b If done, report the unadjusted association between each candidate predictor and outcome. 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). 15b Explain how to the use the prediction model. 18 Discuss any limitations of the study (such as nonrepresentative sample, few events per predictor, missing data). 19b Give an overall interpretation of the results, considering objectives, limitations, and results from similar	Imputation) with details of any imputation method.Imputation method.10aDescribe how predictors were handled in the analyses.N/A10bSpecify type of model, all model-building procedures (including any predictor selection), and method for internal validation.N/A10dSpecify all measures used to assess model performance and, if relevant, to compare multiple models.Page.5-6/L.145-17411Provide details on how risk groups were created, if done.N/A13aDescribe 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.N/A13bDescribe the characteristics of the participants (basic demographics, clinical features, available predictors), including the number of participants with missing data for predictors and outcome.N/A14aSpecify the number of participants and outcome events in each analysis.Page.6-7/Linc.177-19814bIf done, report the unadjusted association between each candidate predictor and outcome.N/A15aPresent 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).Page.6-7/Linc.177-19816Report performance measures (with Cls) for the prediction model.N/AN/AImplemention of the results, considering objectives, limitations, and results from similar ata).16Give an overall interpretation of the results, considering objectives, limitations, and results from similar ata).Page.9-10/Line.271-277

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