Article Information

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Article ID

Article available for data extraction

Endnote number





Background and objectives

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Article ID	
The background and rationale are presented	○ Yes ○ No
Reference to existing models is included (or stated that there are no existing models)	○ Yes ○ No
Any description of why ML techniques are being used to address the objective is reported	○ Yes ○ No
If yes, please provide the statement below	
It is stated whether the study describes development and/or validation and/or incremental (added) value	○ Yes ○ No
Any additional comment about the "background and objectives" section of this article?	
	(If there is something in the "Background" that

does not fit into the questions of this form please use this space to detail. Also use this space to detail anything you are unsure about.)



Reviewer Information

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Article ID

Reviewer Information

Reviewer Name

(Provide your initials)

Date of extraction



06-21-2020 01:50

General Information

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Article ID	
General study information	
Title	
laurnal	
Journal	
Publication year	
Number of authors listed	
Name of the first author	
	(Initials, Surname (e.g. EW Steyerberg))
What is the affiliation of the 1st author?	Clinical Epidemiology
	Health informatics
	Data sciences Other
If other, please specify	
What is the clinical area being investigated?	Oncology
	 Cardiovascular medicine Critical care
	 Healthcare services Geriatric
	 Hepatology
	O Psychiatry
	\bigcirc Neonatology
	O Nutrition
	 Obstetrics & Gynaecology Physical medicine
	O Primary care
	Surgery
	 Neurology
	Ophthalmology
	O Medical imaging
If other please specify	

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uential	Page 2 of 2
What is the type of study?	 Diagnosis Prognosis
What is the type of study?	 Classification Risk prediction Unclear (We need to distinguish if the problem is a prediction problem or a classification problem. A classification problem is about predicting a label and a prediction problems is about predicting a predicting a quantity.)
What is the purpose of the article?	 Clinical use Simulation/tutoring Contest/Challenge (If tutoring, please finish this form and don't follow with the extraction.)
What is the study design?	 Development only (including internal validation Development with external validation (same m Development with external validation (different model) External validation only
What is the primary outcome for the model?	
	(Please include timing of primary outcome. Extrac on primary outcome only.)
What is the format of the primary outcome?	 Continuous Binary Ordinal Multinomial Time to event Count Other
If other, please specify	

(If there is information in "General Information" that does not fit into the questions of this form - please use this space to detail. Also use this space to detail anything you are unsure about.)



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Methods

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Article ID		
Is adherence to a reporting guideline mentioned?	○ Yes ○ No	
If yes, to which guideline?	 ☐ TRIPOD ☐ CONSORT ☐ STROBE ☐ Other 	
If other, please specify		
What type of study is reported?	 Development (including internal validation) Development with external validation (same model) Development with external validation (different model) External validation only (This questions is repeated due to branching logic. Please answer again.) 	
What is the type of external validation?	 Temporal Geographical Independent data Fully independent Unclear Other 	
If other, please specify		
	(E.g. different setting, different participants population (pediatric/adult))	
Differences or similarities in definitions with the development study are described	 Yes No NA (Mentioning of any differences in all four (setting, eligibility criteria, predictors and outcome) is required to score Yes. If it is explicitly mentioned that there were no differences in setting, eligibility criteria, predictors and outcomes, score Yes. For incremental value reports, in case additional predictors are not added to a previously developed prediction model but rather added to conventional predictors in a newly fitted model, score Not applicable.) 	
In which domains are differences?	 Setting Eligibility criteria Predictors Outcomes No differences were reported Other 	

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	Page 2 c
If other, please specify	
	(list using (;) to separate if more than 1.)
Is there a diagram/draw to clarify the methods used?	○ Yes ○ No
Source of data	
Development The study design or source of data is reported	 Yes No (E.g. Prospectively designed, existing cohort, existing RCT, registry/medical records, case control, case series. This needs to be explicitly reported; reference to this information in another article alone is insufficient.)
External validation The study design or source of data is reported	 Yes No (E.g. Prospectively designed, existing cohort, existing RCT, registry/medical records, case control, case series. This needs to be explicitly reported; reference to this information in another article alone is insufficient.)
External validation The study design or source of data is reported	 Yes No (E.g. Prospectively designed, existing cohort, existing RCT, registry/medical records, case control, case series. This needs to be explicitly reported; reference to this information in another article alone is insufficient.)
Development If yes, what was the data source origin?	 RCT Prospective cohort Retrospective cohort Registry Electronic medical records Case-control/case-cohort study Individual patient data - meta analysis Claims Other (Multiples answers are possible)
External validation If yes, what was the data source origin?	 RCT Prospective cohort Retrospective cohort Registry Electronic medical records Case-control/case-cohort study Individual patient data - meta analysis Claims Other (Multiples answers are possible)





	Page 3	
External validation If yes, what was the data source origin?	 RCT Prospective cohort Retrospective cohort Registry Electronic medical records Case-control/case-cohort study Individual patient data - meta analysis Claims Other (Multiples answers are possible) 	
If other, please specify		
If other, please specify		
If other, please specify		
Development The starting date of accrual is reported	○ Yes ○ No	
External validation The starting date of accrual is reported	○ Yes ○ No	
External validation The starting date of accrual is reported	⊖ Yes ⊖ No	
If yes, what is the start data of data collection?		
If yes, what is the start data of data collection?		
If yes, what is the start data of data collection?		
Development The end date of accrual is reported	○ Yes ○ No	
External validation The end date of accrual is reported	○ Yes ○ No	
External validation The end date of accrual is reported	⊖ Yes ⊖ No	
If yes, what is the end date of data collection?		
If yes, what is the end date of data collection?		
If yes, what is the end date of data collection?		



Confidential Page 4 of 45 Development ⊖ Yes O No The length of follow-up is reported Ó NA (E.g. "Patients were followed from baseline for 10 years" and "10-year prediction of..."; notably for prognostic studies with long term follow-up. If this is not applicable for an article (i.e. diagnostic study or no follow-up), then score Not applicable.) External validation ⊖ Yes The length of follow-up is reported 🔿 No Ó NA (E.g. "Patients were followed from baseline for 10 years" and "10-year prediction of ... "; notably for prognostic studies with long term follow-up. If this is not applicable for an article (i.e. diagnostic study or no follow-up), then score Not applicable.) External validation ⊖ Yes The length of follow-up is reported ⊙ No \bigcirc NA (E.g. "Patients were followed from baseline for 10 years" and "10-year prediction of..."; notably for prognostic studies with long term follow-up. If this is not applicable for an article (i.e. diagnostic study or no follow-up), then score Not applicable.) If yes, what is the length of follow up? If yes, what is the length of follow up? If yes, what is the length of follow up? ⊖ Yes Development The length of the prediction horizon/time frame is ⊖ No reported \bigcirc NA External validation ○ Yes The length of the prediction horizon/time frame is ◯ No reported \bigcirc NA ⊖ Yes External validation The length of the prediction horizon/time frame is \bigcirc No Õ NA reported If yes, what is the length of the prediction horizon/time frame? If yes, what is the length of the prediction horizon/time frame? If yes, what is the length of the prediction horizon/time frame?



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Participants	
Development The study setting is reported	 Yes No (E.g.: 'surgery for endometrial cancer patients' is considered to be enough information about the study setting.)
External validation The study setting is reported	 Yes No (E.g.: 'surgery for endometrial cancer patients' is considered to be enough information about the study setting.)
External validation The study setting is reported	 Yes No (E.g.: 'surgery for endometrial cancer patients' is considered to be enough information about the study setting.)
Development What is the setting for the model?	 Primary care Secondary care Tertiary care General population Other (Primary care = GPs, dentists and pharmacists (often first point of care). Secondary care = hospital or clinic based care - can be planned (e.g., cataract operation) or emergency (e.g., fracture). Tertiary care = highly specialised treatments (e.g., transplant, hip replacement).)
External validation What is the setting for the model?	 Primary care Secondary care Tertiary care General population Other (Primary care = GPs, dentists and pharmacists (often first point of care). Secondary care = hospital or clinic based care - can be planned (e.g., cataract operation) or emergency (e.g., fracture). Tertiary care = highly specialised treatments (e.g., transplant, hip replacement).)
External validation What is the setting for the model?	 Primary care Secondary care Tertiary care General population Other (Primary care = GPs, dentists and pharmacists (often first point of care). Secondary care = hospital or clinic based care - can be planned (e.g., cataract operation) or emergency (e.g., fracture). Tertiary care = highly specialised treatments (e.g., transplant, hip replacement).)
If other, please specify	
If other, please specify	
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	Page 6 o
If other, please specify	
Development The number of centres involved is reported	 Yes No (If the number is not reported explicitly, but can be concluded from the name of the centre/cent or if clearly a single centre study, score Yes.)
External validation The number of centres involved is reported	 Yes No (If the number is not reported explicitly, but can be concluded from the name of the centre/cent or if clearly a single centre study, score Yes.)
External validation The number of centres involved is reported	 Yes No (If the number is not reported explicitly, but can be concluded from the name of the centre/cent or if clearly a single centre study, score Yes.)
How many centres involved?	
How many centres involved?	
How many centres involved?	
Development The geographical location (at least country) of centres involved is reported	 Yes No (If no geographical location is specified, but the location can be concluded from the name of the centre(s), score Yes.)
External validation The geographical location (at least country) of centres involved is reported	 Yes No (If no geographical location is specified, but the location can be concluded from the name of the centre(s), score Yes.)
External validation The geographical location (at least country) of centres involved is reported	 Yes No (If no geographical location is specified, but the location can be concluded from the name of the centre(s), score Yes.)
If yes, what was the geographic location of the data collection?	 Europe North America Latin America Asia Africa Oceania (Multiples answers are possible)

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If yes, what was the geographic location of the data collection?	 Europe North America Latin America Asia Africa Oceania (Multiples answers are possible) 	
If yes, what was the geographic location of the data collection?	 Europe North America Latin America Asia Africa Oceania (Multiples answers are possible) 	
Eligibility criteria		
Development In-/exclusion criteria are stated	 Yes No (These should explicitly be stated. Reasons for exclusion only described in a participant flow is not sufficient.) 	
External validation In-/exclusion criteria are stated	 Yes No (These should explicitly be stated. Reasons for exclusion only described in a participant flow is not sufficient.) 	
External validation In-/exclusion criteria are stated	 Yes No (These should explicitly be stated. Reasons for exclusion only described in a participant flow is not sufficient.) 	
Development What was the participant population?		
External validation What was the participant population?		
External validation What was the participant population?		
Development Details of any treatments received are described	 Yes No NA (This item is notably for prognostic modelling studies and is about treatment at baseline or during follow-up. The 'if relevant' judgment of treatment requires clinical knowledge and interpretation. If you are certain that treatment was not relevant, e.g. in some diagnostic model studies, score Not applicable 	





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External validation Details of any treatments received are described	 Yes No NA (This item is notably for prognostic modelling studies and is about treatment at baseline or during follow-up. The 'if relevant' judgment of treatment requires clinical knowledge and interpretation. If you are certain that treatment was not relevant, e.g. in some diagnostic model studies, score Not applicable 	
External validation Details of any treatments received are described	 Yes No NA (This item is notably for prognostic modelling studies and is about treatment at baseline or during follow-up. The 'if relevant' judgment of treatment requires clinical knowledge and interpretation. If you are certain that treatment was not relevant, e.g. in some diagnostic model studies, score Not applicable) 	
Outcome		
Development The outcome definition is clearly presented	○ Yes ○ No	
External validation The outcome definition is clearly presented	○ Yes ○ No	
External validation The outcome definition is clearly presented	○ Yes ○ No	
Development What is the type of primary outcome?	 Death Complications Recurrence Survival Other 	
External validation What is the type of primary outcome?	 Death Complications Recurrence Survival Other 	
External validation What is the type of primary outcome?	 Death Complications Recurrence Survival Other 	
If other, please specify		
If other, please specify		
If other, please specify		
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Development It is described how outcome was assessed	 Yes No (Including all elements of any composite, for example CVD [e.g. MI, HF, stroke])
External validation It is described how outcome was assessed	 Yes No (Including all elements of any composite, for example CVD [e.g. MI, HF, stroke])
External validation It is described how outcome was assessed	 Yes No (Including all elements of any composite, for example CVD [e.g. MI, HF, stroke])
Development It is described when the outcome was assessed (time point(s) since T0)	○ Yes ○ No
External validation It is described when the outcome was assessed (time point(s) since T0)	○ Yes ○ No
External validation It is described when the outcome was assessed (time point(s) since T0)	○ Yes ○ No
Development Actions to blind assessment of outcome to be predicted are reported	 Yes No (If it is clearly a non-issue (e.g. all-cause mortality or an outcome not requiring interpretation), score Yes. In all other instances, an explicit mention is expected.)
External validation Actions to blind assessment of outcome to be predicted are reported	 Yes No (If it is clearly a non-issue (e.g. all-cause mortality or an outcome not requiring interpretation), score Yes. In all other instances, an explicit mention is expected.)
External validation Actions to blind assessment of outcome to be predicted are reported	 Yes No (If it is clearly a non-issue (e.g. all-cause mortality or an outcome not requiring interpretation), score Yes. In all other instances, an explicit mention is expected.)



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Predictors	
Development All predictors are reported	 Yes No (For development, "all predictors" refers to all predictors that potentially could have been included in the 'final' model (including those considered in any univariable analyses).)
External validation All predictors are reported	 Yes No (For validation, "all predictors" means the predictors in the model being evaluated.)
External validation All predictors are reported	 Yes No (For validation, "all predictors" means the predictors in the model being evaluated.)
Development Number of candidate predictors considered	(If the number of candidate predictors is unclear, please fill this question with "UN")
External validation External validation Number of candidate predictors considered	(If the number of candidate predictors is unclear, please fill this question with 'UN')
Development What are the categories of the candidate predictors?	 Demography Clinical history Physical examination Blood and Urine parameters Imaging Genetic Risk Score Pathology Scale Score (e.g. pain, wellbeing, QoL) Questionnaires Other (Multiples answers are possible)
External validation External validation What are the categories of the candidate predictors?	 Demography Clinical history Physical examination Blood and Urine parameters Imaging Genetic Risk Score Pathology Scale Score (e.g. pain, wellbeing, QoL) Questionnaires Other (Multiples answers are possible)
If other, please specify	
	(list using (;) to separate if more than 1)
If other, please specify	(list using (;) to separate if more than 1)
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Development Were a-priori predictors considered/forced into the model?	 Yes No Unclear (E.g. Clinical reasoning, literature review, money constraints)
Development Predictor definitions are clearly presented	○ Yes ○ No
External validation Predictor definitions are clearly presented	○ Yes ○ No
External validation Predictor definitions are clearly presented	○ Yes ○ No
Development It is clearly described how the predictors were measured	○ Yes ○ No
External validation It is clearly described how the predictors were measured	○ Yes ○ No
External validation It is clearly described how the predictors were measured	○ Yes ○ No
Development It is clearly described when the predictors were measured	○ Yes ○ No
External validation It is clearly described when the predictors were measured	○ Yes ○ No
External validation It is clearly described when the predictors were measured	○ Yes ○ No
Development It is clearly described whether predictor assessments were blinded for outcome	 Yes No (For predictors for which it is clearly a non-issue (e.g. automatic blood pressure measurement, age, sex) and for instances where the predictors were clearly assessed before outcome assessment, score Yes. For all other predictors an explicit mention is expected.)
External validation It is clearly described whether predictor assessments were blinded for outcome	 Yes No (For predictors for which it is clearly a non-issue (e.g. automatic blood pressure measurement, age, sex) and for instances where the predictors were clearly assessed before outcome assessment, score Yes. For all other predictors an explicit mention is expected.)





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External validation It is clearly described whether predictor assessments were blinded for outcome	 Yes No (For predictors for which it is clearly a non-issue (e.g. automatic blood pressure measurement, age, sex) and for instances whe the predictors were clearly assessed before outcome assessment, score Yes. For all other predictors an explicit mention is expected.)
Development It is clearly described whether predictor assessments were blinded for the other predictors	○ Yes ○ No
External validation It is clearly described whether predictor assessments were blinded for the other predictors	○ Yes ○ No
External validation It is clearly described whether predictor assessments were blinded for the other predictors	⊖ Yes ○ No
Sample Size	
Development It is explained how the sample size was arrived at	 Yes No (Is there any mention of sample size, e.g. whet this was done on statistical grounds or practical/logistical grounds (e.g. an existing study cohort or data set of a RCT was used)?)
External validation It is explained how the sample size was arrived at	 Yes No (Is there any mention of sample size, e.g. whet this was done on statistical grounds or practical/logistical grounds (e.g. an existing study cohort or data set of a RCT was used)?)
External validation It is explained how the sample size was arrived at	 Yes No (Is there any mention of sample size, e.g. whet this was done on statistical grounds or practical/logistical grounds (e.g. an existing study cohort or data set of a RCT was used)?)
Development What is the reason for the sample?	 Power Justified time interval Size of existing/available data Events per variable Other
External validation What is the reason for the sample?	 Power Justified time interval Size of existing/available data Events per variable Other

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	Page 1
External validation What is the reason for the sample?	 Power Justified time interval Size of existing/available data Events per variable Other
Development If other, please specify	
External validation If other, please specify	
External validation If other, please specify	
Development What was the initial sample size of the study?	(If unclear, please fill this with "UN")
Development What was the final sample size of the study?	(If unclear, please fill this with "UN")
External validation What was the initial sample size of the study?	(If unclear, please fill this with "UN")
External validation What was the final sample size of the study?	(If unclear, please fill this with "UN")
External validation What was the initial sample size of the study?	(If unclear, please fill this with "UN")
External validation What was the final sample size of the study?	(If unclear, please fill this with "UN")
Missing Data	
Development Was missingness an explicit exclusion criterion for the data?	 ○ Yes ○ No ○ Unclear
External validation Was missingness an explicit exclusion criterion for the data?	 ○ Yes ○ No ○ Unclear
External validation Was missingness an explicit exclusion criterion for the data?	 ○ Yes ○ No ○ Unclear
Development If yes, how many patients were excluded due to missing data?	(If not reported, please fill this with "NR")



	Page 14 o
External validation If yes, how many patients were excluded due to missing data?	(If not reported, please fill this with "NR")
External validation If yes, how many patients were excluded due to missing data?	(If not reported, fill this with "NR")
Development The method for handling missing data (predictors and outcome) is mentioned	 Yes No (E.g. Complete case (explicit mention that individuals with missing values have been excluded), single imputation, multiple imputation, mean/median imputation. If there i no missing data, there should be an explicit mention that there is no missing data for all predictors and outcome. If so, score Yes. If it is unclear whether there is missing data (from e.g. the reported methods or results), score No If it is clear there is missing data is unclear, score No.)
External validation The method for handling missing data (predictors and outcome) is mentioned	 Yes No (E.g. Complete case (explicit mention that individuals with missing values have been excluded), single imputation, multiple imputation, mean/median imputation. If there i no missing data, there should be an explicit mention that there is no missing data for all predictors and outcome. If so, score Yes. If it is unclear whether there is missing data (from e.g. the reported methods or results), score No If it is clear there is missing data is unclear, score No.)
External validation The method for handling missing data (predictors and outcome) is mentioned	 Yes No (E.g. Complete case (explicit mention that individuals with missing values have been excluded), single imputation, multiple imputation, mean/median imputation. If there i no missing data, there should be an explicit mention that there is no missing data for all predictors and outcome. If so, score Yes. If it is unclear whether there is missing data (from e.g. the reported methods or results), score No If it is clear there is missing data, but the method for handling missing data is unclear, score No.)







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Development In presence of missing data, how was this dealt with?	 No missing data No imputation Multiple Imputation Complete case analysis Mean imputation Median imputation Other (If there is any discrepancy between how missing values were handled for outcome and predictors, please specify so it in the comments below the methods form.)
External validation In presence of missing data, how was this dealt with?	 No missing data No imputation Multiple Imputation Complete case analysis Mean imputation Median imputation Other (If there is any discrepancy between how missing values were handled for outcome and predictors, please specify so it in the comments below the methods form.)
External validation In presence of missing data, how was this dealt with?	 No missing data No imputation Multiple Imputation Complete case analysis Mean imputation Median imputation Other (If there is any discrepancy between how missing values were handled for outcome and predictors, please specify so it in the comments below the methods form. Report here imputation for predictors .)
If other, please specify	
If other, please specify	
If other, please specify	
Development If missing data were imputed, details of the software used are given	 ○ Yes ○ No ○ NA
External validation If missing data were imputed, details of the software used are given	 Yes No NA
External validation If missing data were imputed, details of the software used are given	 ○ Yes ○ No ○ NA

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	Page 16
Development If missing data were imputed, a description of which variables were included in the imputation procedure is given.	 ○ Yes ○ No ○ NA
External validation If missing data were imputed, a description of which variables were included in the imputation procedure is given.	 ○ Yes ○ No ○ NA
External validation If missing data were imputed, a description of which variables were included in the imputation procedure is given.	 ○ Yes ○ No ○ NA
Development If multiple imputation was used, the number of imputations is reported	○ Yes ○ No
External validation If multiple imputation was used, the number of imputations is reported	○ Yes ○ No
External validation If multiple imputation was used, the number of imputations is reported	○ Yes ○ No
Development How is missing data presented in the paper or supplemental material?	 Overall By all candidate variables By all final model variables By number of variables Not summarised
External validation How is missing data presented in the paper or supplemental material?	 Overall By all candidate variables By all final model variables By number of variables Not summarised
External validation How is missing data presented in the paper or supplemental material?	 Overall By all candidate variables By all final model variables By number of variables Not summarised
Development If missing data is presented/summarised, what is the percentage/number of individuals have missing data (overall)	(If this is unclear, please fill this with "UN")
External validation If missing data is presented/summarised, what is the percentage/number of individuals have missing data (overall)	(If this is unclear, please fill this with "UN")





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External validation If missing data is presented/summarised, what is the percentage/number of individuals have missing data (overall)	(If this is unclear, please fill this with "UN")
Statistical Analysis	
Data pre-processing	
Did the candidate predictors include continuous variables?	 ○ Yes ○ No ○ Unclear
For continuous predictors it is described whether they were modelled as linear, nonlinear (type of transformation specified) or categorized.	 Yes No Unclear (A general statement is sufficient, no need to describe this for each predictors.)
How were continuous predictors handled?	 Linear (no change) Non-linear (explicity/planned) Non-linear (implicitly/unplanned) Categorised (some) Categorised (all) Other (Non linear terms may not be explicitly reported/planned but handled within the model building process for ML and maybe implicit/unplanned. If an ML model is being evaluated and non-linear terms are not explicitly reported - choose 'Yes (implicit/unplanned)'. A general statement is sufficient, no need to describe this for each predictor separately.)
If other, please specify	
For categorical or categorized predictors, the cut-points were reported	○ Yes ○ No
For categorized predictors the method to choose the cut-points was clearly described	 Yes No NA (If no categorized predictors, score Not applicable.)
If categorised/dichotomised, how was this done?	 Quantiles Data dependent Mixture No rationale Based on previous literature or/and standarizatio



dential	Page 18 o
Is there any other data pre-processing methods used?	 Cleaning Aggregation Transformation Sampling Standardization Integration Reduction Other No (Multiples answers are possible. E.g. data aggregation (calculating predictors from other collected data), other predictor transformations data sampling (only using part of a dataset), predictor standardisation.)
If other, please specify	
	(list using (;) to separate if more then 1)
ls class imbalance addressed?	 Yes No NA (There is a disproportionate ratio of observatior in each class/group -most machine learning algorithms work best when the number of samp in each class are about equal.)
If yes, how?	
	(list using (;) to separate if more than 1)
Is class imbalance handling justified?	○ Yes ○ No
Is there any data reduction techniques used?	 Yes No (E.g. Missing values ratio, Low variance filter, High correlation filter, Random Forest / Ensembles tress, Principal Component Analysis (PCA), Backward feature elimination, Forward feature construction, autoencoder, Non-negativ matrix factorization, Kernel PCA, Graph-based kernel PCA, etc.)
If yes, what techniques were used?	
	(list using (;) to separate if more then 1)
Is data reduction justified?	○ Yes





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Was collinearity assessed?	 Yes, implicitly Yes, explicitly No NA (Collinearity may not be explicitly reported/planned but handled within the model building process for ML and maybe implicit/unplanned. If an ML model is being evaluated and collinearity are not explicitly reported - choose 'Yes (implicit/unplanned)'. A general statement is sufficient, no need to describe this for each predictor separately.)
Model building Instructions -Please extract the models in the order they -If more than 10 models were developed for -If a comparison with logistic regression was count and extract information.	y are presented in the article. The main outcome, only refer to the first 10. Is made, please included this model in the final
How many models were developed for the primary outcome?	(This should reflect the number of models you a going to extract on - primary outcome and prim timepoint (If more than 10 models were develop only refer to the first 10). If a comparison with logistic regression was made, please included this model in the final count.)
	this model in the final count.)
External validation It is described how predictions for individuals (in the validation set) were obtained from the model being validated	 Yes No (E.g. Using the original reported model coefficients with or without the intercept, and/or using updated or refitted model coefficients, or using a nomogram, spreadsheet web calculator.)
External validation It is described how predictions for individuals (in the validation set) were obtained from the model being validated	 Yes No (E.g. Using the original reported model coefficients with or without the intercept, and/or using updated or refitted model coefficients, or using a nomogram, spreadsheet web calculator.)
External validation It is described how predictions for individuals (in the validation set) were obtained from the model being validated Model 1 The type of statistical modelling approach is reported	 Yes No (E.g. Using the original reported model coefficients with or without the intercept, and/or using updated or refitted model coefficients, or using a nomogram, spreadsheet web calculator.) Yes No (E.g. Neural Network)
External validation It is described how predictions for individuals (in the validation set) were obtained from the model being validated Model 1 The type of statistical modelling approach is reported What is the ML technique being used?	 Yes No (E.g. Using the original reported model coefficients with or without the intercept, and/or using updated or refitted model coefficients, or using a nomogram, spreadsheet web calculator.) Yes No (E.g. Neural Network) Neural network Random forest Classification and regression tree (CART) Support vector machine Gradient boosting machine Logistic regression Other



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Do they clearly state why the technique is selected?	
	(E.g. R code availability, clinical question, previous literature, comparison. If not reported, fill this with "NR")
The approach used for predictor selection before modelling is described	 Yes No NA (Before modelling' means before any univariable multivariable analysis of predictor-outcome associations. If no predictor selection before modelling is done, score Not applicable. If it is unclear whether predictor selection before modelling is done, score No. If it is clear there was predictor selection before modelling but the method was not described, score No.)
The approach used for predictor selection during modelling is described	 Yes No NA (E.g. Univariable analysis, stepwise selection, bootstrap, Lasso. 'During modelling' includes both univariable or multivariable analysis of predictor-outcome associations. If no predictor selection during modelling is done (so-called full model approach), score Not applicable. If it is unclear whether predictor selection during modelling is done, score No. If it is clear there was predictor selection during modelling but the method was not described, score No.)
What was the model building strategy?	 Stepwise Forward selection Backward selection All predictors All significant in univariable Data-driven Other Unclear
If other, please specify	
	(list using (;) to separate if more than 1)
Are hyper-parameters tuning reported?	 Yes No NA (Answer yes if any information is giving about how the models were set-up. Term as number of layer node, optimization, hyperparameters, etc.)
Is predictor importance assessed?	 Yes No (E.g. Importance, Mean decrease/Increase in accuracy, Mean decrease Gini, Gini Index, Averag impurity decrease, etc.)





If yes, how?	
	(list using (;) to separate if more then 1)
Is there any ML technique being used for	⊖ Yes
predictors/feature selection?	\bigcirc No (E a SV(M))
	(E.g. SVM)
If yes, which technique?	
	(list using (;) to separate if more then 1)
Testing of interaction terms is described	○ Yes, explicitly.
	O Yes, implicitly.
	(Interaction terms may not be explicitly
	reported/planned but handled within the mode
	building process for ML and maybe
	Implicit/unplanned. If an ML model is being evaluated and interactions are not explicitly
	reported - choose 'Yes (implicit/unplanned)'. A
	general statement is sufficient, no need to
	describe this for each predictor separately.)
If applicable, how was censoring accounted for?	
	(If not applicable, fill this with "NA")
Testing of the proportionality of hazards in survival	∩ Yes
models is described	○ No
	Not applicable.)
What shrinkage/negalisation methods were used?	
what shimkage/penalisation methods were used:	O Uniform shrinkage
	\bigcirc Penalised estimation
	⊖ Other
If other, please specify	
Model 2	
The type of statistical modelling approach is	⊖ Yes
reported	() NO (E.g. Neural Notwork)
	(L.g. Neural Network)
What is the ML technique being used?	O Neural network
	\bigcirc Classification and regression tree (CAPT)
	\bigcirc Support vector machine
	O Gradient boosting machine
	O Logistic regression
	() Other



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Do they clearly state why the technique is selected?	
	(E.g. R code availability, clinical question, previous literature, comparison. If not reported, fill this with "NR")
The approach used for predictor selection before modelling is described	 Yes No NA (Before modelling' means before any univariable or multivariable analysis of predictor-outcome associations. If no predictor selection before modelling is done, score Not applicable. If it is unclear whether predictor selection before modelling is done, score No. If it is clear there was predictor selection before modelling but the method was not described, score No.)
The approach used for predictor selection during modelling is described	 Yes No NA (E.g. Univariable analysis, stepwise selection, bootstrap, Lasso. 'During modelling' includes both univariable or multivariable analysis of predictor-outcome associations. If no predictor selection during modelling is done (so-called full model approach), score Not applicable. If it is unclear whether predictor selection during modelling is done, score No. If it is clear there was predictor selection during modelling but the method was not described, score No.)
What was the model building strategy?	 Stepwise Forward selection Backward selection All predictors All significant in univariable Data-driven Other Unclear
If other, please specify	
	(list using (;) to separate if more than 1)
Are hyper-parameters tuning reported?	 Yes No NA (Answer yes if any information is giving about how the models were set-up. Term as number of layers node, optimization, hyperparameters, etc.)
Is predictor importance assessed?	 Yes No (E.g. Importance, Mean decrease/Increase in accuracy, Mean decrease Gini, Gini Index, Average impurity decrease etc.)





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If yes, how?	
	(list using (;) to separate if more than 1)
Is there any ML technique being used for predictors/feature selection?	○ Yes ○ No (E.g. SVM)
If yes, which technique?	
	(list using (;) to separate if more than 1)
Testing of interaction terms is described	 Yes, explicitly. Yes, implicitly. No (Interaction terms may not be explicitly reported/planned but handled within the mode building process for ML and maybe implicit/unplanned. If an ML model is being evaluated and interactions are not explicitly reported - choose 'Yes (implicit/unplanned)'. A general statement is sufficient, no need to describe this for each predictor separately.)
If applicable, how was censoring accounted for?	
	(If not applicable, fill this with "NA")
Testing of the proportionality of hazards in survival models is described	 ○ Yes ○ No ○ NA
What shrinkage/penalisation methods were used?	 None Uniform shrinkage Penalised estimation Other
If other, please specify	
Model 3	
The type of statistical modelling approach is reported	○ Yes ○ No (E.g. Neural Network)
What is the ML technique being used?	 Neural network Random forest Classification and regression tree (CART) Support vector machine Gradient boosting machine Logistic regression Other



Do they clearly state why the technique is selected?	
	(E.g. R code availability, clinical question, previous literature, comparison. If not reported, fill this with "NR")
The approach used for predictor selection before modelling is described	 Yes No NA (Before modelling' means before any univariable or multivariable analysis of predictor-outcome associations. If no predictor selection before modelling is done, score Not applicable. If it is unclear whether predictor selection before modelling is done, score No. If it is clear there was predictor selection before modelling but the method was not described, score No.)
The approach used for predictor selection during modelling is described	 Yes No NA (E.g. Univariable analysis, stepwise selection, bootstrap, Lasso. 'During modelling' includes both univariable or multivariable analysis of predictor-outcome associations. If no predictor selection during modelling is done (so-called full model approach), score Not applicable. If it is unclear whether predictor selection during modelling is done, score No. If it is clear there was predictor selection during modelling but the method was not described, score No.)
What was the model building strategy?	 Stepwise Forward selection Backward selection All predictors All significant in univariable Data-driven Other Unclear
If other, please specify	
	(list using (;) to separate if more than 1)
Are hyper-parameters tuning reported?	 Yes No NA (Answer yes if any information is giving about how the models were set-up. Term as number of layers node, optimization, hyperparameters, etc.)
Is predictor importance assessed?	 Yes No (E.g. Importance, Mean decrease/Increase in accuracy, Mean decrease Gini, Gini Index, Average impurity decrease, etc.)





If yes, how?	
	(list using (;) to separate if more than 1)
Is there any ML technique being used for predictors/feature selection?	○ Yes ○ No (E.g. SVM)
If yes, which technique?	
	(list using (;) to separate if more than 1)
Testing of interaction terms is described	 Yes, explicitly. Yes, implicitly. No (Interaction terms may not be explicitly reported/planned but handled within the model building process for ML and maybe implicit/unplanned. If an ML model is being evaluated and interactions are not explicitly reported - choose 'Yes (implicit/unplanned)'. A general statement is sufficient, no need to describe this for each predictor separately.)
If applicable, how was censoring accounted for?	
	(If not applicable, fill this with "NA")
Testing of the proportionality of hazards in survival models is described	 ○ Yes ○ No ○ NA
What shrinkage/penalisation methods were used?	 None Uniform shrinkage Penalised estimation Other
If other, please specify	
Model 4	
The type of statistical modelling approach is reported	 ○ Yes ○ No (E.g. Neural Network)
What is the ML technique being used?	 Neural network Random forest Classification and regression tree (CART) Support vector machine Gradient boosting machine Logistic regression Other

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Do they clearly state why the technique is selected?	
	(E.g. R code availability, clinical question, previous literature, comparison. If not reported, fill this with "NR")
The approach used for predictor selection before modelling is described	 Yes No NA (Before modelling' means before any univariable multivariable analysis of predictor-outcome associations. If no predictor selection before modelling is done, score Not applicable. If it is unclear whether predictor selection before modelling is done, score No. If it is clear there was predictor selection before modelling but the method was not described, score No.)
The approach used for predictor selection during modelling is described	 Yes No NA (E.g. Univariable analysis, stepwise selection, bootstrap, Lasso. 'During modelling' includes both univariable or multivariable analysis of predictor-outcome associations. If no predictor selection during modelling is done (so-called full model approach), score Not applicable. If it is unclear whether predictor selection during modelling is done, score No. If it is clear there was predictor selection during modelling but the method was not described, score No.)
What was the model building strategy?	 Stepwise Forward selection Backward selection All predictors All significant in univariable Data-driven Other Unclear
If other, please specify	
	(list using (;) to separate if more than 1)
Are hyper-parameters tuning reported?	 Yes No NA (Answer yes if any information is giving about how the models were set-up. Term as number of layer node, optimization, hyperparameters, etc.)
Is predictor importance assessed?	 Yes No (E.g. Importance, Mean decrease/Increase in accuracy, Mean decrease Gini, Gini Index, Averag impurity decrease, etc.)





If yes, how?	
	(list using (;) to separate if more than 1)
Is there any ML technique being used for predictors/feature selection?	○ Yes ○ No (E.g. SVM)
If yes, which technique?	
	(list using (;) to separate if more than 1)
Testing of interaction terms is described	 Yes, explicitly. Yes, implicitly. No (Interaction terms may not be explicitly reported/planned but handled within the mode building process for ML and maybe implicit/unplanned. If an ML model is being evaluated and interactions are not explicitly reported - choose 'Yes (implicit/unplanned)'. A general statement is sufficient, no need to describe this for each predictor separately.)
If applicable, how was censoring accounted for?	
	(If not applicable, fill this with "NA")
Testing of the proportionality of hazards in survival models is described	 ○ Yes ○ No ○ NA
What shrinkage/penalisation methods were used?	 None Uniform shrinkage Penalised estimation Other
If other, please specify	
Model 5	
The type of statistical modelling approach is reported	 ○ Yes ○ No (E.g. Neural Network)
What is the ML technique being used?	 Neural network Random forest Classification and regression tree (CART) Support vector machine Gradient boosting machine Logistic regression Other



Do they clearly state why the technique is selected?	
	(E.g. R code availability, clinical question, previous literature, comparison. If not reported, fill this with "NR")
The approach used for predictor selection before modelling is described	 Yes No NA (Before modelling' means before any univariable of multivariable analysis of predictor-outcome associations. If no predictor selection before modelling is done, score Not applicable. If it is unclear whether predictor selection before modelling is done, score No. If it is clear there was predictor selection before modelling but the method was not described, score No.)
The approach used for predictor selection during modelling is described	 Yes No NA (E.g. Univariable analysis, stepwise selection, bootstrap, Lasso. 'During modelling' includes both univariable or multivariable analysis of predictor-outcome associations. If no predictor selection during modelling is done (so-called full model approach), score Not applicable. If it is unclear whether predictor selection during modelling is done, score No. If it is clear there was predictor selection during modelling but the method was not described, score No.)
What was the model building strategy?	 Stepwise Forward selection Backward selection All predictors All significant in univariable Data-driven Other Unclear
If other, please specify	
	(list using (;) to separate if more than 1)
Are hyper-parameters tuning reported?	 Yes No NA (Answer yes if any information is giving about how the models were set-up. Term as number of layers node, optimization, hyperparameters, etc.)
Is predictor importance assessed?	 Yes No (E.g. Importance, Mean decrease/Increase in accuracy, Mean decrease Gini, Gini Index, Average impurity decrease, etc.)





If yes, how?	
	(list using (;) to separate if more than 1)
Is there any ML technique being used for predictors/feature selection?	○ Yes ○ No (E.g. SVM)
If yes, which technique?	
	(list using (;) to separate if more than 1)
Testing of interaction terms is described	 Yes, explicitly. Yes, implicitly. No (Interaction terms may not be explicitly reported/planned but handled within the mode building process for ML and maybe implicit/unplanned. If an ML model is being evaluated and interactions are not explicitly reported - choose 'Yes (implicit/unplanned)'. A general statement is sufficient, no need to describe this for each predictor separately.)
If applicable, how was censoring accounted for?	
	(If not applicable, fill this with "NA")
Testing of the proportionality of hazards in survival models is described	 ○ Yes ○ No ○ NA
What shrinkage/penalisation methods were used?	 None Uniform shrinkage Penalised estimation Other
If other, please specify	
Model C	
	Q ¥22
reported	O Yes O No (E.g. Neural Network)
What is the ML technique being used?	 Neural network Random forest Classification and regression tree (CART) Support vector machine Gradient boosting machine Logistic regression Other



Do they clearly state why the technique is selected?	
	(E.g. R code availability, clinical question, previous literature, comparison. If not reported, fill this with "NR")
The approach used for predictor selection before modelling is described	 Yes No NA (Before modelling' means before any univariable of multivariable analysis of predictor-outcome associations. If no predictor selection before modelling is done, score Not applicable. If it is unclear whether predictor selection before modelling is done, score No. If it is clear there was predictor selection before modelling but the method was not described, score No.)
The approach used for predictor selection during modelling is described	 Yes No NA (E.g. Univariable analysis, stepwise selection, bootstrap, Lasso. 'During modelling' includes both univariable or multivariable analysis of predictor-outcome associations. If no predictor selection during modelling is done (so-called full model approach), score Not applicable. If it is unclear whether predictor selection during modelling is done, score No. If it is clear there was predictor selection during modelling but the method was not described, score No.)
What was the model building strategy?	 Stepwise Forward selection Backward selection All predictors All significant in univariable Data-driven Other Unclear
If other, please specify	
	(list using (;) to separate if more than 1)
Are hyper-parameters tuning reported?	 Yes No NA (Answer yes if any information is giving about how the models were set-up. Term as number of layers node, optimization, hyperparameters, etc.)
Is predictor importance assessed?	 Yes No (E.g. Importance, Mean decrease/Increase in accuracy, Mean decrease Gini, Gini Index, Averagi impurity decrease, etc.)





If yes, how?	
	(list using (;) to separate if more than 1)
Is there any ML technique being used for predictors/feature selection?	○ Yes ○ No (E.g. SVM)
If yes, which technique?	
	(list using (;) to separate if more than 1)
Testing of interaction terms is described	 Yes, implicitly. Yes, explicitly. No (Interaction terms may not be explicitly reported/planned but handled within the mode building process for ML and maybe implicit/unplanned. If an ML model is being evaluated and interactions are not explicitly reported - choose 'Yes (implicit/unplanned)'. A general statement is sufficient, no need to describe this for each predictor separately.)
If applicable, how was censoring accounted for?	
	(If not applicable, fill this with "NA")
Testing of the proportionality of hazards in survival models is described	 ○ Yes ○ No ○ NA
What shrinkage/penalisation methods were used?	 None Uniform shrinkage Penalised estimation Other
If other, please specify	
Model 7	
The type of statistical modelling approach is reported	 ○ Yes ○ No (E.g. Neural Network)
What is the ML technique being used?	 Neural network Random forest Classification and regression tree (CART) Support vector machine Gradient boosting machine Logistic regression Other


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Do they clearly state why the technique is selected?	
	(E.g. R code availability, clinical question, previous literature, comparison. If not reported, fill this with "NR")
The approach used for predictor selection before modelling is described	 Yes No NA (Before modelling' means before any univariable multivariable analysis of predictor-outcome associations. If no predictor selection before modelling is done, score Not applicable. If it is unclear whether predictor selection before modelling is done, score No. If it is clear there was predictor selection before modelling but the method was not described, score No.)
The approach used for predictor selection during modelling is described	 Yes No NA (E.g. Univariable analysis, stepwise selection, bootstrap, Lasso. 'During modelling' includes both univariable or multivariable analysis of predictor-outcome associations. If no predictor selection during modelling is done (so-called full model approach), score Not applicable. If it is unclear whether predictor selection during modelling is done, score No. If it is clear there was predictor selection during modelling but the method was not described, score No.)
What was the model building strategy?	 Stepwise Forward selection Backward selection All predictors All significant in univariable Data-driven Other Unclear
If other, please specify	
	(list using (;) to separate if more than 1)
Are hyper-parameters tuning reported?	 Yes No NA (Answer yes if any information is giving about he models were set-up. Term as number of laye node, optimization, hyperparameters, etc.)
Is predictor importance assessed?	 Yes No (E.g. Importance, Mean decrease/Increase in accuracy, Mean decrease Gini, Gini Index, Avera impurity decrease, etc.)



If yes, how?	
	(list using (;) to separate if more than 1)
Is there any ML technique being used for predictors/feature selection?	○ Yes ○ No (E.g. SVM)
If yes, which technique?	
	(list using (;) to separate if more than 1)
Testing of interaction terms is described	 Yes, implicitly. Yes, explicitly. No (Interaction terms may not be explicitly reported/planned but handled within the mode building process for ML and maybe implicit/unplanned. If an ML model is being evaluated and interactions are not explicitly reported - choose 'Yes (implicit/unplanned)'. A general statement is sufficient, no need to describe this for each predictor separately.)
If applicable, how was censoring accounted for?	
	(If not applicable, fill this with "NA")
Testing of the proportionality of hazards in survival models is described	 ○ Yes ○ No ○ NA
What shrinkage/penalisation methods were used?	 None Uniform shrinkage Penalised estimation Other
If other, please specify	
Model 8	
The type of statistical modelling approach is reported	 ○ Yes ○ No (E.g. Neural Network)
What is the ML technique being used?	 Neural network Random forest Classification and regression tree (CART) Support vector machine Gradient boosting machine Logistic regression Other



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Do they clearly state why the technique is selected?	
	(E.g. R code availability, clinical question, previous literature, comparison. If not reported, fill this with "NR")
The approach used for predictor selection before modelling is described	 Yes No NA (Before modelling' means before any univariable multivariable analysis of predictor-outcome associations. If no predictor selection before modelling is done, score Not applicable. If it is unclear whether predictor selection before modelling is done, score No. If it is clear there was predictor selection before modelling but the method was not described, score No.)
The approach used for predictor selection during modelling is described	 Yes No NA (E.g. Univariable analysis, stepwise selection, bootstrap, Lasso. 'During modelling' includes both univariable or multivariable analysis of predictor-outcome associations. If no predictor selection during modelling is done (so-called full model approach), score Not applicable. If it is unclear whether predictor selection during modelling is done, score No. If it is clear there was predictor selection during modelling but the method was not described, score No.)
What was the model building strategy?	 Stepwise Forward selection Backward selection All predictors All significant in univariable Data-driven Other Unclear
If other, please specify	
	(list using (;) to separate if more than 1)
Are hyper-parameters tuning reported?	 Yes No NA (Answer yes if any information is giving about how the models were set-up. Term as number of layer node, optimization, hyperparameters, etc.)
Is predictor importance assessed?	 Yes No (E.g. Importance, Mean decrease/Increase in accuracy, Mean decrease Gini, Gini Index, Averag impurity decrease, etc.)





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If yes, how?	
	(list using (;) to separate if more than 1)
Is there any ML technique being used for predictors/feature selection?	○ Yes ○ No (E.g. SVM)
If yes, which technique?	
	(list using (;) to separate if more than 1)
Testing of interaction terms is described	 Yes, implicitly. Yes, explicitly. No (Interaction terms may not be explicitly reported/planned but handled within the mode building process for ML and maybe implicit/unplanned. If an ML model is being evaluated and interactions are not explicitly reported - choose 'Yes (implicit/unplanned)'. A general statement is sufficient, no need to describe this for each predictor separately.)
If applicable, how was censoring accounted for?	
	(If not applicable, fill this with "NA")
Testing of the proportionality of hazards in survival models is described	 ○ Yes ○ No ○ NA
What shrinkage/penalisation methods were used?	 None Uniform shrinkage Penalised estimation Other
If other, please specify	
Model 9	
The type of statistical modelling approach is reported	 ○ Yes ○ No (E.g. Neural Network)
What is the ML technique being used?	 Neural network Random forest Classification and regression tree (CART) Support vector machine Gradient boosting machine Logistic regression Other



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Do they clearly state why the technique is selected?	
	(E.g. R code availability, clinical question, previous literature, comparison. If not reported, fill this with "NR")
The approach used for predictor selection before modelling is described	 Yes No NA (Before modelling' means before any univariable multivariable analysis of predictor-outcome associations. If no predictor selection before modelling is done, score Not applicable. If it is unclear whether predictor selection before modelling is done, score No. If it is clear there was predictor selection before modelling but the method was not described, score No.)
The approach used for predictor selection during modelling is described	 Yes No NA (E.g. Univariable analysis, stepwise selection, bootstrap, Lasso. 'During modelling' includes both univariable or multivariable analysis of predictor-outcome associations. If no predictor selection during modelling is done (so-called full model approach), score Not applicable. If it is unclear whether predictor selection during modelling is done, score No. If it is clear there was predictor selection during modelling but the method was not described, score No.)
What was the model building strategy?	 Stepwise Forward selection Backward selection All predictors All significant in univariable Data-driven Other Unclear
If other, please specify	
	(list using (;) to separate if more than 1)
Are hyper-parameters tuning reported?	 Yes No NA (Answer yes if any information is giving about how the models were set-up. Term as number of layer node, optimization, hyperparameters, etc.)
Is predictor importance assessed?	 Yes No (E.g. Importance, Mean decrease/Increase in accuracy, Mean decrease Gini, Gini Index, Averag impurity decrease, etc.)





If yes, how?	
	(list using (;) to separate if more than 1)
Is there any ML technique being used for predictors/feature selection?	○ Yes ○ No (E.g. SVM)
If yes, which technique?	
	(list using (;) to separate if more than 1)
Testing of interaction terms is described	 Yes, implicitly. Yes, explicitly. No (Interaction terms may not be explicitly reported/planned but handled within the mode building process for ML and maybe implicit/unplanned. If an ML model is being evaluated and interactions are not explicitly reported - choose 'Yes (implicit/unplanned)'. A general statement is sufficient, no need to describe this for each predictor separately.)
If applicable, how was censoring accounted for?	
	(If not applicable, fill this with "NA")
Testing of the proportionality of hazards in survival models is described	 ○ Yes ○ No ○ NA
What shrinkage/penalisation methods were used?	 None Uniform shrinkage Penalised estimation Other
If other, please specify	
Model 10	
The type of statistical modelling approach is reported	 ○ Yes ○ No (E.g. Neural Network)
What is the ML technique being used?	 Neural network Random forest Classification and regression tree (CART) Support vector machine Gradient boosting machine Logistic regression Other



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Do they clearly state why the technique is selected?	
	(E.g. R code availability, clinical question, previous literature, comparison. If not reported, fill this with "NR")
The approach used for predictor selection before modelling is described	 Yes No NA (Before modelling' means before any univariable multivariable analysis of predictor-outcome associations. If no predictor selection before modelling is done, score Not applicable. If it is unclear whether predictor selection before modelling is done, score No. If it is clear there was predictor selection before modelling but the method was not described, score No.)
The approach used for predictor selection during modelling is described	 Yes No NA (E.g. Univariable analysis, stepwise selection, bootstrap, Lasso. 'During modelling' includes both univariable or multivariable analysis of predictor-outcome associations. If no predictor selection during modelling is done (so-called full model approach), score Not applicable. If it is unclear whether predictor selection during modelling is done, score No. If it is clear there was predictor selection during modelling but the method was not described, score No.)
What was the model building strategy?	 Stepwise Forward selection Backward selection All predictors All significant in univariable Data-driven Other Unclear
If other, please specify	
	(list using (;) to separate if more than 1)
Are hyper-parameters tuning reported?	 Yes No NA (Answer yes if any information is giving about how the models were set-up. Term as number of layer node, optimization, hyperparameters, etc.)
Is predictor importance assessed?	 Yes No (E.g. Importance, Mean decrease/Increase in accuracy, Mean decrease Gini, Gini Index, Averag impurity decrease, etc.)





lf yes, how?	
	(list using (;) to separate if more than 1)
Is there any ML technique being used for predictors/feature selection?	○ Yes ○ No (E.g. SVM)
If yes, which technique?	
	(list using (;) to separate if more than 1)
Testing of interaction terms is described	 Yes, implicitly. Yes, explicitly. No (Interaction terms may not be explicitly reported/planned but handled within the mode building process for ML and maybe implicit/unplanned. If an ML model is being evaluated and interactions are not explicitly reported - choose 'Yes (implicit/unplanned)'. A general statement is sufficient, no need to describe this for each predictor separately.)
If applicable, how was censoring accounted for?	
	(If not applicable, fill this with "NA")
Testing of the proportionality of hazards in survival models is described	 ○ Yes ○ No ○ NA
What shrinkage/penalisation methods were used?	 None Uniform shrinkage Penalised estimation Other
If other, please specify	
Internal validation	
Internal validation is reported	 Yes No (If the use of internal validation is clearly a non-issue (e.g. in case of very large data sets), score Yes. For all other situations an explicit mention is expected.)
How is the model internally validated?	 Split sample Bootstrapping Cross-validation Other Unclear
If split sample, what % split was used for the development?	(If not reported please fill this with "UN")



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If split sample, was it a random or non-random split?	 ○ Random ○ Non-random ○ Unclear
If non-random split was used, how?	
	(E.g. Temporal splitting)
If bootstrap method, how many were performed?	
	(If not reported, please fill this with "UN")
If bootstrap method, were selection of variables included in the bootstrap?	○ Yes ○ No
If cross validation, please specify the method used	
	(E.g. ten-fold. If not reported, fill this with "UN")
Model Performance	
Development Measures for model discrimination are described	○ Yes ○ No
External validation Measures for model discrimination are described	○ Yes ○ No
External validation Measures for model discrimination are described	○ Yes ○ No
Development How was discrimination assessed?	 AUC/ AUROC/Area under the curve C-statistic Harrell's C-index D-Statistic Other
External validation How was discrimination assessed?	 AUC/ AUROC/Area under the curve C-statistic Harrell's C-index D-Statistic Other
External validation How was discrimination assessed?	 AUC/ AUROC/Area under the curve C-statistic Harrell's C-index D-Statistic Other
If other, please specify	
	(list using (;) to separate if more than 1)
If other, please specify	
	(list using (;) to separate if more than 1)

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If other, please specify	
	(list using (;) to separate if more than 1)
Development Measures for model calibration are described	⊖ Yes ⊖ No
External validation Measures for model calibration are described	⊖ Yes ⊖ No
External validation Measures for model calibration are described	⊖ Yes ⊖ No
Development How was calibration assessed?	 H-L Calibration plot Calibration slope Calibration intercept Calibration in the large Calibration table Kappa Observed/expected ratio Other
External validation How was calibration assessed?	 H-L Calibration plot Calibration slope Calibration intercept Calibration in the large Calibration table Kappa Observed/expected ratio Other
External validation How was calibration assessed?	 H-L Calibration plot Calibration slope Calibration intercept Calibration in the large Calibration table Kappa Observed/expected ratio Other
If other, please specify	
If other, please specify	



	Page 42
Development Other performance measures are described	 Yes No (E.g. R2, Brier score, predictive values, sensitivity, specificity, AUC difference, decision curve analysis, net reclassification improvement, integrated discrimination improvement, AIC)
External validation Other performance measures are described	 Yes No (E.g. R2, Brier score, predictive values, sensitivity, specificity, AUC difference, decision curve analysis, net reclassification improvement, integrated discrimination improvement, AIC)
External validation Other performance measures are described	 Yes No (E.g. R2, Brier score, predictive values, sensitivity, specificity, AUC difference, decision curve analysis, net reclassification improvement, integrated discrimination improvement, AIC)
Development lf yes, please specify	 R2 Brier score predictive values sensitivity specificity AUC difference decision curve analysis net reclassification improvement integrated discrimination improvement AIC Other
External validation If yes, please specify	 R2 Brier score predictive values sensitivity specificity AUC difference decision curve analysis net reclassification improvement integrated discrimination improvement AIC Other
External validation If yes, please specify	 R2 Brier score predictive values sensitivity specificity AUC difference decision curve analysis net reclassification improvement integrated discrimination improvement AIC Other



If other, please specify	
If other, please specify	
If other, please specify	
Model Updating	
A description of model-updating is given	 Yes No NA (E.g. Intercept recalibration, regression coefficient recalibration, refitting the whole model, adding a new predictor If updating wa done, it should be clear which updating methor was applied to score Yes. If it is not explicitly mentioned that updating was applied the study, score this item as 'Not applicable'.
lf yes, please specify	
	(E.g. Intercept recalibration, regression coefficient recalibration, refitting the whole model, adding a new predictor.)
Risk groups	
Development Were risk groups created?	○ Yes ○ No
External validation Were risk groups created?	○ Yes ○ No
External validation Were risk groups created?	○ Yes ○ No
How many risk groups were created?	
How many risk groups were created?	
How many risk groups were created?	
Development What method was used to create these risk groups?	 Count factors present Data driven Equal size Other data dependent



External validation What method was used to create these risk groups?	 Count factors present Data driven Equal size Other data dependent Unclear
External validation What method was used to create these risk groups?	 Count factors present Data driven Equal size Other data dependent Unclear
Development If risk groups were created, risk group boundaries (risk thresholds) are specified	 Yes No (Score this item separately for development ar validation if a study includes both developmen and validation.)
External validation If risk groups were created, risk group boundaries (risk thresholds) are specified	 Yes No (Score this item separately for development ar validation if a study includes both developmen and validation.)
External validation If risk groups were created, risk group boundaries (risk thresholds) are specified	 Yes No (Score this item separately for development ar validation if a study includes both developmen and validation.)
Development Is any subgroup analysis prespecified?	<pre>○ Yes ○ No</pre>
External validation Is any subgroup analysis prespecified?	○ Yes ○ No
External validation Is any subgroup analysis prespecified?	○ Yes ○ No
If yes, how many subgroup criteria were defined?	
If yes, how many subgroup criteria were defined?	
If yes, how many subgroup criteria were defined?	
Development Is any sensitivity/subpopulation analysis prespecified?	○ Yes ○ No
External validation Is any sensitivity/subpopulation analysis prespecified?	○ Yes ○ No

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External validation Is any sensitivity/subpopulation analysis prespecified?	○ Yes ○ No	
Is yes, how many subpopulation criteria area defined?		
Is yes, how many subpopulation criteria area defined?		
Is yes, how many subpopulation criteria area defined?		
Comments		
Any additional comment about the methods section of this article?		

(If there is something in the "Methods" section that does not fit into the questions of this form - please use this space to detail. Also use this space to detail anything you are unsure about.)



Results	JC - Machine Learning Systematic Revie Page 1 of 55
Article ID	
What type of study is reported?	 Development (including internal validation) Development with external validation (same model) External validation only (This questions is repeated due to branching logic. Please answer again.)
Is there any diagram/draw to clarify the results?	○ Yes ○ No
Participants	
Development The flow of participants is reported	○ Yes ○ No
External validation The flow of participants is reported	○ Yes ○ No
External validation The flow of participants is reported	○ Yes ○ No
Development The number of participants with and without the outcome is reported	 Yes No NA (If outcomes are continuous, score Not applicable.)
External validation The number of participants with and without the outcome is reported	 Yes No NA (If outcomes are continuous, score Not applicable.)
External validation The number of participants with and without the outcome is reported	 Yes No NA (If outcomes are continuous, score Not applicable.)
Development A summary of follow-up time is presented	 Yes (median) Yes (average) Other No NA (This notably applies to prognosis studies and diagnostic studies with follow-up as diagnostic outcome. If this is not applicable for an article (i.e. diagnostic study or no follow-up), then score Not applicable.)



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External validation A summary of follow-up time is presented	 Yes (median) Yes (average) Other No
	NA (This notably applies to prognosis studies and diagnostic studies with follow-up as diagnostic outcome. If this is not applicable for an article (i.e. diagnostic study or no follow-up), then score Not applicable.)
External validation A summary of follow-up time is presented	 Yes (median) Yes (average) Other No NA (This notably applies to prognosis studies and diagnostic studies with follow-up as diagnostic outcome. If this is not applicable for an article (i.e. diagnostic study or no follow-up), then score Not applicable.)
If other, please specify	
	(list using (;) to separate if more than 1)
If other, please specify	
	(list using (;) to separate if more than 1)
If other, please specify	
	(list using (;) to separate if more than 1)
Development For time-to-event endpoints, do the authors report how many have X-years of follow-up?	 ○ Yes ○ No ○ NA
External validation For time-to-event endpoints, do the authors report how many have X-years of follow-up?	 ○ Yes ○ No ○ NA
Development Basic demographics are reported	\bigcirc Yes \bigcirc No (At least age and sex are reported.)
External validation Basic demographics are reported	\bigcirc Yes \bigcirc No (At least age and sex are reported.)
External validation Basic demographics are reported	 ○ Yes ○ No (At least age and sex are reported.)
Development Number of predictors in the final model	(If the final predictors are unclear, please fill

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External validation Number of predictors in the final model	(If the final predictors are unclear, please fill this question with a 'UN')
External validation Number of predictors in the final model	(If the final predictors are unclear, please fill this question with a 'UN')
Development Summary information is provided for all predictors included in the final developed/validated model	○ Yes ○ No
External validation Summary information is provided for all predictors included in the final developed/validated model	○ Yes ○ No
External validation Summary information is provided for all predictors included in the final developed/validated model	○ Yes ○ No
Development The number of participants with missing data for predictors is reported	 Yes No NA (When no missing values is reported, fill this with "NA")
External validation The number of participants with missing data for predictors is reported	 Yes No NA (When no missing values is reported, fill this with "NA")
External validation The number of participants with missing data for predictors is reported	 Yes No NA (When no missing values is reported, fill this with "NA")
Development Final number of models developed/validated reported	(Please provide the number. If this is unclear, please fill this with "UN")
External validation Final number of models developed/validated reported	(Please provide the number. If this is unclear, please fill this with "UN")
External validation Final number of models developed/validated reported	(Please provide the number. If this is unclear,



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External validation Demographic characteristics (at least age and gender) of the validation study participants are reported along with those of the original development study	 Yes No NA (For incremental value reports, in case addition predictors are not added to a previously developed prediction model but rather added to conventional predictors in a newly fitted model, score Not applicable.)
External validation Distributions of predictors in the model of the validation study participants are reported along with those of the original development study	 Yes No NA (For incremental value reports, in case addition predictors are not added to a previously developed prediction model but rather added to conventional predictors in a newly fitted model, score Not applicable.)
External validation Outcomes of the validation study participants are reported along with those of the original development study	 Yes No NA (For incremental value reports, in case additions predictors are not added to a previously developed prediction model but rather added to conventional predictors in a newly fitted model, score Not applicable)
Model development	
The number of participants in each analysis is specified	\bigcirc Yes \bigcirc No (e.g. in the analysis of each model if more than one model is developed)
What is the number of participants (included in the analysis) reported in the main model?	(If the number of participants is not reported, please fill this with 'NR'.)
The number of outcome events in each analysis is specified	 Yes No NA (e.g. in the analysis of each model if more than one model is developed. If outcomes are continuous, score Not applicable.)
What is the number of events (initial) reported in the main outcome?	(If the number of events is not reported, please fill this with 'NR'.)
What is the number of events (included in the analysis) reported in the main model?	(If the number of events is not reported, please fill this with 'NR'.)



The unadjusted associations between each predictor and outcome are reported	 Yes No NA (If any univariable analysis is mentioned in the methods but not in the results, score No. If nothing on univariable analysis (in methods or results) is reported, score this item as Not applicable)
Model specification	
Development The regression coefficient (or a derivative such as hazard ratio, odds ratio, risk ratio) for each predictor in the model is reported	 ○ Yes ○ No ○ NA
Development The intercept or the cumulative baseline hazard (or baseline survival) for at least one time point is reported	 ○ Yes ○ No ○ NA
Development An explanation (e.g. a simplified scoring rule, chart, nomogram of the model, reference to online calculator, or worked example) is provided to explain how to use the model for individualised predictions.	○ Yes ○ No
Development Is there enough information to calculate the risk of the outcome in a new individual?	○ Yes ○ No

-If more than 10 models were developed for the main outcome, only refer to the first 10.

-If a comparison with logistic regression was made, please include this model in the final

count and extract information.

How many models were developed for the primary outcome?

(This should reflect the number of models you are going to extract on - primary outcome and primary timepoint. If more than 10 models were developed, please refer to the first 10 models. If a Logistic regression model was performed, please also extract data from this model)



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Model 1	
Is this the recommended model?	\bigcirc Yes \bigcirc No (If there is only one model, please state 'Yes')
Development A discrimination measure is presented	 ○ Yes ○ No (E.g. C-index / area under the ROC curve)
External validation A discrimination measure is presented	\bigcirc Yes \bigcirc No (E.g. C-index / area under the ROC curve)
Development Which discrimination measures are described?	 AUC/AUROC/Area under the curve C-statistic Harrell's c-index D-statistic
External validation Which discrimination measures are described?	 AUC/AUROC/Area under the curve C-statistic Harrell's c-index D-statistic
The confidence interval (or standard error) of the AUC/AUROC/Area under the curve is presented	⊖ Yes ⊖ No
The confidence interval (or standard error) of the AUC/AUROC/Area under the curve is presented	○ Yes ○ No
What was the AUC/AUROC/Area under the curve apparent discrimination estimate (+95% Cl, if given)?	(Please use the following format - estimate (lower Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the AUC/AUROC/Area under the curve apparent discrimination estimate (+95% CI, if given)?	(Please use the following format - estimate (lower Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the AUC/AUROC/Area under the curve corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected AUC/AUROC/Area under the curve (+95% CI, if given)?	(Please use the following format - estimate (lower Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the C-statistic is presented	○ Yes ○ No
The confidence interval (or standard error) of the C-statistic is presented	○ Yes ○ No

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What was the C-statistic apparent discrimination	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the C-statistic apparent discrimination	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the C-statistic corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected C-statistic (+95% CI,	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the Harrell's c-index is presented	○ Yes ○ No
The confidence interval (or standard error) of the Harrell's c-index is presented	○ Yes ○ No
What was the Harrell's c-index apparent	
	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the Harrell's c-index apparent	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the Harrell's c-index corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected Harrell's c-index (+95% Cl, if given)?	
	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the d-statistic presented	○ Yes ○ No
The confidence interval (or standard error) of the	O Yes

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What was the d-statistic apparent discrimination estimate (+95% Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the d-statistic apparent discrimination estimate (+95% CI, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the d-statistic corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected d-statistic (+95% Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Development Measures for model calibration are described	 Yes No (E.g. calibration plot, calibration slope or intercept, calibration table, Hosmer Lemeshow test, O/E ratio.)
External validation Measures for model calibration are described	 Yes No (E.g. calibration plot, calibration slope or intercept, calibration table, Hosmer Lemeshow test, O/E ratio.)
Development How was calibration assessed?	 Calibration plot Calibration slope Calibration intercept Calibration in the large Calibration table Kappa Observed/expected ratio H-L
External validation How was calibration assessed?	 Calibration plot Calibration slope Calibration intercept Calibration in the large Calibration table Kappa Observed/expected ratio H-L
What was the apparent [gen_methods_103:checked] estimate (+95% Cl, if given)?	(list using (;) to separate if more than 1. If not reported, fill this with "NR")

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What was the apparent [gen_methods_117:checked] estimate (+95% Cl, if given)?	
	(list using (;) to separate if more than 1. If not reported, fill this with "NR")
Was the [gen_methods_103:checked] measure corrected for optimism?	○ Yes○ No○ Unclear
What was the optimism corrected [gen_methods_103:checked] estimate (+95% Cl, if given)?	(list using (;) to separate if more than 1. If measures were corrected for optimism but not reported, please fill this with "NR".)
Development Other model performance measures are presented	 Yes No (E.g. R2, Brier score, predictive values, sensitivity, specificity, AUC difference, decision curve analysis, net reclassification improvement, integrated discrimination improvement, AIC.)
External validation Other model performance measures are presented	 Yes No (E.g. R2, Brier score, predictive values, sensitivity, specificity, AUC difference, decision curve analysis, net reclassification improvement, integrated discrimination improvement, AIC.)
Development lf yes, please specify	 R2 Brier score predictive values sensitivity specificity AUC difference decision curve analysis net reclassification improvement integrated discrimination improvement AIC Accuracy Other (Multiples answers are possible)
External validation f yes, please specify	 R2 Brier score predictive values sensitivity specificity AUC difference decision curve analysis net reclassification improvement integrated discrimination improvement AIC Accuracy Other (Multiples answers are possible)



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If other, please specify	
	(list the names using (;) to separate if more that 1)
If other, please specify	
	(list the names using (;) to separate if more that 1)
What was the apparent [gen_methods_104:checked] estimate (+95% Cl, if given)?	
	(list using (;) to separate if more than 1. If not reported, fill this with "NR")
What was the apparent [gen_methods_118:checked] estimate (+95% Cl, if given)?	
	(list using (;) to separate if more than 1. If not reported, fill this with "NR")
Was the [gen_methods_104:checked] measure corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected [gen_methods_104:checked] estimate (+95% Cl, if given)?	(list using (;) to separate if more than 1. If not reported, fill this with "NR")
Model 2	
Is this the recommended model?	 ○ Yes ○ No (If there is only one model, please state 'Yes')
Development A discrimination measure is presented	\bigcirc Yes \bigcirc No (E.g. C-index / area under the ROC curve)
External validation A discrimination measure is presented	\bigcirc Yes \bigcirc No (E.g. C-index / area under the ROC curve)
Development Which performance measures are described?	 AUC/AUROC/Area under the curve C-statistic Harrell's c-index D-statistic (Multiples answers are possible)
External validation Which performance measures are described?	 AUC/AUROC/Area under the curve C-statistic Harrell's c-index D-statistic (Multiples answers are possible)



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The confidence interval (or standard error) of the AUC/AUROC/Area under the curve is presented	⊖ Yes ⊖ No
The confidence interval (or standard error) of the AUC/AUROC/Area under the curve is presented	○ Yes ○ No
What was the AUC/AUROC/Area under the curve apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported, fill this with "NR")
What was the AUC/AUROC/Area under the curve apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the AUC/AUROC/Area under the curve corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected AUC/AUROC/Area under the curve (95% CI, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the C-statistics is presented	○ Yes ○ No
The confidence interval (or standard error) of the C-statistics is presented	○ Yes ○ No
What was the C-statistics apparent discrimination estimate (+95%CI, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the C-statistics apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the C-statistics corrected for optimism?	○ Yes○ No○ Unclear
What was the optimism corrected C-statistic (95% Cl, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the Harrel's c-index is presented	⊖ Yes ⊖ No

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The confidence interval (or standard error) of the Harrel's c-index is presented	⊖ Yes ⊖ No
What was the Harrel's c-index apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the Harrel's c-index apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the Harrel's c-index corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected Harrel's c-index (95% Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the D-statistic is presented	○ Yes ○ No
The confidence interval (or standard error) of the D-statistic is presented	○ Yes ○ No
What was the D-statistics apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the D-statistics apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the D-statistics corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected D-statistics (95% Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Development Measures for model calibration are described	 Yes No (E.g. calibration plot, calibration slope or intercept, calibration table, Hosmer Lemeshow test, O/E ratio.)

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External validation Measures for model calibration are described	 Yes No (E.g. calibration plot, calibration slope or intercept, calibration table, Hosmer Lemeshow test, O/E ratio.)
Development How was calibration assessed?	 Calibration plot Calibration slope Calibration intercept Calibration in the large Calibration table Kappa Observed/expected ratio H-L
External validation How was calibration assessed?	 Calibration plot Calibration slope Calibration intercept Calibration in the large Calibration table Kappa Observed/expected ratio H-L
What was the apparent [gen_methods_119:checked] estimate (+95% Cl, if given)?	(list using (;) to separate if more than 1. If not reported, fill this with "NR")
What was the apparent [gen_methods_120:checked] estimate (+95% Cl, if given)?	(list using (;) to separate if more than 1. If not reported, fill this with "NR")
Was the [gen_methods_119:checked] measure corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected [gen_methods_119:checked] estimate (+95% Cl, if given)?	(list using (;) to separate if more than 1. If not reported, fill this with "NR")
Development Other model performance measures are presented	 Yes No (E.g. R2, Brier score, predictive values, sensitivity, specificity, AUC difference, decision curve analysis, net reclassification improvement, integrated discrimination improvement, AIC.)
External validation Other model performance measures are presented	 Yes No (E.g. R2, Brier score, predictive values, sensitivity, specificity, AUC difference, decision curve analysis, net reclassification improvement, integrated discrimination

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Development If yes, please specify	 R2 Brier score predictive values sensitivity specificity AUC difference decision curve analysis pet reclassification improvement
	 integrated discrimination improvement AIC Accuracy Other (Multiples answers are possible)
External validation If yes, please specify	 R2 Brier score predictive values sensitivity specificity AUC difference decision curve analysis net reclassification improvement integrated discrimination improvement AIC Accuracy Other (Multiples answers are possible)
If other, please specify	
	(list using (;) to separate if more than 1)
If other, please specify	
	(list using (;) to separate if more than 1)
What was the apparent [gen_methods_121:checked] estimate (+95% CI, if given)?	
	(list using (;) to separate if more than 1. If not reported, fill this with "NR")
What was the apparent [gen_methods_122:checked] estimate (+95% CI, if given)?	
	(list using (;) to separate if more than 1. If not reported, fill this with "NR")
Was the [gen_methods_121:checked] measure corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected [gen_methods_121:checked] estimate (+95% CI, if given)?	
g	(list using (;) to separate if more than 1. If not reported, fill this with "NR")

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Model 3	
Is this the recommended model?	\bigcirc Yes \bigcirc No (If there is only one model, please state 'Yes')
Development A discrimination measure is presented	\bigcirc Yes \bigcirc No (E.g. C-index / area under the ROC curve)
External validation A discrimination measure is presented	\bigcirc Yes \bigcirc No (E.g. C-index / area under the ROC curve)
Development Which performance measures are described?	 AUC/AUROC/Area under the curve C-statistic Harrell's c-index D-statistic (Multiples answers are possible)
External validation Which performance measures are described?	 AUC/AUROC/Area under the curve C-statistic Harrell's c-index D-statistic (Multiples answers are possible)
The confidence interval (or standard error) of the AUC/AUROC/Area under the curve is presented	○ Yes ○ No
The confidence interval (or standard error) of the AUC/AUROC/Area under the curve is presented	○ Yes ○ No
What was the AUC/AUROC/Area under the curve apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (lower Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the AUC/AUROC/Area under the curve apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (lower Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the AUC/AUROC/Area under the curve corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected AUC/AUROC/Area under the curve (95% CI, if given)?	(Please use the following format - estimate (lower Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the C-statistics is presented	○ Yes ○ No

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The confidence interval (or standard error) of the C-statistics is presented	○ Yes ○ No
What was the C-statistics apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81).
	If not reported fill this with "NR")
What was the C-statistics apparent discrimination estimate (+95%Cl, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the C-statistics corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected C-statistics (95% CI,	
ir given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the Harrel's c-index is presented	○ Yes ○ No
The confidence interval (or standard error) of the Harrel's c-index is presented	○ Yes ○ No
What was the Harrel's c-index apparent discrimination estimate (+95%CL if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the Harrel's c-index apparent discrimination	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the Harrel's c-index corrected for optimism?	○ Yes○ No○ Unclear
What was the optimism corrected Harrel's c-index (95%	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the D-statistics is presented	○ Yes ○ No

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The confidence interval (or standard error) of the D-statistics is presented	○ Yes ○ No
What was the D-statistics apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (lov
	Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the D-statistics apparent discrimination estimate (+95%Cl, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the D-statistics corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected D-statistics (95% CI, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Development Measures for model calibration are described	 Yes No (E.g. calibration plot, calibration slope or intercept, calibration table, Hosmer Lemeshow test, O/E ratio.)
External validation Measures for model calibration are described	⊖ Yes ○ No
	(E.g. calibration plot, calibration slope or intercept, calibration table, Hosmer Lemeshow test, O/E ratio.)
Development How was calibration assessed?	 Calibration plot Calibration slope
	 Calibration intercept Calibration in the large Calibration table
	 Calibration table Kappa Observed/expected ratio H-L
External validation How was calibration assessed?	Calibration plot
	☐ Calibration intercept ☐ Calibration in the large ☐ Calibration table
	☐ Kappa ☐ Observed/expected ratio

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What was the apparent [gen_methods_139:checked] estimate (+95% CI, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
What was the apparent [gen_methods_140:checked] estimate (+95% Cl, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Was the [gen_methods_139:checked] measure corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected [gen_methods_139:checked] estimate (+95% CI, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Development Other model performance measures are presented	 Yes No (E.g. R2, Brier score, predictive values, sensitivity, specificity, AUC difference, decision curve analysis, net reclassification improvement, integrated discrimination improvement, AIC.)
External validation Other model performance measures are presented	 Yes No (E.g. R2, Brier score, predictive values, sensitivity, specificity, AUC difference, decision curve analysis, net reclassification improvement, integrated discrimination improvement, AIC.)
Development If yes, please specify	 R2 Brier score predictive values sensitivity specificity AUC difference decision curve analysis net reclassification improvement integrated discrimination improvement AIC Accuracy Other (Multiples answers are possible)





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External validation If yes, please specify	 R2 Brier score predictive values sensitivity specificity AUC difference decision curve analysis net reclassification improvement integrated discrimination improvement AIC Accuracy Other (Multiples answers are possible)
If other, please specify	
	(list using (;) to separate if more than 1)
If other, please specify	
	(list using (;) to separate if more than 1)
What was the apparent [gen_methods_123:checked] estimate (+95% Cl, if given)?	
	(list using (;) to separate if more than 1. If not reported, fill this with "NR")
What was the apparent [gen_methods_123:checked] estimate (+95% Cl, if given)?	
	(list using (;) to separate if more than 1. If not reported, fill this with "NR")
Was the [gen_methods_123:checked] measure corrected for optimism?	○ Yes○ No○ Unclear
What was the optimism corrected [gen_methods_123:checked] estimate (+95% Cl, if given)?	
	(list using (;) to separate if more than 1. If measures were corrected for optimism but not reported, please fill this with "NR".)
Model 4	
Is this the recommended model?	\bigcirc Yes \bigcirc No (If there is only one model, please state 'Yes')
Development A discrimination measure is presented	\bigcirc Yes \bigcirc No (E.g. C-index / area under the ROC curve)
External validation A discrimination measure is presented	 ○ Yes ○ No (E.g. C-index / area under the BOC curve)

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Development Which performance measures are described?	 AUC/AUROC/Area under the curve C-statistic Harrell's c-index D-statistic (Multiples answers are possible)
External validation Which performance measures are described?	 AUC/AUROC/Area under the curve C-statistic Harrell's c-index D-statistic (Multiples answers are possible)
The confidence interval (or standard error) of the AUC/AUROC/Area under the curve is presented	○ Yes ○ No
The confidence interval (or standard error) of the AUC/AUROC/Area under the curve is presented	○ Yes ○ No
What was the AUC/AUROC/Area under the curve apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (lowe Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the AUC/AUROC/Area under the curve apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (lowe Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the AUC/AUROC/Area under the curve corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected AUC/AUROC/Area under the curve (95% CI, if given)?	(Please use the following format - estimate (lowe Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the C-statistics is presented	○ Yes ○ No
The confidence interval (or standard error) of the C-statistics is presented	○ Yes ○ No
What was the C-statistics apparent discrimination estimate (+95%CI, if given)?	(Please use the following format - estimate (lowe CI - upper CI). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")



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What was the C-statistics apparent discrimination	
estimate (+95%Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the C-statistics corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected C-statistics (95% Cl, if given)?	
	(Please use the following format - estimate (Iov Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the Harrel's c-index is presented	○ Yes ○ No
The confidence interval (or standard error) of the Harrel's c-index is presented	○ Yes ○ No
What was the Harrel's c-index apparent discrimination estimate (+95%CI, if given)?	
	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the Harrel's c-index apparent discrimination estimate (+95%CL if given)?	
	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the Harrel's c-index corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected Harrel's c-index (95%	
Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the D-statistic is presented	○ Yes ○ No
The confidence interval (or standard error) of the D-statistic is presented	○ Yes ○ No
What was the D-statistic apparent discrimination	
	(Please use the following format - estimate (low CI - upper CI). For example 0.79 (0.75 - 0.81).



What was the D-statistic apparent discrimination estimate (+95%CI, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the D-statistic corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected D-statistic (95% CI, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Development Measures for model calibration are described	⊖ Yes ⊖ No
	(E.g. calibration plot, calibration slope or intercept, calibration table, Hosmer Lemeshow test, O/E ratio.)
External validation	⊖ Yes ⊖ No
	(E.g. calibration plot, calibration slope or intercept, calibration table, Hosmer Lemeshow test, O/E ratio.)
Development How was calibration assessed?	Calibration plot Calibration slope
	Calibration intercept
	Calibration table
	 Observed/expected ratio H-L
External validation	Calibration plot
	Calibration intercept
	Kappa Observed/expected ratio H-L
What was the apparent [gen_methods_141:checked] estimate (+95% CI, if given)?	
- ,, <u>-</u> ,-	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
What was the apparent [gen_methods_142:checked]	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")


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Was the [gen_methods_141:checked] measure corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected [gen_methods_141:checked] estimate (+95% CI, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Development Other model performance measures are presented	 Yes No (E.g. R2, Brier score, predictive values, sensitivity, specificity, AUC difference, decision curve analysis, net reclassification improvement, integrated discrimination improvement, AIC.)
External validation Other model performance measures are presented	 Yes No (E.g. R2, Brier score, predictive values, sensitivity, specificity, AUC difference, decision curve analysis, net reclassification improvement, integrated discrimination improvement, AIC.)
Development lf yes, please specify	 R2 Brier score predictive values sensitivity specificity AUC difference decision curve analysis net reclassification improvement integrated discrimination improvement AIC Accuracy Other (Multiples answers are possible)
External validation If yes, please specify	 R2 Brier score predictive values sensitivity specificity AUC difference decision curve analysis net reclassification improvement integrated discrimination improvement AIC Accuracy Other (Multiples answers are possible)
If other please specify	

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If other, please specify	
	(list using (;) to separate if more than 1.)
What was the apparent [gen_methods_125:checked] estimate (+95% Cl, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
What was the apparent [gen_methods_126:checked] estimate (+95% CI, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Was the [gen_methods_125:checked] measure corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected [gen_methods_125:checked] estimate (+95% Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Model 5	
Is this the recommended model?	\bigcirc Yes \bigcirc No (If there is only one model, please state 'Yes')
Development A discrimination measure is presented	\bigcirc Yes \bigcirc No (E.g. C-index / area under the ROC curve)
External validation A discrimination measure is presented	\bigcirc Yes \bigcirc No (E.g. C-index / area under the ROC curve)
Development Which performance measures are described?	 AUC/AUROC/Area under the curve C-statistic Harrell's c-index D-statistic (Multiples answers are possible)
External validation Which performance measures are described?	 AUC/AUROC/Area under the curve C-statistic Harrell's c-index D-statistic (Multiples approximate are pessible)

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The confidence interval (or standard error) of the AUC/AUROC/Area under the curve is presented	○ Yes ○ No
The confidence interval (or standard error) of the AUC/AUROC/Area under the curve is presented	○ Yes ○ No
What was the AUC/AUROC/Area under the curve apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the AUC/AUROC/Area under the curve apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the AUC/AUROC/Area under the curve corrected for optimism?	○ Yes○ No○ Unclear
What was the optimism corrected AUC/AUROC/Area under the curve (95% CI, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the C-statistic is presented	○ Yes ○ No
The confidence interval (or standard error) of the C-statistic is presented	○ Yes ○ No
What was the C-statistic apparent discrimination estimate (+95%CI, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the C-statistic apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the C-statistic corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected C-statistic (95% Cl, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the Harrel's c-index is presented	⊖ Yes ⊖ No

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The confidence interval (or standard error) of the Harrel's c-index is presented	⊖ Yes ⊖ No
What was the Harrel's c-index apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the Harrel's c-index apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the Harrel's c-index corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected Harrel's c-index (95% Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the D-statistic is presented	○ Yes ○ No
The confidence interval (or standard error) of the D-statistic is presented	○ Yes ○ No
What was the D-statistic apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the D-statistic apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the D-statistic corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected D-statistic (95% Cl, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Development Measures for model calibration are described	 Yes No (E.g. calibration plot, calibration slope or intercept, calibration table, Hosmer Lemeshow test, O/E ratio.)

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External validation Measures for model calibration are described	 Yes No (E.g. calibration plot, calibration slope or intercept, calibration table, Hosmer Lemeshow test, O/E ratio.)
Development How was calibration assessed?	 Calibration plot Calibration slope Calibration intercept Calibration in the large Calibration table Kappa Observed/expected ratio H-L
External validation How was calibration assessed?	 Calibration plot Calibration slope Calibration intercept Calibration in the large Calibration table Kappa Observed/expected ratio H-L
What was the apparent [gen_methods_143:checked] estimate (+95% CI, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
What was the apparent [gen_methods_144:checked] estimate (+95% Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Was the [gen_methods_143:checked] measure corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected [gen_methods_143:checked] estimate (+95% CI, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Development Other model performance measures are presented	 Yes No (E.g. R2, Brier score, predictive values, sensitivity, specificity, AUC difference, decision curve analysis, net reclassification improvement, integrated discrimination improvement, AIC.)



External validation Other model performance measures are presented	 Yes No (E.g. R2, Brier score, predictive values, sensitivity, specificity, AUC difference, decision curve analysis, net reclassification improvement, integrated discrimination improvement, AIC.)
Development If yes, please specify	 R2 Brier score predictive values sensitivity specificity AUC difference decision curve analysis net reclassification improvement integrated discrimination improvement AIC Accuracy Other (Multiples answers are possible)
External validation If yes, please specify	 R2 Brier score predictive values sensitivity specificity AUC difference decision curve analysis net reclassification improvement integrated discrimination improvement AIC Accuracy Other (Multiples answers are possible)
If other, please specify	
	(list using (;) to separate if more than 1)
If other, please specify	
	(list using (;) to separate if more than 1)
What was the apparent [gen_methods_127:checked] estimate (+95% CI, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
What was the apparent [gen_methods_128:checked] estimate (+95% CI, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")

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Was the [gen_methods_127:checked] measure corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected [gen_methods_127:checked] estimate (+95% Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported,
	fill this with "NR")
Model 6	
Is this the recommended model?	\bigcirc Yes \bigcirc No (If there is only one model, please state 'Yes')
Development A discrimination measure is presented	 ○ Yes ○ No (E.g. C-index / area under the ROC curve)
External validation A discrimination measure is presented	\bigcirc Yes \bigcirc No (E.g. C-index / area under the ROC curve)
Development Which performance measures are described?	 AUC/AUROC/Area under the curve C-statistic Harrell's c-index D-statistic (Multiples answers are possible)
External validation Which performance measures are described?	 AUC/AUROC/Area under the curve C-statistic Harrell's c-index D-statistic (Multiples answers are possible)
The confidence interval (or standard error) of the AUC/AUROC/Area under the curve is presented	○ Yes ○ No
The confidence interval (or standard error) of the AUC/AUROC/Area under the curve is presented	○ Yes ○ No
What was the AUC/AUROC/Area under the curve apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the AUC/AUROC/Area under the curve apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")

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Was the AUC/AUROC/Area under the curve corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected AUC/AUROC/Area under the curve (95% CI, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the C-statistic is presented	○ Yes ○ No
The confidence interval (or standard error) of the C-statistic is presented	○ Yes ○ No
What was the C-statistic apparent discrimination estimate (+95%CI, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the C-statistic apparent discrimination estimate (+95%CI, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the C-statistic corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected C-statistic (95% Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the Harrel's c-index is presented	○ Yes ○ No
The confidence interval (or standard error) of the Harrel's c-index is presented	○ Yes ○ No
What was the Harrel's c-index apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the Harrel's c-index apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")

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Was the Harrel's c-index corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected Harrel's c-index (95% Cl, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the D-statistic is presented	○ Yes ○ No
The confidence interval (or standard error) of the D-statistic is presented	○ Yes ○ No
What was the D-statistic apparent discrimination estimate (+95%Cl, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the D-statistic apparent discrimination estimate (+95%Cl, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the D-statistic corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected D-statistic (95% Cl,	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Development Measures for model calibration are described	○ Yes ○ No
	(E.g. calibration plot, calibration slope or intercept, calibration table, Hosmer Lemeshow test, O/E ratio.)
External validation Measures for model calibration are described	○ Yes ○ No
	(E.g. calibration plot, calibration slope or intercept, calibration table, Hosmer Lemeshow test, O/E ratio.)
Development How was calibration assessed?	Calibration plot Calibration slope
	 Calibration intercept Calibration in the large
	☐ Calibration table □ Kappa
	Observed/expected ratio

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External validation How was calibration assessed?	 Calibration plot Calibration slope Calibration intercept Calibration in the large Calibration table Kappa Observed/expected ratio H-L
What was the apparent [gen_methods_145:checked] estimate (+95% Cl, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
What was the apparent [gen_methods_146:checked] estimate (+95% Cl, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Was the [gen_methods_145:checked] measure corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected [gen_methods_145:checked] estimate (+95% Cl, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Development Other model performance measures are presented	 Yes No (E.g. R2, Brier score, predictive values, sensitivity, specificity, AUC difference, decision curve analysis, net reclassification improvement, integrated discrimination improvement, AIC.)
External validation Other model performance measures are presented	 Yes No (E.g. R2, Brier score, predictive values, sensitivity, specificity, AUC difference, decision curve analysis, net reclassification improvement, integrated discrimination improvement, AIC.)





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Development If yes, please specify	 R2 Brier score predictive values sensitivity specificity AUC difference decision curve analysis net reclassification improvement integrated discrimination improvement AIC Accuracy Other (Multiples answers are possible)
External validation If yes, please specify	 R2 Brier score predictive values sensitivity specificity AUC difference decision curve analysis net reclassification improvement integrated discrimination improvement AIC Accuracy Other (Multiples answers are possible)
If other, please specify	(list using (·) to separate if more than 1)
If other, please specify	
	(list using (;) to separate if more than 1)
What was the apparent [gen_methods_129:checked] estimate (+95% CI, if given)?	
	(Please use the following format - estimate (low CI - upper CI). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
What was the apparent [gen_methods_138:checked] estimate (+95% CI, if given)?	
	(Please use the following format - estimate (low CI - upper CI). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Was the [gen_methods_129:checked] measure corrected for optimism?	 ○ Yes ○ No ○ Unclear





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What was the optimism corrected [gen_methods_129:checked] estimate (+95% CI, if given)?	
given).	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Model 7	
Is this the recommended model?	\bigcirc Yes \bigcirc No (If there is only one model, please state 'Yes')
Development A discrimination measure is presented	\bigcirc Yes \bigcirc No (E.g. C-index / area under the ROC curve)
External validation A discrimination measure is presented	 ○ Yes ○ No (E.g. C-index / area under the ROC curve)
Development Which performance measures are described?	 AUC/AUROC/Area under the curve C-statistic Harrell's c-index D-statistic (Multiples answers are possible)
External validation Which performance measures are described?	 AUC/AUROC/Area under the curve C-statistic Harrell's c-index D-statistic (Multiples answers are possible)
The confidence interval (or standard error) of the AUC/AUROC/Area under the curve is presented	○ Yes ○ No
The confidence interval (or standard error) of the AUC/AUROC/Area under the curve is presented	○ Yes ○ No
What was the AUC/AUROC/Area under the curve apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the AUC/AUROC/Area under the curve apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the AUC/AUROC/Area under the curve corrected for optimism?	 ○ Yes ○ No ○ Unclear

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What was the optimism corrected AUC/AUROC/Area under the curve (95% CI, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the C-statistic is presented	○ Yes ○ No
The confidence interval (or standard error) of the C-statistic is presented	○ Yes ○ No
What was the C-statistic apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the C-statistic apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the C-statistic corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected C-statistic (95% CI, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the Harrel's c-index is presented	○ Yes ○ No
The confidence interval (or standard error) of the Harrel's c-index is presented	○ Yes ○ No
What was the Harrel's c-index apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the Harrel's c-index apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the Harrel's c-index corrected for optimism?	 ○ Yes ○ No ○ Unclear

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What was the optimism corrected Harrel's c-index (95% Cl, if given)?	(Please use the following format - estimate (low
	Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the D-statistic is presented	○ Yes ○ No
The confidence interval (or standard error) of the D-statistic is presented	○ Yes ○ No
What was the D-statistic apparent discrimination estimate (+95%CI, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the D-statistic apparent discrimination estimate (+95%CL if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the D-statistic corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected D-statistic (95% Cl,	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Development Measures for model calibration are described	○ Yes ○ No
	(E.g. calibration plot, calibration slope or intercept, calibration table, Hosmer Lemeshow test, O/E ratio.)
External validation Measures for model calibration are described	○ Yes ○ No
	(E.g. calibration plot, calibration slope or intercept, calibration table, Hosmer Lemeshow test, O/E ratio.)
Development How was calibration assessed?	Calibration plot Calibration slope
	Calibration intercept Calibration in the large
	Calibration table
	Observed/expected ratio H-L

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External validation How was calibration assessed?	 Calibration plot Calibration slope Calibration intercept Calibration in the large Calibration table Kappa Observed/expected ratio H-L
What was the apparent [gen_methods_147:checked] estimate (+95% Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
What was the apparent [gen_methods_148:checked] estimate (+95% Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Was the [gen_methods_147:checked] measure corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected [gen_methods_147:checked] estimate (+95% Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Development Other model performance measures are presented	 Yes No (E.g. R2, Brier score, predictive values, sensitivity, specificity, AUC difference, decision curve analysis, net reclassification improvement, integrated discrimination improvement, AIC.)
External validation Other model performance measures are presented	 Yes No (E.g. R2, Brier score, predictive values, sensitivity, specificity, AUC difference, decision curve analysis, net reclassification improvement, integrated discrimination improvement. AIC.)





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Development If yes, please specify	 R2 Brier score predictive values sensitivity specificity AUC difference decision curve analysis net reclassification improvement integrated discrimination improvement AIC Accuracy Other (Multiples answers are possible)
External validation If yes, please specify	 R2 Brier score predictive values sensitivity specificity AUC difference decision curve analysis net reclassification improvement integrated discrimination improvement AIC Accuracy Other (Multiples answers are possible)
If other, please specify	
	(list using (;) to separate if more than 1.)
If other, please specify	
	(list using (;) to separate if more than 1.)
What was the apparent [gen_methods_130:checked] estimate (+95% CI, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
What was the apparent [gen_methods_137:checked] estimate (+95% Cl, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Was the [gen_methods_130:checked] measure corrected for optimism?	 ○ Yes ○ No ○ Unclear



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What was the optimism corrected [gen_methods_130:checked] estimate (+95% CI, if	
given):	(Please use the following format - estimate (low CI - upper CI). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Model 8	
Is this the recommended model?	 ○ Yes ○ No (If there is only one model, please state 'Yes')
Development A discrimination measure is presented	\bigcirc Yes \bigcirc No (E.g. C-index / area under the ROC curve)
External validation A discrimination measure is presented	\bigcirc Yes \bigcirc No (E.g. C-index / area under the ROC curve)
Development Which performance measures are described?	 AUC/AUROC/Area under the curve C-statistic Harrell's c-index D-statistic (Multiples answers are possible)
External validation Which performance measures are described?	 AUC/AUROC/Area under the curve C-statistic Harrell's c-index D-statistic (Multiples answers are possible)
The confidence interval (or standard error) of the AUC/AUROC/Area under the curve is presented	○ Yes ○ No
The confidence interval (or standard error) of the AUC/AUROC/Area under the curve is presented	○ Yes ○ No
What was the AUC/AUROC/Area under the curve apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the AUC/AUROC/Area under the curve apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the AUC/AUROC/Area under the curve corrected for optimism?	 ○ Yes ○ No ○ Unclear

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What was the optimism corrected AUC/AUROC/Area under the curve (95% CI, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the C-statistic is presented	○ Yes ○ No
The confidence interval (or standard error) of the C-statistic is presented	○ Yes ○ No
What was the C-statistic apparent discrimination estimate (+95%Cl. if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the C-statistic apparent discrimination estimate (+95%Cl. if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the C-statistic corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected C-statistic (95% CI,	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the Harrel's c-index is presented	○ Yes ○ No
The confidence interval (or standard error) of the Harrel's c-index is presented	○ Yes ○ No
What was the Harrel's c-index apparent discrimination estimate (+95%Cl, if given)?	
· _ ·	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the Harrel's c-index apparent discrimination estimate (+95%Cl. if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the Harrel's c-index corrected for optimism?	O Yes O No

Was the Harrel's c-index corrected for optimism?	○ Yes○ No○ Unclear
What was the optimism corrected Harrel's c-index (95% Cl, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the optimism corrected Harrel's c-index (95% Cl. if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the D-statistic is presented	○ Yes ○ No
The confidence interval (or standard error) of the D-statistic is presented	○ Yes ○ No
What was the D-statistic apparent discrimination estimate (+95%Cl, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the D-statistic apparent discrimination estimate (+95%CL if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the D-statistic corrected for optimism?	○ Yes ○ No
	⊖ Unclear
What was the optimism corrected D-statistic (95% CI, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Development Measures for model calibration are described	○ Yes ○ No
	(E.g. calibration plot, calibration slope or intercept, calibration table, Hosmer Lemeshow test, O/E ratio.)
External validation Measures for model calibration are described	○ Yes ○ No
	(E.g. calibration plot, calibration slope or intercept, calibration table, Hosmer Lemeshow test. Q/F ratio.)

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Development How was calibration assessed?	 Calibration plot Calibration slope Calibration intercept Calibration in the large Calibration table Kappa Observed/expected ratio H-L
External validation How was calibration assessed?	 Calibration plot Calibration slope Calibration intercept Calibration in the large Calibration table Kappa Observed/expected ratio H-L
What was the apparent [gen_methods_149:checked] estimate (+95% CI, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
What was the apparent [gen_methods_150:checked] estimate (+95% Cl, if given)?	(Please use the following format - estimate (lov CI - upper CI). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Was the [gen_methods_149:checked] measure corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected [gen_methods_149:checked] estimate (+95% CI, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Development Other model performance measures are presented	 Yes No (E.g. R2, Brier score, predictive values, sensitivity, specificity, AUC difference, decision curve analysis, net reclassification improvement, integrated discrimination improvement, AIC.)
External validation Other model performance measures are presented	 Yes No (E.g. R2, Brier score, predictive values, sensitivity, specificity, AUC difference, decision curve analysis, net reclassification improvement, integrated discrimination improvement, AIC.)



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Development If yes, please specify	 R2 Brier score predictive values sensitivity specificity AUC difference decision curve analysis net reclassification improvement integrated discrimination improvement AIC Accuracy Other (Multiples answers are possible)
External validation If yes, please specify	 R2 Brier score predictive values sensitivity specificity AUC difference decision curve analysis net reclassification improvement integrated discrimination improvement AIC Accuracy Other (Multiples answers are possible)
If other, please specify	(list using (:) to separate if more than 1.)
If other, please specify	
	(list using (;) to separate if more than 1.)
What was the apparent [gen_methods_131:checked] estimate (+95% Cl, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
What was the apparent [gen_methods_136:checked] estimate (+95% CI, if given)?	
	(Please use the following format - estimate (low CI - upper CI). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Was the [gen_methods_131:checked] measure corrected for optimism?	 ○ Yes ○ No ○ Unclear





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What was the optimism corrected [gen_methods_131:checked] estimate (+95% Cl, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Model 9	
Is this the recommended model?	\bigcirc Yes \bigcirc No (If there is only one model, please state 'Yes')
Development A discrimination measure is presented	 ○ Yes ○ No (E.g. C-index / area under the ROC curve)
External validation A discrimination measure is presented	 ○ Yes ○ No (E.g. C-index / area under the ROC curve)
Development Which performance measures are described?	 AUC/AUROC/Area under the curve C-statistic Harrell's c-index D-statistic (Multiples answers are possible)
External validation Which performance measures are described?	 AUC/AUROC/Area under the curve C-statistic Harrell's c-index D-statistic (Multiples answers are possible)
The confidence interval (or standard error) of the AUC/AUROC/Area under the curve is presented	⊖ Yes ⊖ No
The confidence interval (or standard error) of the AUC/AUROC/Area under the curve is presented	○ Yes ○ No
What was the AUC/AUROC/Area under the curve apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the AUC/AUROC/Area under the curve apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the AUC/AUROC/Area under the curve corrected for optimism?	 ○ Yes ○ No ○ Uncloar

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What was the optimism corrected AUC/AUROC/Area under the curve (95% CI, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the C-statistic is presented	○ Yes ○ No
The confidence interval (or standard error) of the C-statistic is presented	○ Yes ○ No
What was the C-statistic apparent discrimination estimate (+95%Cl, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the C-statistic apparent discrimination estimate (+95%CL if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the C-statistic corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected C-statistic (95% CI,	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the Harrel's c-index is presented	○ Yes ○ No
The confidence interval (or standard error) of the Harrel's c-index is presented	○ Yes ○ No
What was the Harrel's c-index apparent discrimination estimate (+95%Cl, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the Harrel's c-index apparent discrimination estimate (+95%Cl. if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the Harrel's c-index corrected for optimism?	○ Yes ○ No

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What was the optimism corrected Harrel's c-index (95%	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the D-statistic is presented	○ Yes ○ No
The confidence interval (or standard error) of the D-statistic is presented	⊖ Yes ⊖ No
What was the D-statistic apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the D statistic apparent discrimination	
estimate (+95%Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the D-statistic corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected D-statistic (95% CI,	
if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Development Measures for model calibration are described	 Yes No (E.g. calibration plot, calibration slope or intercept, calibration table, Hosmer Lemeshow test, O/E ratio.)
External validation Measures for model calibration are described	 Yes No (E.g. calibration plot, calibration slope or intercept, calibration table, Hosmer Lemeshow test, O/E ratio.)
Development How was calibration assessed?	 Calibration plot Calibration slope Calibration intercept Calibration in the large Calibration table Kappa Observed/expected ratio H-L

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External validation How was calibration assessed?	 Calibration plot Calibration slope Calibration intercept Calibration in the large Calibration table Kappa Observed/expected ratio H-L
What was the apparent [gen_methods_155:checked] estimate (+95% Cl, if given)?	(Please use the following format - estimate (lor CI - upper CI). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
What was the apparent [gen_methods_166:checked] estimate (+95% Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Was the [gen_methods_155:checked] measure corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected [gen_methods_155:checked] estimate (+95% Cl, if given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Development Other model performance measures are presented	 Yes No (E.g. R2, Brier score, predictive values, sensitivity, specificity, AUC difference, decision curve analysis, net reclassification improvement, integrated discrimination improvement, AIC.)
External validation Other model performance measures are presented	 Yes No (E.g. R2, Brier score, predictive values, sensitivity, specificity, AUC difference, decision curve analysis, net reclassification improvement, integrated discrimination improvement, AIC.)





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Development If yes, please specify	 R2 Brier score predictive values sensitivity specificity AUC difference decision curve analysis net reclassification improvement integrated discrimination improvement AIC Accuracy Other (Multiples answers are possible)
External validation If yes, please specify	 R2 Brier score predictive values sensitivity specificity AUC difference decision curve analysis net reclassification improvement integrated discrimination improvement AIC Accuracy Other (Multiples answers are possible)
If other, please specify	
	(list using (;) to separate if more than 1.)
If other, please specify	
	(list using (;) to separate if more than 1.)
What was the apparent [gen_methods_132:checked] estimate (+95% Cl, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
What was the apparent [gen_methods_135:checked] estimate (+95% CI, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Was the [gen_methods_132:checked] measure corrected for optimism?	 ○ Yes ○ No ○ Unclear

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What was the optimism corrected [gen_methods_132:checked] estimate (+95% Cl, if given)?	
g	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Model 10	
Is this the recommended model?	 Yes No (If there is only one model, please state 'Yes')
Development A discrimination measure is presented	 ○ Yes ○ No (E.g. C-index / area under the ROC curve)
External validation A discrimination measure is presented	 ○ Yes ○ No (E.g. C-index / area under the ROC curve)
Development Which performance measures are described?	 AUC/AUROC/Area under the curve C-statistic Harrell's c-index D-statistic (Multiples answers are possible)
External validation Which performance measures are described?	 AUC/AUROC/Area under the curve C-statistic Harrell's c-index D-statistic (Multiples answers are possible)
The confidence interval (or standard error) of the AUC/AUROC/Area under the curve is presented	○ Yes ○ No
The confidence interval (or standard error) of the AUC/AUROC/Area under the curve is presented	○ Yes ○ No
What was the AUC/AUROC/Area under the curve apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the AUC/AUROC/Area under the curve apparent discrimination estimate (+95%Cl, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the AUC/AUROC/Area under the curve corrected for optimism?	 ○ Yes ○ No ○ Unclear

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What was the optimism corrected AUC/AUROC/Area under the curve (95% CL if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the C-statistic is presented	○ Yes ○ No
The confidence interval (or standard error) of the C-statistic is presented	○ Yes ○ No
What was the C-statistic apparent discrimination	
	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the C-statistic apparent discrimination	
	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the C-statistic corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected C-statistic (95% CI,	
ir given)?	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the Harrel's c-index is presented	○ Yes ○ No
The confidence interval (or standard error) of the Harrel's c-index is presented	○ Yes ○ No
What was the Harrel's c-index apparent discrimination	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the Harrel's c-index apparent discrimination	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the Harrel's c-index corrected for optimism?	⊖ Yes ⊖ No

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What was the optimism corrected Harrel's c-index (95% CI, if given)?	(Diasco uso the following formation optimate (low
	Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
The confidence interval (or standard error) of the D-statistic is presented	○ Yes ○ No
The confidence interval (or standard error) of the D-statistic is presented	○ Yes ○ No
What was the D-statistic apparent discrimination estimate (+95%Cl, if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
What was the D-statistic apparent discrimination estimate (+95%CL if given)?	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Was the D-statistic corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected D-statistic (95% CI,	
	(Please use the following format - estimate (low Cl - upper Cl). For example 0.79 (0.75 - 0.81). If not reported fill this with "NR")
Development Measures for model calibration are described	○ Yes ○ No
	(E.g. calibration plot, calibration slope or intercept, calibration table, Hosmer Lemeshow test, O/E ratio.)
External validation Measures for model calibration are described	○ Yes ○ No
	(E.g. calibration plot, calibration slope or intercept, calibration table, Hosmer Lemeshow test, O/E ratio.)
Development How was calibration assessed?	Calibration plot Calibration slope
	 Calibration intercept Calibration in the large
	 ☐ Calibration table ☐ Kappa
	Observed/expected ratio

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External validation How was calibration assessed?	 Calibration plot Calibration slope Calibration intercept Calibration in the large Calibration table Kappa Observed/expected ratio H-L
What was the apparent [gen_methods_167:checked] estimate (+95% Cl, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
What was the apparent [gen_methods_168:checked] estimate (+95% Cl, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Was the [gen_methods_167:checked] measure corrected for optimism?	 ○ Yes ○ No ○ Unclear
What was the optimism corrected [gen_methods_167:checked] estimate (+95% Cl, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Development Other model performance measures are presented	 Yes No (E.g. R2, Brier score, predictive values, sensitivity, specificity, AUC difference, decision curve analysis, net reclassification improvement, integrated discrimination improvement, AIC.)
External validation Other model performance measures are presented	 Yes No (E.g. R2, Brier score, predictive values, sensitivity, specificity, AUC difference, decision curve analysis, net reclassification improvement, integrated discrimination improvement, AIC.)





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Development If yes, please specify	 R2 Brier score predictive values sensitivity specificity AUC difference decision curve analysis net reclassification improvement integrated discrimination improvement AIC Accuracy Other (Multiples answers are possible)
External validation If yes, please specify	 R2 Brier score predictive values sensitivity specificity AUC difference decision curve analysis net reclassification improvement integrated discrimination improvement AIC Accuracy Other (Multiples answers are possible)
If other, please specify	
	(list using (;) to separate if more than 1.)
If other, please specify	
	(list using (;) to separate if more than 1.)
What was the apparent [gen_methods_133:checked] estimate (+95% Cl, if given)?	
	(Please use the following format - estimate (lowe Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
What was the apparent [gen_methods_134:checked] estimate (+95% Cl, if given)?	
	(Please use the following format - estimate (lowe Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
Was the [gen_methods_133:checked] measure corrected for optimism?	 ○ Yes ○ No ○ Unclear



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What was the optimism corrected [gen_methods_133:checked] estimate (+95% Cl, if given)?	(Please use the following format - estimate (lov Cl - upper Cl). For example 0.79 (0.75 - 0.81). List using (;) if more than 1. If not reported, fill this with "NR")
SPIN	
Is there use of leading words/strong statement in the results section to describe model and/or model performance/accuracy/effectiveness?	 Yes No (The prediction estimate is described with a va judgement like "statistically significant", "significant")
If yes, please specify the leading word/strong statement	 Novel Excellent Accurate Optimal Perfect Significant Other
If other, please specify	(list using (;) to separate if more than 1)
If yes, please copy the statement below	
ls at least ONE non-significant/non relevant model reported?	 Yes No NA (Select NA when only 1 model was developed)
If yes, did the authors make use of leading word to reject those non predictive models reported?	 Yes No (E.g. The effect is said to be significant, although the 95% confidence interval of the adjusted odds ratio crosses 1; OR Words like "trend" or "borderline, "significance", "statistically significant" are used)
If yes, please copy the statement below	
Is the prediction model defined in multiples ways?	 Yes No (E.g. Different thresholds of categorization AND continuous, or absolute value and relative value





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Presence of spin in the presentation of tables and figures	 Yes No (E.g. Non-significant p values adjusted for multiple comparisons are written below the tak whereas significant unadjusted p values are highly visible)
Model Updating	
The updated regression coefficients for each predictor in the model are reported	 Yes No NA (If model updating was described as 'not neede score Yes)
The updated intercept or cumulative baseline hazard or baseline survival (for at least one time point) is reported	 Yes No NA (If model updating was described as 'not needed score Yes)
The discrimination of the updated model is reported	○ Yes ○ No
The confidence interval (or standard error) of the discrimination measure of the updated model is reported	○ Yes ○ No
The calibration of the updated model is reported	○ Yes ○ No
The confidence interval (or standard error) of the calibration measure of the updated model is reported	○ Yes ○ No

Any additional comment about the results section of this article?

(If there is something in the "Results" section that does not fit into the questions of this form - please use this space to detail. Also use this space to detail anything you are unsure about.)



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Discussion / Conclusion

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 Yes No (Discuss any limitations of the study is sufficient (such as non-representative sample, few events per predictor, missing data).)
○ Yes ○ No
 Yes No Some outcomes in favour and not in favour for others Unclear
 Yes No (Considering objectives, limitations, results from similar studies, and other relevant evidence.)
○ Yes ○ No
○ Yes ○ No
 Novel Excellent Accurate Optimal Perfect Significant Other (Multiples answers are possible)
(list using (;) to separate if more than 1)



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	Page 2 c
ls at least (1) non-significant/non relevant model discussed?	 ○ Yes ○ No ○ NA (Select NA when only 1 model was developed)
Implications	
The potential clinical use is discussed	 Yes No (An explicit description of the context in which the prediction model is to be used (e.g. to identify high risk groups to help direct treatment, or to triage patients for referral to subsequent care).)
Does the recommendation include using the model in a difference clinical setting/population?	○ Yes ○ No
If yes, please provide the statement below	
Implications for future research are discussed	 Yes No (E.g. a description of what the next stage of investigation of the prediction model should be such as "We suggest further external validation
If yes, please provide the statement below	
Were there any other perspectives addressed in the discussion section?	 Yes No (E.g. unexpected finding from the analyses.)
If yes, please provide the statement below or a short description	
Is uncertainty reported in the discussion?	 Yes No (The use of any verbs as "may" or "could", or a words as "likely to" or "maybe")
Is there any other misleading strategy reported in the discussion/conclusion section?	○ Yes ○ No



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Other Information	
Information about supplementary resources is provided	○ Yes ○ No
If yes, what type of supplementary material?	 Web calculator repository for dataset repository for R code extra results main results missing data predictors Protocol Details on statistics Other (Multiples answers are possible. Protocol inclu registered/published protocol as well.)
If other, please specify	
	(list using (;) to separate if more than 1)
Funding	
The source of funding is reported or there is explicit mention that there was no external funding involved	○ Yes ○ No
If yes, which type of funding?	 Profit Non-Profit Both Unclear
The role of funders is reported	 ○ Yes ○ No ○ NA (If there is no external funding, please select "NA")
Is there a "Disclosure of authors' potential conflicts of interest (COI)" section in the journal?	○ Yes ○ No
If yes, are COI reported?	○ Yes ○ No
If yes, how many authors declared COI?	
	(Number only the ones who declare having conflicts)
Any additional comment about the discussion/conclusion section of this article?	
	(If there is something in the "Discussion/conclusion" section that does not into the questions of this form - please use this space to detail. Also use this space to detail


Title	Page 1
Article ID	
The words developing/development, validation/validating, incremental/added value (or synonyms) are reported in the title	○ Yes ○ No
Is the title supportive of the clinical relevance of the model, despite the study reporting non-significant/relevant measures?	\bigcirc Yes \bigcirc No (Title is inconsistent with the study results)
Is there use of leading words/strong statement in the title to describe model and/or model performance?	 Yes No (E.g. Novel, excellent, acurrate, significant, promising, breakthrough, etc.)
If yes, please select the leading word(s)/strong statement(s) used	 Novel Excellent Accurate Optimal Perfect Significant Other (Multiples answers are possible)
If other, please specify	
	(list using (;) to separate if more than 1)
The words prediction, risk prediction, prediction model, risk models, prognostic models, prognostic indices, risk scores (or synonyms) are reported in the title	○ Yes ○ No
The target population is reported in the title	○ Yes ○ No
The outcome to be predicted is reported in the title	○ Yes ○ No

(If there is something in the "Title" section that does not fit into the questions of this form please use this space to detail. Also use this space to detail anything you are unsure about.)



Abstract

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Article ID	
The objectives are reported in the abstract	○ Yes ○ No
Sources of data are reported in the abstract	 Yes No (E.g. Prospective cohort, registry data, RCT data.)
The setting is reported in the abstract	 Yes No (E.g. Primary care, secondary care, general population, adult care, or paediatric care. The setting should be reported for both the development and validation datasets, if applicable.)
A general definition of the study participants is reported in the abstract	 Yes No (E.g. patients with suspicion of certain disease, patients with a specific disease, or general eligibility criteria.)
The overall sample size is reported in the abstract	○ Yes ○ No
The number of events (or % outcome together with overall sample size) is reported in the abstract	 Yes No NA (If a continuous outcome was studied, score Not applicable)
Predictors included in the final model are reported in the abstract. For validation studies of well-known models, at least the name/acronym of the validated model is reported	 Yes No (Broad descriptions are sufficient, e.g. 'all information from patient history and physical examination'. Check in the main text whether all predictors of the final model are indeed reported in the abstract.)
The outcome is reported in the abstract	⊖ Yes ⊖ No
Statistical methods are described in the abstract	 Yes No (For model development, at least the type of statistical model should be reported. For validation studies a quote like "model's discrimination and calibration was assessed" is considered adequate. If done, methods of updating should be reported.)
ML techniques that will be used are reported in the abstract	○ Yes ○ No



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Results for model discrimination are reported in the abstract	 Yes No (This should be reported separately for development and validation if a study includes both development and validation.)
If yes, which measures are described in the abstract?	
	(list the names using (;) to separate if more than 1)
If yes, are confidence interval reported?	○ Yes ○ No
Results for model calibration are reported in the abstract	 Yes No (This should be reported separately for development and validation if a study includes both development and validation.)
If yes, which measures are described in the abstract?	
	(list the names using (;) to separate if more than 1)
If yes, are precision estimates reported?	○ Yes ○ No
Is there any other type of measures reported?	○ Yes ○ No
If yes, please specify	
	(list the names using (;) to separate if more than 1)
Conclusions are reported in the abstract	 Yes No (In publications addressing both model developmer and validation, there is no need for separate conclusions for both; one conclusion is sufficient.)
If yes, are the conclusion consistent with the reported results in the abstract section?	○ Yes ○ No
Does the conclusion statement focus solely on significant/relevant results?	 Yes No (absence of non signicant results reports)
Is there emphasis on model relevance in the conclusion section of the abstract while there is not enough information given to concluded results are predictive?	○ Yes ○ No

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	Page 3 o
Is there discrepancies between the full-text (discussion) and abstract (conclusion) explanations of the study findings?	\bigcirc Yes \bigcirc No (The discussion is consistent with the study findings, whereas the abstract conclusion is no [+/-]; OR The discussion is not consistent with the study findings, whereas the abstract conclusion is [-/+])
If yes, please copy the statement below	
What is the recommended next step for the prediction model?	 To be used in clinical practice Validate the models in a different setting/population Other recommendations for further studies None reported (Multiples answers are possible)
If other recommendations, please specify	
	(list using (;) to separate if more than 1)
Is there any reference to previous prediction model in literature in the abstract section?	○ Yes ○ No
If yes, please copy the statement below	
Is there use of leading words/strong statement in the abstract section to describe model and/or model performance/accuracy/effectiveness?	 Yes No (E.g. Novel, excellent, acurrate, significant, promising, breakthrough, etc.)
If yes, which leading words/strong statement?	 Novel Excellent Accurate Optimal Perfect Significant Other (Multiples answers are possible)
If other, please specify	
	(list using (;) to separate if more than 1)
Please copy the statement below	
Is uncertainty reported in the abstract?	 Yes No (The use of any verbs as "may" or "could", nor words as "likely to" or "maybe")
Limitations are reported in the abstract	⊖ Yes ⊖ No
	⊖ No

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Please copy the statement below

Any additional comment about the "abstract" section of this article?

(If there is something in the "Abstract" section that does not fit into the questions of this form - please use this space to detail. Also use this space to detail anything you are unsure about.)



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Confi	dential

Probast

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Article ID

Instructions When assessing ROB using PROBAST, please refer to the "best performance" model for the primary outcome suggested by the authors

What type of study is reported?	 Development (including internal validation) Development with external validation (same model) Development with external validation (different model) External validation only (This questions is repeated due to branching logic. Please answer again.)
Domain 1 : Participants	
Development Were appropriate data sources used, e.g. cohort, RCT or nested case-control study data?	 Yes / Probably yes No / Probably no No information
External validation Were appropriate data sources used, e.g. cohort, RCT or nested case-control study data?	 Yes / Probably yes No / Probably no No information
Development Were all inclusions and exclusions of participants appropriate?	 Yes / Probably yes No / Probably no No information (Yes/probably yes: If inclusion and exclusion of participants was appropriate, so participants correspond to unselected participants of interest. No/probably no: If participants are included who would already have been identified as having the outcome and so are no longer participants at suspicion of disease (diagnostic studies) or at risk of developing outcome (prognostic studies), or if specific subgroups are excluded that may have altered the performance of the prediction model for the intended target population. No information: When there is no information on whether inappropriate inclusions or exclusions took place.)



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External validation Were all inclusions and exclusions of participants appropriate?	 Yes / Probably yes No / Probably no No information (Yes/probably yes: If inclusion and exclusion of participants was appropriate, so participants correspond to unselected participants of interest. No/probably no: If participants are included who would already have been identified as having the outcome and so are no longer participants at suspicion of disease (diagnostic studies) or at risk of developing outcome (prognostic studies), or if specific subgroups are excluded that may have altered the performance of the prediction model for the intended target population. No information: When there is no information on whether inappropriate inclusions or exclusions took place.)
Development Risk of bias introduced by Participants	 Low ROB High ROB Unclear ROB (Low risk of bias: If the answer to all signaling questions is "Yes" or "Probably yes," then risk of bias can be considered low. If ≥1 of the answers is "No" or "Probably no," the judgment could still be "Low risk of bias" but specific reasons should be provided why the risk of bias can be considered low. High risk of bias: If the answer to any of the signaling questions is "No" or "Probably no," there is a potential for bias, except if defined at low risk of bias above. Unclear risk of bias: If relevant information is missing for some of the signaling questions is judged to put this domain at high risk of bias.)
External validation Risk of bias introduced by Participants	 Low ROB High ROB Unclear ROB (Low risk of bias: If the answer to all signaling questions is "Yes" or "Probably yes," then risk of bias can be considered low. If ≥1 of the answers is "No" or "Probably no," the judgment could still be "Low risk of bias" but specific reasons should be provided why the risk of bias can be considered low. High risk of bias: If the answer to any of the signaling questions is "No" or "Probably no," there is a potential for bias, except if defined at low risk of bias above. Unclear risk of bias: If relevant information is missing for some of the signaling questions and none of the signaling questions is judged to put this domain at high risk of bias.)
Development Support for Judgement	
External validation Support for Judgement	

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Domain 2 : Predictors	
Development Were predictors defined and assessed in a similar way for all participants?	 Yes / Probably yes No / Probably No No information (Yes/probably yes: If definitions of predictors and their assessment were similar for all participants. No/probably no: If different definitions were used for the same predictor or if predictors requiring subjective interpretation were assessed by differently experienced assessors. No information: If there is no information on how predictors were defined or assessed.)
External validation Were predictors defined and assessed in a similar way for all participants?	 Yes / Probably yes No / Probably No No information (Yes/probably yes: If definitions of predictors and their assessment were similar for all participants. No/probably no: If different definitions were used for the same predictor or if predictors requiring subjective interpretation were assessed by differently experienced assessors. No information: If there is no information on how predictors were defined or assessed.)
Development Were predictor assessments made without knowledge of outcome data?	 Yes / Probably yes No / Probably no No information
External validation Were predictor assessments made without knowledge of outcome data?	 Yes / Probably yes No / Probably no No information
Development Are all predictors available at the time the model is intended to be used?	 Yes / Probably Yes No / Probably No No information
External validation Are all predictors available at the time the model is intended to be used?	 Yes / Probably Yes No / Probably No No information



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Development Risk of bias introduced by predictors	 Low ROB High ROB Unclear ROB (Low risk of bias: If the answer to all signaling questions is "Yes" or "Probably Yes," then risk of bias can be considered low. If ≥1 of the answers is "No" or "Probably no," the judgment could still be "Low risk of bias" but specific reasons should be provided why the risk of bias can be considered low, e.g., use of objective predictors not requiring subjective interpretation. High risk of bias: If the answer to any of the signaling questions is "No" or "Probably no," there is a potential for bias. Unclear risk of bias: If relevant information is missing for some of the signaling questions is judged to put the domain at high risk of bias.)
External validation Risk of bias introduced by predictors	 Low ROB High ROB Unclear ROB (Low risk of bias: If the answer to all signaling questions is "Yes" or "Probably Yes," then risk of bias can be considered low. If ≥1 of the answers is "No" or "Probably no," the judgment could still be "Low risk of bias" but specific reasons should be provided why the risk of bias can be considered low, e.g., use of objective predictors not requiring subjective interpretation. High risk of bias: If the answer to any of the signaling questions is "No" or "Probably no," there is a potential for bias. Unclear risk of bias: If relevant information is missing for some of the signaling questions is judged to put the domain at high risk of bias.)
Development Support for Judgement	
External validation Support for Judgement	
Domain 3 : Outcome	
Development Was the outcome determined appropriately?	 Yes / Probably yes No / Probably no No information
External validation Was the outcome determined appropriately?	 Yes / Probably yes No / Probably no No information
Development Was a pre-specified or standard outcome definition used?	 Yes / Probably yes No / Probably no No information

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External validation Was a pre-specified or standard outcome definition used?	 Yes / Probably yes No / Probably no No information
Development Were predictors excluded from the outcome definition?	 Yes / Probably yes No / Probably no No information
External validation Were predictors excluded from the outcome definition?	 Yes / Probably yes No / Probably no No information
Development Was the outcome defined and determined in a similar way for all participants?	 Yes / Probably yes No / Probably no No information
External validation Was the outcome defined and determined in a similar way for all participants?	 Yes / Probably yes No / Probably no No information
Development Was the outcome determined without knowledge of predictor information?	 Yes / Probably yes No / Probably no No information
External validation Was the outcome determined without knowledge of predictor information?	 Yes / Probably yes No / Probably no No information
Development Was the time interval between predictor assessment and outcome determination appropriate?	 Yes / Probably yes No / Probably no No information
External validation Was the time interval between predictor assessment and outcome determination appropriate?	 Yes / Probably yes No / Probably no No information
Development Risk of bias introduced by the outcome	 Low ROB High ROB Unclear ROB Unclear ROB (Low risk of bias: If the answer to all signaling questions is "Yes" or "Probably yes," then risk of bias can be considered low. If ≥1 of the answers is "No" or "Probably no," the judgment could still be low risk of bias, but specific reasons should be provided why the risk of bias can be considered low, e.g., when the outcome was determined with knowledge of predictor information but the outcome assessment did not require much interpretation by the assessor (e.g., death regardless of cause). High risk of bias: If the answer to any of the signaling questions is "No" or "Probably no," there is a potential for bias. Unclear risk of bias: If relevant information about the outcome is missing for some of the signaling questions and none of the signaling questions is judged to put this domain at high risk of bias.)

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External validation Risk of bias introduced by the outcome	 Low ROB High ROB Unclear ROB (Low risk of bias: If the answer to all signaling questions is "Yes" or "Probably yes," then risk of bias can be considered low. If ≥1 of the answers is "No" or "Probably no," the judgment could still be low risk of bias, but specific reasons should be provided why the risk of bias can be considered low, e.g., when the outcome was determined with knowledge of predictor information but the outcome assessment did not require much interpretation by the assessor (e.g., death regardless of cause). High risk of bias: If the answer to any of the signaling questions is "No" or "Probably no," there is a potential for bias. Unclear risk of bias: If relevant information about the outcome is missing
	relevant information about the outcome is missing for some of the signaling questions and none of the signaling questions is judged to put this

Development Support for Judgement

External validation Support for Judgement

Domain 4 : Analysis

Development Were there a reasonable number of participants with the outcome?

○ Yes / Probably yes ○ No / Probably no \bigcirc No information (Yes/probably yes: For model development studies, if the number of participants with the outcome relative to the number of candidate predictor parameters is ≥ 20 (EPV ≥ 20).* For model validation studies, if the number of participants with the outcome is ≥ 100 . No/probably no: For model development studies, if the number of participants with the outcome relative to the number of candidate predictor parameters is < 10(EPV < 10).* For model validation studies, if the number of participants with the outcome is < 100. No information: For model development studies, no information on the number of candidate predictor parameters or number of participants with the outcome, such that the EPV cannot be calculated. For model validation studies, no information on the number of participants with the outcome.)

domain at high risk of bias.)





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External validation Were there a reasonable number of participants with the outcome?	○ Yes / Probably yes ○ No / Probably no ○ No information (Yes/probably yes: For model development studies, if the number of participants with the outcome relative to the number of candidate predictor parameters is ≥20 (EPV ≥20).* For model validation studies, if the number of participants with the outcome is ≥100. No/probably no: For model development studies, if the number of participants with the outcome relative to the number of candidate predictor parameters is < 10 (EPV < 10).* For model validation studies, if the number of participants with the outcome is < 100. No information: For model development studies, no information on the number of candidate predictor parameters or number of participants with the outcome, such that the EPV cannot be calculated. For model validation studies, no information on the number of participants with the outcome.)
Development Were continuous and categorical handled appropriately?	 Yes / Probably yes No / Probably no No information (Yes/probably yes: If continuous predictors are not converted into ≥2 categories when included in the model (i.e., dichotomized or categorized), or if continuous predictors are examined for nonlinearity using, for example, fractional polynomials or restricted cubic splines, or if categorical predictor groups are defined using a prespecified method. For model validation studies, if continuous predictors are included using the same definitions or transformations, and categorical variables are categorized using the same cut points, as compared with the development study. No/probably no: If categorical predictor group definitions do not use a prespecified method. For model development studies, if continuous predictors are converted into ≥2 categories when included in the model. For model validation studies, if continuous predictors are converted into ≥2 categories when included in the model. For model validation studies, or categorical variables are categorical variables are converted into ≥2 categories when included in the model. For model validation studies, if continuous predictors are converted into ≥2 categorized using different definitions or transformations, or categorical variables are categorized using different cut points, as compared with the development study. No information: No information on whether continuous predictors are examined for nonlinearity and no information on how categorical predictor groups are defined. For model validation studies, no information on whether the same definitions or transformations on the model validation studies, no information on whether the same definitions or transformations and the same cut points are used, as compared with the development study.)

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External validation Were continuous and categorical handled appropriately?	 Yes / Probably yes No / Probably no No information (Yes/probably yes: If continuous predictors are not converted into ≥2 categories when included in the model (i.e., dichotomized or categorized), or if continuous predictors are examined for nonlinearity using, for example, fractional polynomials or restricted cubic splines, or if categorical predictor groups are defined using a prespecified method. For model validation studies, if continuous predictors are included using the same definitions or transformations, and categorical variables are categorized using the same cut points, as compared with the development study. No/probably no: If categorical predictor group definitions do not use a prespecified method. For model development studies, if continuous predictors are converted into ≥2 categories when included in the model. For model validation studies, if continuous predictors are converted into ≥2 categories when included in the model. For model validation studies, if continuous predictors are converted into ≥2 categories when included in the model. For model validation studies, if continuous predictors are converted into ≥2 categories when included in the model. For model validation studies, if continuous predictors are converted into ≥2 categories when included in the model. No information: No information on whether continuous predictors are examined for nonlinearity and no information on whether continuous predictors are examined for nonlinearity and no information on whether the same definitions or transformations and the same cut points are used, as compared with the development study.)
Development Were enrolled participants included in the analysis?	 Yes / Probably yes No / Probably no No information
External validation Were enrolled participants included in the analysis?	 Yes / Probably yes No / Probably no No information
Development Were participants with missing data handled appropriately?	 Yes / Probably yes No / Probably no No information
External validation Were participants with missing data handled appropriately?	 Yes / Probably yes No / Probably no No information
Development Was selection of predictors based on univariable analysis avoided?	 Yes / Probably yes No / Probably no No information
Development Were complexities in the data (e.g., censoring, competing risks, sampling of control participants) accounted appropriately?	 Yes / Probably yes No / Probably no No information

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External validation Were complexities in the data (e.g., censoring, competing risks, sampling of control participants) accounted appropriately?	 Yes / Probably yes No / Probably no No information
Development Were relevant model performance measures evaluated appropriately?	 Yes / Probably yes No / Probably no No information (Yes/probably yes: If both calibration and discrimination are evaluated appropriately (including relevant measures tailored for mode predicting survival outcomes). No/probably no both calibration and discrimination are not evaluated, or if only goodness-of-fit tests, such as the Hosmer-Lemeshow test, are used to eva calibration, or if for models predicting survival outcomes performance measures accounting f censoring are not used, or if classification measures (like sensitivity, specificity, or predictive values) were presented using predict probability thresholds derived from the data set at hand. No information: Either calibration or discrimination are not reported, or no information is provided as to whether appropriate performance measures for survival outcomes as used (e.g., references to relevant literature or specific mention of methods, such as using Kaplan-Meier estimates), or no information on thresholds for estimating classification measures is given.)
External validation Were relevant model performance measures evaluated appropriately?	 Yes / Probably yes No / Probably no No information (Yes/probably yes: If both calibration and discrimination are evaluated appropriately (including relevant measures tailored for mode predicting survival outcomes). No/probably no both calibration and discrimination are not evaluated, or if only goodness-of-fit tests, such as the Hosmer-Lemeshow test, are used to eva calibration, or if for models predicting survival outcomes performance measures accounting f censoring are not used, or if classification measures (like sensitivity, specificity, or predictive values) were presented using predicting robability thresholds derived from the data set at hand. No information: Either calibration or discrimination are not reported, or no information is provided as to whether approprime performance measures for survival outcomes are used (e.g., references to relevant literature or specific mention of methods, such as using Kaplan-Meier estimates), or no information on thresholds for estimating classification measures is given.)
Development Were model overfitting, under-fitting, and optimism in model performance accounted for?	 Yes / Probably yes No / Probably no No information

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 Yes / Probably yes No / Probably no No information
 Yes / Probably yes No / Probably no No information
 Yes / Probably yes No / Probably no No information
 Low ROB High ROB Unclear ROB (Low risk of bias: If the answer to all signaling questions is "Yes" or "Probably yes," then risk of bias can be considered low. If ≥1 of the answers is "No" or "Probably no," the judgment could still be low risk of bias, but specific reasons should be provided why the risk of bias: can be considered low. High risk of bias: If the answer to any of the signaling questions is "No or "Probably no," there is a potential for bias. Unclear risk of bias: If relevant information about the analysis is missing for some of the signaling question answers is judged to put the analysis high risk of bias)
 Low ROB High ROB Unclear ROB Unclear ROB (Low risk of bias: If the answer to all signaling questions is "Yes" or "Probably yes," then risk of bias can be considered low. If ≥1 of the answers is "No" or "Probably no," the judgment could still be low risk of bias, but specific reasons should be provided why the risk of bias: can be considered low. High risk of bias: If the answer to any of the signaling questions is "No or "Probably no," there is a potential for bias. Unclear risk of bias: If relevant information about the analysis is missing for some of the signaling question answers is judged to put the analysis high risk of bias)



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Overall assessment of ROB	
Development Overall risk of bias	 Low risk of bias High risk of bias Unclear risk of bias (Low ROB: If all domains were rated low risk of bias. If a prediction model was developed without any external validation, and it was rated as low risk of bias for all domains, consider downgrading to high risk of bias. Such a model evaluation can only be considered as low risk of bias, if the development was based on a very large data set and included some form of internal validation. High ROB: If ≥1 domain is judged to be at high risk of bias. Unclear ROB: If an unclear risk of bias was noted in ≥1 domain and it was low risk for all other domains.)
External validation Overall risk of bias	 Low risk of bias High risk of bias Unclear risk of bias Unclear risk of bias (Low ROB: If all domains were rated low risk of bias. If a prediction model was developed without any external validation, and it was rated as low risk of bias for all domains, consider downgrading to high risk of bias. Such a model evaluation can only be considered as low risk of bias, if the development was based on a very large data set and included some form of internal validation. High ROB: If ≥1 domain is judged to be at high risk of bias. Unclear ROB: If an unclear risk of bias was noted in ≥1 domain and it was low risk for all other domains.)

Any additional comments about ROB on this article?

(If there is something in the "PROBAST" that does not fit into the questions of this form - please use this space to detail. Also use this space to detail anything you are unsure about.)

