

Title     1     D:V     Identify the study as developing and/or validating a multivariable prediction model, the target population, Abstract     Title       Abstract     2     D:V     Provide a summary of objectives, study design, setting, participants, sample size, predictors, outcome, statistical analysis, results, and conclusions.     Abstract       Introduction     Explain the medical context (including whether diagnostic or prognostic) and rationale for developing or validating the multivariable prediction model, including references to existing models.     Introduction, paragraph 1-4       Background and objectives     3a     D:V     Specify the objectives, including whether the study describes the development or validation of the model or both.     Introduction, paragraph 1-4       We therefore aimed to investigate if urinary LAM detection, along with other clinical variables readily available in high-burden settings, could be used to pedictive patients admitted to hospital and diagnosed with TB patients were at high risk of early mortality in, and to externally validate the predictive tool."       Methods     4a     D:V     Describe the study design or source of data (e.g., randomized trial, cohort, or registry data), separately for the development and validation data sets, if applicable.     Methods, paragraph 1 (development), paragraph 10 (validation)       Source of data     5a     D:V     Specify the key study dates, including start of accrual; and, if applicable, end of follow- up.     Methods, paragraph 1 (development	Section/Topic	Item		Checklist Item	Page
Title     1     UV     and the outcome to be predicied.     Title       Abstract     2     D/V     Provide a summary of objectives, study design, setting, participants, sample size, predictors, outcome, and the outcome to be predicide.     Abstract     Abstract       Inroduction     3a     D/V     Explain the medical context (including whether diagnostic or prognostic) and rationale for developing or validating the multivariable prediction model, including references to existing models.     Introduction, paragraph 14       Background and objectives     3b     D/V     Specify the objectives, including whether the study describes the development or validation of the model or oboth.     Wet therefore sime do investigate if urinary LAM detection, along with other clinical variables readily araitable in high-hurden stering, could be tend to or oboth.       Methods     For the development and validation data sets, if applicable.     Methods, paragraph 1 (development), paragraph 10 (validate).       Source of data     4a     D/V     Describe the study design or source of data (e.g., randomized trial, cohort, or registry data), separately (validation).     Methods, paragraph 1 (development), paragraph 11 (LAM RCT recriment occurred between January 2013 and October 2015 and September 2017) and Methods, paragraph 1 (cobote 2015 and September 2017) and Methods, paragraph 1 (cobote 2015 and September 2017) and Methods, paragraph 1 (cobote 2015).       Participants     5b     D/V     Specify t	Title and abstract				
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Source of data     4     D:V     validating the multivariable prediction model, including references to existing models.     Introduction, paragraph 1       Background and objectives     3b     D:V     Specify the objectives, including whether the study describes the development or validation of the model     Introduction, paragraph 1       Methods	Introduction				
Background and objectives     Jb     D:V     Specify the objectives, including whether the study describes the development or validation of the model or both.     "We therefore aimed to investigate if urinary LAM detection, along with other clinical variables readily waiiable in high-burden settings, could be used to predict which HU-positive patients admitted to hospital and diagnosed with TB patients were at high risk of early mortality in, and to externally validate the predictive tool."       Methods     Describe the study design or source of data (e.g., randomized trial, cohort, or registry data), separately for the development, paragraph 1 (development), paragraph 1 (development), paragraph 1 (r the development and validation data sets, if applicable.     Methods, paragraph 1 (development), (validate the predictive tool."       Source of data     4a     D:V     Specify the key study dates, including start of accrual; and if applicable, end of follow- up.     Methods, paragraph 1 (development), paragraph 1 (Development), paragraph 11 (LAM RCT recruitment occurred between January 2013 and October 2014, and the MSF cohor to between October 2013 and August 2015.)       Participants     5a     D:V     including number and location of centres.     Methods, paragraph 1 (hoctob, paragraph 1 (Methods, paragraph 3)     Methods, paragraph 3       Outcome     6a     D:V     Clearly define the outcome that is predicted by the prediction model, including how and when assessed.     Methods, paragraph 3.     Methods, paragraph 3.       Predictors     6b     D:V		3a	D;V	Explain the medical context (including whether diagnostic or prognostic) and rationale for developing or validating the multivariable prediction model, including references to existing models.	Introduction, paragraphs 1-4
background and objectives     3b     D <sub>V</sub> Specify the objectives, including whether the study describes the development or validation of the model or both.     detection, along with other clinical variables readily predict which HIV-positive patients admitted to predict which HIV-positive patients admitted to predict which HIV-positive patients were at high risk of early mortality in, and to externally validate the predictive tool."       Methods     A     D <sub>V</sub> Describe the study design or source of data (e.g., randomized trial, cohort, or registry data), separately for the development and validation data sets, if applicable.     Methods, paragraph 1 (development), paragraph 10 (validation)       Source of data     4a     D <sub>V</sub> V     Specify the key study dates, including start of accrual; end of accrual; and, if applicable, end of follow- up.     Methods, paragraph 1 (October 2015 and September 2017) and Methods, paragraph 11 (LAM RCT recruitment occurred between January 2013 up.       Participants     5a     D <sub>V</sub> V     Specify key elements of the study setting (e.g., primary care, secondary care, general population) including number and location of centres.     Methods, paragraph 1 (hospital cohort)       Outcome     6a     D <sub>V</sub> V     Clearly define the outcome that is predicted by the prediction model, including how and when assessed.     Methods, paragraph 3 Methods, paragraph 3.       Outcome     6a     D <sub>V</sub> V     Report any actions to blind assessment of predictors were for predictors.     Methods, paragraph 2.3 Predictors wer					Introduction, paragraph 4
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analysis methods 100 D internal validation.		10a	D		Methods, paragraph 6-7
		10b	D		Methods, paragraph 6-7
		10c	V	For validation, describe how the predictions were calculated.	Methods, paragraph 12



	10d	D;V	Specify all measures used to assess model performance and, if relevant, to compare multiple models.	Methods, paragraph 8-9, 13
	10e	V	Describe any model updating (e.g., recalibration) arising from the validation, if done.	None done
Risk groups	11	D;V	Provide details on how risk groups were created, if done.	Methods, paragraph 7 "High-, medium- and low-risk groups for mortality were then arbitrarily defined after plotting risk score against observed mortality, so the high-risk group accounted for most (>50%) deaths, and low-risk group accounted for as few deaths as possible"
Development vs. validation	12	V	For validation, identify any differences from the development data in setting, eligibility criteria, outcome, and predictors.	Methods, paragraph 11
Results				
Participants	13a	D;V	Describe the flow of participants through the study, including the number of participants with and without the outcome and, if applicable, a summary of the follow-up time. A diagram may be helpful.	Results, paragraph 1; Figure 1
	13b	D;V	Describe the characteristics of the participants (basic demographics, clinical features, available predictors), including the number of participants with missing data for predictors and outcome.	Results, paragraph 1; Table 1
	13c	V	For validation, show a comparison with the development data of the distribution of important variables (demographics, predictors and outcome).	Table 1, Figure 4
Model	14a	D	Specify the number of participants and outcome events in each analysis.	Results, paragraph 1;, Figure 1
development	14b	D	If done, report the unadjusted association between each candidate predictor and outcome.	Table 2
Model	15a	D	Present the full prediction model to allow predictions for individuals (i.e., all regression coefficients, and model intercept or baseline survival at a given time point).	Table 2
specification	15b	D	Explain how to the use the prediction model.	Results, paragraph 4-6;
Model performance	16	D;V	Report performance measures (with CIs) for the prediction model.	Figures 3 & 4; Results, paragraph 4, 6, 8-9
Model-updating	17	V	If done, report the results from any model updating (i.e., model specification, model performance).	NA
Discussion				
Limitations	18	D;V	Discuss any limitations of the study (such as nonrepresentative sample, few events per predictor, missing data).	Discussion, paragraphs 9-10
Interpretation	19a	V	For validation, discuss the results with reference to performance in the development data, and any other validation data.	Discussion, paragraph 11
	19b	D;V	Give an overall interpretation of the results, considering objectives, limitations, results from similar studies, and other relevant evidence.	Discussion, paragraphs 1-6
Implications	20	D;V	Discuss the potential clinical use of the model and implications for future research.	Discussion, paragraphs 5-6
Other information				
Supplementary information	21	D;V	Provide information about the availability of supplementary resources, such as study protocol, Web calculator, and data sets.	Data statement, supplement
Funding	22	D;V	Give the source of funding and the role of the funders for the present study.	Financial Disclosure Statement

\*Items relevant only to the development of a prediction model are denoted by D, items relating solely to a validation of a prediction model are denoted by V, and items relating to both are denoted D;V. We recommend using the TRIPOD Checklist in conjunction with the TRIPOD Explanation and Elaboration document.