## TRIPOD Checklist: Prediction Model Development and Validation

| Section                   | Item |     | Checklist description  | Reported on Page<br>Number/Line<br>Number | Reported on<br>Section/Paragraph |
|---------------------------|------|-----|--|---|----------------------------------|
| Title and abstract        |      |     |  |   |                                  |
| Title                     | 1    | D;V | Identify the study as developing and/or validating a multivariable prediction model, the target population, and the outcome to be predicted.   | Page.1/Line.1-3                           | Title                            |
| Abstract                  | 2    | D;V | Provide a summary of objectives, study design, setting, participants, sample size, predictors, outcome, statistical analysis, results, and conclusions.  | Page.1-2/Line.27-56                       | Abstract                         |
| Introduction              |      |     |  |   |                                  |
| Background and objectives | 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. | Page.3-4/Line.77-114                      | Introduction/Paragraph.1-2       |
|                           | 3b   | D;V | Specify the objectives, including whether the study describes the development or validation of the model or both.  | Page.4/Line.115-119                       | Introduction/Paragraph.3         |
| Methods                   |      |     |  |   |                                  |
| Source of data            | 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, ifapplicable.                           | Page.4-6/Line.122-172                     | Methods/Paragraph.1-10           |
|                           | 4b   | D;V | Specify the key study dates, including start of accrual; end of accrual; and, if applicable, end of follow-up.   | N/A                                       | N/A                              |
| Participants              | 5a   | D;V | Specify key elements of the study setting (e.g., primary care, secondary care, general population) including number and location of centres.   | Page.4/Line.122-128                       | Methods/Paragraph.1-2            |
|                           | 5b   | D;V | Describe eligibility criteria for participants.  | N/A                                       | N/A                              |
|                           | 5c   | D;V | Give details of treatments received, if relevant.  | N/A                                       | N/A                              |
| Outcome                   | 6a   | D;V | Clearly define the outcome that is predicted by the prediction model, including how and when assessed.   | Page.4-5/Line.130-142                     | Methods/Paragraph.3-5            |
|                           | 6b   | D;V | Report any actions to blind assessment of the outcome to be predicted.   | N/A                                       | N/A                              |
| Predictors                | 7a   | D;V | Clearly define all predictors used in developing or validating the multivariable prediction model, including how and when they were measured.  | Page.5-6/L.144-172                        | Methods/Paragraph.6-10           |
|                           | 7b   | D;V | Report any actions to blind assessment of predictors for the outcome and other predictors.   | N/A                                       | N/A                              |
| Sample size               | 8    | D;V | Explain how the study size was arrived at.   | Page.4-6/Line.122-172                     | Methods/Paragraph.1-10           |

| Missing data                 | 9   | D;V | Describe how missing data were handled (e.g., complete-case analysis, single imputation, multiple imputation) with details of any imputation method.  | N/A                   | N/A                    |
|------------------------------|-----|-----|---|-----------------------|------------------------|
| Statistical analysis methods | 10a | D   | Describe how predictors were handled in the analyses.   | Page.6/L.174-176      | Methods/Paragraph.11   |
|                              | 10b | D   | Specify type of model, all model-building procedures (including any predictor selection), and method for internal validation.   | N/A                   | N/A                    |
|                              | 10c | V   | For validation, describe how the predictions were calculated.   | Page.6/L.174-176      | Methods/Paragraph.11   |
|                              | 10d | D;V | Specify all measures used to assess model performance and, if relevant, to compare multiple models.   | Page.6/L.174-176      | Methods/Paragraph.11   |
|                              | 10e | V   | Describe any model updating (e.g., recalibration) arising from the validation, if done.   | N/A                   | N/A                    |
| Risk groups                  | 11  | D;V | Provide details on how risk groups were created, if done.   | N/A                   | N/A                    |
| Development vs. validation   | 12  | V   | For validation, identify any differences from the development data in setting, eligibility criteria, outcome, and predictors.   | N/A                   | N/A                    |
| 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. | N/A                   | N/A                    |
|                              | 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.    | N/A                   | N/A                    |
|                              | 13c | V   | For validation, show a comparison with the development data of the distribution of important variables (demographics, predictors and outcome).  | N/A                   | N/A                    |
| Model<br>development         | 14a | D   | Specify the number of participants and outcome events in each analysis.   | N/A                   | N/A                    |
|                              | 14b | D   | If done, report the unadjusted association between each candidate predictor and outcome.  | N/A                   | N/A                    |
| Model<br>specification       | 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).                           | Page.6/Line.179-198   | Results/Paragraph.1-3  |
|                              | 15b | D   | Explain how to the use the prediction model.  | Page.6-9/Line.200-282 | Results/Paragraph.4-12 |
| Model<br>performance         | 16  | D;V | Report performance measures (with CIs) for the prediction model.  | N/A                   | N/A                    |
| Model-updating               | 17  | V   | If done, report the results from any model updating (i.e., model specification, model performance).   | N/A                   | N/A                    |
| Discussion                   |     |     |   | •                     | 1                      |
| Limitations                  | 18  | D;V | Discuss any limitations of the study (such as nonrepresentative sample, few events per predictor, missing data).  | Page.11/Line.346-353  | Conclusion             |
|                              |     |     |   | 1                     | 1                      |

| Interpretation            | 19a | V   | For validation, discuss the results with reference to performance in the development data, and any other validation data.                          | Page.11/Line.346-353 | Conclusion |  |  |  |
|---------------------------|-----|-----|--|----------------------|------------|--|--|--|
|                           | 19b | D;V | Give an overall interpretation of the results, considering objectives, limitations, and results from similar studies, and other relevant evidence. | Page.11/Line.346-353 | Conclusion |  |  |  |
| Implications              | 20  | D;V | Discuss the potential clinical use of the model and implications for future research.  | Page.11/Line.346-353 | Conclusion |  |  |  |
| Other information         |     |     |  |                      |            |  |  |  |
| Supplementary information | 21  | D;V | Provide information about the availability of supplementary resources, such as study protocol, Web calculator, and data sets.                      | N/A                  | N/A        |  |  |  |
| Funding                   | 22  | D;V | Give the source of funding and the role of the funders for the present study.  | N/A                  | N/A        |  |  |  |

<sup>\*</sup> 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.

Article information: https://dx.doi.org/10.21037/jtd-23-542

<sup>\*</sup>As the checklist was provided upon initial submission, the page number/line number reported may be changed due to copyediting and may not be referable in the published version. In this case, the section/paragraph may be used as an alternative reference.