## The REMARK checklist

| Item to be reported                                                                 | Reported on Page Number/Line Number | Reported on Section/Paragraph |
|-------------------------------------------------------------------------------------|------------------------------------|------------------------------|
| **INTRODUCTION**                                                                    |                                    |                              |
| 1 State the marker examined, the study objectives, and any pre-specified hypotheses. | page 6/ line 129-138              | Introduction/ para 3          |
| **MATERIALS AND METHODS**                                                           |                                    |                              |
| **Patients**                                                                        |                                    |                              |
| 2 Describe the characteristics (e.g., disease stage or co-morbidities) of the study patients, including their source and inclusion and exclusion criteria. | page 6-7/ line 141-151            | Patients and Methods - Patients and study design/ para 1 |
| 3 Describe treatments received and how chosen (e.g., randomized or rule-based).      | page 6/ line 143-145              | Patients and Methods          |
| **Specimen characteristics**                                                       |                                    |                              |
| 4 Describe type of biological material used (including control samples) and methods of preservation and storage. | page 7/ line 147-150              | Patients and Methods          |
| **Assay methods**                                                                  |                                    |                              |
| 5 Specify the assay method used and provide (or reference) a detailed protocol, including specific reagents or kits used, quality control procedures, reproducibility assessments, quantitation methods, and scoring and reporting protocols. Specify whether and how assays were performed blinded to the study endpoint. | page 7/ line 147-150              | Patients and Methods - Patients and study design/ para 1 |
| **Study design**                                                                   |                                    |                              |
| 6 State the method of case selection, including whether prospective or retrospective and whether stratification or matching (e.g., by stage of disease or age) was used. Specify the time period from which cases were taken, the end of the follow-up period, and the median follow-up time. | page 6-7/ line 141-151            | Patients and Methods - Patients and study design/ para 1 |
| 7 Precisely define all clinical endpoints examined.                                | page 8/ line 178-190              | Patients and Methods          |
| 8 List all candidate variables initially examined or considered for inclusion in models. | page 7-8/ line 161-177            | Patients and Methods          |
| 9 Give rationale for sample size; if the study was designed to detect a specified effect size, give the target power and effect size. | NA for the enrollment of          | NA for the enrollment of      |
| **Statistical analysis methods**                                                    |                                    |                              |
| 10 Specify all statistical methods, including details of any variable selection procedures and other model-building issues, how model assumptions were verified, and how missing data were handled. | page 7-8/ line 153-190            | Patients and Methods - Clinical outcome variables |
| 11 Clarify how marker values were handled in the analyses; if relevant, describe methods used for cutpoint determination. | page 7-8/ line 161-177            | Patients and Methods          |
RESULTS

Data

| 12 | Describe the flow of patients through the study, including the number of patients included in each stage of the analysis (a diagram may be helpful) and reasons for dropout. Specifically, both overall and for each subgroup extensively examined report the numbers of patients and the number of events. | page 10/ line 216-217, 226-227, 237-238 | Results-Baseline immune blood biomarkers and survival outcomes/ para 3-5 |
| 13 | Report distributions of basic demographic characteristics (at least age and sex), standard (disease-specific) prognostic variables, and tumor marker, including numbers of missing values. | page 9/ line 193-201 Table 2 | Results-Patients characteristics/ para 1 |

Analysis and presentation

| 14 | Show the relation of the marker to standard prognostic variables. | page 9/ line 203-215 | Results-Baseline immune |
| 15 | Present univariable analyses showing the relation between the marker and outcome, with the estimated effect (e.g., hazard ratio and survival probability). Preferably provide similar analyses for all other variables being analyzed. For the effect of a tumor marker on a time-to-event outcome, a Kaplan-Meier plot is recommended. | page 10-11/line 216-247 page 12-13/ line 269-293, 301-308 Table 2, 5 | Results-Baseline immune blood biomarkers and survival outcomes/ para 3-5 |
| 16 | For key multivariable analyses, report estimated effects (e.g., hazard ratio) with confidence intervals for the marker and, at least for the final model, all other variables in the model. | page 11/ line 248-260 page 13-14/ 309-314 | Results-Baseline immune blood biomarkers and survival outcomes/ para 3-5 |
| 17 | Among reported results, provide estimated effects with confidence intervals from an analysis in which the marker and standard prognostic variables are included, regardless of their statistical significance. | Table 3 | Table 3 |
| 18 | If done, report results of further investigations, such as checking assumptions, sensitivity analyses, and internal validation. | page 11/ line 52-55 | Results-Baseline immune |

DISCUSSION

| 19 | Interpret the results in the context of the pre-specified hypotheses and other relevant studies; include a discussion of limitations of the study. | page 15-20/ line 344-364, | Discussion/ para 3, 5, 6, 8 |
| 20 | Discuss implications for future research and clinical value. | page 19/ line 438-441 | Discussion/ para 8, 9 |

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*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.