The REMARK checklist

Item	to be reported	Reported on Page Number/Line Number	Reported on Section/Paragraph
INTR	ODUCTION		·
1	State the marker examined, the study objectives, and any pre-specified hypotheses.	Page3/Line96-98	Introduction/Paragraph4
MATE	ERIALS AND METHODS		
Patier	nts		
2	Describe the characteristics (e.g., disease stage or co-morbidities) of the study patients, including their source and inclusion and exclusion criteria.	Page4/Line110-118	Method/Paragraph1
3	Describe treatments received and how chosen (e.g., randomized or rule-based).	N/A	N/A
Speci	imen characteristics		
4	Describe type of biological material used (including control samples) and methods of preservation and storage.	N/A	N/A
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.	N/A	N/A
Study	/ design	1	1
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.	Page4/Line123-132	Method/Paragraph3
7	Precisely define all clinical endpoints examined.	N/A	N/A
8	List all candidate variables initially examined or considered for inclusion in models.	N/A	N/A
9	Give rationale for sample size; if the study was designed to detect a specified effect size, give the target power and effect size.	Page4/ILine123-132	Method/Paragraph3
Statis	tical 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.	Page5/Line151-158	Method/Paragraph6
11	Clarify how marker values were handled in the analyses; if relevant, describe methods used for cutpoint determination.	Page5/line151-155	Method/Paragraph6

RESU	ILTS		
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.	N/A	N/A
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.	Page5/Line162-167 Pgae6/Line171-175	Results/Paragraph1-2
Analy	sis and presentation		
14	Show the relation of the marker to standard prognostic variables.	Page7/Line217-226	Results/Paragraph7
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.	Page7/Line217-226 Page22/Line466-467 Page22/Line469-470	Results/Paregraph7 Figure/Paragraph1 Figure/Paragraph1
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.	N/A	N/A
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.	N/A	N/A
18	If done, report results of further investigations, such as checking assumptions, sensitivity analyses, and internal validation.	N/A	N/A
DISC	USSION		
19	Interpret the results in the context of the pre-specified hypotheses and other relevant studies; include a discussion of limitations of the study.	Page9/Line298-304	Discussion/Paragraph6
20	Discuss implications for future research and clinical value.	Page9/Line304-306	Discussion/Paragraph6

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