The REMARK checklist

Item	to be reported	Reported on Page Number/Line Number	Reported on Section/Paragraph
INTR	DDUCTION		
1	State the marker examined, the study objectives, and any pre-specified hypotheses.	Page4-5	Introduction/Paragraph3
MATE	RIALS AND METHODS		
Patier	its		
2	Describe the characteristics (e.g., disease stage or co-morbidities) of the study patients, including their source and inclusion and exclusion criteria.	Page5/Line 20-25	Method/Paragraph1
3	Describe treatments received and how chosen (e.g., randomized or rule-based).	Page5/Line 20-25	Method/Paragraph1
Speci	men characteristics		
4	Describe type of biological material used (including control samples) and methods of preservation and storage.	Page5/Line 20-25	Method/Paragrhp1
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.	N/A	N/A
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.	N/A	N/A
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.	N/A	N/A
11	Clarify how marker values were handled in the analyses; if relevant, describe methods used for cutpoint determination.	Page8	Method/Paragraph7

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.	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.	N/A	N/A	
Analysis and presentation				
14	Show the relation of the marker to standard prognostic variables.	Page8-9	Results/Paragraph1-2	
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.	Page9	Results/Paragraph2	
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.	Page9	Results/Paragraph2	
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.	Page9	Results/Paragraph2	
18	If done, report results of further investigations, such as checking assumptions, sensitivity analyses, and internal validation.	N/A	N/A	
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.	Page12-13	Discussion/Paragraph1-2	
20	Discuss implications for future research and clinical value.	Page14/Line16-22	Conclusions/Paragraph1	

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