The REMARK checklist

em to be reported	Reported on Page Number/Line Number	Reported on Section/Paragraph		
INTRODUCTION				
State the marker examined, the study objectives, and any pre-specified hypotheses.	Page3/Li ne110-116	Introduction/P5		
ATERIALS AND METHODS				
tients				
Describe the characteristics (e.g., disease stage or co-morbidities) of the study patients, including their source and inclusion and exclusion criteria.	Page3/Li ne128–141	Met hods/P1, 2		
Describe treatments received and how chosen (e.g., randomized or rule-based).	N/A	N/A		
ecimen characteristics				
Describe type of biological material used (including control samples) and methods of preservation and storage.	Page5/Li ne169	Met t hods/P7		
say methods				
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.	Page4/Li ne168-221	Met hods/P8, 9, 10, 11		
udy design				
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/Li ne129	Met hods/P1		
Precisely define all clinical endpoints examined.	Page5/Li ne153-155	Met hods/P4		
List all candidate variables initially examined or considered for inclusion in models.	Page8/Li ne262-263	Results/P4		
Give rationale for sample size; if the study was designed to detect a specified effect size, give the target power and effect size.	Page4/Li ne128-130	Met hods/P1		
atistical analysis methods				
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.	Page7/Li ne224-228	Met hods/P1, 2		
Clarify how marker values were handled in the analyses; if relevant, describe methods used for cutpoint determination.	Page5/Li ne153	Met hods/P4		

RESULTS Data				
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.	Page7/Li ne232-239	Pesul ts/P1	
Analy	sis and presentation			
14	Show the relation of the marker to standard prognostic variables.	Page8/Li ne256-266	Results/P4	
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.	Page8/249-253	Results/P3	
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.	Page8/Li ne 256-266	Pesul ts/P4	
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.	Page15/Li ne462-465	Fi gur e/P5	
18	If done, report results of further investigations, such as checking assumptions, sensitivity analyses, and internal validation.	NA	NA	
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.	Page9-10/Li ne302-331	Di scussi on/p2	
20	Discuss implications for future research and clinical value.	Page10/Li ne332-333	Di scussi on/P5	

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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.