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.	Line 44 - 80	INTRODUCTION
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.	Line 97 - 103	MATERIALS AND METHODS
3	Describe treatments received and how chosen (e.g., randomized or rule-based).	This study does not in	MATERIALS AND METHODS
Specimen characteristics			
4	Describe type of biological material used (including control samples) and methods of preservation and storage.	f biological materials	MATERIALS 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.	Line 144 - 148 This study is a bioinformatics analysis, and the main analysis and	MATERIALS AND METHODS
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.	data of public databas e, which does not invo lve the selection of s amples.	MATERIALS AND METHODS
7	Precisely define all clinical endpoints examined.	This is an analytical	RESULTS
8	List all candidate variables initially examined or considered for inclusion in models.	riable.	MATERIALS 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.	Due to the limitation	MATERIALS AND METHODS
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.	hypothesis. Only case s with complete inform ation were included.	MATERIALS AND METHODS/ RESULTS
11	Clarify how marker values were handled in the analyses; if relevant, describe methods used for cutpoint determination.	Line 97- 148	MATERIALS 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.	The case data in this study came from public database, and there was no flow of samples.	RESULTS
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.	This study is a pancancer analysis. We have shown the expression of TPGS2	RESULTS
Analysis and presentation			
14	Show the relation of the marker to standard prognostic variables.	This study involved	RESULTS
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.	Line 211 - 225; Figure 5-7	RESULTS
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.	This project does not involve multivariate analysis.	RESULTS
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.	Line 211 - 225; Figure 5-7	RESULTS
18	If done, report results of further investigations, such as checking assumptions, sensitivity analyses, and internal validation.	ayses, and internal validation.	RESULTS
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.	Line 308 - 398	DISCUSSION
20	Discuss implications for future research and clinical value.	Line 308 - 398	CONCLUSION

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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.