Matrix Metalloproteinase-3 -1171 5A/6A Polymorphism (rs35068180) is Associated with Risk of Periodontitis

Cheng Ding¹, Xing Chen¹, Peng-tao Zhang¹, Jin-ping Huang², Yan Xu³, Ning Chen^{4,*} & Liang-jun Zhong^{1,5,*}

Criteria items	Score
Citetia tems	(0 to 40)
Title and Abstract	\square 0 \square 1
Title and Abstract: Indicate the study design (case-control or cohort study) in the title or the abstract	□ 0 □ 1
Abstract: Provide an informative and balanced summary of the study	\square 0 \square 1
Introduction: Explain the scientific background and rationale for the investigation	\square 0 \square 1
Introduction: State specific objectives, including any prespecified hypotheses	\square 0 \square 1
Methods	\square 0 \square 1
Study Design: Present key elements of study design	\square 0 \square 1
Setting: Describe the setting, locations, and relevant dates, including periods of recruitment,	
exposure, follow-up, and data collection	\square 0 \square 1
Participants: Give the eligibility criteria of case	\square 0 \square 1
Participants: Give the sources and methods of case ascertainment and control selection	\square 0 \square 1
Participants: Give matching criteria and the number of controls	\square 0 \square 1
Variables: Clearly define all outcomes, exposures, predictors, potential confounders, effect	
modifiers	\square 0 \square 1
Data sources/Measurement: Give sources of data and details of methods of assessment	\square 0 \square 1
Data sources/Measurement: Describe comparability of assessment methods	\square 0 \square 1
Bias: Describe any efforts to address potential sources of bias	\square 0 \square 1
Study size: Explain and describe the estimation of the study size	\square 0 \square 1
Quantitative variables: Explain how quantitative variables were handled in the analyses	\square 0 \square 1
Quantitative variables: Give group included criteria in the analyses	\square 0 \square 1
Statistical methods: Describe all statistical methods, including those used to control for	□ 0 □ 1
confounding	
Statistical methods: Describe any methods used to examine subgroups and interactions	\square 0 \square 1
Statistical methods: Explain how missing data were addressed	\square 0 \square 1
Statistical methods: Explain how matching of cases and controls was addressed	\square 0 \square 1
Statistical methods: Describe any sensitivity analyses	\square 0 \square 1
Hardy-Weinberg equilibrium: HWE was assessed	\square 0 \square 1
Hardy-Weinberg equilibrium: HWE of control group was assessed	\square 0 \square 1
Results	
Participants: Report the numbers of individuals at each stage of the study, such as numbers	
potentially eligible, examined for eligibility, confirmed eligible, included in the study,	□ 0 □ 1
completing follow-up and analyzed	
Participants: Give reasons for non-participation at each stage	$\sqcup 0 \sqcup 1$
Participants: Give a flow diagram Descriptive data: Give abstractoristics of study participants (e.g. demographic clinical)	$\sqcup 0 \sqcup 1$
Descriptive data: Give characteristics of study participants (e.g. demographic, clinical diagnosis, ethnicity, sex ratio, etc.)	\square 0 \square 1
	\square 0 \square 1
Descriptive data: Indicate the number of participants with missing data Outcome data: Papart numbers in each exposure actorogy, or summers measures of exposure	
Outcome data: Report numbers in each exposure category, or summary measures of exposure	$\sqcup 0 \sqcup 1$
Main results: Give unadjusted estimates and confounder-adjusted estimates and their 95%	\square 0 \square 1

confidence intervals	
Main results: Make clear which confounders were adjusted for and why they were included	\square 0 \square 1
Main results: If relevant, consider translating estimates of relative risk into absolute risk for a meaningful time period	□ 0 □ 1
Other analyses: Report other analyses such subgroups, interactions, and sensitivity analyses	\square 0 \square 1
Discussion	\square 0 \square 1
Key results: Summarize key results with reference to study objectives	\square 0 \square 1
Main results: Report category boundaries when continuous variables were categorized	\square 0 \square 1
Limitations: Discuss both direction and magnitude of any potential bias	\square 0 \square 1
Interpretation: Give a cautious overall interpretation of results considering objectives, limitations, multiplicity of analyses, results from similar studies, and other relevant evidence	□ 0 □ 1
Generalizability: Discuss the generalizability (external validity) of the study results	\square 0 \square 1
Other	\square 0 \square 1
Funding: Give the source of funding and the role of the funders for the present study	\square 0 \square 1

Supplementary Table S1. The modified STROBE quality score systems.