

Decreased risk of developing lung cancer in subjects carrying the *CLPTM1L* rs401681 (G>A) polymorphism: evidence from a meta-analysis

X.L. Zhang^{1,2*}, X.J. Zhang^{1*}, Y. Chen^{4*}, Y.M. Zhang³, Q. Zhang¹, C.X. Cao¹, D.Y. Gu¹, J.F. Shi¹, Y.L. Gong¹, J.F. Chen¹ and C.J. Tang¹

¹Department of Oncology, Nanjing First Hospital, Nanjing Medical University, Nanjing, China

²Department of Oncology, Nantong Tumor Hospital, Nantong, Jiangsu, China ³Department of Oncology, Xuzhou Hospital Affiliated to Medical College of Southeast University and Xuzhou Central Hospital, Xuzhou, China ⁴Medical School of Southeast University, Nanjing, China

*These authors contributed equally to this study. Corresponding authors: J.F. Chen / C.J. Tang E-mail: jinfeichen@sohu.com / tangcuiju@tom.com

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ABSTRACT. A genome-wide association study revealed that a single nucleotide polymorphism, CLPTMIL - rs401681 (G>A), located at the 5p15.33 locus was significantly associated with increased risk of various cancers; however, its association with lung cancer is currently inconclusive. In order to explore the relationship between this polymorphism and lung cancer risk more precisely, we performed a meta-analysis of eight eligible studies involving 9935 cases and 11,261 controls. The pooled odds ratio (OR) and the 95% confidence interval (CI) were calculated using a fixed-or random-effect models. Results indicated that this polymorphism was significantly associated with lung cancer risk in all genetic models (GA vs GG: OR = 0.88, 95%CI = 0.83-0.94; AA vs GG: OR = 0.81, 95%CI = 0.70-0.93; AA/GA vs GG: OR = 0.86, 95%CI = 0.81-0.91; AA vs GA/GG:

OR = 0.86, 95%CI = 0.76-0.99). An analysis stratified by ethnicity and source of controls revealed a significantly decreased risk among European groups and population-based studies in all genetic models, and among Asian populations only in the dominant model comparison. Additionally, in a subgroup analysis by histology type, the *CLPTM1L* rs401681 polymorphism was found to significantly decrease the risks of both adenocarcinoma and squamous cell carcinoma of the lung in all genetic models. In conclusion, our study indicated that the *CLPTM1L* - rs401681 (G>A) polymorphism was significantly associated with decreased lung cancer risk, especially among European populations. Due to some minor limitations, our findings should be confirmed in further studies.

Key words: Genetic polymorphism; *TERT-CLPTM1L*; rs401681 (G>A); Lung cancer; Meta-analysis