

Letter to the Editor

Comment on "IL1-RN variable number of tandem repeats polymorphism with osteoarthritis risk"

Beuv Joob¹ , Viroi Wiwanitkit^{2, 3}

¹Sanitation 1 Medical Academic Center, Bangkok, Thailand ²Dr D Y Patil University, Pune, India ³Joseph Ayobabalola University, Ikeiji-Araeji, Nigeria

Article history: Received 1 November 2019 Accepted 27 May 2020

ORCID iDs of the authors: B.J. 0000-0002-5281-0369; V.W. 0000-0003-1039-3728.

Corresponding Author: Beuv Joob beuyjoob@hotmail.com



Content of this journal is licensed under a Creative Commons Attribution-NonCommercial 4.0 International License.

A R T I C L E I N F O Dear Editor, we read the publication from China on "Meta-analysis of the association of IL1-RN variable number of tandem repeats polymorphism with osteoarthritis risk" with a great interest (1). Xu et al. concluded that "IL1-RN VNTR polymorphism may increase the susceptibility to OA (1)." We would like to share ideas on this interesting report. First, the clinical association of IL and medical problem is widely mentioned. The effect on clinical phenotypic manifestation in IL1-RN variable number of tandem repeats polymorphism is explainable by molecular pathophysiology. As observed in other IL polymorphisms, the molecular change is resulted from genetic mutation and can further result in altered phenotypic expression (2). In IL1-RN variable number of tandem repeats polymorphism, a significant molecular change due to variation of the number of tandem repeats is detectable. However, in the present study, Xu et al. focused on single genetic polymorphism. There are also other possible genetic polymorphisms that possibly relate to OA risk. The good examples of those polymorphisms are IL17A, CD52, and CCL2 polymorphisms (3-6). Therefore, there should be further studies to cover other genetic polymorphisms that might be associated with OA risk.

> Conflict of Interest: The authors have no conflicts of interest to declare.

> Financial Disclosure: The authors declared that this study has received no financial support.

References

- 1. Xu B, Shi XQ, Xing RL, Xiao YC, Wu P, Wang PM. Meta-analysis of the association of IL1-RN variable number of tandem repeats polymorphism with osteoarthritis risk. Acta Orthop Traumatol Turc 2019; 53: 497-501. [Crossref]
- Srriwijitalai W, Wiwanitkit V. Interleukin-6 2. -174G/C polymorphism and end-stage renal disease: Is there any role? Saudi J Kidney Dis Transpl 2018; 29: 747-8. [Crossref]
- 3. Gao S, Mao C, Cheng J, Deng Q, Sheng W. Association of IL-17A-197G/A and IL-17F-7488T/C polymorphisms and osteoarthritis susceptibility: A meta-analysis. Int J Rheum Dis 2020; 23: 37-46 [Crossref]
- Xu Z, Li J, Yang H, et al. Association of CCL2 4. gene variants with osteoarthritis. Arch Med Res 2019; 50: 86-90. [Crossref]
- 5. Wang Y, Zhang X, Niu X, Xu Y, Lu L, Li H. The genetic relationship of SOX9 polymorphisms with osteoarthritis risk in Chinese population: A case-control study. Medicine (Baltimore) 2019; 98: e14096. doi: 10.1097/ MD.000000000014096. [Crossref]
- 6. Shang H, Hao Y, Hu W, Hu X, Jin Q. CDH2 gene rs11564299 polymorphism is a risk factor for knee osteoarthritis in a Chinese population: A case-control study. J Orthop Surg Res 2019; 14: 208. doi: 10.1186/s13018-019-1256-0. [Crossref]

Author's response:

"Author's reply to the comment couldn't obtained despite all attempts."

Cite this article as: Joob B, Wiwanitkit V. Comment on "IL1-RN variable number of tandem repeats polymorphism with osteoarthritis risk". Acta Orthop Traumatol Turc 2020; 54(4): 472.