



Letter to the Editor

Trimethylamine N-oxide: A new therapeutic target for atrial fibrillation?

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ARTICLE INFO

Article history:

Received 13 July 2018

Received in revised form 19 July 2018

Accepted 31 July 2018

Keywords:

Atrial fibrillation

Trimethylamine N-oxide

Dear Editor,

We have read with great interest of the recently published article entitled "A potential relationship between gut microbes and atrial fibrillation: Trimethylamine N-oxide, a gut microbe-derived metabolite, facilitates the progression of atrial fibrillation" by Yu et al. [1]. Atrial fibrillation (AF) is a common cardiac arrhythmia, especially in aged population. In Yu's paper, trimethylamine N-oxide (TMAO) was reported to increase the electric remodeling in canine AF models via activating inflammation in ganglionated plexi [1]. Interestingly, circulating TMAO elevation was also found in AF patients [2,3]. These raise the question that whether lowering the blood TMAO level is beneficial for AF patients. TMAO is a common gut microbe's metabolite, the concentration of TMAO in the blood increases after consuming food containing carnitine or lecithin [4]. These indicate that TMAO level can be reduced with diet control. TMAO is also associated with other cardiovascular disease, a recent report shows that TMAO level is associated with the vulnerability

of atherosclerotic plaques in coronary artery disease patients [5]. A large number of AF patients also have other cardiovascular disease. Taken together, a diet control to lower blood TMAO may benefit AF patients, especially in those comorbid with other cardiovascular disease.

Conflict of interest

None.

Acknowledgments

This authors acknowledge the grant supported by National Natural Science Foundation of China (No. 81700426).

References

- [1] L. Yu, G. Meng, B. Huang, X. Zhou, S. Stavrakis, M. Wang, et al., A potential relationship between gut microbes and atrial fibrillation: Trimethylamine N-oxide, a gut microbe-derived metabolite, facilitates the progression of atrial fibrillation, *Int. J. Cardiol.* 255 (2018) 92–98.
- [2] G.F.T. Svingen, H. Zuo, P.M. Ueland, R. Seifert, K.H. Loland, E.R. Pedersen, et al., Increased plasma trimethylamine-N-oxide is associated with incident atrial fibrillation, *Int. J. Cardiol.* 267 (2018) 100–106.
- [3] R.S. Mishima, A.D. Elliott, P. Sanders, D. Linz, Microbiome and atrial fibrillation, *Int. J. Cardiol.* 255 (2018) 103–104.
- [4] W.H. Tang, Z. Wang, B.S. Levison, R.A. Koeth, E.B. Britt, X. Fu, et al., Intestinal microbial metabolism of phosphatidylcholine and cardiovascular risk, *N. Engl. J. Med.* 368 (2013) 1575–1584.
- [5] X. Liu, Z. Xie, M. Sun, X. Wang, J. Li, J. Cui, et al., Plasma trimethylamine N-oxide is associated with vulnerable plaque characteristics in CAD patients as assessed by optical coherence tomography, *Int. J. Cardiol.* 265 (2018) 18–23.

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