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Increased Resting Heart Rate and Genetic Variability

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This is an Open Access article distributed under the terms of the Creative Commons Attribution Non-Commercial License (http:// creativecommons.org/licenses/bync/4.0/) which permits unrestricted non-commercial use, distribution, and reproduction in any medium, provided the original work is properly cited. Copyright© 2019 Mongolian National University of Medical Sciences Increased resting heart rate is associated with various cardiovascular events and diseases, such as atherosclerotic lesions in the coronary arteries, atrial fibrillation, acute myocardial infarction, and hypertension [1]. These associations have been shown not only in patients with cardiovascular diseases but also in the general population [2]. It has been reported that an increased resting HR is an independent risk factor for cardiovascular mortality, including sudden cardiac death and total mortality even after adjusting for other potential risk factors [3]. However, the association between increased resting heart rate and both cardiovascular diseases and mortality has been shown in a large number of studies. However, their molecular mechanism is still not well understood.

The heart rate is a complex trait influenced by multiple genetic and environmental factors and their interactions. Several environmental risk factors for increased heart rate are already known, and the heritability of heart rate has been estimated up to 32-63 %, which could result from common or rare genetic variants [4,5]. Although heritable factors play an extensive role in regulating heart rate, evidence about its specific genetic determinants is still limited, even though it may provide additional insight into the pathophysiology of cardiac function and help explain the molecular mechanisms and genetic basis for disorders associated with its regulation.

Much effort has been devoted to mapping the genes that are associated with heart rate regulation, including linkage and genome-wide linkage studies as well as candidate-gene approaches; however, the results have been inconsistent. In recent years, genome-wide association studies [6] have been a successful strategy for identifying highly compelling candidate genes that influence common human traits, including those for various electrocardiogram parameters such as PR, QT and RR intervals [7-11]. The hope is that through a better understanding of genetic factors impacting resting heart rate, patients can be better advised regarding their risks for sudden cardiac death, and perhaps someday gene-based treatments may be offered.

Editorial

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