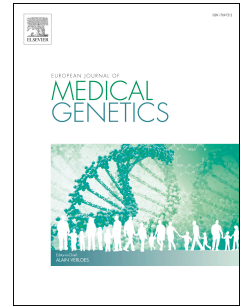


Accepted Manuscript

Whole-exome sequencing identifies rare compound heterozygous mutations in the *MYBPC3* gene associated with severe familial hypertrophic cardiomyopathy

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PII: S1769-7212(17)30728-0

DOI: [10.1016/j.ejmg.2018.03.001](https://doi.org/10.1016/j.ejmg.2018.03.001)

Reference: EJMG 3429

To appear in: *European Journal of Medical Genetics*

Received Date: 31 October 2017

Revised Date: 1 March 2018

Accepted Date: 2 March 2018

Please cite this article as: N. Zhou, S. Qin, Y. Liu, L. Tang, W. Zhao, C. Pan, Z. Qiu, X. Wang, X. Shu, Whole-exome sequencing identifies rare compound heterozygous mutations in the *MYBPC3* gene associated with severe familial hypertrophic cardiomyopathy, *European Journal of Medical Genetics* (2018), doi: 10.1016/j.ejmg.2018.03.001.

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Whole-exome Sequencing Identifies Rare Compound Heterozygous Mutations in the *MYBPC3* Gene Associated with Severe Familial Hypertrophic Cardiomyopathy

Running head: *MYBPC3* in hypertrophic cardiomyopathy

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