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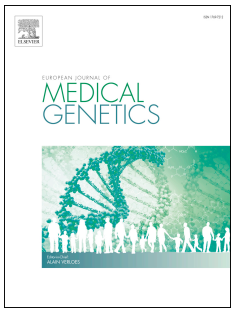

Whole-exome sequencing identifies rare compound heterozygous mutations in the *MYBPC3* gene associated with severe familial hypertrophic cardiomyopathy

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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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