

# CITATION REPORT

List of articles citing

**Biallelic C1QBP Mutations Cause Severe Neonatal-,  
Childhood-, or Later-Onset Cardiomyopathy  
Associated with Combined Respiratory-Chain Deficiencies**

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**American Journal of Human Genetics, 2017, 101, 525-538.**

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#	Paper	IF	Citations
53	In vitro and in vivo studies of the ALS-FTLD protein CHCHD10 reveal novel mitochondrial topology and protein interactions. <i>Human Molecular Genetics</i> , <b>2018</b> , 27, 160-177	5.6	39
52	Mitochondrial p32/C1qbp Is a Critical Regulator of Dendritic Cell Metabolism and Maturation. <i>Cell Reports</i> , <b>2018</b> , 25, 1800-1815.e4	10.6	35
51	Mitochondrial disease genetics update: recent insights into the molecular diagnosis and expanding phenotype of primary mitochondrial disease. <i>Current Opinion in Pediatrics</i> , <b>2018</b> , 30, 714-724	3.2	27
50	The Expression Pattern of in Sheep Muscle and Its Role in Differentiation, Cell Proliferation, and Apoptosis of Myoblasts. <i>International Journal of Molecular Sciences</i> , <b>2019</b> , 20,	6.3	1
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