

Michaela Yuen

List of Publications by Year in descending order

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Version: 2024-02-01

7
papers

348
citations

1307543

7
h-index

1720014

7
g-index

7
all docs

7
docs citations

7
times ranked

686
citing authors

#	ARTICLE	IF	CITATIONS
1	Leiomodin-3 dysfunction results in thin filament disorganization and nemaline myopathy. <i>Journal of Clinical Investigation</i> , 2014, 124, 4693-4708.	8.2	153
2	Mutation-specific effects on thin filament length in thin filament myopathy. <i>Annals of Neurology</i> , 2016, 79, 959-969.	5.3	54
3	Variants in <i>SLC18A3</i> , vesicular acetylcholine transporter, cause congenital myasthenic syndrome. <i>Neurology</i> , 2016, 87, 1442-1448.	1.1	46
4	Muscle weakness in <i>TPM3</i> -myopathy is due to reduced Ca ²⁺ -sensitivity and impaired acto-myosin cross-bridge cycling in slow fibres. <i>Human Molecular Genetics</i> , 2015, 24, 6278-6292.	2.9	38
5	KBTBD13 is an actin-binding protein that modulates muscle kinetics. <i>Journal of Clinical Investigation</i> , 2020, 130, 754-767.	8.2	25
6	Nebulin: big protein with big responsibilities. <i>Journal of Muscle Research and Cell Motility</i> , 2020, 41, 103-124.	2.0	23
7	Neonatal-lethal dilated cardiomyopathy due to a homozygous <i>LMOD2</i> donor splice-site variant. <i>European Journal of Human Genetics</i> , 2022, 30, 450-457.	2.8	9