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List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Imaging neural activity in the ventral nerve cord of behaving adult Drosophila. Nature Communications, 2018, 9, 4390.	12.8	62
2	Pseudo-acetylation of K326 and K328 of actin disrupts Drosophila melanogaster indirect flight muscle structure and performance. Frontiers in Physiology, 2015, 6, 116.	2.8	33
3	Profilin modulates sarcomeric organization and mediates cardiomyocyte hypertrophy. Cardiovascular Research, 2016, 110, 238-248.	3.8	31
4	Modest overexpression of <i><scp>FOXO</scp></i> maintains cardiac proteostasis and ameliorates ageâ€associated functional decline. Aging Cell, 2017, 16, 93-103.	6.7	31
5	Distortion of the Actin A-Triad Results in Contractile Disinhibition and Cardiomyopathy. Cell Reports, 2017, 20, 2612-2625.	6.4	26
6	Prolonged cross-bridge binding triggers muscle dysfunction in a Drosophila model of myosin-based hypertrophic cardiomyopathy. ELife, 2018, 7, .	6.0	26
7	<i>TNNT2</i> mutations in the tropomyosin binding region of TNT1 disrupt its role in contractile inhibition and stimulate cardiac dysfunction. Proceedings of the National Academy of Sciences of the United States of America, 2020, 117, 18822-18831.	7.1	21
8	CaMKII oxidation is a critical performance/disease trade-off acquired at the dawn of vertebrate evolution. Nature Communications, 2021, 12, 3175.	12.8	19
9	Myosin storage myopathy mutations yield defective myosin filament assembly in vitro and disrupted myofibrillar structure and function in vivo. Human Molecular Genetics, 2017, 26, 4799-4813.	2.9	16
10	A role for actin flexibility in thin filament-mediated contractile regulation and myopathy. Nature Communications, 2020, 11, 2417.	12.8	16
11	Cardiac-Restricted Expression of VCP/TER94 RNAi or Disease Alleles Perturbs Drosophila Heart Structure and Impairs Function. Journal of Cardiovascular Development and Disease, 2016, 3, 19.	1.6	14
12	Conservation of cardiac L-type Ca2+ channels and their regulation in Drosophila: A novel genetically-pliable channelopathic model. Journal of Molecular and Cellular Cardiology, 2018, 119, 64-74.	1.9	9
13	A Restrictive Cardiomyopathy Mutation in an Invariant Proline at the Myosin Head/Rod Junction Enhances Head Flexibility and Function, Yielding Muscle Defects in Drosophila. Journal of Molecular Biology, 2016, 428, 2446-2461.	4.2	8
14	Myosin dilated cardiomyopathy mutation S532P disrupts actomyosin interactions, leading to altered muscle kinetics, reduced locomotion, and cardiac dilation in <i>Drosophila</i> . Molecular Biology of the Cell, 2021, 32, 1690-1706.	2.1	8
15	Quantifying Tissue-Specific Overexpression of FOXO in Drosophila via mRNA Fluorescence In Situ Hybridization Using Branched DNA Probe Technology. Methods in Molecular Biology, 2019, 1890, 171-190.	0.9	3
16	The R369 Myosin Residue within Loop 4 Is Critical for Actin Binding and Muscle Function in Drosophila. International Journal of Molecular Sciences, 2022, 23, 2533.	4.1	1