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List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Effect of actin C-terminal modification on tropomyosin isoforms binding and thin filament regulation. Biochimica Et Biophysica Acta - Proteins and Proteomics, 2009, 1794, 237-243.	2.3	20
2	Congenital myopathyâ€related mutations in tropomyosin disrupt regulatory function through altered actin affinity and tropomodulin binding. FEBS Journal, 2019, 286, 1877-1893.	4.7	14
3	Functional effects of substitutions I92T and V95A in actin-binding period 3 of tropomyosin. Biochimica Et Biophysica Acta - Proteins and Proteomics, 2018, 1866, 558-568.	2.3	13
4	Different positions of tropomyosin isoforms on actin filament are determined by specific sequences of endâ€ŧoâ€end overlaps. Cytoskeleton, 2011, 68, 300-312.	2.0	10
5	Tropomyosin isoforms regulate cofilin 1 activity by modulating actin filament conformation. Archives of Biochemistry and Biophysics, 2020, 682, 108280.	3.0	10
6	Mutations Q93H and E97K in TPM2 Disrupt Ca-Dependent Regulation of Actin Filaments. International Journal of Molecular Sciences, 2021, 22, 4036.	4.1	7
7	Regulation of Actin Filament Length by Muscle Isoforms of Tropomyosin and Cofilin. International Journal of Molecular Sciences, 2020, 21, 4285.	4.1	6
8	Structural Effects of Disease-Related Mutations in Actin-Binding Period 3 of Tropomyosin. Molecules, 2021, 26, 6980.	3.8	4
9	Abnormal movement of tropomyosin and response of myosin heads and actin during the ATPase cycle caused by the Arg167His, Arg167Cly and Lys168Glu mutations in TPM1 gene. Archives of Biochemistry and Biophysics, 2016, 606, 157-166.	3.0	3
10	Structural differences between C-terminal regions of tropomyosin isoforms. PeerJ, 2013, 1, e181.	2.0	2
11	The primary cause of muscle disfunction associated with substitutions E240K and R244G in tropomyosin is aberrant behavior of tropomyosin and response of actin and myosin during ATPase cycle. Archives of Biochemistry and Biophysics, 2018, 644, 17-28.	3.0	0