

Qiuping Zhang

List of Publications by Year in descending order

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Version: 2024-02-01

30
papers

3,909
citations

393982

19
h-index

454577

30
g-index

31
all docs

31
docs citations

31
times ranked

4768
citing authors

#	ARTICLE	IF	CITATIONS
1	SUN1/2 Are Essential for RhoA/ROCK-Regulated Actomyosin Activity in Isolated Vascular Smooth Muscle Cells. <i>Cells</i> , 2020, 9, 132.	1.8	22
2	Nesprin-1-alpha2 associates with kinesin at myotube outer nuclear membranes, but is restricted to neuromuscular junction nuclei in adult muscle. <i>Scientific Reports</i> , 2019, 9, 14202.	1.6	14
3	Nesprin-1/2: roles in nuclear envelope organisation, myogenesis and muscle disease. <i>Biochemical Society Transactions</i> , 2018, 46, 311-320.	1.6	36
4	Mouse models of nesprin-related diseases. <i>Biochemical Society Transactions</i> , 2018, 46, 669-681.	1.6	9
5	Novel nesprin-1 mutations associated with dilated cardiomyopathy cause nuclear envelope disruption and defects in myogenesis. <i>Human Molecular Genetics</i> , 2017, 26, 2258-2276.	1.4	91
6	Selumetinib, an Oral Anti-Neoplastic Drug, May Attenuate Cardiac Hypertrophy via Targeting the ERK Pathway. <i>PLoS ONE</i> , 2016, 11, e0159079.	1.1	20
7	Identification and Validation of Putative Nesprin Variants. <i>Methods in Molecular Biology</i> , 2016, 1411, 211-220.	0.4	4
8	N-terminal nesprin-2 variants regulate β -catenin signalling. <i>Experimental Cell Research</i> , 2016, 345, 168-179.	1.2	17
9	Specific localization of nesprin-1 β 2, the short isoform of nesprin-1 with a KASH domain, in developing, fetal and regenerating muscle, using a new monoclonal antibody. <i>BMC Cell Biology</i> , 2016, 17, 26.	3.0	26
10	Nesprin-2-dependent ERK1/2 compartmentalisation regulates the DNA damage response in vascular smooth muscle cell ageing. <i>Cell Death and Differentiation</i> , 2015, 22, 1540-1550.	5.0	35
11	Nesprins: Tissue-Specific Expression of Epsilon and Other Short Isoforms. <i>PLoS ONE</i> , 2014, 9, e94380.	1.1	72
12	Bidirectional cross-regulation between the endothelial nitric oxide synthase and β -catenin signalling pathways. <i>Cardiovascular Research</i> , 2014, 104, 116-126.	1.8	21
13	Multiple Novel Nesprin-1 and Nesprin-2 Variants Act as Versatile Tissue-Specific Intracellular Scaffolds. <i>PLoS ONE</i> , 2012, 7, e40098.	1.1	93
14	Nesprin-1 and actin contribute to nuclear and cytoskeletal defects in lamin A/C-deficient cardiomyopathy. <i>Journal of Molecular and Cellular Cardiology</i> , 2011, 50, 479-486.	0.9	29
15	Paternal uniparental isodisomy of chromosome 6 causing a complex syndrome including complete IFN γ receptor 1 deficiency. <i>American Journal of Medical Genetics, Part A</i> , 2010, 152A, 622-629.	0.7	22
16	Novel Nuclear Nesprin-2 Variants Tether Active Extracellular Signal-regulated MAPK1 and MAPK2 at Promyelocytic Leukemia Protein Nuclear Bodies and Act to Regulate Smooth Muscle Cell Proliferation. <i>Journal of Biological Chemistry</i> , 2010, 285, 1311-1320.	1.6	47
17	Epidermal growth factor receptor gene polymorphisms, R497K, but not (CA) n repeat, is associated with dilated cardiomyopathy. <i>Clinica Chimica Acta</i> , 2009, 403, 184-187.	0.5	8
18	Cell Nuclei Spin in the Absence of Lamin B1. <i>Journal of Biological Chemistry</i> , 2007, 282, 20015-20026.	1.6	83

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19	Nesprin-1 and -2 are involved in the pathogenesis of Emeryâ€Dreifuss muscular dystrophy and are critical for nuclear envelope integrity. <i>Human Molecular Genetics</i> , 2007, 16, 2816-2833.	1.4	461
20	Distinct functional domains in nesprin-1 [±] and nesprin-2 [±] bind directly to emerin and both interactions are disrupted in X-linked Emeryâ€Dreifuss muscular dystrophy. <i>Experimental Cell Research</i> , 2007, 313, 2845-2857.	1.2	84
21	Nesprin-2 is a multi-isomeric protein that binds lamin and emerin at the nuclear envelope and forms a subcellular network in skeletal muscle. <i>Journal of Cell Science</i> , 2005, 118, 673-687.	1.2	236
22	Nesprins: intracellular scaffolds that maintain cell architecture and coordinate cell function?. <i>Expert Reviews in Molecular Medicine</i> , 2005, 7, 1-15.	1.6	1,274
23	Osteo/Chondrocytic Transcription Factors and Their Target Genes Exhibit Distinct Patterns of Expression in Human Arterial Calcification. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2003, 23, 489-494.	1.1	479
24	The Nesprins Are Giant Actin-Binding Proteins, Orthologous to <i>Drosophila melanogaster</i> Muscle Protein MSP-300. <i>Genomics</i> , 2002, 80, 473-481.	1.3	182
25	The nesprins are giant actin-binding proteins, orthologous to <i>Drosophila melanogaster</i> muscle protein MSP-300. <i>Genomics</i> , 2002, 80, 473-81.	1.3	93
26	Nesprins: a novel family of spectrin-repeat-containing proteins that localize to the nuclear membrane in multiple tissues. <i>Journal of Cell Science</i> , 2001, 114, 4485-4498.	1.2	348
27	Common Genetic Variants of Lipoprotein Lipase and Apolipoproteins AIâ€CIII That Relate to Coronary Artery Disease: A Study in Chinese and European Subjects. <i>Molecular Genetics and Metabolism</i> , 1998, 64, 177-183.	0.5	19
28	Mutations at the Lipoprotein Lipase Gene Locus in Subjects with Diabetes Mellitus, Obesity and Lipaemia. <i>Clinical Science</i> , 1997, 93, 335-341.	1.8	15
29	Lipid Transport Genes and Their Relation to the Syndrome of Insulin Resistance. <i>Annals of the New York Academy of Sciences</i> , 1997, 827, 100-109.	1.8	1
30	Common genetic variants of lipoprotein lipase that relate to lipid transport in patients with premature coronary artery disease. <i>Clinical Genetics</i> , 1995, 48, 293-298.	1.0	44