

Gaynor Miller

List of Publications by Year in descending order

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Version: 2024-02-01

15
papers

473
citations

840776

11
h-index

996975

15
g-index

15
all docs

15
docs citations

15
times ranked

943
citing authors

#	ARTICLE	IF	CITATIONS
1	Appearances can be deceiving: phenotypes of knockout mice. <i>Briefings in Functional Genomics & Proteomics</i> , 2007, 6, 91-103.	3.8	175
2	A targeted deletion of the C-terminal end of titin, including the titin kinase domain, impairs myofibrillogenesis. <i>Journal of Cell Science</i> , 2003, 116, 4811-4819.	2.0	46
3	Preventing phosphorylation of dystroglycan ameliorates the dystrophic phenotype in mdx mouse. <i>Human Molecular Genetics</i> , 2012, 21, 4508-4520.	2.9	33
4	Altered Macrophage Polarization Induces Experimental Pulmonary Hypertension and Is Observed in Patients With Pulmonary Arterial Hypertension. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2021, 41, 430-445.	2.4	33
5	PyMT-Maclow: A novel, inducible, murine model for determining the role of CD68 positive cells in breast tumor development. <i>PLoS ONE</i> , 2017, 12, e0188591.	2.5	33
6	Specific and Potent RNA Interference in Terminally Differentiated Myotubes. <i>Journal of Biological Chemistry</i> , 2003, 278, 934-939.	3.4	32
7	Structural and functional analysis of the sarcoglycan-sarcospan subcomplex. <i>Experimental Cell Research</i> , 2007, 313, 639-651.	2.6	26
8	Disrupted mechanical stability of the dystrophin-glycoprotein complex causes severe muscular dystrophy in sarcospan transgenic mice. <i>Journal of Cell Science</i> , 2007, 120, 996-1008.	2.0	25
9	ENU Mutagenesis Reveals a Novel Phenotype of Reduced Limb Strength in Mice Lacking Fibrillin 2. <i>PLoS ONE</i> , 2010, 5, e9137.	2.5	19
10	Heterologous expression of wild-type and mutant β -cardiac myosin changes the contractile kinetics of cultured mouse myotubes. <i>Journal of Physiology</i> , 2003, 548, 167-174.	2.9	14
11	Over-expression of Microspan, a novel component of the sarcoplasmic reticulum, causes severe muscle pathology with triad abnormalities. <i>Journal of Muscle Research and Cell Motility</i> , 2006, 27, 545-558.	2.0	13
12	<i>In Vivo</i> Fiber Optic Raman Spectroscopy of Muscle in Preclinical Models of Amyotrophic Lateral Sclerosis and Duchenne Muscular Dystrophy. <i>ACS Chemical Neuroscience</i> , 2021, 12, 1768-1776.	3.5	12
13	Generation of a novel mouse model for the inducible depletion of macrophages in vivo. <i>Genesis</i> , 2013, 51, 41-49.	1.6	6
14	N232S, G741R and D778G β -cardiac myosin mutants, implicated in familial hypertrophic cardiomyopathy, do not disrupt myofibrillar organisation in cultured myotubes. <i>FEBS Letters</i> , 2000, 486, 325-327.	2.8	5
15	Nanospan, an alternatively spliced isoform of sarcospan, localizes to the sarcoplasmic reticulum in skeletal muscle and is absent in limb girdle muscular dystrophy 2F. <i>Skeletal Muscle</i> , 2017, 7, 11.	4.2	1