

Jeffrey J Widrick

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/5075204/publications.pdf>

Version: 2024-02-01

20
papers

1,524
citations

759233

12
h-index

794594

19
g-index

20
all docs

20
docs citations

20
times ranked

2848
citing authors

#	ARTICLE	IF	CITATIONS
1	In vivo gene editing in dystrophic mouse muscle and muscle stem cells. <i>Science</i> , 2016, 351, 407-411.	12.6	889
2	Directed evolution of a family of AAV capsid variants enabling potent muscle-directed gene delivery across species. <i>Cell</i> , 2021, 184, 4919-4938.e22.	28.9	193
3	MicroRNA-486-dependent modulation of DOCK3/PTEN/AKT signaling pathways improves muscular dystrophy-associated symptoms. <i>Journal of Clinical Investigation</i> , 2014, 124, 2651-2667.	8.2	128
4	Concurrent muscle and bone deterioration in a murine model of cancer cachexia. <i>Physiological Reports</i> , 2013, 1, e00144.	1.7	38
5	Dystrophic muscle improvement in zebrafish via increased heme oxygenase signaling. <i>Human Molecular Genetics</i> , 2014, 23, 1869-1878.	2.9	38
6	RNA helicase, DDX27 regulates skeletal muscle growth and regeneration by modulation of translational processes. <i>PLoS Genetics</i> , 2018, 14, e1007226.	3.5	34
7	Muscle dysfunction in a zebrafish model of Duchenne muscular dystrophy. <i>Physiological Genomics</i> , 2016, 48, 850-860.	2.3	29
8	Whole Body Periodic Acceleration Is an Effective Therapy to Ameliorate Muscular Dystrophy in mdx Mice. <i>PLoS ONE</i> , 2014, 9, e106590.	2.5	25
9	Evaluation of Electrical Impedance as a Biomarker of Myostatin Inhibition in Wild Type and Muscular Dystrophy Mice. <i>PLoS ONE</i> , 2015, 10, e0140521.	2.5	21
10	Discovery of Novel Therapeutics for Muscular Dystrophies using Zebrafish Phenotypic Screens. <i>Journal of Neuromuscular Diseases</i> , 2019, 6, 271-287.	2.6	21
11	Gait characteristics of adults with Down syndrome explain their greater metabolic rate during walking. <i>Gait and Posture</i> , 2015, 41, 180-184.	1.4	19
12	A limb-girdle muscular dystrophy 2l model of muscular dystrophy identifies corrective drug compounds for dystroglycanopathies. <i>JCI Insight</i> , 2018, 3, .	5.0	17
13	Transgenic zebrafish model of DUX4 misexpression reveals a developmental role in FSHD pathogenesis. <i>Human Molecular Genetics</i> , 2019, 28, 320-331.	2.9	14
14	An open source microcontroller based flume for evaluating swimming performance of larval, juvenile, and adult zebrafish. <i>PLoS ONE</i> , 2018, 13, e0199712.	2.5	13
15	IMP2 Increases Mouse Skeletal Muscle Mass and Voluntary Activity by Enhancing Autocrine Insulin-Like Growth Factor 2 Production and Optimizing Muscle Metabolism. <i>Molecular and Cellular Biology</i> , 2019, 39, .	2.3	12
16	miR-486 is essential for muscle function and suppresses a dystrophic transcriptome. <i>Life Science Alliance</i> , 2022, 5, e202101215.	2.8	10
17	Tetraspanin CD82 is necessary for muscle stem cell activation and supports dystrophic muscle function. <i>Skeletal Muscle</i> , 2020, 10, 34.	4.2	9
18	PDE10A Inhibition Reduces the Manifestation of Pathology in DMD Zebrafish and Represses the Genetic Modifier PTPNA. <i>Molecular Therapy</i> , 2021, 29, 1086-1101.	8.2	9

#	ARTICLE	IF	CITATIONS
19	Dynamin-2 reduction rescues the skeletal myopathy of a SPEG-deficient mouse model. JCI Insight, 2022, 7, .	5.0	5
20	Skeletal Muscle Dysfunction in Experimental Pulmonary Hypertension Kosmas Kosmas ^{1,2} , Zoe Michael ^{2,3} , Fotios Spyropoulos ^{1,2} , Jeffrey Widrick ^{2,4} , Ravi Jasuja ² , Aimilia Papathanasiou ^{1,2} , Helen Christou ^{1,2} ¹ Department of Pediatric New. FASEB Journal, 2022, 36, .	0.5	0