

Jeffrey J Widrick

List of Publications by Year in descending order

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

#	ARTICLE	IF	CITATIONS
19	Dystrophic muscle improvement in zebrafish via increased heme oxygenase signaling. Human Molecular Genetics, 2014, 23, 1869-1878.	2.9	38
20	Concurrent muscle and bone deterioration in a murine model of cancer cachexia. Physiological Reports, 2013, 1, e00144.	1.7	38