Marco Morsch

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/2147880/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Brainâ€Targeted Aggregationâ€Inducedâ€Emission Nanoparticles with Nearâ€Infrared Imaging at 1550Ânm Boosts Orthotopic Glioblastoma Theranostics. Advanced Materials, 2022, 34, e2106082.	11.1	75
2	Enhanced Antioxidant Effects of the Anti-Inflammatory Compound Probucol When Released from Mesoporous Silica Particles. Pharmaceutics, 2022, 14, 502.	2.0	5
3	Brain-Targeted Codelivery of Bcl-2/Bcl-xl and Mcl-1 Inhibitors by Biomimetic Nanoparticles for Orthotopic Glioblastoma Therapy. ACS Nano, 2022, 16, 6293-6308.	7.3	40
4	TDP-43 is a ubiquitylation substrate of the SCFcyclin F complex. Neurobiology of Disease, 2022, 167, 105673.	2.1	11
5	Aspect Ratio of PEGylated Upconversion Nanocrystals Affects the Cellular Uptake In Vitro and In Vivo. Acta Biomaterialia, 2022, 147, 403-413.	4.1	11
6	In vivo Validation of Bimolecular Fluorescence Complementation (BiFC) to Investigate Aggregate Formation in Amyotrophic Lateral Sclerosis (ALS). Molecular Neurobiology, 2021, 58, 2061-2074.	1.9	5
7	ALS/FTD-causing mutation in cyclin F causes the dysregulation of SFPQ. Human Molecular Genetics, 2021, 30, 971-984.	1.4	16
8	Unbiased Label-Free Quantitative Proteomics of Cells Expressing Amyotrophic Lateral Sclerosis (ALS) Mutations in CCNF Reveals Activation of the Apoptosis Pathway: A Workflow to Screen Pathogenic Gene Mutations. Frontiers in Molecular Neuroscience, 2021, 14, 627740.	1.4	12
9	Nanotechnologyâ€Based Strategies for Early Diagnosis of Central Nervous System Disorders. Advanced NanoBiomed Research, 2021, 1, 2100008.	1.7	16
10	Riluzole does not ameliorate disease caused by cytoplasmic TDPâ€43 in a mouse model of amyotrophic lateral sclerosis. European Journal of Neuroscience, 2021, 54, 6237-6255.	1.2	15
11	Splicing factor proline and glutamine rich intron retention, reduced expression and aggregate formation are pathological features of amyotrophic lateral sclerosis. Neuropathology and Applied Neurobiology, 2021, 47, 990-1003.	1.8	11
12	Polymeric nanoparticle mediated inhibition of miR-21 with enhanced miR-124 expression for combinatorial glioblastoma therapy. Biomaterials, 2021, 276, 121036.	5.7	29
13	Cytokine Signalling at the Microglial Penta-Partite Synapse. International Journal of Molecular Sciences, 2021, 22, 13186.	1.8	13
14	A Robust Intrinsically Green Fluorescent Poly(Amidoamine) Dendrimer for Imaging and Traceable Central Nervous System Delivery in Zebrafish. Small, 2020, 16, 2003654.	5.2	8
15	Foreword to the special issue on zebrafish imaging: Emerging techniques and methodologies. Micron, 2020, 136, 102877.	1.1	0
16	Observation and characterisation of macrophages in zebrafish liver. Micron, 2020, 132, 102851.	1.1	7
17	Muscle specific kinase protects dystrophic <i>mdx</i> mouse muscles from eccentric contractionâ€induced loss of forceâ€producing capacity. Journal of Physiology, 2019, 597, 4831-4850.	1.3	11
18	Using proteomics to identify ubiquitin ligase–substrate pairs: how novel methods may unveil therapeutic targets for neurodegenerative diseases. Cellular and Molecular Life Sciences, 2019, 76, 2499-2510.	2.4	18

Marco Morsch

#	Article	IF	CITATIONS
19	Albumin uptake and distribution in the zebrafish liver as observed via correlative imaging. Experimental Cell Research, 2019, 374, 162-171.	1.2	8
20	Aurora kinase B regulates axonal outgrowth and regeneration in the spinal motor neurons of developing zebrafish. Cellular and Molecular Life Sciences, 2018, 75, 4269-4285.	2.4	17
21	DNA nanoclew templated spherical nucleic acids for siRNA delivery. Chemical Communications, 2018, 54, 3609-3612.	2.2	50
22	Selective Spatiotemporal Vulnerability of Central Nervous System Neurons to Pathologic TAR DNA-Binding Protein 43 in Aged Transgenic Mice. American Journal of Pathology, 2018, 188, 1447-1456.	1.9	8
23	Cannabinoid-induced increase of quantal size and enhanced neuromuscular transmission. Scientific Reports, 2018, 8, 4685.	1.6	17
24	Pathogenic mutation in the ALS/FTD gene, CCNF, causes elevated Lys48-linked ubiquitylation and defective autophagy. Cellular and Molecular Life Sciences, 2018, 75, 335-354.	2.4	44
25	The mouse passiveâ€ŧransfer model of MuSK myasthenia gravis: disrupted MuSK signaling causes synapse failure. Annals of the New York Academy of Sciences, 2018, 1412, 54-61.	1.8	8
26	Real-time visualization of oxidative stress-mediated neurodegeneration of individual spinal motor neurons in vivo. Redox Biology, 2018, 19, 226-234.	3.9	41
27	Nucleo-cytoplasmic transport of TDP-43 studied in real time: impaired microglia function leads to axonal spreading of TDP-43 in degenerating motor neurons. Acta Neuropathologica, 2018, 136, 445-459.	3.9	66
28	Utility and reliability of nonâ€invasive muscle function tests in highâ€fatâ€fed mice. Experimental Physiology, 2017, 102, 773-778.	0.9	17
29	Expression of ALS/FTD-linked mutant CCNF in zebrafish leads to increased cell death in the spinal cord and an aberrant motor phenotype. Human Molecular Genetics, 2017, 26, 2616-2626.	1.4	44
30	Relocation is the key to successful correlative fluorescence and scanning electron microscopy. Methods in Cell Biology, 2017, 140, 215-244.	0.5	5
31	Multifunctional Hybrid Nanoparticles for Traceable Drug Delivery and Intracellular Microenvironmentâ€Controlled Multistage Drugâ€Release in Neurons. Small, 2017, 13, 1603966.	5.2	21
32	A versatile upconversion surface evaluation platform for bio–nano surface selection for the nervous system. Nanoscale, 2017, 9, 13683-13692.	2.8	13
33	Triggering Cell Stress and Death Using Conventional UV Laser Confocal Microscopy. Journal of Visualized Experiments, 2017, , .	0.2	13
34	Casein kinase II phosphorylation of cyclin F at serine 621 regulates the Lys48-ubiquitylation E3 ligase activity of the SCF (cyclin F) complex. Open Biology, 2017, 7, 170058.	1.5	29
35	A Tol2 Gateway-Compatible Toolbox for the Study of the Nervous System and Neurodegenerative Disease. Zebrafish, 2017, 14, 69-72.	0.5	56
36	Improving the Delivery of SOD1 Antisense Oligonucleotides to Motor Neurons Using Calcium Phosphate-Lipid Nanoparticles. Frontiers in Neuroscience, 2017, 11, 476.	1.4	53

Marco Morsch

#	Article	IF	CITATIONS
37	Ultrastructural Mapping of the Zebrafish Gastrointestinal System as a Basis for Experimental Drug Studies. BioMed Research International, 2016, 2016, 1-13.	0.9	14
38	The effects of high-fat feeding on physical function and skeletal muscle extracellular matrix. Nutrition and Diabetes, 2015, 5, e187-e187.	1.5	24
39	Forced expression of muscle specific kinase slows postsynaptic acetylcholine receptor loss in a mouse model of MuSK myasthenia gravis. Physiological Reports, 2015, 3, e12658.	0.7	13
40	Assessment of neuro-muscular function tests in mouse models of obesity and diabetes. Journal of the Neurological Sciences, 2015, 357, e203.	0.3	0
41	In vivo characterization of microglial engulfment of dying neurons in the zebrafish spinal cord. Frontiers in Cellular Neuroscience, 2015, 9, 321.	1.8	91
42	The established and emerging roles of astrocytes and microglia in amyotrophic lateral sclerosis and frontotemporal dementia. Frontiers in Cellular Neuroscience, 2015, 9, 414.	1.8	90
43	Electrophysiological analysis of neuromuscular synaptic function in myasthenia gravis patients and animal models. Experimental Neurology, 2015, 270, 41-54.	2.0	43
44	Muscleâ€specific kinase (MuSK) autoantibodies suppress the MuSK pathway and ACh receptor retention at the mouse neuromuscular junction. Journal of Physiology, 2014, 592, 2881-2897.	1.3	29
45	Clinical and scientific aspects of muscle-specific tyrosine kinase-related myasthenia gravis. Current Opinion in Neurology, 2014, 27, 558-565.	1.8	26
46	The Neuromuscular Junction: Measuring Synapse Size, Fragmentation and Changes in Synaptic Protein Density Using Confocal Fluorescence Microscopy. Journal of Visualized Experiments, 2014, , .	0.2	24
47	Effects of the ß2-Adrenoceptor Agonist, Albuterol, in a Mouse Model of Anti-MuSK Myasthenia Gravis. PLoS ONE, 2014, 9, e87840.	1.1	44
48	Pyridostigmine but not 3,4â€diaminopyridine exacerbates ACh receptor loss and myasthenia induced in mice by muscleâ€specific kinase autoantibody. Journal of Physiology, 2013, 591, 2747-2762.	1.3	63
49	Sequence of Age-Associated Changes to the Mouse Neuromuscular Junction and the Protective Effects of Voluntary Exercise. PLoS ONE, 2013, 8, e67970.	1.1	63
50	Muscle specific kinase autoantibodies cause synaptic failure through progressive wastage of postsynaptic acetylcholine receptors. Experimental Neurology, 2012, 237, 286-295.	2.0	50
51	Muscle Specific Kinase: Organiser of synaptic membrane domains. International Journal of Biochemistry and Cell Biology, 2011, 43, 295-298.	1.2	60
52	Modulation of the Ca2+ conductance of nicotinic acetylcholine receptors by Lypd6. European Neuropsychopharmacology, 2009, 19, 670-681.	0.3	49