

Brett M Morrison

List of Publications by Year in descending order

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Version: 2024-02-01

28
papers

3,094
citations

394421

19
h-index

610901

24
g-index

30
all docs

30
docs citations

30
times ranked

4516
citing authors

#	ARTICLE	IF	CITATIONS
1	MCT1 Deletion in Oligodendrocyte Lineage Cells Causes Late-Onset Hypomyelination and Axonal Degeneration. <i>Cell Reports</i> , 2021, 34, 108610.	6.4	65
2	Metabolic support of tumour-infiltrating regulatory T cells by lactic acid. <i>Nature</i> , 2021, 591, 645-651.	27.8	492
3	Macrophage monocarboxylate transporter 1 promotes peripheral nerve regeneration after injury in mice. <i>Journal of Clinical Investigation</i> , 2021, 131, .	8.2	29
4	Metabolic Transporters in the Peripheral Nerve—What, Where, and Why?. <i>Neurotherapeutics</i> , 2021, 18, 2185-2199.	4.4	5
5	Monocarboxylate transporter 1 in Schwann cells contributes to maintenance of sensory nerve myelination during aging. <i>Glia</i> , 2020, 68, 161-177.	4.9	46
6	Absence of Survival and Motor Deficits in 500 Repeat C9ORF72 BAC Mice. <i>Neuron</i> , 2020, 108, 775-783.e4.	8.1	33
7	Reducing monocarboxylate transporter MCT1 worsens experimental diabetic peripheral neuropathy. <i>Experimental Neurology</i> , 2020, 333, 113415.	4.1	11
8	Lactate Transporters Mediate Glia-Neuron Metabolic Crosstalk in Homeostasis and Disease. <i>Frontiers in Cellular Neuroscience</i> , 2020, 14, 589582.	3.7	35
9	Surprising New Players in Glia-Neuron Crosstalk: Role in CNS Regeneration. <i>Cell Metabolism</i> , 2020, 32, 695-696.	16.2	1
10	Glia-neuron energy metabolism in health and diseases: New insights into the role of nervous system metabolic transporters. <i>Experimental Neurology</i> , 2018, 309, 23-31.	4.1	123
11	Neuromuscular Diseases. <i>Seminars in Neurology</i> , 2016, 36, 409-418.	1.4	59
12	Motor neuron disease, TDP-43 pathology, and memory deficits in mice expressing ALS/FTD-linked <i>UBQLN2</i> mutations. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2016, 113, E7580-E7589.	7.1	77
13	Deficiency in monocarboxylate transporter 1 (MCT1) in mice delays regeneration of peripheral nerves following sciatic nerve crush. <i>Experimental Neurology</i> , 2015, 263, 325-338.	4.1	71
14	Oligodendroglia: metabolic supporters of axons. <i>Trends in Cell Biology</i> , 2013, 23, 644-651.	7.9	196
15	Amyotrophic Lateral Sclerosis and Novel Therapeutic Strategies. <i>Neurology Research International</i> , 2012, 2012, 1-3.	1.3	2
16	Medication, Toxic, and Vitamin-Related Neuropathies. <i>CONTINUUM Lifelong Learning in Neurology</i> , 2012, 18, 139-160.	0.8	5
17	Oligodendroglia metabolically support axons and contribute to neurodegeneration. <i>Nature</i> , 2012, 487, 443-448.	27.8	1,287
18	Expanding the spectrum of monoclonal light chain deposition disease in muscle. <i>Muscle and Nerve</i> , 2012, 45, 755-761.	2.2	15

#	ARTICLE	IF	CITATIONS
19	Approach to the Patient with Abnormal Cerebrospinal Fluid Glucose Content. , 2009, , 281-285.		0
20	A soluble activin type IIB receptor improves function in a mouse model of amyotrophic lateral sclerosis. <i>Experimental Neurology</i> , 2009, 217, 258-268.	4.1	75
21	Magnetic resonance imaging of mouse skeletal muscle to measure denervation atrophy. <i>Experimental Neurology</i> , 2008, 212, 448-457.	4.1	58
22	Genetically Decreased Spinal Cord Copper Concentration Prolongs Life in a Transgenic Mouse Model of Amyotrophic Lateral Sclerosis. <i>Journal of Neuroscience</i> , 2004, 24, 7945-7950.	3.6	50
23	Early and Selective Pathology of Light Chain Neurofilament in the Spinal Cord and Sciatic Nerve of G86R Mutant Superoxide Dismutase Transgenic Mice. <i>Experimental Neurology</i> , 2000, 165, 207-220.	4.1	27
24	Amyotrophic lateral sclerosis associated with mutations in superoxide dismutase: a putative mechanism of degeneration. <i>Brain Research Reviews</i> , 1999, 29, 121-135.	9.0	78
25	Time course of neuropathology in the spinal cord of G86R superoxide dismutase transgenic mice. <i>Journal of Comparative Neurology</i> , 1998, 391, 64-77.	1.6	91
26	Light and electron microscopic distribution of the AMPA receptor subunit, GluR2, in the spinal cord of control and G86R mutant superoxide dismutase transgenic mice. , 1998, 395, 523-534.		57
27	Superoxide dismutase and neurofilament transgenic models of amyotrophic lateral sclerosis. <i>The Journal of Experimental Zoology</i> , 1998, 282, 32-47.	1.4	23
28	Quantitative immunocytochemical analysis of the spinal cord in G86R superoxide dismutase transgenic mice: Neurochemical correlates of selective vulnerability. , 1996, 373, 619-631.		83