

Josep E Esquerda

List of Publications by Year in descending order

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60
papers

1,744
citations

257357

24
h-index

315616

38
g-index

60
all docs

60
docs citations

60
times ranked

1921
citing authors

#	ARTICLE	IF	CITATIONS
1	Accumulation of misfolded SOD1 outlines distinct patterns of motor neuron pathology and death during disease progression in a SOD1 ^{G93A} mouse model of amyotrophic lateral sclerosis. <i>Brain Pathology</i> , 2022, 32, .	2.1	6
2	Microglial recruitment and mechanisms involved in the disruption of afferent synaptic terminals on spinal cord motor neurons after acute peripheral nerve injury. <i>Glia</i> , 2021, 69, 1216-1240.	2.5	22
3	Cover Image, Volume 69, Issue 5. <i>Glia</i> , 2021, 69, C1.	2.5	0
4	Beneficial effects of dietary supplementation with green tea catechins and cocoa flavanols on aging-related regressive changes in the mouse neuromuscular system. <i>Aging</i> , 2021, 13, 18051-18093.	1.4	4
5	Motoneuron deafferentation and gliosis occur in association with neuromuscular regressive changes during ageing in mice. <i>Journal of Cachexia, Sarcopenia and Muscle</i> , 2020, 11, 1628-1660.	2.9	21
6	Localization and dynamic changes of neuregulin-1 at C-type synaptic boutons in association with motor neuron injury and repair. <i>FASEB Journal</i> , 2019, 33, 7833-7851.	0.2	30
7	The Y172 Monoclonal Antibody Against p-c-Jun (Ser63) Is a Marker of the Postsynaptic Compartment of C-Type Cholinergic Afferent Synapses on Motoneurons. <i>Frontiers in Cellular Neuroscience</i> , 2019, 13, 582.	1.8	1
8	Glial Activation and Central Synapse Loss, but Not Motoneuron Degeneration, Are Prevented by the Sigma-1 Receptor Agonist PRE-084 in the Smn2B ^{+/+} Mouse Model of Spinal Muscular Atrophy. <i>Journal of Neuropathology and Experimental Neurology</i> , 2018, 77, 577-597.	0.9	30
9	Neuregulin 1-ErbB module in C-bouton synapses on somatic motor neurons: molecular compartmentation and response to peripheral nerve injury. <i>Scientific Reports</i> , 2017, 7, 40155.	1.6	32
10	Chronic Treatment with the AMPK Agonist AICAR Prevents Skeletal Muscle Pathology but Fails to Improve Clinical Outcome in a Mouse Model of Severe Spinal Muscular Atrophy. <i>Neurotherapeutics</i> , 2016, 13, 198-216.	2.1	27
11	Adverse effects of a SOD1-peptide immunotherapy on SOD1G93A mouse slow model of amyotrophic lateral sclerosis. <i>Neuroscience</i> , 2015, 310, 38-50.	1.1	7
12	Accumulation of Misfolded SOD1 in Dorsal Root Ganglion Degenerating Proprioceptive Sensory Neurons of Transgenic Mice with Amyotrophic Lateral Sclerosis. <i>BioMed Research International</i> , 2014, 2014, 1-13.	0.9	38
13	Mechanisms Involved in Spinal Cord Central Synapse Loss in a Mouse Model of Spinal Muscular Atrophy. <i>Journal of Neuropathology and Experimental Neurology</i> , 2014, 73, 519-535.	0.9	57
14	Neuregulin-1 is concentrated in the postsynaptic subsurface cistern of C-bouton inputs to motoneurons and altered during motoneuron diseases. <i>FASEB Journal</i> , 2014, 28, 3618-3632.	0.2	65
15	Chronic treatment with lithium does not improve neuromuscular phenotype in a mouse model of severe spinal muscular atrophy. <i>Neuroscience</i> , 2013, 250, 417-433.	1.1	8
16	Synaptic defects in type I spinal muscular atrophy in human development. <i>Journal of Pathology</i> , 2013, 229, 49-61.	2.1	77
17	Immunodetection of Disease-Associated Conformers of Mutant Cu/Zn Superoxide Dismutase 1 Selectively Expressed in Degenerating Neurons in Amyotrophic Lateral Sclerosis. <i>Journal of Neuropathology and Experimental Neurology</i> , 2013, 72, 646-661.	0.9	17
18	Sera from amyotrophic lateral sclerosis patients induce the non-canonical activation of NMDA receptors <i>in vitro</i> . <i>Neurochemistry International</i> , 2011, 59, 954-964.	1.9	16

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19	Defective Neuromuscular Junction Organization and Postnatal Myogenesis in Mice With Severe Spinal Muscular Atrophy. <i>Journal of Neuropathology and Experimental Neurology</i> , 2011, 70, 444-461.	0.9	68
20	Increased intramuscular nerve branching and inhibition of programmed cell death of chick embryo motoneurons by immunoglobulins from patients with motoneuron disease. <i>Journal of Neuroimmunology</i> , 2010, 229, 157-168.	1.1	3
21	Neurotoxic Species of Misfolded SOD1 ^{G93A} Recognized by Antibodies Against the P2X ₄ Subunit of the ATP Receptor Accumulate in Damaged Neurons of Transgenic Animal Models of Amyotrophic Lateral Sclerosis. <i>Journal of Neuropathology and Experimental Neurology</i> , 2010, 69, 176-187.	0.9	23
22	Lithium prevents excitotoxic cell death of motoneurons in organotypic slice cultures of spinal cord. <i>Neuroscience</i> , 2010, 165, 1353-1369.	1.1	58
23	Excitotoxic motoneuron degeneration induced by glutamate receptor agonists and mitochondrial toxins in organotypic cultures of chick embryo spinal cord. <i>Journal of Comparative Neurology</i> , 2009, 516, 277-290.	0.9	21
24	Development of microglia in the chick embryo spinal cord: Implications in the regulation of motoneuronal survival and death. <i>Journal of Neuroscience Research</i> , 2009, 87, 2447-2466.	1.3	48
25	Caspase-independent type III programmed cell death in chronic lymphocytic leukemia: the key role of the F-actin cytoskeleton. <i>Haematologica</i> , 2009, 94, 507-517.	1.7	26
26	Strong P2X ₄ purinergic receptor-like immunoreactivity is selectively associated with degenerating neurons in transgenic rodent models of amyotrophic lateral sclerosis. <i>Journal of Comparative Neurology</i> , 2008, 506, 75-92.	0.9	65
27	The rescue of developing avian motoneurons from programmed cell death by a selective inhibitor of the fetal muscle-specific nicotinic acetylcholine receptor. <i>Developmental Neurobiology</i> , 2008, 68, 972-980.	1.5	15
28	7-Bromoindirubin-3-oxime uncovers a serine protease-mediated paradigm of necrotic cell death. <i>Biochemical Pharmacology</i> , 2008, 76, 39-52.	2.0	22
29	Drp1 Mediates Caspase-Independent Type III Cell Death in Normal and Leukemic Cells. <i>Molecular and Cellular Biology</i> , 2007, 27, 7073-7088.	1.1	98
30	Survival and death of mature avian motoneurons in organotypic slice culture: Trophic requirements for survival and different types of degeneration. <i>Journal of Comparative Neurology</i> , 2007, 501, 669-690.	0.9	30
31	Excitotoxic motoneuron disease in chick embryo evolves with autophagic neurodegeneration and deregulation of neuromuscular innervation. <i>Journal of Neuroscience Research</i> , 2007, 85, 2726-2740.	1.3	15
32	Protein retention in the endoplasmic reticulum, blockade of programmed cell death and autophagy selectively occur in spinal cord motoneurons after glutamate receptor-mediated injury. <i>Molecular and Cellular Neurosciences</i> , 2005, 29, 283-298.	1.0	45
33	Rescue of developing spinal motoneurons from programmed cell death by the GABA _A agonist muscimol acts by blockade of neuromuscular activity and increased intramuscular nerve branching. <i>Molecular and Cellular Neurosciences</i> , 2003, 22, 331-343.	1.0	31
34	c-Jun-like Immunoreactivity in Apoptosis Is the Result of a Crossreaction with Neoantigenic Sites Exposed by Caspase-3-mediated Proteolysis. <i>Journal of Histochemistry and Cytochemistry</i> , 2002, 50, 961-972.	1.3	11
35	In Vivo Analysis of Schwann Cell Programmed Cell Death in the Embryonic Chick: Regulation by Axons and Glial Growth Factor. <i>Journal of Neuroscience</i> , 2002, 22, 4509-4521.	1.7	36
36	Occurrence of glutamate receptor subunit 1-containing aggresome-like structures during normal development of rat spinal cord interneurons. <i>Journal of Comparative Neurology</i> , 2002, 442, 23-34.	0.9	17

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37	Long-Lasting Aberrant Tubulovesicular Membrane Inclusions Accumulate in Developing Motoneurons after a Sublethal Excitotoxic Insult: A Possible Model for Neuronal Pathology in Neurodegenerative Disease. <i>Journal of Neuroscience</i> , 2001, 21, 8072-8081.	1.7	25
38	Antibodies against c-Jun N-terminal peptide cross-react with neo-epitopes emerging after caspase-mediated proteolysis during apoptosis. <i>Journal of Neurochemistry</i> , 2001, 77, 904-915.	2.1	14
39	c-Jun regulation in rat neonatal motoneurons postaxotomy. <i>Journal of Neuroscience Research</i> , 2001, 63, 469-479.	1.3	21
40	Induction of reactive astrocytosis and prevention of motoneuron cell death by the I2 -imidazoline receptor ligand LSL 60101. <i>British Journal of Pharmacology</i> , 2000, 130, 1767-1776.	2.7	28
41	Opposing Effects of Excitatory Amino Acids on Chick Embryo Spinal Cord Motoneurons: Excitotoxic Degeneration or Prevention of Programmed Cell Death. <i>Journal of Neuroscience</i> , 1999, 19, 10803-10812.	1.7	43
42	Specific association of c-Jun-like immunoreactivity but not c-Jun p39 with normal and induced programmed cell death in the chick embryo. , 1999, 38, 171-190.		27
43	Nitric oxide synthase in rat neuromuscular junctions and in nerve terminals of Torpedo electric organ: its role as regulator of acetylcholine release. , 1998, 51, 90.		51
44	Effects of excitatory amino acids on neuromuscular development in the chick embryo. , 1997, 387, 73-95.		36
45	The carbohydrate N-acetylglucosamine is involved in the guidance of neurites from chick ciliary ganglion neurons through the extracellular matrix of rat skeletal muscle fiber. <i>Neuroscience Letters</i> , 1996, 207, 81-84.	1.0	6
46	Prevention by lamotrigine, MK-801 and N ^ω -nitro-l-arginine methyl ester of motoneuron cell death after neonatal axotomy. <i>Neuroscience</i> , 1996, 71, 313-325.	1.1	60
47	Intramuscular nerve sprouting induced by CNTF is associated with increases in CGRP content in mouse motor nerve terminals. <i>Neuroscience Letters</i> , 1996, 219, 60-64.	1.0	18
48	Schwann Cell Apoptosis during Normal Development and after Axonal Degeneration Induced by Neurotoxins in the Chick Embryo. <i>Journal of Neuroscience</i> , 1996, 16, 3979-3990.	1.7	75
49	Regulation of Motoneuronal Calcitonin Gene-related Peptide (CGRP) During Axonal Growth and Neuromuscular Synaptic Plasticity Induced by Botulinum Toxin in Rats. <i>European Journal of Neuroscience</i> , 1996, 8, 829-836.	1.2	63
50	S-laminin and N-acetylgalactosamine located at the synaptic basal lamina of skeletal muscle are involved in synaptic recognition by growing neurites. <i>Journal of Neurocytology</i> , 1995, 24, 903-915.	1.6	6
51	Evidence for calcium regulation of spinal cord motoneuron death in the chick embryo in vivo. <i>Developmental Brain Research</i> , 1995, 86, 167-179.	2.1	12
52	Calcitonin gene-related peptide in rat spinal cord motoneurons: Subcellular distribution and changes induced by axotomy. <i>Neuroscience</i> , 1992, 48, 449-461.	1.1	66
53	Treatment with digestive agents reveals several glycoconjugates specifically associated with rat neuromuscular junction. <i>Histochemistry</i> , 1992, 97, 125-131.	1.9	16
54	Absence of histochemical immunoreactivity to calcitonin gene-related peptide (CGRP) in spinal cord motoneurons from (+)-tubocurarine-treated chick embryos. <i>Neuroscience Letters</i> , 1989, 105, 1-6.	1.0	18

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55	Synaptic localization of a 66-kDa soluble protein from skeletal muscle: Evidence for its developmental and neural regulation. <i>Experimental Neurology</i> , 1989, 105, 211-218.	2.0	0
56	Phylogenetic polymorphism on lectin binding to junctional and non-junctional basal lamina at the vertebrate neuromuscular junction. <i>Histochemistry</i> , 1987, 87, 301-307.	1.9	12
57	Receptors to agglutinin from <i>Dolichus biflorus</i> (DBA) at the synaptic basal lamina of rat neuromuscular junction. <i>Cell and Tissue Research</i> , 1987, 248, 111-117.	1.5	14
58	Structural changes at pure cholinergic synaptosomes during the transmitter release induced by A-23187 in <i>Torpedo marmorata</i> . <i>Cell and Tissue Research</i> , 1987, 248, 207-214.	1.5	21
59	Binding of 125 I-bungarotoxin to <i>Torpedo</i> electric organ synaptosomes. A high resolution autoradiographic study. <i>Neuroscience</i> , 1982, 7, 751-758.	1.1	16
60	Ionic dependence of adenosine uptake by isolated nerve endings from <i>Torpedo</i> electric organ. <i>Neurochemistry International</i> , 1982, 4, 513-521.	1.9	6