

Alberto Garc a-Redondo

List of Publications by Year in descending order

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Version: 2024-02-01

20
papers

1,492
citations

623734

14
h-index

888059

17
g-index

20
all docs

20
docs citations

20
times ranked

2893
citing authors

#	ARTICLE	IF	CITATIONS
1	Molecular Alterations in Sporadic and SOD1-ALS Immortalized Lymphocytes: Towards a Personalized Therapy. <i>International Journal of Molecular Sciences</i> , 2021, 22, 3007.	4.1	16
2	Type XIX collagen: a promising biomarker from the basement membranes. <i>Neural Regeneration Research</i> , 2020, 15, 988.	3.0	13
3	Collagen XIX Alpha 1 Improves Prognosis in Amyotrophic Lateral Sclerosis. , 2019, 10, 278.		18
4	Circulating Cytokines Could Not Be Good Prognostic Biomarkers in a Mouse Model of Amyotrophic Lateral Sclerosis. <i>Frontiers in Immunology</i> , 2019, 10, 801.	4.8	16
5	Mutations in the vesicular trafficking protein annexin A11 are associated with amyotrophic lateral sclerosis. <i>Science Translational Medicine</i> , 2017, 9, .	12.4	129
6	Comparative study of hematopoietic stem and progenitor cells between sexes in mice under physiological conditions along time. <i>Cell Biology International</i> , 2017, 41, 1399-1405.	3.0	0
7	Granulocyte Colony-Stimulating Factor Ameliorates Skeletal Muscle Dysfunction in Amyotrophic Lateral Sclerosis Mice and Improves Proliferation of SOD1-G93A Myoblasts in vitro. <i>Neurodegenerative Diseases</i> , 2017, 17, 1-13.	1.4	11
8	Hematopoietic stem and progenitor cells as novel prognostic biomarkers of longevity in a murine model for amyotrophic lateral sclerosis. <i>American Journal of Physiology - Cell Physiology</i> , 2016, 311, C910-C919.	4.6	0
9	Neuregulin-1 promotes functional improvement by enhancing collateral sprouting in SOD1G93A ALS mice and after partial muscle denervation. <i>Neurobiology of Disease</i> , 2016, 95, 168-178.	4.4	44
10	NEK1 variants confer susceptibility to amyotrophic lateral sclerosis. <i>Nature Genetics</i> , 2016, 48, 1037-1042.	21.4	218
11	CCNF mutations in amyotrophic lateral sclerosis and frontotemporal dementia. <i>Nature Communications</i> , 2016, 7, 11253.	12.8	174
12	Analysis of the <i>CHCHD10</i> gene in patients with frontotemporal dementia and amyotrophic lateral sclerosis from Spain. <i>Brain</i> , 2015, 138, e400-e400.	7.6	56
13	MicroRNA-206: A Potential Circulating Biomarker Candidate for Amyotrophic Lateral Sclerosis. <i>PLoS ONE</i> , 2014, 9, e89065.	2.5	154
14	Characterization of the repeat expansion size in C9orf72 in amyotrophic lateral sclerosis and frontotemporal dementia. <i>Human Molecular Genetics</i> , 2014, 23, 749-754.	2.9	98
15	Exome-wide Rare Variant Analysis Identifies TUBA4A Mutations Associated with Familial ALS. <i>Neuron</i> , 2014, 84, 324-331.	8.1	308
16	Genetic Biomarkers for ALS Disease in Transgenic SOD1G93A Mice. <i>PLoS ONE</i> , 2012, 7, e32632.	2.5	53
17	Altered Expression of Myogenic Regulatory Factors in the Mouse Model of Amyotrophic Lateral Sclerosis. <i>Neurodegenerative Diseases</i> , 2011, 8, 386-396.	1.4	39
18	Oxidative stress in skin fibroblasts cultures from patients with Parkinson's disease. <i>BMC Neurology</i> , 2010, 10, 95.	1.8	37

#	ARTICLE	IF	CITATIONS
19	Oxidative Stress in Skin Fibroblasts Cultures of Patients with Huntington's Disease. <i>Neurochemical Research</i> , 2006, 31, 1103-1109.	3.3	57
20	Early onset multisystem mitochondrial disorder caused by a nonsense mutation in the mitochondrial DNA Cytochrome C oxidase II gene. <i>Annals of Neurology</i> , 2001, 50, 409-413.	5.3	51