Maite Mendioroz

List of Publications by Year
in descending order

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49 papers	1,402 citations	377584 21 h-index	406436 35 g-index
57 all docs	57 docs citations	57 times ranked	2708 citing authors

#	Article	IF	Citations
1	NXN Gene Epigenetic Changes in an Adult Neurogenesis Model of Alzheimerâ€™s Disease. Cells, 2022, 11, 1069.	1.8	3
2	Role of Biomarkers for the Diagnosis of Prion Diseases: A Narrative Review. Medicina (Lithuania), 2022, 58, 473.	0.8	6
3	Profile of TREM2-Derived circRNA and mRNA Variants in the Entorhinal Cortex of Alzheimerâ€™s Disease Patients. International Journal of Molecular Sciences, 2022, 23, 7682.	1.8	6
4	Modificaciones epigenÃ©ticas en las cefaleas. NeurologÃa, 2021, 36, 369-376.	0.3	8
5	The Participation of Microglia in Neurogenesis: A Review. Brain Sciences, 2021, 11, 658.	1.1	29
6	Gender-Dependent Deregulation of Linear and Circular RNA Variants of HOMER1 in the Entorhinal Cortex of Alzheimerâ€™s Disease. International Journal of Molecular Sciences, 2021, 22, 9205.	1.8	13
7	Telomere length correlates with subtelomeric DNA methylation in long-term mindfulness practitioners. Scientific Reports, 2020, 10, 4564.	1.6	21
8	Early epigenetic changes of Alzheimerâ€™s disease in the human hippocampus. Epigenetics, 2020, 15, 1083-1092.	1.3	11
9	Microgliaâ€Related Gene Triggering Receptor Expressed in Myeloid Cells 2 (TREM2) Is Upregulated in the Substantia Nigra of Progressive Supranuclear Palsy. Movement Disorders, 2020, 35, 885-890.	2.2	11
10	Globular glial tauopathy caused by MAPT P301T mutation: clinical and neuropathological findings. Journal of Neurology, 2019, 266, 2396-2405.	1.8	22
11	DNA methylation signature of human hippocampus in Alzheimerâ€™s disease is linked to neurogenesis. Clinical Epigenetics, 2019, 11, 91.	1.8	67
12	Hippocampal LMNA Gene Expression is Increased in Late-Stage Alzheimerâ€™s Disease. International Journal of Molecular Sciences, 2019, 20, 878.	1.8	17
13	PATJ Low Frequency Variants Are Associated With Worse Ischemic Stroke Functional Outcome. Circulation Research, 2019, 124, 114-120.	2.0	49
14	Epigenetic Response to Mindfulness in Peripheral Blood Leukocytes Involves Genes Linked to Common Human Diseases. Mindfulness, 2018, 9, 1146-1159.	1.6	30
15	PLD3 epigenetic changes in the hippocampus of Alzheimerâ€™s disease. Clinical Epigenetics, 2018, 10, 116.	1.8	21
16	Mass Spectrometry-Based Proteomic Profiling of Thrombotic Material Obtained by Endovascular Thrombectomy in Patients with Ischemic Stroke. International Journal of Molecular Sciences, 2018, 19, 498.	1.8	32
17	Evaluation of Chitotriosidase and CC-Chemokine Ligand 18 as Biomarkers of Microglia Activation in Amyotrophic Lateral Sclerosis. Neurodegenerative Diseases, 2018, 18, 208-215.	0.8	17
18	Liquid biopsy: a new source of candidate biomarkers in amyotrophic lateral sclerosis. Annals of Clinical and Translational Neurology, 2018, 5, 763-768.	1.7	14

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19	GRECOS Project (Genotyping Recurrence Risk of Stroke). Stroke, 2017, 48, 1147-1153.	1.0	23
20	Vitamin D receptor gene is epigenetically altered and transcriptionally up-regulated in multiple sclerosis. PLoS ONE, 2017, 12, e0174726.	1.1	26
21	CRTC1 gene is differentially methylated in the human hippocampus in Alzheimerâ€™s disease. Alzheimer's Research and Therapy, 2016, 8, 15.	3.0	28
22	TREM2 upregulation correlates with 5-hydroxymethycytosine enrichment in Alzheimerâ€™s disease hippocampus. Clinical Epigenetics, 2016, 8, 37.	1.8	68
23	Trans effects of chromosome aneuploidies on DNA methylation patterns in human Down syndrome and mouse models. Genome Biology, 2015, 16, 263.	3.8	68
24	Genes involved in hemorrhagic transformations that follow recombinant t-PA treatment in stroke patients. Pharmacogenomics, 2013, 14, 495-504.	0.6	18
25	Brain Natriuretic Peptide Is Associated with Worsening and Mortality in Acute Stroke Patients but Adds No Prognostic Value to Clinical Predictors of Outcome. Cerebrovascular Diseases, 2012, 34, 240-245.	0.8	32
26	Role of the MMP9 Gene in Hemorrhagic Transformations After Tissue-Type Plasminogen Activator Treatment in Stroke Patients. Stroke, 2012, 43, 1398-1400.	1.0	13
27	IL1BandVWFVariants Are Associated With Fibrinolytic Early Recanalization in Patients With Ischemic Stroke. Stroke, 2012, 43, 2659-2665.	1.0	28
28	A predictive clinicalâ€“genetic model of tissue plasminogen activator response in acute ischemic stroke. Annals of Neurology, 2012, 72, 716-729.	2.8	39
29	Differentiating ischemic from hemorrhagic stroke using plasma biomarkers: The S100B/RAGE pathway. Journal of Proteomics, 2012, 75, 4758-4765.	1.2	68
30	The gender gap in stroke: a meta-analysis. Acta Neurologica Scandinavica, 2012, 125, 83-90.	1.0	70
31	ACE variants and risk of intracerebral hemorrhage recurrence in amyloid angiopathy. Neurobiology of Aging, 2011, 32, 551.e13-551.e22.	1.5	22
32	Update on the Serum Biomarkers and Genetic Factors Associated with Safety and Efficacy of rt-PA Treatment in Acute Stroke Patients. Stroke Research and Treatment, 2011, 2011, 1-10.	0.5	3
33	A panel of biomarkers including caspase-3 and D-dimer may differentiate acute stroke from stroke-mimicking conditions in the emergency department. Journal of Internal Medicine, 2011, 270, 166-174.	2.7	61
34	Leukoaraiosis is associated with genes regulating blood-brain barrier homeostasis in ischaemic stroke patients. European Journal of Neurology, 2011, 18, 826-835.	1.7	24
35	No evidence of APP point mutation and locus duplication in individuals with cerebral amyloid angiopathy. European Journal of Neurology, 2011, 18, 1279-1281.	1.7	8
36	Osteopontin predicts long-term functional outcome among ischemic stroke patients. Journal of Neurology, 2011, 258, 486-493.	1.8	23

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37	The I/D polymorphism of the ACE1 gene is not associated with ischaemic stroke in Spanish individuals. European Journal of Neurology, 2010, 17, 1390-1392.	1.7	18
38	Association of a Genetic Variant in the <i>ALOX5AP</i> with Higher Risk of Ischemic Stroke: A Case-Control, Meta-Analysis and Functional Study. Cerebrovascular Diseases, 2010, 29, 528-537.	0.8	54
39	PAI-1 4G/5G Polymorphism is Associated with Brain Vessel Reocclusion After Successful Fibrinolytic Therapy in Ischemic Stroke Patients. International Journal of Neuroscience, 2010, 120, 245-251.	0.8	17
40	Stroke after prolonged air travel associated with a pulmonary arteriovenous malformation. Journal of the Neurological Sciences, 2010, 292, 99-100.	0.3	9
41	A missense HTRA1 mutation expands CARASIL syndrome to the Caucasian population. Neurology, 2010, 75, 2033-2035.	1.5	66
42	KCNK17 genetic variants in ischemic stroke. Atherosclerosis, 2010, 208, 203-209.	0.4	22
43	CD40-1C>T polymorphism (rs1883832) is associated with brain vessel reocclusion after fibrinolysis in ischemic stroke. Pharmacogenomics, 2010, 11, 763-772.	0.6	16
44	Lower concentrations of thrombin-antithrombin complex (TAT) correlate to higher recanalisation rates among ischaemic stroke patients treated with t-PA. Thrombosis and Haemostasis, 2009, 102, 759-764.	1.8	19
45	Endogenous Activated Protein C Predicts Hemorrhagic Transformation and Mortality after Tissue Plasminogen Activator Treatment in Stroke Patients. Cerebrovascular Diseases, 2009, 28, 143-150.	0.8	23
46	CADASIL management or what to do when there is little one can do. Expert Review of Neurotherapeutics, 2009, 9, 197-210.	1.4	20
47	Caspase-3 is related to infarct growth after human ischemic stroke. Neuroscience Letters, 2008, 430, 1-6.	1.0	36
48	Genetics of stroke: a review of recent advances. Expert Review of Molecular Diagnostics, 2008, 8, 495-513.	1.5	49
49	Influence of thrombinâ€activatable fibrinolysis inhibitor and plasminogen activator inhibitorâ€1 gene polymorphisms on tissueâ€type plasminogen activatorâ€induced recanalization in ischemic stroke patients. Journal of Thrombosis and Haemostasis, 2007, 5, 1862-1868.	1.9	49