Martin Medrano

List of Publications by Year
in descending order

15	456	933447 10	1125743 13
papers	citations	h-index	g-index

15	15	15	1323
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Progranulin mutations in clinical and neuropathological Alzheimer's disease. Alzheimer's and Dementia, 2022, 18, 2458-2467.	0.8	12
2	Younger age of stroke in lowâ€middle income countries is related to healthcare access and quality. Annals of Clinical and Translational Neurology, 2022, 9, 415-427.	3.7	17
3	FMNL2 regulates gliovascular interactions and is associated with vascular risk factors and cerebrovascular pathology in Alzheimerâ€™s disease. Acta Neuropathologica, 2022, 144, 59-79.	7.7	19
4	Vascular mild cognitive impairment and its relationship to hemoglobin A1c levels and apolipoprotein E genotypes in the Dominican Republic. Dementia E Neuropsychologia, 2021, 15, 69-78.	0.8	0
5	Rare Variants Imputation in Admixed Populations: Comparison Across Reference Panels and Bioinformatics Tools. Frontiers in Genetics, 2019, 10, 239.	2.3	20
6	Cerebrovascular Disease and Neurodegeneration in Alzheimerâ€™s Disease with and without a Strong Family History: A Pilot Magnetic Resonance Imaging Study in Dominican Republic. Journal of Alzheimer's Disease, 2018, 66, 1519-1528.	2.6	3
7	Linkage analysis of multiplex Caribbean Hispanic families loaded for unexplained earlyâ€onset cases identifies novel Alzheimer's disease loci. Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring, 2018, 10, 554-562.	2.4	8
8	An Alzheimer's Disease-Linked Loss-of-Function CLN5 Variant Impairs Cathepsin D Maturation, Consistent with a Retromer Trafficking Defect. Molecular and Cellular Biology, 2018, 38, .	2.3	34
9	Fâ€box/ <scp>LRR</scp> â€repeat protein 7 is genetically associated with Alzheimer's disease. Annals of Clinical and Translational Neurology, 2015, 2, 810-820.	3.7	54
10	Rare coding mutations identified by sequencing of <scp>A</scp>lzheimer disease genomeâ€wide association studies loci. Annals of Neurology, 2015, 78, 487-498.	5.3	126
11	Inbreeding among Caribbean Hispanics from the Dominican Republic and its effects on risk of Alzheimer disease. Genetics in Medicine, 2015, 17, 639-643.	2.4	20
12	Genetic Modifiers of Age at Onset in Carriers of the G206A Mutation inPSEN1With Familial Alzheimer Disease Among Caribbean Hispanics. JAMA Neurology, 2015, 72, 1043.	9.0	50
13	Diseaseâ€related mutations among Caribbean Hispanics with familial dementia. Molecular Genetics & Genomic Medicine, 2014, 2, 430-437.	1.2	36
14	P3-016: SORBS2, SH3RF3, AND NPHP1 MODIFY AGE AT ONSET IN CARRIERS OF THE G206A MUTATION IN PSEN1 WITH FAMILIAL ALZHEIMER'S DISEASE. , 2014, 10, P632-P632.		2
15	Familial Alzheimer Disease Among Caribbean Hispanics. Archives of Neurology, 2002, 59, 87.	4.5	55