

Eduardo Prez-Palma

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

20
papers

324
citations

11
h-index

17
g-index

23
ext. papers

521
ext. citations

9.2
avg, IF

2.98
L-index

#	Paper	IF	Citations
20	Wnt/βcatenin signaling in Alzheimer's disease. <i>CNS and Neurological Disorders - Drug Targets</i> , 2014 , 13, 745-54	2.6	58
19	Overrepresentation of glutamate signaling in Alzheimer's disease: network-based pathway enrichment using meta-analysis of genome-wide association studies. <i>PLoS ONE</i> , 2014 , 9, e95413	3.7	42
18	A novel functional low-density lipoprotein receptor-related protein 6 gene alternative splice variant is associated with Alzheimer's disease. <i>Neurobiology of Aging</i> , 2013 , 34, 1709.e9-18	5.6	27
17	Predicting functional effects of missense variants in voltage-gated sodium and calcium channels. <i>Science Translational Medicine</i> , 2020 , 12,	17.5	27
16	Spectrum of neurodevelopmental disease associated with the GNAO1 guanosine triphosphate-binding region. <i>Epilepsia</i> , 2019 , 60, 406-418	6.4	26
15	Spectrum of GABAA receptor variants in epilepsy. <i>Current Opinion in Neurology</i> , 2019 , 32, 183-190	7.1	24
14	Simple ClinVar: an interactive web server to explore and retrieve gene and disease variants aggregated in ClinVar database. <i>Nucleic Acids Research</i> , 2019 , 47, W99-W105	20.1	21
13	Variants in ABCG8 and TRAF3 genes confer risk for gallstone disease in admixed Latinos with Mapuche Native American ancestry. <i>Scientific Reports</i> , 2019 , 9, 772	4.9	16
12	Early Transcriptional Changes Induced by Wnt/-Catenin Signaling in Hippocampal Neurons. <i>Neural Plasticity</i> , 2016 , 2016, 4672841	3.3	16
11	Heterogeneous contribution of microdeletions in the development of common generalised and focal epilepsies. <i>Journal of Medical Genetics</i> , 2017 , 54, 598-606	5.8	14
10	Identification of pathogenic variant enriched regions across genes and gene families. <i>Genome Research</i> , 2020 , 30, 62-71	9.7	14
9	Current knowledge of SLC6A1-related neurodevelopmental disorders. <i>Brain Communications</i> , 2020 , 2, fcaa170	4.5	11
8	Whole Genome Sequence, Variant Discovery and Annotation in Mapuche-Huilliche Native South Americans. <i>Scientific Reports</i> , 2019 , 9, 2132	4.9	6
7	Variant Score Ranker-a web application for intuitive missense variant prioritization. <i>Bioinformatics</i> , 2019 , 35, 4478-4479	7.2	5
6	Intestinal-Cell Kinase and Juvenile Myoclonic Epilepsy. <i>New England Journal of Medicine</i> , 2019 , 380, e24	59.2	4
5	Assessment of genetic variant burden in epilepsy-associated brain lesions. <i>European Journal of Human Genetics</i> , 2019 , 27, 1738-1744	5.3	4
4	Genome-Wide Analysis of Copy Number Variation in Latin American Parkinson's Disease Patients. <i>Movement Disorders</i> , 2021 , 36, 434-441	7	4

3	Development and Validation of a Prediction Model for Early Diagnosis of -Related Epilepsies.. <i>Neurology</i> , 2022 ,	6.5	2
2	Copy number variants in lipid metabolism genes are associated with gallstones disease in men. <i>European Journal of Human Genetics</i> , 2020 , 28, 264-273	5.3	2
1	Duplications at 19q13.33 in patients with neurodevelopmental disorders. <i>Neurology: Genetics</i> , 2018 , 4, e210	3.8	1