Eleanna Kara

List of Publications by Year
in descending order

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17 papers	2,566 citations	686830 13 h-index	887659 17 g-index
19 all docs	19 docs citations	19 times ranked	6874 citing authors

#	Article	IF	Citations
1	Integrating protein networks and machine learning for disease stratification in the Hereditary Spastic Paraplegias. IScience, 2021, 24, 102484.	1.9	8
2	An integrated genomic approach to dissect the genetic landscape regulating the cell-to-cell transfer of Î±-synuclein. Cell Reports, 2021, 35, 109189.	2.9	8
3	Sex-dependent calcium hyperactivity due to lysosomal-related dysfunction in astrocytes from APOE4 versus APOE3 gene targeted replacement mice. Molecular Neurodegeneration, 2020, 15, 35.	4.4	35
4	Toxic Protein Spread in Neurodegeneration: Reality versus Fantasy. Trends in Molecular Medicine, 2018, 24, 1007-1020.	3.5	26
5	A flow cytometryâ€“based in vitro assay reveals that formation of apolipoprotein E (ApoE)â€“amyloid beta complexes depends on ApoE isoform and cell type. Journal of Biological Chemistry, 2018, 293, 13247-13256.	1.6	11
6	Isoform- and cell type-specific structure of apolipoprotein E lipoparticles as revealed by a novel Forster resonance energy transfer assay. Journal of Biological Chemistry, 2017, 292, 14720-14729.	1.6	20
7	Pathogenic PS1 phosphorylation at Ser367. ELife, 2017, 6, .	2.8	18
8	Genetic and phenotypic characterization of complex hereditary spastic paraplegia. Brain, 2016, 139, 1904-1918.	3.7	170
9	Mutations in SLC39A14 disrupt manganese homeostasis and cause childhood-onset parkinsonismâ€“dystonia. Nature Communications, 2016, 7, 11601.	5.8	233
10	A 6.4 Mb Duplication of the Î±-Synuclein Locus Causing Frontotemporal Dementia and Parkinsonism. JAMA Neurology, 2014, 71, 1162.	4.5	60
11	Large-scale meta-analysis of genome-wide association data identifies six new risk loci for Parkinson's disease. Nature Genetics, 2014, 46, 989-993.	9.4	1,685
12	Assessment of Parkinson's disease risk loci in Greece. Neurobiology of Aging, 2014, 35, 442.e9-442.e16.	1.5	18
13	Kjellin syndrome: hereditary spastic paraplegia with pathognomonic macular appearance. Practical Neurology, 2014, 14, 278-279.	0.5	6
14	Î±-Synuclein mutations cluster around a putative protein loop. Neuroscience Letters, 2013, 546, 67-70.	1.0	36
15	Genetic Analysis of Inherited Leukodystrophies. JAMA Neurology, 2013, 70, 875.	4.5	147
16	The pallidopyramidal syndromes. Current Opinion in Neurology, 2013, 26, 381-394.	1.8	25
17	The MAPT p.A152T variant is a risk factor associated with tauopathies with atypical clinical and neuropathological features. Neurobiology of Aging, 2012, 33, 2231.e7-2231.e14.	1.5	60