

Eleanna Kara

List of Publications by Year in descending order

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17
papers

2,566
citations

687363

13
h-index

888059

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. <i>IScience</i> , 2021, 24, 102484.	4.1	8
2	An integrated genomic approach to dissect the genetic landscape regulating the cell-to-cell transfer of Î±-synuclein. <i>Cell Reports</i> , 2021, 35, 109189.	6.4	8
3	Sex-dependent calcium hyperactivity due to lysosomal-related dysfunction in astrocytes from APOE4 versus APOE3 gene targeted replacement mice. <i>Molecular Neurodegeneration</i> , 2020, 15, 35.	10.8	35
4	Toxic Protein Spread in Neurodegeneration: Reality versus Fantasy. <i>Trends in Molecular Medicine</i> , 2018, 24, 1007-1020.	6.7	26
5	A flow cytometry-based in vitro assay reveals that formation of apolipoprotein E (ApoE) amyloid beta complexes depends on ApoE isoform and cell type. <i>Journal of Biological Chemistry</i> , 2018, 293, 13247-13256.	3.4	11
6	Isoform- and cell type-specific structure of apolipoprotein E lipoparticles as revealed by a novel Forster resonance energy transfer assay. <i>Journal of Biological Chemistry</i> , 2017, 292, 14720-14729.	3.4	20
7	Pathogenic PS1 phosphorylation at Ser367. <i>ELife</i> , 2017, 6, .	6.0	18
8	Genetic and phenotypic characterization of complex hereditary spastic paraplegia. <i>Brain</i> , 2016, 139, 1904-1918.	7.6	170
9	Mutations in SLC39A14 disrupt manganese homeostasis and cause childhood-onset parkinsonism-dystonia. <i>Nature Communications</i> , 2016, 7, 11601.	12.8	233
10	A 6.4 Mb Duplication of the Î±-Synuclein Locus Causing Frontotemporal Dementia and Parkinsonism. <i>JAMA Neurology</i> , 2014, 71, 1162.	9.0	60
11	Large-scale meta-analysis of genome-wide association data identifies six new risk loci for Parkinson's disease. <i>Nature Genetics</i> , 2014, 46, 989-993.	21.4	1,685
12	Assessment of Parkinson's disease risk loci in Greece. <i>Neurobiology of Aging</i> , 2014, 35, 442.e9-442.e16.	3.1	18
13	Kjellin syndrome: hereditary spastic paraplegia with pathognomonic macular appearance. <i>Practical Neurology</i> , 2014, 14, 278-279.	1.1	6
14	Î±-Synuclein mutations cluster around a putative protein loop. <i>Neuroscience Letters</i> , 2013, 546, 67-70.	2.1	36
15	Genetic Analysis of Inherited Leukodystrophies. <i>JAMA Neurology</i> , 2013, 70, 875.	9.0	147
16	The pallidopyramidal syndromes. <i>Current Opinion in Neurology</i> , 2013, 26, 381-394.	3.6	25
17	The MAPT p.A152T variant is a risk factor associated with tauopathies with atypical clinical and neuropathological features. <i>Neurobiology of Aging</i> , 2012, 33, 2231.e7-2231.e14.	3.1	60