

Emil K Gustavsson

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.

23
papers

580
citations

12
h-index

24
g-index

30
ext. papers

862
ext. citations

9
avg, IF

2.94
L-index

#	Paper	IF	Citations
23	DNAJC13 mutations in Parkinson disease. <i>Human Molecular Genetics</i> , 2014 , 23, 1794-801	5.6	209
22	Comparative study of Parkinson's disease and leucine-rich repeat kinase 2 p.G2019S parkinsonism. <i>Neurobiology of Aging</i> , 2014 , 35, 1125-31	5.6	64
21	DNM3 and genetic modifiers of age of onset in LRRK2 Gly2019Ser parkinsonism: a genome-wide linkage and association study. <i>Lancet Neurology</i> , 2016 , 15, 1248-1256	24.1	50
20	Altered dopamine release and monoamine transporters in Vps35 p.D620N knock-in mice. <i>Npj Parkinsons Disease</i> , 2018 , 4, 27	9.7	38
19	DNAJC13 genetic variants in parkinsonism. <i>Movement Disorders</i> , 2015 , 30, 273-8	7	32
18	Genome sequencing analysis identifies new loci associated with Lewy body dementia and provides insights into its genetic architecture. <i>Nature Genetics</i> , 2021 , 53, 294-303	36.3	31
17	DCTN1 p.K56R in progressive supranuclear palsy. <i>Parkinsonism and Related Disorders</i> , 2016 , 28, 56-61	3.6	24
16	A Case of Parkinson's Disease with No Lewy Body Pathology due to a Homozygous Exon Deletion in. <i>Case Reports in Neurological Medicine</i> , 2018 , 2018, 6838965	0.7	24
15	Genetic variability of the retromer cargo recognition complex in parkinsonism. <i>Movement Disorders</i> , 2015 , 30, 580-4	7	20
14	The influence of mitonuclear genetic variation on personality in seed beetles. <i>Proceedings of the Royal Society B: Biological Sciences</i> , 2014 , 281, 20141039	4.4	20
13	TDP-43 loss and ALS-risk SNPs drive mis-splicing and depletion of UNC13A.. <i>Nature</i> , 2022 , 603, 131-137	50.4	14
12	Genetic Identification in Early Onset Parkinsonism among Norwegian Patients. <i>Movement Disorders Clinical Practice</i> , 2017 , 4, 499-508	2.2	13
11	Parkinson's disease, genetic variability and the Faroe Islands. <i>Parkinsonism and Related Disorders</i> , 2015 , 21, 75-8	3.6	9
10	Genomewide Association Studies of LRRK2 Modifiers of Parkinson's Disease. <i>Annals of Neurology</i> , 2021 , 90, 76-88	9.4	9
9	Neuropathological findings in PINK1-associated Parkinson's disease. <i>Parkinsonism and Related Disorders</i> , 2020 , 78, 105-108	3.6	7
8	Novel LRRK2 mutations in Parkinsonism. <i>Parkinsonism and Related Disorders</i> , 2015 , 21, 1119-21	3.6	6
7	DNAJC13 p.Asn855Ser, implicated in familial parkinsonism, alters membrane dynamics of sorting nexin 1. <i>Neuroscience Letters</i> , 2019 , 706, 114-122	3.3	5

6	Family with primary periodic paralysis and a mutation in MCM3AP, a gene implicated in mRNA transport. <i>Muscle and Nerve</i> , 2019 , 60, 311-314	3.4	2
5	Human-lineage-specific genomic elements are associated with neurodegenerative disease and APOE transcript usage. <i>Nature Communications</i> , 2021 , 12, 2076	17.4	1
4	Genome-wide association study of REM sleep behavior disorder identifies novel loci with distinct polygenic and brain expression effects		1
3	Novel variants broaden the phenotypic spectrum of PLEKHG5-associated neuropathies. <i>European Journal of Neurology</i> , 2021 , 28, 1344-1355	6	0
2	Deep brain stimulation in a Parkinson's disease patient with calcifications and a mutation in the SLC20A2 gene.. <i>Parkinsonism and Related Disorders</i> , 2022 , 96, 88-90	3.6	
1	Genomic features specific to the human lineage are associated with neurological diseases and intelligence. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2022 , 93, A97.1-A97	5.5	