Jennifer A Ruskey

List of Publications by Year
in descending order

27	784	567281 15	580821 25
papers	citations	h-index	g-index

31	31	31	1286
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Rare PSAP Variants and Possible Interaction with GBA in REM Sleep Behavior Disorder. Journal of Parkinson's Disease, 2022, 12, 333-340.	2.8	3
2	Comprehensive Analysis of Familial Parkinsonism Genes in Rapidâ€Eyeâ€Movement Sleep Behavior Disorder. Movement Disorders, 2021, 36, 235-240.	3.9	11
3	Analysis of Heterozygous <scp>PRKN</scp> Variants and Copyâ€Number Variations in Parkinson's Disease. Movement Disorders, 2021, 36, 178-187.	3.9	39
4	Analysis of DNM3 and VAMP4 as genetic modifiers of LRRK2 Parkinsonâ€™s disease. Neurobiology of Aging, 2021, 97, 148.e17-148.e24.	3.1	16
5	Evidence for Nonâ€Mendelian Inheritance in Spastic Paraplegia 7. Movement Disorders, 2021, 36, 1664-1675.	3.9	11
6	LRRK2 p.M1646T is associated with glucocerebrosidase activity and with Parkinson's disease. Neurobiology of Aging, 2021, 103, 142.e1-142.e5.	3.1	11
7	Common Xâ€Chromosome Variants Are Associated with Parkinson Disease Risk. Annals of Neurology, 2021, 90, 22-34.	5.3	28
8	Association study of DNAJC13, UCHL1, HTRA2, GIGYF2, and EIF4G1 with Parkinson's disease. Neurobiology of Aging, 2021, 100, 119.e7-119.e13.	3.1	19
9	Investigation of Autosomal Genetic Sex Differences in Parkinson's Disease. Annals of Neurology, 2021, 90, 35-42.	5.3	29
10	Fine mapping of the HLA locus in Parkinsonâ€™s disease in Europeans. Npj Parkinson's Disease, 2021, 7, 84.	5.3	31
11	Novel Associations of BST1 and LAMP3 With REM Sleep Behavior Disorder. Neurology, 2021, 96, e1402-e1412.	1.1	12
12	Genetic and epidemiological characterization of restless legs syndrome in QuÃ©bec. Sleep, 2020, 43, .	1.1	9
13	Genetic, Structural, and Functional Evidence Link TMEM175 to Synucleinopathies. Annals of Neurology, 2020, 87, 139-153.	5.3	65
14	Genetic modifiers of risk and age at onset in GBA associated Parkinsonâ€™s disease and Lewy body dementia. Brain, 2020, 143, 234-248.	7.6	149
15	GBA variants in REM sleep behavior disorder. Neurology, 2020, 95, e1008-e1016.	1.1	45
16	Fineâ€Mapping of SNCA in Rapid Eye Movement Sleep Behavior Disorder and Overt Synucleinopathies. Annals of Neurology, 2020, 87, 584-598.	5.3	39
17	Variants in the Niemannâ€“Pick type C gene NPC1 are not associated with Parkinson's disease. Neurobiology of Aging, 2020, 93, 143.e1-143.e4.	3.1	13
18	Analysis of common and rare VPS13C variants in late-onset Parkinson disease. Neurology: Genetics, 2020, 6, 385.	1.9	19

#	Article	IF	Citations
19	SMPD1 variants do not have a major role in rapid eye movement sleep behavior disorder. Neurobiology of Aging, 2020, 93, 142.e5-142.e7.	3.1	4
20	Increased yield of full GBA sequencing in Ashkenazi Jews with Parkinson's disease. European Journal of Medical Genetics, 2019, 62, 65-69.	1.3	49
21	Glucocerebrosidase mutations and phenoconversion of REM sleep behavior disorder to parkinsonism and dementia. Parkinsonism and Related Disorders, 2019, 65, 230-233.	2.2	26
22	Common and rare GCH1 variants are associated with Parkinson'sÂdisease. Neurobiology of Aging, 2019, 73, 231.e1-231.e6.	3.1	20
23	Carriers of both GBA and LRRK2 mutations, compared to carriers of either, in Parkinson's disease: Risk estimates and genotype-phenotype correlations. Parkinsonism and Related Disorders, 2019, 62, 179-184.	2.2	58
24	CAPN1 mutations: Expanding the CAPN1-related phenotype: From hereditary spastic paraparesis to spastic ataxia. European Journal of Medical Genetics, 2019, 62, 103605.	1.3	21
25	TOX3 Variants Are Involved in Restless Legs Syndrome and Parkinsonâ€™s Disease with Opposite Effects. Journal of Molecular Neuroscience, 2018, 64, 341-345.	2.3	11
26	LRRK2 protective haplotype and full sequencing study in REM sleep behavior disorder. Parkinsonism and Related Disorders, 2018, 52, 98-101.	2.2	25
27	Sequencing of the GBA coactivator, Saposin C, in Parkinson disease. Neurobiology of Aging, 2018, 72, 187.e1-187.e3.	3.1	16