Rebecca R Valentino

List of Publications by Year
in descending order

13	175	1307594 7	1281871 11
papers	citations	h-index	g-index

14	14	14	319
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Genetic determinants of survival in progressive supranuclear palsy: a genome-wide association study. Lancet Neurology, The, 2021, 20, 107-116.	10.2	62
2	Association of MAPT Subhaplotypes With Risk of Progressive Supranuclear Palsy and Severity of Tau Pathology. JAMA Neurology, 2019, 76, 710.	9.0	39
3	Association of MAPT H1 subhaplotypes with neuropathology of lewy body disease. Movement Disorders, 2019, 34, 1325-1332.	3.9	15
4	Screening non-MAPT genes of the Chr17q21 H1 haplotype in Parkinson's disease. Parkinsonism and Related Disorders, 2020, 78, 138-144.	2.2	12
5	Frequency of spinocerebellar ataxia mutations in patients with multiple system atrophy. Clinical Autonomic Research, 2021, 31, 117-125.	2.5	10
6	Associations of mitochondrial genomic variation with corticobasal degeneration, progressive supranuclear palsy, and neuropathological tau measures. Acta Neuropathologica Communications, 2020, 8, 162.	5.2	9
7	MAPT subhaplotypes in corticobasal degeneration: assessing associations with disease risk, severity of tau pathology, and clinical features. Acta Neuropathologica Communications, 2020, 8, 218.	5.2	8
8	Fine-mapping of the non-coding variation driving the Caucasian LRRK2 GWAS signal in Parkinson's disease. Parkinsonism and Related Disorders, 2021, 83, 22-30.	2.2	7
9	Association of Tripartite Motif Containing 11 rs564309 With Tau Pathology in Progressive Supranuclear Palsy. Movement Disorders, 2020, 35, 890-894.	3.9	6
10	Association of mitochondrial genomic background with risk of Multiple System Atrophy. Parkinsonism and Related Disorders, 2020, 81, 200-204.	2.2	4
11	Investigating ELOVL7 coding variants in multiple system atrophy. Neuroscience Letters, 2021, 749, 135723.	2.1	2
12	Association of Mitochondrial DNA Genomic Variation With Risk of Pick Disease. Neurology, 2021, 96, e1755-e1760.	1.1	1
13	Mitochondrial genomic variation in dementia with Lewy bodies: association with disease risk and neuropathological measures. Acta Neuropathologica Communications, 2022, 10, .	5.2	0