Mie Rizig

List of Publications by Year
in descending order

27 papers	1,966 citations	759233 12 h-index	610901 24 g-index
31 all docs	31 docs citations	31 times ranked	3405 citing authors

#	Article	IF	Citations
1	Identification of novel risk loci, causal insights, and heritable risk for Parkinson's disease: a meta-analysis of genome-wide association studies. Lancet Neurology, The, 2019, 18, 1091-1102.	10.2	1,414
2	Identification of Candidate Parkinson Disease Genes by Integrating Genome-Wide Association Study, Expression, and Epigenetic Data Sets. JAMA Neurology, 2021, 78, 464.	9.0	95
3	The endocytic membrane trafficking pathway plays a major role in the risk of Parkinson's disease. Movement Disorders, 2019, 34, 460-468.	3.9	66
4	Penetrance of Parkinson's Disease in LRRK2 p.G2019S Carriers Is Modified by a Polygenic Risk Score. Movement Disorders, 2020, 35, 774-780.	3.9	57
5	PDXK mutations cause polyneuropathy responsive to pyridoxal 5â€²â€phosphate supplementation. Annals of Neurology, 2019, 86, 225-240.	5.3	54
6	A threonine to isoleucine missense mutation in the pericentriolar material 1 gene is strongly associated with schizophrenia. Molecular Psychiatry, 2010, 15, 615-628.	7.9	50
7	The Genetic Architecture of Parkinson Disease in Spain: Characterizing Populationâ€Specific Risk, Differential Haplotype Structures, and Providing Etiologic Insight. Movement Disorders, 2019, 34, 1851-1863.	3.9	47
8	Investigation of Autosomal Genetic Sex Differences in Parkinson's Disease. Annals of Neurology, 2021, 90, 35-42.	5.3	29
9	Comparing fluid biomarkers of Alzheimer's disease between African American or Black African and white groups: A systematic review and meta-analysis. Journal of the Neurological Sciences, 2021, 421, 117270.	0.6	19
10	Leucine rich repeat kinase 2 (LRRK2) GLY2019SER mutation is absent in a second cohort of Nigerian Africans with Parkinson disease. PLoS ONE, 2018, 13, e0207984.	2.5	18
11	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
12	The Nigeria Parkinson Disease Registry: Process, Profile, and Prospects of a Collaborative Project. Movement Disorders, 2020, 35, 1315-1322.	3.9	14
13	The International Parkinson Disease Genomics Consortium Africa. Lancet Neurology, The, 2021, 20, 335.	10.2	14
14	Novel fluid biomarkers to differentiate frontotemporal dementia and dementia with Lewy bodies from Alzheimer's disease: A systematic review. Journal of the Neurological Sciences, 2020, 415, 116886.	0.6	13
15	WARS2 mutations cause dopa-responsive early-onset parkinsonism and progressive myoclonus ataxia. Parkinsonism and Related Disorders, 2022, 94, 54-61.	2.2	13
16	The effect of clozapine on mRNA expression for genes encoding G protein-coupled receptors and the protein components of clathrin-mediated endocytosis. Psychiatric Genetics, 2013, 23, 153-162.	1.1	10
17	Negative screening for 12 rare LRRK2 pathogenic variants in a cohort of Nigerians with Parkinson's disease. Neurobiology of Aging, 2021, 99, 101.e15-101.e19.	3.1	6
18	A Crossâ€Sectional Comprehensive Assessment of the Profile and Burden of Nonâ€motor Symptoms in Relation to Motor Phenotype in the Nigeria Parkinson Disease Registry Cohort. Movement Disorders Clinical Practice, 2021, 8, 1206-1215.	1.5	6

#	Article	IF	Citations
19	Parkinsonâ€™s Disease in Central Asian and Transcaucasian Countries: A Review of Epidemiology, Genetics, Clinical Characteristics, and Access to Care. Parkinson's Disease, 2019, 2019, 1-7.	1.1	5
20	COVIDâ€19 and the state of African neurology. European Journal of Neurology, 2020, 27, e48-e49.	3.3	5
21	LRRK2 Mutations and Asian Disease-Associated Variants in the First Parkinsonâ€™s Disease Cohort from Kazakhstan. Parkinson's Disease, 2020, 2020, 1-10.	1.1	3
22	Prevalence of Fabry Disease among Patients with Parkinsonâ€™s Disease. Parkinson's Disease, 2022, 2022, 1-8.	1.1	3
23	Spectrum of movement disorders: Experience of a one and half year of existence of the first specialized center in Senegal. Parkinsonism and Related Disorders, 2022, 98, 13-15.	2.2	3
24	Parkinsonâ€™s Disease in Kazakhstan: Clinico-Demographic Description of a Large Cohort. Journal of Parkinson's Disease, 2020, 10, 707-709.	2.8	1
25	LRRK2 mutations in Parkinson's disease patients from Central Europe: A case control study. Parkinsonism and Related Disorders, 2021, 83, 110-112.	2.2	1
26	DEEP PHENOTYPING OF THE G2019S LRRK2 MUTATION IN PARKINSON'S DISEASE: UCL COHORT. Journal of Neurology, Neurosurgery and Psychiatry, 2016, 87, e1.16-e1.	1.9	0
27	PARKINSON'S FAMILIES PROJECT: RECRUITMENT OF FAMILIAL PD PATIENTS VIA THE BNSU. Journal of Neurology, Neurosurgery and Psychiatry, 2016, 87, e1.20-e1.	1.9	0