# Mie Rizig

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/8339740/publications.pdf

Version: 2024-02-01

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

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | WARS2 mutations cause dopa-responsive early-onset parkinsonism and progressive myoclonus ataxia. Parkinsonism and Related Disorders, 2022, 94, 54-61. | 2.2 | 13 |
| 2 | Prevalence of Fabry Disease among Patients with Parkinsonâ€™s Disease. Parkinson's Disease, 2022, 2022, 1-8. | 1.1 | 3 |
| 3 | 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 |
| 4 | 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 |
| 5 | 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 |
| 6 | 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 |
| 7 | 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 |
| 8 | 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 |
| 9 | Investigation of Autosomal Genetic Sex Differences in Parkinson's Disease. Annals of Neurology, 2021, 90, 35-42. | 5.3 | 29 |
| 10 | The International Parkinson Disease Genomics Consortium Africa. Lancet Neurology, The, 2021, 20, 335. | 10.2 | 14 |
| 11 | 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 |
| 12 | 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 |
| 13 | The Nigeria Parkinson Disease Registry: Process, Profile, and Prospects of a Collaborative Project. Movement Disorders, 2020, 35, 1315-1322. | 3.9 | 14 |
| 14 | COVIDâ€19 and the state of African neurology. European Journal of Neurology, 2020, 27, e48-e49. | 3.3 | 5 |
| 15 | 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 |
| 16 | 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 |
| 17 | Parkinsonâ€™s Disease in Kazakhstan: Clinico-Demographic Description of a Large Cohort. Journal of Parkinson's Disease, 2020, 10, 707-709. | 2.8 | 1 |
| 18 | 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 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | 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 |
| 20 | 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 |
| 21 | 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 |
| 22 | PDXK mutations cause polyneuropathy responsive to pyridoxal 5â€²â€phosphate supplementation. Annals of Neurology, 2019, 86, 225-240. | 5.3 | 54 |
| 23 | 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 |
| 24 | 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 |
| 25 | 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 |
| 26 | 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 |
| 27 | 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 |