# Emil K Gustavsson

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/7440140/publications.pdf

Version: 2024-02-01

| 24 | 1,196 | 623699 14 | 677123 22 |
|---|---|---|---|
| papers | citations | h-index | g-index |

| 30 | 30 | 30 | 1740 |
|---|---|---|---|
| all docs | docs citations | times ranked | citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | DNAJC13 mutations in Parkinson disease. Human Molecular Genetics, 2014, 23, 1794-1801. | 2.9 | 258 |
| 2 | Genome sequencing analysis identifies new loci associated with Lewy body dementia and provides insights into its genetic architecture. Nature Genetics, 2021, 53, 294-303. | 21.4 | 198 |
| 3 | TDP-43 loss and ALS-risk SNPs drive mis-splicing and depletion of UNC13A. Nature, 2022, 603, 131-137. | 27.8 | 188 |
| 4 | A comparative study of Parkinson's disease and leucine-rich repeat kinase 2 p.G2019S parkinsonism. Neurobiology of Aging, 2014, 35, 1125-1131. | 3.1 | 83 |
| 5 | ggtranscript: an R package for the visualization and interpretation of transcript isoforms usingggplot2. Bioinformatics, 2022, 38, 3844-3846. | 4.1 | 76 |
| 6 | DNM3 and genetic modifiers of age of onset in LRRK2 Gly2019Ser parkinsonism: a genome-wide linkage and association study. Lancet Neurology, The, 2016, 15, 1248-1256. | 10.2 | 69 |
| 7 | Altered dopamine release and monoamine transporters in Vps35 p.D620N knock-in mice. Npj Parkinson's Disease, 2018, 4, 27. | 5.3 | 51 |
| 8 | DNAJC13 genetic variants in parkinsonism. Movement Disorders, 2015, 30, 273-278. | 3.9 | 42 |
| 9 | A Case of Parkinsonâ€™s Disease with No Lewy Body Pathology due to a Homozygous Exon Deletion in Parkin. Case Reports in Neurological Medicine, 2018, 2018, 1-4. | 0.4 | 33 |
| 10 | Genomewide Association Studies of <scp>LRRK2</scp> Modifiers of Parkinson's Disease. Annals of Neurology, 2021, 90, 76-88. | 5.3 | 30 |
| 11 | DCTN1 p.K56R in progressive supranuclear palsy. Parkinsonism and Related Disorders, 2016, 28, 56-61. | 2.2 | 27 |
| 12 | The influence of mitonuclear genetic variation on personality in seed beetles. Proceedings of the Royal Society B: Biological Sciences, 2014, 281, 20141039. | 2.6 | 25 |
| 13 | Genetic Identification in Early Onset Parkinsonism among Norwegian Patients. Movement Disorders Clinical Practice, 2017, 4, 499-508. | 1.5 | 25 |
| 14 | Genetic variability of the retromer cargo recognition complex in parkinsonism. Movement Disorders, 2015, 30, 580-584. | 3.9 | 23 |
| 15 | Neuropathological findings in PINK1-associated Parkinson's disease. Parkinsonism and Related Disorders, 2020, 78, 105-108. | 2.2 | 14 |
| 16 | Parkinson's disease, genetic variability and the Faroe Islands. Parkinsonism and Related Disorders, 2015, 21, 75-78. | 2.2 | 11 |
| 17 | Human-lineage-specific genomic elements are associated with neurodegenerative disease and APOE transcript usage. Nature Communications, 2021, 12, 2076. | 12.8 | 9 |
| 18 | Novel LRRK2 mutations in Parkinsonism. Parkinsonism and Related Disorders, 2015, 21, 1119-1121. | 2.2 | 8 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | DNAJC13 p.Asn855Ser, implicated in familial parkinsonism, alters membrane dynamics of sorting nexin 1. Neuroscience Letters, 2019, 706, 114-122. | 2.1 | 8 |
| 20 | Novel variants broaden the phenotypic spectrum of PLEKHG5 â€associated neuropathies. European Journal of Neurology, 2021, 28, 1344-1355. | 3.3 | 4 |
| 21 | Family with primary periodic paralysis and a mutation in MCM3AP, a gene implicated in mRNA transport. Muscle and Nerve, 2019, 60, 311-314. | 2.2 | 3 |
| 22 | Polygenic risk of Alzheimerâ€™s disease in the Faroe Islands. European Journal of Neurology, 2022, , . | 3.3 | 1 |
| 23 | Deep brain stimulation in a Parkinson's disease patient with calcifications and a mutation in the SLC20A2 gene. Parkinsonism and Related Disorders, 2022, 96, 88-90. | 2.2 | 0 |
| 24 | Genomic features specific to the human lineage are associated with neurological diseases and intelligence. Journal of Neurology, Neurosurgery and Psychiatry, 2022, 93, A97.1-A97. | 1.9 | 0 |