## Anneke Ja Kievit

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/1883715/publications.pdf

Version: 2024-02-01

1163117 1281871 12 366 8 11 citations h-index g-index papers 13 13 13 789 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	<i>D</i> <scp><i>NAJC</i></scp> <i>6</i> Mutations <scp>A</scp> ssociated <scp>W</scp> ith <scp>E</scp> arlyâ€ <scp>O</scp> nset <scp>P</scp> arkinson's <scp>D</scp> isease. Annals of Neurology, 2016, 79, 244-256.	5.3	148
2	Paroxysmal exercise-induced dystonia within the phenotypic spectrum of <i>ECHS1</i> deficiency. Movement Disorders, 2016, 31, 1041-1048.	3.9	58
3	Panel-Based Exome Sequencing for Neuromuscular Disorders as a Diagnostic Service. Journal of Neuromuscular Diseases, 2019, 6, 241-258.	2.6	32
4	Lessons learned from 40 novel <i>PIGA</i> patients and a review of the literature. Epilepsia, 2020, 61, 1142-1155.	5.1	32
5	Cysteinyl-tRNA Synthetase Mutations Cause a Multi-System, Recessive Disease That Includes Microcephaly, Developmental Delay, and Brittle Hair and Nails. American Journal of Human Genetics, 2019, 104, 520-529.	6.2	31
6	Mutations inTMEM230are not a common cause of Parkinson's disease. Movement Disorders, 2017, 32, 302-304.	3.9	14
7	Deciphering the natural history of SCA7 in children. European Journal of Neurology, 2020, 27, 2267-2276.	3.3	12
8	A new alpha-synuclein missense variant (Thr72Met) in two Turkish families with Parkinson's disease. Parkinsonism and Related Disorders, 2021, 89, 63-72.	2.2	11
9	Early onset Xâ€inked female limited high myopia in three multigenerational families caused by novel mutations in the <i>ARR3</i> gene. Human Mutation, 2022, 43, 380-388.	2.5	11
10	The Effect of Predictive Testing in Adultâ€Onset Neurodegenerative Diseases on Social and Personal Life. Journal of Genetic Counseling, 2018, 27, 947-954.	1.6	10
11	Experience in Genetic Counseling for GBA1 Variants in Parkinson's Disease. Movement Disorders Clinical Practice, 2021, 8, 33-36.	1.5	5
12	Modelling the cascade of biomarker changes in progranulinâ€related frontotemporal dementia. Alzheimer's and Dementia, 2020, 16, e040934.	0.8	0