Ebba Lohmann

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/1770873/publications.pdf

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27 papers	1,454 citations	933447 10 h-index	642732 23 g-index
30 all docs	30 docs citations	30 times ranked	3600 citing authors

#	Article	IF	Citations
1	PHACTR1 genetic variability is not critical in small vessel ischemic disease patients and PcomA recruitment in C57BL/6J mice. Scientific Reports, 2021, 11, 6072.	3.3	2
2	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
3	Peripheral TREM2 mRNA levels in early and late-onset Alzheimer diseaseâ€™s patients. Molecular Biology Reports, 2020, 47, 5903-5909.	2.3	8
4	Analysis of copy number variation in a Turkish dementia cohort. Alzheimer's and Dementia, 2020, 16, e044868.	0.8	1
5	Characterization of Recessive Parkinson Disease in a Large Multicenter Study. Annals of Neurology, 2020, 88, 843-850.	5.3	40
6	Event-related potential changes due to early-onset Parkinsonâ€™s disease in parkin (PARK2) gene mutation carriers and non-carriers. Clinical Neurophysiology, 2020, 131, 1444-1452.	1.5	7
7	Association between selected cholesterol-related gene polymorphisms and Alzheimerâ€™s disease in a Turkish cohort. Molecular Biology Reports, 2019, 46, 1701-1707.	2.3	6
8	Patients with Lately Diagnosed Cerebrotendinous Xanthomatosis. Neurodegenerative Diseases, 2019, 19, 218-224.	1.4	11
9	Single Molecule Molecular Inversion Probes for High Throughput Germline Screenings in Dystonia. Frontiers in Neurology, 2019, 10, 1332.	2.4	2
10	Role of LRRK2 and SNCA in autosomal dominant Parkinson's disease in Turkey. Parkinsonism and Related Disorders, 2018, 48, 34-39.	2.2	8
11	Mutations in TYROBP are not a common cause of dementia in a Turkish cohort. Neurobiology of Aging, 2017, 58, 240.e1-240.e3.	3.1	6
12	Needs and Requirements of Modern Biobanks on the Example of Dystonia Syndromes. Frontiers in Neurology, 2017, 8, 9.	2.4	6
13	Mutation Frequency of the Major Frontotemporal Dementia Genes, MAPT, GRN and C9ORF72 in a Turkish Cohort of Dementia Patients. PLoS ONE, 2016, 11, e0162592.	2.5	19
14	Vitamin D deficiency might pose a greater risk for ApoEÉ›4 non-carrier Alzheimerâ€™s disease patients. Neurological Sciences, 2016, 37, 1633-1643.	1.9	36
15	A novel homozygous DJ1 mutation causes parkinsonism and ALS in a Turkish family. Parkinsonism and Related Disorders, 2016, 29, 117-120.	2.2	23
16	Loss of VPS13C Function in Autosomal-Recessive Parkinsonism Causes Mitochondrial Dysfunction and Increases PINK1/Parkin-Dependent Mitophagy. American Journal of Human Genetics, 2016, 98, 500-513.	6.2	333
17	PLA2G6 Mutations Related to Distinct Phenotypes: A New Case with Early-onset Parkinsonism. Tremor and Other Hyperkinetic Movements, 2016, 6, 363.	2.0	11
18	P3-097: Serum interleukin 1 alpha and alpha 2 macroglobulin levels in patients with early- or late-onset Alzheimer's disease or mild cognitive impairment. , 2015, 11, P657-P657.		2

#	Article	IF	Citations
19	The interleukin 1 alpha, interleukin 1 beta, interleukin 6 and alpha-2-macroglobulin serum levels in patients with early or late onset Alzheimer's disease, mild cognitive impairment or Parkinson's disease. Journal of Neuroimmunology, 2015, 283, 50-57.	2.3	197
20	P3-082: Compromised regulation of serum cytokine levels and BDNF due to low levels of vitamin d in patients with early- or late-onset Alzheimer's disease or parkinson's disease. , 2015, 11, P649-P650.		0
21	A new Fâ€box protein 7 gene mutation causing typical Parkinson's disease. Movement Disorders, 2015, 30, 1130-1133.	3.9	59
22	TREM2 mutations implicated in neurodegeneration impair cell surface transport and phagocytosis. Science Translational Medicine, 2014, 6, 243ra86.	12.4	600
23	Unusual variability of PRRT2 linked phenotypes within a family. European Journal of Paediatric Neurology, 2014, 18, 540-542.	1.6	9
24	P1-287: NEUROIMAGING FINDINGS OF NASU-HAKOLA DISEASE. , 2014, 10, P415-P415.		0
25	The Impact of Familial Structure on Parkinson's Disease in Istanbul Medical School, Turkey. International Journal of Neuroscience, 2012, 122, 102-105.	1.6	0
26	Identification of PSEN1 and PSEN2 gene mutations and variants in Turkish dementia patients. Neurobiology of Aging, 2012, 33, 1850.e17-1850.e27.	3.1	44
27	A homozygous frameshift mutation of sepiapterin reductase gene causing parkinsonism with onset in childhood. Parkinsonism and Related Disorders, 2012, 18, 191-193.	2.2	13