

Elisabeth StÄjgmann

List of Publications by Year in descending order

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Version: 2024-02-01

58
papers

12,490
citations

279487

23
h-index

161609

54
g-index

62
all docs

62
docs citations

62
times ranked

20783
citing authors

#	ARTICLE	IF	CITATIONS
1	User experience and acceptance of a device assisting persons with dementia in daily life: a multicenter field study. <i>Aging Clinical and Experimental Research</i> , 2022, 34, 869-879.	1.4	6
2	Usability testing of the first prototype of the Memento system: a technological device to promote an independent living in people with dementia. <i>Disability and Rehabilitation: Assistive Technology</i> , 2022, , 1-10.	1.3	1
3	Reduction of physical activity during the COVID-19 pandemic is related to increased neuropsychiatric symptoms in memory clinic patients. <i>Clinical Medicine</i> , 2022, 22, 177-180.	0.8	4
4	No effect of thyroid hormones on 5-year mortality in patients with subjective cognitive decline, mild cognitive disorder, and Alzheimer's disease. <i>Journal of Neuroendocrinology</i> , 2022, 34, e13107.	1.2	5
5	Serum NfL in Alzheimer Dementia: Results of the Prospective Dementia Registry Austria. <i>Medicina (Lithuania)</i> , 2022, 58, 433.	0.8	5
6	Mapping genomic loci implicates genes and synaptic biology in schizophrenia. <i>Nature</i> , 2022, 604, 502-508.	13.7	929
7	AI-Based Predictive Modelling of the Onset and Progression of Dementia. <i>Smart Cities</i> , 2022, 5, 700-714.	5.5	2
8	Individual cognitive changes in subjective cognitive decline, mild cognitive impairment and Alzheimer's disease using the reliable change index methodology. <i>Wiener Klinische Wochenschrift</i> , 2021, 133, 1064-1069.	1.0	5
9	A NOTCH3 homozygous nonsense mutation in familial Sneddon syndrome with pediatric stroke. <i>Journal of Neurology</i> , 2021, 268, 810-816.	1.8	11
10	Co-incident <i>C9orf72</i> expansion mutation-related frontotemporal lobar degeneration pathology and sporadic Creutzfeldt-Jakob disease. <i>European Journal of Neurology</i> , 2021, 28, 1009-1015.	1.7	2
11	Genetics of Alzheimer's disease. <i>Wiener Medizinische Wochenschrift</i> , 2021, 171, 249-256.	0.5	11
12	Utilization of occupational therapy services and relation to survival in people taking dementia-specific medication in Austria: A retrospective population-based study with a 13-year observation period. <i>International Journal of Geriatric Psychiatry</i> , 2021, 36, 1179-1187.	1.3	0
13	A Comparison of Ten Polygenic Score Methods for Psychiatric Disorders Applied Across Multiple Cohorts. <i>Biological Psychiatry</i> , 2021, 90, 611-620.	0.7	103
14	Depressive symptoms and olfactory function in patients with subjective cognitive decline, mild cognitive impairment and Alzheimer's disease. <i>Brain Disorders</i> , 2021, 2, 100014.	1.1	1
15	Epidemiology of dementia: the epidemic we saw coming. <i>Wiener Medizinische Wochenschrift</i> , 2021, 171, 247-248.	0.5	2
16	Genotype-guided diagnostic reassessment after exome sequencing in neuromuscular disorders: experiences with a two-step approach. <i>European Journal of Neurology</i> , 2020, 27, 51-61.	1.7	21
17	Awareness of Olfactory Dysfunction in Subjective Cognitive Decline, Mild Cognitive Decline, and Alzheimer's Disease. <i>Chemosensory Perception</i> , 2020, 13, 59-70.	0.7	4
18	Frequency of comedication in patients with dementia. <i>Alzheimer's and Dementia</i> , 2020, 16, e039493.	0.4	0

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19	Prescription patterns of antidementives in a high income country: A pharmacoepidemiologic study. <i>Alzheimer's and Dementia: Translational Research and Clinical Interventions</i> , 2020, 6, e12014.	1.8	5
20	Argyrophilic grain disease in individuals younger than 75 years: clinical variability in an underrecognized limbic tauopathy. <i>European Journal of Neurology</i> , 2020, 27, 1856-1866.	1.7	13
21	Multidimensional Design Research for Dementia and Its Methodological Opportunities for Cross-Disciplinary Consortia. <i>Design Journal</i> , 2020, 23, 597-619.	0.5	2
22	Anatomical and functional changes in the retina in patients with Alzheimer's disease and mild cognitive impairment. <i>Acta Ophthalmologica</i> , 2020, 98, e914-e921.	0.6	33
23	Diagnostic exome sequencing in non-acquired focal epilepsies highlights a major role of GATOR1 complex genes. <i>Journal of Medical Genetics</i> , 2020, 57, 624-633.	1.5	16
24	Increased risk of death associated with the use of proton pump inhibitors in patients with dementia and controls – a pharmacoepidemiological claims data analysis. <i>European Journal of Neurology</i> , 2020, 27, 1422-1428.	1.7	3
25	Identification of odors, faces, cities and naming of objects in patients with subjective cognitive decline, mild cognitive impairment and Alzheimer's disease: a longitudinal study. <i>International Psychogeriatrics</i> , 2019, 31, 537-549.	0.6	22
26	Differences regarding the five-factor personality model in patients with subjective cognitive decline and mild cognitive impairment. <i>Neuropsychiatrie</i> , 2019, 33, 35-45.	1.3	9
27	Memento for Living, Working and Caring: An Archetypal Object for Being with Dementia. <i>Communications in Computer and Information Science</i> , 2019, , 114-127.	0.4	2
28	Macrophagic scavenging of A β . , 2019, 38, 48-50.		0
29	Estimation of Genetic Correlation via Linkage Disequilibrium Score Regression and Genomic Restricted Maximum Likelihood. <i>American Journal of Human Genetics</i> , 2018, 102, 1185-1194.	2.6	119
30	Analysis of shared heritability in common disorders of the brain. <i>Science</i> , 2018, 360, .	6.0	1,085
31	Semantic memory and depressive symptoms in patients with subjective cognitive decline, mild cognitive impairment, and Alzheimer's disease. <i>International Psychogeriatrics</i> , 2017, 29, 1123-1135.	0.6	17
32	Hereditary spastic paraplegia caused by compound heterozygous mutations outside the motor domain of the <i>KIF1A</i> gene. <i>European Journal of Neurology</i> , 2017, 24, 741-747.	1.7	25
33	Manifestations of neurological symptoms and thromboembolism in adults with MTHFR-deficiency. <i>Journal of the Neurological Sciences</i> , 2017, 383, 123-127.	0.3	9
34	Activities of Daily Living and Depressive Symptoms in Patients with Subjective Cognitive Decline, Mild Cognitive Impairment, and Alzheimer's Disease. <i>Journal of Alzheimer's Disease</i> , 2016, 49, 1043-1050.	1.2	48
35	The impact of depressive symptoms on health-related quality of life in patients with subjective cognitive decline, mild cognitive impairment, and Alzheimer's disease. <i>International Psychogeriatrics</i> , 2016, 28, 2045-2054.	0.6	21
36	Self-reported and informant-reported memory functioning and awareness in patients with mild cognitive impairment and Alzheimer's disease. <i>Neuropsychiatrie</i> , 2016, 30, 103-112.	1.3	13

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37	Genetic influences on schizophrenia and subcortical brain volumes: large-scale proof of concept. <i>Nature Neuroscience</i> , 2016, 19, 420-431.	7.1	204
38	Modeling Linkage Disequilibrium Increases Accuracy of Polygenic Risk Scores. <i>American Journal of Human Genetics</i> , 2015, 97, 576-592.	2.6	1,098
39	Partitioning Heritability of Regulatory and Cell-Type-Specific Variants across 11 Common Diseases. <i>American Journal of Human Genetics</i> , 2014, 95, 535-552.	2.6	569
40	Biological insights from 108 schizophrenia-associated genetic loci. <i>Nature</i> , 2014, 511, 421-427.	13.7	6,934
41	Reply: Autosomal recessive epilepsy associated with contactin 2 mutation is different from familial cortical tremor, myoclonus and epilepsy. <i>Brain</i> , 2013, 136, e254-e254.	3.7	1
42	Epilepsy, hippocampal sclerosis and febrile seizures linked by common genetic variation around SCN1A. <i>Brain</i> , 2013, 136, 3140-3150.	3.7	168
43	Autosomal recessive cortical myoclonic tremor and epilepsy: association with a mutation in the potassium channel associated gene CNTN2. <i>Brain</i> , 2013, 136, 1155-1160.	3.7	137
44	Lack of association between <i>ABCC2</i> gene variants and treatment response in epilepsy. <i>Pharmacogenomics</i> , 2012, 13, 185-190.	0.6	33
45	Central serotonin 1A receptor binding in temporal lobe epilepsy: A [carbonyl-11C]WAY-100635 PET study. <i>Epilepsy and Behavior</i> , 2010, 19, 467-473.	0.9	37
46	A novel mutation in the VCP gene (G157R) in a german family with inclusion body myopathy with paget disease of bone and frontotemporal dementia. <i>Muscle and Nerve</i> , 2009, 39, 389-391.	1.0	60
47	A novel mutation in the MFSD8 gene in late infantile neuronal ceroid lipofuscinosis. <i>Neurogenetics</i> , 2009, 10, 73-77.	0.7	33
48	A splice site variant in the sodium channel gene <i>SCN1A</i> confers risk of febrile seizures. <i>Neurology</i> , 2009, 72, 974-978.	1.5	50
49	A functional polymorphism in the <i>SCN1A</i> gene is not associated with carbamazepine dosages in Austrian patients with epilepsy. <i>Epilepsia</i> , 2008, 49, 1108-1109.	2.6	48
50	Sequence analysis of the complete SLITRK1 gene in Austrian patients with Tourette's disorder. <i>Psychiatric Genetics</i> , 2008, 18, 308-309.	0.6	58
51	Clinical Seizure Lateralization in Frontal Lobe Epilepsy. <i>Epilepsia</i> , 2007, 48, 517-523.	2.6	63
52	Lack of Association between a GABAB Receptor 1 Gene Polymorphism and Temporal Lobe Epilepsy. <i>Epilepsia</i> , 2006, 47, 437-439.	2.6	10
53	Mutations in the CLCN2 gene are a rare cause of idiopathic generalized epilepsy syndromes. <i>Neurogenetics</i> , 2006, 7, 265-268.	0.7	22
54	Andreas Rett and benign familial neonatal convulsions revisited. <i>Neurology</i> , 2006, 67, 864-866.	1.5	25

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55	Idiopathic generalized epilepsy phenotypes associated with different EFHC1 mutations. <i>Neurology</i> , 2006, 67, 2029-2031.	1.5	63
56	Association of an ABCB1 gene haplotype with pharmaco-resistance in temporal lobe epilepsy. <i>Neurology</i> , 2004, 63, 1087-1089.	1.5	207
57	A functional polymorphism in the prodynorphin gene promotor is associated with temporal lobe epilepsy. <i>Annals of Neurology</i> , 2002, 51, 260-263.	2.8	94
58	Long-term Olfactory Functions in Patients with Subjective Cognitive Decline and Mild Cognitive Impairment. <i>Chemosensory Perception</i> , 0, , 1.	0.7	0