Anne Sieben

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/7792985/publications.pdf

Version: 2024-02-01

59 papers	2,529 citations	393982 19 h-index	205818 48 g-index
65 all docs	65 docs citations	65 times ranked	3979 citing authors

#	Article	IF	Citations
1	Contribution of rare homozygous and compound heterozygous VPS13C missense mutations to dementia with Lewy bodies and Parkinsonâ€™s disease. Acta Neuropathologica Communications, 2021, 9, 25.	2.4	23
2	Hippocampal Sclerosis in Frontotemporal Dementia: When Vascular Pathology Meets Neurodegeneration. Journal of Neuropathology and Experimental Neurology, 2021, 80, 313-324.	0.9	5
3	Cerebellar ataxia in progressive supranuclear palsy: a clinico-pathological case report. Acta Neurologica Belgica, 2021, 121, 599-602.	0.5	0
4	The Electrophysiological Correlates of Phoneme Perception in Primary Progressive Aphasia: A Preliminary Case Series. Frontiers in Human Neuroscience, 2021, 15, 618549.	1.0	6
5	Family-based exome sequencing identifies RBM45 as a possible candidate gene for frontotemporal dementia and amyotrophic lateral sclerosis. Neurobiology of Disease, 2021, 156, 105421.	2.1	2
6	A case series of verbal semantic processing in primary progressive aphasia: Evidence from the N400 effect. International Journal of Language and Communication Disorders, 2021, 56, 1165-1189.	0.7	2
7	Diagnostic Performance of Automated MRI Volumetry by icobrain dm for Alzheimerâ€™s Disease in a Clinical Setting: A REMEMBER Study. Journal of Alzheimer's Disease, 2021, 83, 623-639.	1.2	7
8	Neurogranin as biomarker in CSF is non-specific to Alzheimer's disease dementia. Neurobiology of Aging, 2021, 108, 99-109.	1.5	13
9	Diagnostic performance of automated MRI volumetry by icobrain DM for Alzheimer's disease in a clinical setting: A REMEMBER study. Alzheimer's and Dementia, 2021, 17, .	0.4	0
10	Multivariate analysis reveals anatomical correlates of naming errors in primary progressive aphasia. Neurobiology of Aging, 2020, 88, 71-82.	1.5	21
11	Amyloid-Î²1â€“43 cerebrospinal fluid levels and the interpretation of APP, PSEN1 and PSEN2 mutations. Alzheimer's Research and Therapy, 2020, 12, 108.	3.0	17
12	ABCA7 PTC mutation carriers present with Alzheimerâ€™s disease pathology and cerebral amyloid angiopathy. Alzheimer's and Dementia, 2020, 16, e041513.	0.4	0
13	A familyâ€based genetic study identifies mutations in TLR9 impairing receptor activation: A role for innate immunity in AD pathogenesis. Alzheimer's and Dementia, 2020, 16, e047212.	0.4	2
14	Dementia, End of Life, and Euthanasia: A Survey Among Dementia Specialists Organized by the Belgian Dementia Council. Journal of Alzheimer's Disease, 2019, 69, 989-1001.	1.2	14
15	Dura mater graft-associated Creutzfeldtâ€“Jakob disease with an incubation period of 30Âyears, mimicking non-convulsive status epilepticus. Acta Neurologica Belgica, 2019, 119, 497-499.	0.5	1
16	Validation of the Erlangen Score Algorithm for Differential Dementia Diagnosis in Autopsy-Confirmed Subjects. Journal of Alzheimer's Disease, 2019, 68, 1151-1159.	1.2	9
17	Loss of DPP6 in neurodegenerative dementia: a genetic player in the dysfunction of neuronal excitability. Acta Neuropathologica, 2019, 137, 901-918.	3.9	37
18	Clinical variability and onset age modifiers in an extended Belgian GRN founder family. Neurobiology of Aging, 2018, 67, 84-94.	1.5	17

#	Article	IF	Citations
19	Teenage-onset progressive myoclonic epilepsy due to a familial C9orf72 repeat expansion. Neurology, 2018, 90, e658-e663.	1.5	9
20	Diagnostic value of cerebrospinal fluid tau, neurofilament, and progranulin in definite frontotemporal lobar degeneration. Alzheimer's Research and Therapy, 2018, 10, 31.	3.0	42
21	Extended FTLD pedigree segregating a Belgian GRN-null mutation: neuropathological heterogeneity in one family. Alzheimer's Research and Therapy, 2018, 10, 7.	3.0	10
22	Rare nonsynonymous variants in SORT1 are associated with increased risk for frontotemporal dementia. Neurobiology of Aging, 2018, 66, 181.e3-181.e10.	1.5	19
23	NEK1 genetic variability in a Belgian cohort of ALS and ALS-FTD patients. Neurobiology of Aging, 2018, 61, 255.e1-255.e7.	1.5	32
24	Monoaminergic impairment in Down syndrome with Alzheimer's disease compared to earlyâ€onset Alzheimer's disease. Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring, 2018, 10, 99-111.	1.2	9
25	A Retrospective Belgian Multi-Center MRI Biomarker Study in Alzheimerâ€™s Disease (REMEMBER). Journal of Alzheimer's Disease, 2018, 63, 1509-1522.	1.2	17
26	Single-word comprehension deficits in the nonfluent variant of primary progressive aphasia. Alzheimer's Research and Therapy, 2018, 10, 68.	3.0	16
27	Distinct [18F]THK5351 binding patterns in primary progressive aphasia variants. European Journal of Nuclear Medicine and Molecular Imaging, 2018, 45, 2342-2357.	3.3	16
28	Investigating the role of ALS genes CHCHD10 and TUBA4A in Belgian FTD-ALS spectrum patients. Neurobiology of Aging, 2017, 51, 177.e9-177.e16.	1.5	60
29	Clinical Evidence of Disease Anticipation in Families Segregating a C9orf72 Repeat Expansion. JAMA Neurology, 2017, 74, 445.	4.5	56
30	Alzheimerâ€™s disease and driving: review of the literature and consensus guideline from Belgian dementia experts and the Belgian road safety institute endorsed by the Belgian Medical Association. Acta Neurologica Belgica, 2017, 117, 811-819.	0.5	15
31	Bilingualism and Cognitive Decline: A Story of Pride and Prejudice. Journal of Alzheimer's Disease, 2017, 60, 1237-1239.	1.2	21
32	[O2â€“08â€“06]: CONTRIBUTION OF RARE DELETERIOUS ABCA7 MUTATIONS TO A BELGIAN EARLYâ€ONSET ALZHEIMER's DISEASE COHORT. Alzheimer's and Dementia, 2017, 13, P573.	0.4	0
33	[P4â€“069]: A PROSPECTIVE NEUROGENETIC STUDY ON EARLYâ€ONSET DEMENTIA IN PATIENTS WITH UNCLEAR INITIAL DIAGNOSIS OF DEGENERATIVE DEMENTIA. Alzheimer's and Dementia, 2017, 13, P1284.	0.4	0
34	Impaired Processing of Serial Order Determines Working Memory Impairments in Alzheimerâ€™s Disease. Journal of Alzheimer's Disease, 2017, 59, 1171-1186.	1.2	6
35	No added diagnostic value of non-phosphorylated tau fraction (p-taurel) in CSF as a biomarker for differential dementia diagnosis. Alzheimer's Research and Therapy, 2017, 9, 49.	3.0	11
36	EEG Dominant Frequency Peak Differentiates Between Alzheimerâ€™s Disease and Frontotemporal Lobar Degeneration. Journal of Alzheimer's Disease, 2016, 55, 53-58.	1.2	13

#	Article	IF	Citations
37	P1â€176: CSF Exploratory Biomarker Study for (DIFFERENTIAL) Diagnosis of Frontotemporal Lobar Degeneration. Alzheimer's and Dementia, 2016, 12, P471.	0.4	0
38	P2-153: Diagnostic Performance of Non-Phosphorylated TAU Fraction (PTAU REL) in CSF as Biomarker for Differential Dementia Diagnosis. , 2016, 12, P672-P673.		0
39	P4â€120: Increased CSF Levels of Biomarkers for Neurodegeneration in FTLDâ€GRN Mutation Carriers. Alzheimer's and Dementia, 2016, 12, P1058.	0.4	0
40	O4â€09â€03: Eeg Dominant Frequency Peak Differentiates Between Alzheimer's Disease and Frontotemporal Lobar Degeneration. Alzheimer's and Dementia, 2016, 12, P354.	0.4	0
41	Phenotypic characteristics of Alzheimer patients carrying an ABCA7 mutation. Neurology, 2016, 86, 2126-2133.	1.5	29
42	Mutated CTSF in adult-onset neuronal ceroid lipofuscinosis and FTD. Neurology: Genetics, 2016, 2, e102.	0.9	21
43	Drosophila screen connects nuclear transport genes to DPR pathology in c9ALS/FTD. Scientific Reports, 2016, 6, 20877.	1.6	239
44	Clinical features ofTBK1carriers compared withC9orf72,GRNand non-mutation carriers in a Belgian cohort. Brain, 2016, 139, 452-467.	3.7	86
45	Motor neuron degeneration in spastic paraplegia 11 mimics amyotrophic lateral sclerosis lesions. Brain, 2016, 139, aww061.	3.7	43
46	Investigating the role of filamin C in Belgian patients with frontotemporal dementia linked to GRN deficiency in FTLD-TDP brains. Acta Neuropathologica Communications, 2015, 3, 68.	2.4	13
47	DT-02-01: Loss-of-function mutations in TBK1 are frequently associated with frontotemporal lobar degeneration in a belgian patient cohort. , 2015, 11, P333-P333.		0
48	Clinical utility and applicability of biomarker-based diagnostic criteria for Alzheimerâ€™s disease: a BeDeCo survey. Acta Neurologica Belgica, 2015, 115, 547-555.	0.5	7
49	Bilingualism delays clinical manifestation of Alzheimer's disease. Bilingualism, 2015, 18, 568-574.	1.0	128
50	Loss of TBK1 is a frequent cause of frontotemporal dementia in a Belgian cohort. Neurology, 2015, 85, 2116-2125.	1.5	151
51	Partial deletion of AFG3L2 causing spinocerebellar ataxia type 28. Neurology, 2014, 82, 2092-2100.	1.5	24
52	Rare mutations in SQSTM1 modify susceptibility to frontotemporal lobar degeneration. Acta Neuropathologica, 2014, 128, 397-410.	3.9	93
53	A Panâ€<scp>E</scp>uropean Study of theC9orf72Repeat Associated with<scp>FTLD</scp>: Geographic Prevalence, Genomic Instability, and Intermediate Repeats. Human Mutation, 2013, 34, 363-373.	1.1	247
54	Distinct Clinical Characteristics of C9orf72 Expansion Carriers Compared With GRN, MAPT, and Nonmutation Carriers in a Flanders-Belgian FTLD Cohort. JAMA Neurology, 2013, 70, 365.	4.5	85

#	Article	IF	Citations
55	The genetics and neuropathology of frontotemporal lobar degeneration. Acta Neuropathologica, 2012, 124, 353-372.	3.9	242
56	Neoplasm Related Encephalopathies. , 2012, , .		0
57	Late-onset post-irradiation vasculopathy of the posterior cerebral vasculature. Acta Neurologica Belgica, 2012, 112, 101-104.	0.5	1
58	A C9orf72 promoter repeat expansion in a Flanders-Belgian cohort with disorders of the frontotemporal lobar degeneration-amyotrophic lateral sclerosis spectrum: a gene identification study. Lancet Neurology, The, 2012, 11, 54-65.	4.9	565
59	Deep brain stimulation of the internal pallidum in multiple system atrophy. Parkinsonism and Related Disorders, 2006, 12, 181-183.	1.1	27