## Adrian Danek

## List of Publications by Year in descending order

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300 papers 12,921 citations

59 h-index 98 g-index

352 all docs

352 docs citations

times ranked

352

13232 citing authors

#	Article	IF	CITATIONS
1	Neuropsychological and psychiatric changes after deep brain stimulation for Parkinson's disease: a randomised, multicentre study. Lancet Neurology, The, 2008, 7, 605-614.	10.2	582
2	Symptom onset in autosomal dominant Alzheimer disease. Neurology, 2014, 83, 253-260.	1.1	391
3	Vestibular cortex lesions affect the perception of verticality. Annals of Neurology, 1994, 35, 403-412.	5.3	386
4	A conserved sorting-associated protein is mutant in chorea-acanthocytosis. Nature Genetics, 2001, 28, 119-120.	21.4	357
5	Motor recovery following capsular stroke. Brain, 1993, 116, 369-382.	7.6	341
6	Frontotemporal dementia and its subtypes: a genome-wide association study. Lancet Neurology, The, 2014, 13, 686-699.	10.2	302
7	Isolation of the gene for McLeod syndrome that encodes a novel membrane transport protein. Cell, 1994, 77, 869-880.	28.9	272
8	A Panâ€∢scp>European Study of the <i>C9orf72</i> Repeat Associated with <scp>FTLD</scp> : Geographic Prevalence, Genomic Instability, and Intermediate Repeats. Human Mutation, 2013, 34, 363-373.	2.5	247
9	McLeod neuroacanthocytosis: Genotype and phenotype. Annals of Neurology, 2001, 50, 755-764.	5.3	244
10	Deactivation of human visual cortex during involuntary ocular oscillations. Brain, 1996, 119, 101-110.	7.6	215
11	Early changes in CSF sTREM2 in dominantly inherited Alzheimer's disease occur after amyloid deposition and neuronal injury. Science Translational Medicine, 2016, 8, 369ra178.	12.4	211
12	Neuroacanthocytosis Syndromes. Orphanet Journal of Rare Diseases, 2011, 6, 68.	2.7	209
13	The anatomy of fronto-occipital connections from early blunt dissections to contemporary tractography. Cortex, 2014, 56, 73-84.	2.4	204
14	Chorein detection for the diagnosis of choreaâ€acanthocytosis. Annals of Neurology, 2004, 56, 299-302.	5.3	186
15	Age at symptom onset and death and disease duration in genetic frontotemporal dementia: an international retrospective cohort study. Lancet Neurology, The, 2020, 19, 145-156.	10.2	175
16	Clinical features and molecular bases of neuroacanthocytosis. Journal of Molecular Medicine, 2002, 80, 475-491.	3.9	174
17	Mutational spectrum of the CHAC gene in patients with chorea-acanthocytosis. European Journal of Human Genetics, 2002, 10, 773-781.	2.8	172
18	Glial Fibrillary Acidic Protein in Serum is Increased in Alzheimer's Disease and Correlates with Cognitive Impairment. Journal of Alzheimer's Disease, 2019, 67, 481-488.	2.6	171

#	Article	lF	Citations
19	Neurologic phenotypes associated with acanthocytosis. Neurology, 2007, 68, 92-98.	1.1	155
20	Imaging cortical anatomy by high-resolution MR at 3.0T: Detection of the stripe of Gennari in visual area 17. Magnetic Resonance in Medicine, 2002, 48, 735-738.	3.0	151
21	The Pattern of Cognitive Performance in CADASIL: A Monogenic Condition Leading to Subcortical Ischemic Vascular Dementia. American Journal of Psychiatry, 2005, 162, 2078-2085.	7.2	141
22	Serum neurofilament light chain in genetic frontotemporal dementia: a longitudinal, multicentre cohort study. Lancet Neurology, The, 2019, 18, 1103-1111.	10.2	128
23	Chorea-Acanthocytosis: Genetic Linkage to Chromosome 9q21. American Journal of Human Genetics, 1997, 61, 899-908.	6.2	126
24	Preclinical trials in autosomal dominant AD: Implementation of the DIAN-TU trial. Revue Neurologique, 2013, 169, 737-743.	1.5	122
25	slan-defined subsets of CD16-positive monocytes: impact of granulomatous inflammation and M-CSF receptor mutation. Blood, 2015, 126, 2601-2610.	1.4	116
26	White matter diffusion alterations precede symptom onset in autosomal dominant Alzheimer's disease. Brain, 2018, 141, 3065-3080.	7.6	116
27	Plasma glial fibrillary acidic protein is raised in progranulin-associated frontotemporal dementia. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 263-270.	1.9	106
28	McLeod syndrome: a distinct form of neuroacanthocytosis. Journal of Neurology, 1992, 239, 302-306.	3.6	98
29	Different neuroinflammatory profile in amyotrophic lateral sclerosis and frontotemporal dementia is linked to the clinical phase. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 4-10.	1.9	96
30	Evaluation of early-phase [ 18 F]-florbetaben PET acquisition in clinical routine cases. NeuroImage: Clinical, 2017, 14, 77-86.	2.7	91
31	Polyâ€∢scp>GP in cerebrospinal fluid links <i>C9orf72</i> â€associated dipeptide repeat expression to the asymptomatic phase of <scp>ALS</scp> / <scp>FTD</scp> . EMBO Molecular Medicine, 2017, 9, 859-868.	6.9	90
32	Chitotriosidase (CHIT1) is increased in microglia and macrophages in spinal cord of amyotrophic lateral sclerosis and cerebrospinal fluid levels correlate with disease severity and progression. Journal of Neurology, Neurosurgery and Psychiatry, 2018, 89, 239-247.	1.9	89
33	Neurological manifestations of autosomal dominant familial Alzheimer's disease: a comparison of the published literature with the Dominantly Inherited Alzheimer Network observational study (DIAN-OBS). Lancet Neurology, The, 2016, 15, 1317-1325.	10.2	87
34	<i>TBK1</i> Mutation Spectrum in an Extended European Patient Cohort with Frontotemporal Dementia and Amyotrophic Lateral Sclerosis. Human Mutation, 2017, 38, 297-309.	2.5	87
35	Serum neurofilament light chain in behavioral variant frontotemporal dementia. Neurology, 2018, 91, e1390-e1401.	1.1	85
36	Motor responses after transcranial electrical stimulation of cerebral hemispheres with a degenerated pyramidal tract. Annals of Neurology, 1991, 29, 646-650.	<b>5.</b> 3	84

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37	Neuroacanthocytosis. Current Opinion in Neurology, 2005, 18, 386-392.	3.6	84
38	Tracing of Neuronal Connections in the Human Brain by Magnetic Resonance Imaging in vivo. European Journal of Neuroscience, 1990, 2, 112-115.	2.6	83
39	Left frontal hub connectivity delays cognitive impairment in autosomal-dominant and sporadic Alzheimer's disease. Brain, 2018, 141, 1186-1200.	7.6	83
40	Neurofilament light chain as a blood biomarker to differentiate psychiatric disorders from behavioural variant frontotemporal dementia. Journal of Psychiatric Research, 2019, 113, 137-140.	3.1	81
41	Quantification of brain lesions using interactive automated software. Behavior Research Methods, 2002, 34, 6-18.	1.3	79
42	Preferential degradation of cognitive networks differentiates Alzheimer's disease from ageing. Brain, 2018, 141, 1486-1500.	7.6	79
43	Cortically evoked motor responses in patients with Xp22.3-linked Kallmann's syndrome and in female gene carriers. Annals of Neurology, 1992, 31, 299-304.	5.3	78
44	Medial Prefrontal and Subcortical Mechanisms Underlying the Acquisition of Motor and Cognitive Action Sequences in Humans. Neuron, 2002, 35, 371-381.	8.1	77
45	Neuroacanthocytosis: new developments in a neglected group of dementing disorders. Journal of the Neurological Sciences, 2005, 229-230, 171-186.	0.6	77
46	Central mechanisms in human enhanced physiological tremor. Neuroscience Letters, 1998, 241, 135-138.	2.1	75
47	Aneurysms and vacuolar degeneration of cerebral arteries in late-onset acid maltase deficiency. Journal of the Neurological Sciences, 1990, 98, 169-183.	0.6	74
48	Erythrocyte membrane changes of chorea-acanthocytosis are the result of altered Lyn kinase activity. Blood, 2011, 118, 5652-5663.	1.4	73
49	Neurofilament as a blood marker for diagnosis and monitoring of primary progressive aphasias. Neurology, 2017, 88, 961-969.	1.1	73
50	Aging is associated with increased collagen type IV accumulation in the basal lamina of human cerebral microvessels. BMC Neuroscience, 2004, 5, 37.	1.9	72
51	Kallman syndrome versus idiopathic hypogonadotropic hypogonadism at MR imaging Radiology, 1994, 191, 53-57.	7.3	71
52	Is improvement in the quality of life after subthalamic nucleus stimulation in Parkinson's disease predictable?. Movement Disorders, 2011, 26, 2516-2521.	3.9	71
53	Genetic Heterogeneity in Alzheimer Disease and Implications for Treatment Strategies. Current Neurology and Neuroscience Reports, 2014, 14, 499.	4.2	70
54	Mirror movements in healthy humans across the lifespan: effects of development and ageing. Developmental Medicine and Child Neurology, 2010, 52, 1106-1112.	2.1	69

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55	Risk factors for executive dysfunction after subthalamic nucleus stimulation in Parkinson's disease. Movement Disorders, 2010, 25, 1583-1589.	3.9	68
56	Specific serum and CSF microRNA profiles distinguish sporadic behavioural variant of frontotemporal dementia compared with Alzheimer patients and cognitively healthy controls. PLoS ONE, 2018, 13, e0197329.	2.5	68
57	Persistent mirror movements: functional MR imaging of the hand motor cortex Radiology, 1997, 203, 545-552.	7.3	67
58	Segregation of functional networks is associated with cognitive resilience in Alzheimer's disease. Brain, 2021, 144, 2176-2185.	7.6	66
59	Geniospasm: Hereditary chin trembling. Movement Disorders, 1993, 8, 335-338.	3.9	64
60	McLeod myopathy revisited: more neurogenic and less benign. Brain, 2007, 130, 3285-3296.	7.6	64
61	<scp>CSF</scp> progranulin increases in the course of Alzheimer's disease and is associated with <scp>sTREM</scp> 2, neurodegeneration and cognitive decline. EMBO Molecular Medicine, 2018, 10, .	6.9	64
62	Alzheimer's disease in Down syndrome: An overlooked population for prevention trials. Alzheimer's and Dementia: Translational Research and Clinical Interventions, 2018, 4, 703-713.	3.7	63
63	Head of the caudate nucleus is most vulnerable in chorea–acanthocytosis: A voxel-based morphometry study. Movement Disorders, 2006, 21, 1728-1731.	3.9	62
64	The BDNFVal66Met SNP modulates the association between beta-amyloid and hippocampal disconnection in Alzheimer's disease. Molecular Psychiatry, 2021, 26, 614-628.	7.9	61
65	McLeod syndrome: a neurohaematological disorder. Vox Sanguinis, 2007, 93, 112-121.	1.5	59
66	Tongue protrusion and feeding dystonia: A hallmark of choreaâ€acanthocytosis. Movement Disorders, 2010, 25, 127-129.	3.9	59
67	Factors Associated With the Onset and Persistence of Post–Lumbar Puncture Headache. JAMA Neurology, 2015, 72, 325.	9.0	59
68	[18F]-THK5351 PET Correlates with Topology and Symptom Severity in Progressive Supranuclear Palsy. Frontiers in Aging Neuroscience, 2017, 9, 440.	3.4	58
69	Neuronal pentraxin 2: a synapse-derived CSF biomarker in genetic frontotemporal dementia. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 612-621.	1.9	55
70	Characteristic head drops and axial extension in advanced choreaâ€acanthocytosis. Movement Disorders, 2010, 25, 1487-1491.	3.9	54
71	First manifestation of multiple sclerosis after immunization with the Pfizer-BioNTech COVID-19 vaccine. Journal of Neurology, 2022, 269, 55-58.	3.6	54
72	Validation of the German Revised Addenbrooke's Cognitive Examination for Detecting Mild Cognitive Impairment, Mild Dementia in Alzheimer's Disease and Frontotemporal Lobar Degeneration. Dementia and Geriatric Cognitive Disorders, 2010, 29, 448-456.	1.5	53

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73	Negative impact of borderline global cognitive scores on quality of life after subthalamic nucleus stimulation in Parkinson's disease. Journal of the Neurological Sciences, 2011, 310, 261-266.	0.6	52
74	Plasma Neurofilament Light for Prediction of Disease Progression in Familial Frontotemporal Lobar Degeneration. Neurology, 2021, 96, e2296-e2312.	1.1	52
75	Recognition of Facial Expressions of Different Emotional Intensities in Patients with Frontotemporal Lobar Degeneration. Behavioural Neurology, 2007, 18, 31-36.	2.1	51
76	Relationship between physical activity, cognition, and Alzheimer pathology in autosomal dominant Alzheimer's disease. Alzheimer's and Dementia, 2018, 14, 1427-1437.	0.8	51
77	Current state of knowledge in Chorea-Acanthocytosis as core Neuroacanthocytosis syndrome. European Journal of Medical Genetics, 2018, 61, 699-705.	1.3	50
78	Shape alterations in the striatum in chorea-acanthocytosis. Psychiatry Research - Neuroimaging, 2011, 192, 29-36.	1.8	49
79	The chorea of McLeod syndrome. Movement Disorders, 2001, 16, 882-889.	3.9	48
80	A new molecular link between defective autophagy and erythroid abnormalities in chorea-acanthocytosis. Blood, 2016, 128, 2976-2987.	1.4	47
81	Predicting sporadic Alzheimer's disease progression via inherited Alzheimer's diseaseâ€informed machineâ€learning. Alzheimer's and Dementia, 2020, 16, 501-511.	0.8	47
82	Cortical [ <scp><sup>18</sup>F</scp> ] <scp>PI</scp> â€2620 Binding Differentiates Corticobasal Syndrome Subtypes. Movement Disorders, 2021, 36, 2104-2115.	3.9	46
83	The neuropsychiatry of neuroacanthocytosis syndromes. Neuroscience and Biobehavioral Reviews, 2011, 35, 1275-1283.	6.1	45
84	Cerebral involvement in McLeod syndrome. Neurology, 1994, 44, 117-117.	1.1	45
85	Developments in neuroacanthocytosis: Expanding the spectrum of choreatic syndromes. Movement Disorders, 2006, 21, 1794-1805.	3.9	44
86	Short and Long Term Outcome of Bilateral Pallidal Stimulation in Chorea-Acanthocytosis. PLoS ONE, 2013, 8, e79241.	2.5	44
87	Anisotropy of transcallosal motor fibres indicates functional impairment in children with periventricular leukomalacia. Developmental Medicine and Child Neurology, 2011, 53, 179-186.	2.1	43
88	Predicting primary progressive aphasias with support vector machine approaches in structural MRI data. Neurolmage: Clinical, 2017, 14, 334-343.	2.7	42
89	Metabolic Correlates of Dopaminergic Loss in Dementia with Lewy Bodies. Movement Disorders, 2020, 35, 595-605.	3.9	42
90	Progression of Behavioral Disturbances and Neuropsychiatric Symptoms in Patients With Genetic Frontotemporal Dementia. JAMA Network Open, 2021, 4, e2030194.	5.9	42

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91	Long-term follow-up in primary progressive aphasia: Clinical course and health care utilisation. Aphasiology, 2014, 28, 981-992.	2.2	41
92	Atrophy in the Thalamus But Not Cerebellum Is Specific for C9orf72 FTD and ALS Patients – An Atlas-Based Volumetric MRI Study. Frontiers in Aging Neuroscience, 2018, 10, 45.	3.4	40
93	4-Aminopyridine improves gait variability in cerebellar ataxia due to CACNA 1A mutation. Journal of Neurology, 2011, 258, 1708-1711.	3.6	39
94	Brain, blood, and iron: Perspectives on the roles of erythrocytes and iron in neurodegeneration. Neurobiology of Disease, 2012, 46, 607-624.	4.4	39
95	A C6orf10/LOC101929163 locus is associated with age of onset in C9orf72 carriers. Brain, 2018, 141, 2895-2907.	7.6	39
96	Reduction of striatal glucose metabolism in McLeod choreoacanthocytosis. Journal of Neurology, Neurosurgery and Psychiatry, 2001, 70, 517-520.	1.9	38
97	Phenotypic variation among brothers with the McLeod neuroacanthocytosis syndrome. Movement Disorders, 2007, 22, 244-247.	3.9	38
98	Seizures in Alzheimer's disease are highly recurrent and associated with a poor disease course. Journal of Neurology, 2020, 267, 2941-2948.	3.6	38
99	Lyme neuroborreliosis disguised as normal pressure hydrocephalus. Neurology, 1996, 46, 1743-1745.	1.1	37
100	White matter abnormalities on MRI in neuroacanthocytosis. Journal of Neurology, Neurosurgery and Psychiatry, 2004, 75, 1200-1201.	1.9	37
101	McLeod phenotype without the McLeod syndrome. Transfusion, 2007, 47, 299-305.	1.6	37
102	In Vivo Assessment of Neuroinflammation in <scp>4â€Repeat</scp> Tauopathies. Movement Disorders, 2021, 36, 883-894.	3.9	37
103	Identification of a VPS13A founder mutation in French Canadian families with chorea-acanthocytosis. Neurogenetics, 2005, 6, 151-158.	1.4	36
104	Lexical retrieval treatment in primary progressive aphasia: An investigation of treatment duration in a heterogeneous case series. Cortex, 2019, 115, 133-158.	2.4	36
105	Clinical, pathophysiological and genetic features of motor symptoms in autosomal dominant Alzheimer's disease. Brain, 2019, 142, 1429-1440.	7.6	36
106	Brain functional network integrity sustains cognitive function despite atrophy in presymptomatic genetic frontotemporal dementia. Alzheimer's and Dementia, 2021, 17, 500-514.	0.8	36
107	Eye Movements in Chorea-Acanthocytosis. , 2005, 46, 1979.		34
108	A Modified Reading the Mind in the Eyes Test Predicts Behavioral Variant Frontotemporal Dementia Better Than Executive Function Tests. Frontiers in Aging Neuroscience, 2018, 10, 11.	3.4	34

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109	"Neuroacanthocytosis―– Overdue for a Taxonomic Update. Tremor and Other Hyperkinetic Movements, 2021, 11, 1.	2.0	34
110	The inner fluctuations of the brain in presymptomatic Frontotemporal Dementia: The chronnectome fingerprint. Neurolmage, 2019, 189, 645-654.	4.2	33
111	Different CSF protein profiles in amyotrophic lateral sclerosis and frontotemporal dementia with <i>C9orf72</i> hexanucleotide repeat expansion. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 503-511.	1.9	33
112	Prosopagnosia after unilateral right cerebral infarction. Journal of Neurology, 2002, 249, 933-935.	3.6	32
113	Neuroacanthocytosis. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2011, 100, 141-151.	1.8	32
114	Apathy in presymptomatic genetic frontotemporal dementia predicts cognitive decline and is driven by structural brain changes. Alzheimer's and Dementia, 2021, 17, 969-983.	0.8	31
115	Deep brain stimulation in chorea acanthocytosis. Movement Disorders, 2009, 24, 1546-1547.	3.9	30
116	PET Imaging of Astrogliosis and Tau Facilitates Diagnosis of Parkinsonian Syndromes. Frontiers in Aging Neuroscience, 2019, 11, 249.	3.4	30
117	Binding characteristics of [ <sup>18</sup> F]Pl-2620 distinguish the clinically predicted tau isoform in different tauopathies by PET. Journal of Cerebral Blood Flow and Metabolism, 2021, 41, 2957-2972.	4.3	30
118	Movement-related cortical potentials in persistent mirror movements. Electroencephalography and Clinical Neurophysiology, 1995, 95, 350-358.	0.3	29
119	DEVELOPMENT OF MESIAL TEMPORAL LOBE EPILEPSY IN CHOREA-ACANTHOCYTOSIS. Neurology, 2009, 73, 1419-1422.	1.1	29
120	Characterizing the Clinical Features and Atrophy Patterns of <i>MAPT</i> -Related Frontotemporal Dementia With Disease Progression Modeling. Neurology, 2021, 97, e941-e952.	1.1	29
121	FDG-PET underscores the key role of the thalamus in frontotemporal lobar degeneration caused by C9ORF72 mutations. Translational Psychiatry, 2019, 9, 54.	4.8	28
122	Differential early subcortical involvement in genetic FTD within the GENFI cohort. NeuroImage: Clinical, 2021, 30, 102646.	2.7	28
123	Automatic striatal volumetry allows for identification of patients with chorea-acanthocytosis at single subject level. Journal of Neural Transmission, 2008, 115, 1393-1400.	2.8	27
124	Cerebral amyloidosis associated with cognitive decline in autosomal dominant Alzheimer disease. Neurology, 2015, 85, 790-798.	1.1	27
125	White matter hyperintensities in progranulin-associated frontotemporal dementia: A longitudinal GENFI study. NeuroImage: Clinical, 2019, 24, 102077.	2.7	27
126	Seizures as an early symptom of autosomal dominant Alzheimer's disease. Neurobiology of Aging, 2019, 76, 18-23.	3.1	27

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127	A data-driven disease progression model of fluid biomarkers in genetic frontotemporal dementia. Brain, 2022, 145, 1805-1817.	7.6	27
128	Social cognition impairment in genetic frontotemporal dementia within the GENFI cohort. Cortex, 2020, 133, 384-398.	2.4	26
129	A novel mutation in CACNA1A associated with hemiplegic migraine, cerebellar dysfunction and late-onset cognitive decline. Journal of the Neurological Sciences, 2011, 300, 160-163.	0.6	25
130	Brain atrophy in primary progressive aphasia involves the cholinergic basal forebrain and Ayala's nucleus. Psychiatry Research - Neuroimaging, 2014, 221, 187-194.	1.8	25
131	Early symptoms in symptomatic and preclinical genetic frontotemporal lobar degeneration. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 975-984.	1.9	25
132	Brain imaging and neuropsychology in late-onset dementia due to a novel mutation (R93C) of valosin-containing protein., 2007, 26, 232-240.		25
133	Adolescent obsessive compulsive disorder heralding choreaâ€acanthocytosis. Movement Disorders, 2008, 23, 422-425.	3.9	24
134	Conceptual framework for the definition of preclinical and prodromal frontotemporal dementia. Alzheimer's and Dementia, 2022, 18, 1408-1423.	0.8	24
135	Phenotypic variability of a distinct deletion in McLeod syndrome. Movement Disorders, 2007, 22, 1358-1361.	3.9	23
136	Cerebral Glucose Metabolism and Dopaminergic Function in Patients with Corticobasal Syndrome. Journal of Neuroimaging, 2017, 27, 255-261.	2.0	23
137	Education modulates brain maintenance in presymptomatic frontotemporal dementia. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 1124-1130.	1.9	23
138	Clinico-genetic findings in 509 frontotemporal dementia patients. Molecular Psychiatry, 2021, 26, 5824-5832.	7.9	23
139	Primary diffuse leptomeningeal gliomatosis: unusual MRI with non-enhancing nodular lesions on the cerebellar surface and spinal leptomeningeal enhancement. Journal of Neurology, Neurosurgery and Psychiatry, 2000, 69, 385-388.	1.9	22
140	Alterations of Red Cell Membrane Properties in Nneuroacanthocytosis. PLoS ONE, 2013, 8, e76715.	2.5	22
141	Quantifying progression in primary progressive aphasia with structural neuroimaging. Alzheimer's and Dementia, 2021, 17, 1595-1609.	0.8	22
142	Atrophy and structural covariance of the cholinergic basal forebrain in primary progressive aphasia. Cortex, 2016, 83, 124-135.	2.4	21
143	The Erythrocyte Sedimentation Rate and Its Relation to Cell Shape and Rigidity of Red Blood Cells from Chorea-Acanthocytosis Patients in an Off-Label Treatment with Dasatinib. Biomolecules, 2021, 11, 727.	4.0	21
144	Stratifying the Presymptomatic Phase of Genetic Frontotemporal Dementia by Serum <scp>NfL</scp> and <scp>pNfH</scp> : A Longitudinal Multicentre Study. Annals of Neurology, 2022, 91, 33-47.	5.3	21

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145	Normal dystrophin in McLeod myopathy. Annals of Neurology, 1990, 28, 720-722.	5.3	20
146	Analysis of brain atrophy and local gene expression in genetic frontotemporal dementia. Brain Communications, 2020, 2, .	3.3	20
147	Persistent mirror movements: force and timing of ?mirroring? are task-dependent. Experimental Brain Research, 1995, 104, 126-34.	1.5	19
148	Quantitative evaluation of mirror movements in adults with focal brain lesions. European Journal of Neurology, 2005, 12, 964-975.	3.3	19
149	Computational Identification of Phospho-Tyrosine Sub-Networks Related to Acanthocyte Generation in Neuroacanthocytosis. PLoS ONE, 2012, 7, e31015.	2.5	19
150	Faster Cortical Thinning and Surface Area Loss in Presymptomatic and Symptomatic <i>C9orf72</i> Repeat Expansion Adult Carriers. Annals of Neurology, 2020, 88, 113-122.	5.3	19
151	Therapeutic targeting of Lyn kinase to treat chorea-acanthocytosis. Acta Neuropathologica Communications, 2021, 9, 81.	5.2	19
152	Olfactory function in patients with hypogonadotropic hypogonadism: an all-or-none phenomenon?. Chemical Senses, 1994, 19, 57-69.	2.0	18
153	Bilateral temporal lobe epilepsy confirmed with intracranial EEG in chorea-acanthocytosis. Seizure: the Journal of the British Epilepsy Association, 2011, 20, 340-342.	2.0	18
154	A modified Camel and Cactus Test detects presymptomatic semantic impairment in genetic frontotemporal dementia within the GENFI cohort. Applied Neuropsychology Adult, 2022, 29, 112-119.	1.2	18
155	Acanthocyte Sedimentation Rate as a Diagnostic Biomarker for Neuroacanthocytosis Syndromes: Experimental Evidence and Physical Justification. Cells, 2021, 10, 788.	4.1	18
156	Unraveling corticobasal syndrome and alien limb syndrome with structural brain imaging. Cortex, 2019, 117, 33-40.	2.4	17
157	Choreaâ€acanthocytosis: Report of two Brazilian cases. Movement Disorders, 2008, 23, 2090-2093.	3.9	16
158	Hippocampal sclerosis and mesial temporal lobe epilepsy in choreaâ€acanthocytosis: a case with clinical, pathologic and genetic evaluation. Neuropathology and Applied Neurobiology, 2017, 43, 542-546.	3.2	16
159	A mathematical model and a computer tool for the Tower of Hanoi and Tower of London puzzles. Information Sciences, 2009, 179, 2934-2947.	6.9	15
160	Autosomal recessive transmission of chorea-acanthocytosis confirmed. Acta Neuropathologica, 2012, 123, 905-906.	7.7	15
161	Additive value of amyloid-PET in routine cases of clinical dementia work-up after FDG-PET. European Journal of Nuclear Medicine and Molecular Imaging, 2017, 44, 2239-2248.	6.4	15
162	Life expectancy and mortality in chorea-acanthocytosis and McLeod syndrome. Parkinsonism and Related Disorders, 2019, 60, 158-161.	2.2	15

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163	The applause sign in frontotemporal lobar degeneration and related conditions. Journal of Neurology, 2019, 266, 330-338.	3.6	15
164	Atypical parkinsonism due to a <i>D202N</i> Gerstmannâ€ <scp>S</scp> trässlerâ€ <scp>S</scp> cheinker prion protein mutation: First in vivo diagnosed case. Movement Disorders, 2013, 28, 241-245.	3.9	14
165	Neuronal injury biomarkers for assessment of the individual cognitive reserve in clinically suspected Alzheimer's disease. Neurolmage: Clinical, 2019, 24, 101949.	2.7	14
166	Erysense, a Lab-on-a-Chip-Based Point-of-Care Device to Evaluate Red Blood Cell Flow Properties With Multiple Clinical Applications. Frontiers in Physiology, 2022, 13, 884690.	2.8	14
167	The Dream King's Psychiatrist Bernhard von Gudden (1824-1886). Archives of Neurology, 1989, 46, 1349.	4.5	13
168	Dominant transmission of chorea-acanthocytosis with VPS13A mutations remains speculative. Acta Neuropathologica, 2009, 117, 95-96.	7.7	13
169	Dual-Phase $\hat{l}^2$ -Amyloid PET Captures Neuronal Injury and Amyloidosis in Corticobasal Syndrome. Frontiers in Aging Neuroscience, 2021, 13, 661284.	3.4	13
170	Cognitive reserve hypothesis in frontotemporal dementia: A FDG-PET study. NeuroImage: Clinical, 2021, 29, 102535.	2.7	13
171	Dipole source analysis in persistent mirror movements. Brain Topography, 1999, 12, 49-60.	1.8	12
172	Clinical and genetic analysis of 29 Brazilian patients with Huntington's disease-like phenotype. Arquivos De Neuro-Psiquiatria, 2011, 69, 419-423.	0.8	12
173	Abnormal pain perception is associated with thalamo-cortico-striatal atrophy in <i>C9orf72</i> expansion carriers in the GENFI cohort. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 1325-1328.	1.9	12
174	Chorea-Acanthocytosis Genotype in the Original Critchley Kentucky Neuroacanthocytosis Kindred. Archives of Neurology, 2011, 68, 1330.	4.5	11
175	The chorea of McLeod syndrome: Progression to hypokinesia. Movement Disorders, 2012, 27, 1701-1702.	3.9	11
176	Subcortical neurodegeneration in chorea: Similarities and differences between chorea-acanthocytosis and Huntington's disease. Parkinsonism and Related Disorders, 2018, 49, 54-59.	2.2	11
177	Impairment of episodic memory in genetic frontotemporal dementia: A GENFI study. Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring, 2021, 13, e12185.	2.4	11
178	First symptom guides diagnosis and prognosis in neurodegenerative diseasesâ€"a retrospective study of autopsy proven cases. European Journal of Neurology, 2021, 28, 1801-1811.	3.3	11
179	XK-Associated McLeod Syndrome: Nonhematological Manifestations and Relation to VPS13A Disease. Transfusion Medicine and Hemotherapy, 2022, 49, 4-12.	1.6	11
180	Diffuse leukoencephalopathy with spheroids: Biopsy findings and a novel mutation. Clinical Neurology and Neurosurgery, 2014, 122, 113-115.	1.4	10

#	Article	IF	CITATIONS
181	Pathoarchitectonics of the cerebral cortex in choreaâ€acanthocytosis and Huntington's disease. Neuropathology and Applied Neurobiology, 2019, 45, 230-243.	3.2	10
182	MRI data-driven algorithm for the diagnosis of behavioural variant frontotemporal dementia. Journal of Neurology, Neurosurgery and Psychiatry, 2021, 92, 608-616.	1.9	10
183	Gene Expression Imputation Across Multiple Tissue Types Provides Insight Into the Genetic Architecture of Frontotemporal Dementia and Its Clinical Subtypes. Biological Psychiatry, 2021, 89, 825-835.	1.3	10
184	Motor speech disorders in the nonfluent, semantic and logopenic variants of primary progressive aphasia. Cortex, 2021, 140, 66-79.	2.4	10
185	Is transient global amnesia related to endogenous benzodiazepines?. Journal of Neurology, 2002, 249, 628-628.	3.6	9
186	Volitional facial palsy after a vascular lesion of the supplementary motor area. Neurology, 2004, 63, 756-757.	1.1	9
187	An episode of geniospasm in sleep: Toward new insights into pathophysiology?. Movement Disorders, 2008, 23, 274-276.	3.9	9
188	The iso-effect: Is there specific learning of Tower of London iso-problems?. Thinking and Reasoning, 2009, 15, 237-249.	3.2	9
189	Huntington's disease: new aspects on phenotype and genotype. Parkinsonism and Related Disorders, 2012, 18, S107-S109.	2.2	9
190	Wildervanck's syndrome and mirror movements: a congenital disorder of axon migration?. Journal of Neurology, 2012, 259, 761-763.	3.6	9
191	The association of aphasia and right-sided motor impairment in corticobasal syndrome. Journal of Neurology, 2015, 262, 2241-2246.	3.6	9
192	Identification of a rare presenilin 1 single amino acid deletion mutation (F175del) with unusual amyloid- $\hat{l}^2$ processing effects. Neurobiology of Aging, 2019, 84, 241.e5-241.e11.	3.1	9
193	Neuroacanthocytosis in China: A Review of Published Reports. Tremor and Other Hyperkinetic Movements, 2020, 4, 248.	2.0	9
194	Neuroacanthocytosis in china: a review of published reports. Tremor and Other Hyperkinetic Movements, 2014, 4, 248.	2.0	9
195	A panel of CSF proteins separates genetic frontotemporal dementia from presymptomatic mutation carriers: a GENFI study. Molecular Neurodegeneration, 2021, 16, 79.	10.8	9
196	Serum <scp>Betaâ€Synuclein</scp> Is Higher in Down Syndrome and Precedes Rise of <scp>pTau181</scp> . Annals of Neurology, 2022, 92, 6-10.	5.3	9
197	Comparative analysis of machine learning algorithms for multi-syndrome classification of neurodegenerative syndromes. Alzheimer's Research and Therapy, 2022, 14, 62.	6.2	9
198	CSF tau proteins in differential diagnosis of dementia. Translational Neuroscience, 2010, 1, 43-48.	1.4	8

#	Article	IF	CITATIONS
199	Three novel presenilin 1 mutations marking the wide spectrum of age at onset and clinical patterns in familial Alzheimer's disease. Journal of Neural Transmission, 2015, 122, 1715-1719.	2.8	8
200	A language-based sum score for the course and therapeutic intervention in primary progressive aphasia. Alzheimer's Research and Therapy, 2018, 10, 41.	6.2	8
201	Disease-related cortical thinning in presymptomatic granulin mutation carriers. Neurolmage: Clinical, 2021, 29, 102540.	2.7	8
202	Cognitive and Neuropsychiatric Findings in McLeod Syndrome and in Chorea-Acanthocytosis. , 2004, , 95-115.		8
203	Loss of word-meaning with spared object semantics in a case of mixed primary progressive aphasia. Brain and Language, 2010, 113, 96-100.	1.6	7
204	Magnetic resonance spectroscopy in two siblings with choreaâ€acanthocytosis. Movement Disorders, 2010, 25, 2894-2897.	3.9	7
205	Diagnostic Value of Subjective Memory Complaints Assessed with a Single Item in Dominantly Inherited Alzheimer's Disease: Results of the DIAN Study. BioMed Research International, 2015, 2015, 1-7.	1.9	7
206	Comparison of clinical rating scales in genetic frontotemporal dementia within the GENFI cohort. Journal of Neurology, Neurosurgery and Psychiatry, 2022, 93, 158-168.	1.9	7
207	Neuropathology of Chorea-Acanthocytosis. , 2008, , 187-195.		7
208	Dataâ€driven staging of genetic frontotemporal dementia using multiâ€modal <scp>MRI</scp> . Human Brain Mapping, 2022, 43, 1821-1835.	3.6	7
209	A longitudinal linguistic analysis of written text production in a case of semantic variant primary progressive aphasia. Journal of Neurolinguistics, 2016, 39, 26-37.	1.1	6
210	Seizure prevalence in neurodegenerative diseasesâ€"a study of autopsy proven cases. European Journal of Neurology, 2022, 29, 12-18.	3.3	6
211	Neurofilament light chain in serum is significantly increased in chorea-acanthocytosis. Parkinsonism and Related Disorders, 2020, 80, 28-31.	2.2	6
212	Bernhard von Gudden, Neuro-Ophthalmology and the Munich School of Neuroanatomy and Psychiatry. Strabismus, 2006, 14, 211-216.	0.7	5
213	Diagnostic evaluation of clinically normal subjects with chronic hyperCKemia. Neurology, 2007, 68, 535-536.	1.1	5
214	Head drops are also observed in advanced Huntington disease. Parkinsonism and Related Disorders, 2013, 19, 569-570.	2.2	5
215	TSPO PET With 18F-GE-180 to Differentiate Variants of Multiple Sclerosis. Clinical Nuclear Medicine, 2020, 45, e447-e448.	1.3	5
216	Identification of two compound heterozygous <i>VPS13A</i> large deletions in choreaâ€acanthocytosis only by protein and quantitative DNA analysis. Molecular Genetics & Enomic Medicine, 2020, 8, e1179.	1.2	5

#	Article	IF	CITATIONS
217	Neuroacanthocytosis Syndromes: What Links Red Blood Cells and Neurons?., 2004, , 1-14.		5
218	Cerebral Involvement in McLeod Syndrome: The First Autopsy Revisited. , 2008, , 205-215.		5
219	Adaptative Up-Regulation of PRX2 and PRX5 Expression Characterizes Brain from a Mouse Model of Chorea-Acanthocytosis. Antioxidants, 2022, $11$ , $76$ .	5.1	5
220	Longitudinal Cognitive Changes in Genetic Frontotemporal Dementia Within the GENFI Cohort. Neurology, 2022, 99, .	1,1	5
221	Polyradiculitis, Perimyocarditis, and Nephrotic Syndrome: Unusual Manifestations of Infection Due to Listeria monocytogenes. Clinical Infectious Diseases, 1999, 28, 152-153.	5.8	4
222	The Challenge of Axonal Path-Finding. Strabismus, 2006, 14, 95-99.	0.7	4
223	Mineral deposition on magnetic resonance imaging in chorea-acanthocytosis: A pathogenic link with pantothenate kinase-associated neurodegeneration?. Neurology India, 2013, 61, 169.	0.4	4
224	Occurrence of the "applause sign―in patients with amyotrophic lateral sclerosis. Clinical Neurology and Neurosurgery, 2015, 137, 8-10.	1.4	4
225	Local feature suppression effect in face and non-face stimuli. Psychological Research, 2015, 79, 194-205.	1.7	4
226	Mendelian randomization implies no direct causal association between leukocyte telomere length and amyotrophic lateral sclerosis. Scientific Reports, 2020, 10, 12184.	3.3	4
227	Low-degree trisomy 21 mosaicism promotes early-onset Alzheimer disease. Neurobiology of Aging, 2021, 103, 147.e1-147.e5.	3.1	4
228	Cardiac manifestation is evident in chorea-acanthocytosis but different from McLeod syndrome. Parkinsonism and Related Disorders, 2021, 88, 90-95.	2.2	4
229	Different rates of cognitive decline in autosomal dominant and lateâ€onset Alzheimer disease. Alzheimer's and Dementia, 2022, 18, 1754-1764.	0.8	4
230	Predicting disease progression in behavioral variant frontotemporal dementia. Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring, 2021, 13, e12262.	2.4	4
231	Cognitive composites for genetic frontotemporal dementia: GENFI-Cog. Alzheimer's Research and Therapy, 2022, 14, 10.	6.2	4
232	Applied multimodal diagnostics in a case of presenile dementia. BMC Neurology, 2016, 16, 131.	1.8	3
233	Dissemination in time and space in presymptomatic granulin mutation carriers: a GENFI spatial chronnectome study. Neurobiology of Aging, 2021, 108, 155-167.	3.1	3
234	Normdaten zur Erg $\tilde{A}$ <b>\mathbf{\mathbf{n}}</b> zung deutscher Wortanf $\tilde{A}$ <b>\mathbf{n}</b> ge. Zeitschrift F $\tilde{A}$ <sup>1</sup> /4r Neuropsychologie = Journal of Neuropsychology, 2010, 21, 17-23.	0.6	3

#	Article	IF	Citations
235	An Automated Toolbox to Predict Single Subject Atrophy in Presymptomatic Granulin Mutation Carriers. Journal of Alzheimer's Disease, 2022, , 1-14.	2.6	3
236	Ockham's razor, not a barber's weapon but a writer's tool. Brain, 2022, 145, 1870-1873.	7.6	3
237	Karl Friedrich Scheid (1906-1945). Der Nervenarzt, 2002, 73, 1130-1131.	5.0	2
238	Postoperative amnesic state with impairment of static visual perception. European Journal of Neurology, 2008, 15, e44-e45.	3.3	2
239	Mutation in the <i>CHAC</i> gene in a family of autosomal dominant chorea–acanthocytosis. Neurology, 2012, 79, 198-199.	1.1	2
240	Detection Gap of Right-Asymmetric Neuronal Degeneration by CERAD Test Battery in Alzheimer's Disease. Frontiers in Aging Neuroscience, 2021, 13, 611595.	3.4	2
241	Neuroacanthocytosis Syndromes — A Current Overview. , 2008, , 3-20.		2
242	Volumetric Neuroimaging in Neuroacanthocytosis. , 2008, , 175-185.		2
243	Eighth International Chorea-Acanthocytosis Symposium: Summary of Workshop Discussion and Action Points. Tremor and Other Hyperkinetic Movements, 2017, 7, 428.	2.0	2
244	Impact of Partial Volume Correction on [18F]GE-180 PET Quantification in Subcortical Brain Regions of Patients with Corticobasal Syndrome. Brain Sciences, 2022, 12, 204.	2.3	2
245	Examining empathy deficits across familial forms of frontotemporal dementia within the GENFI cohort. Cortex, 2022, 150, 12-28.	2.4	2
246	Patterns and implications of neurological examination findings in autosomal dominant Alzheimer disease. Alzheimer's and Dementia, $0$ , , .	0.8	2
247	The Meaning of Sherlock Holmes. Archives of Neurology, 1992, 49, 350-350.	4.5	1
248	Kognitive Neurologie und Neuropsychologie., 2005,, 41-82.		1
249	Antisocial behaviour and neuroacanthocytosis. International Journal of Clinical Practice, 2007, 61, 1419-1419.	1.7	1
250	Reply: Choreaâ€acanthocytosis: Report of two Brazilian cases. Movement Disorders, 2009, 24, 1254-1254.	3.9	1
251	O5â€02â€02: Personality Characteristics and Alterations in Familial Alzheimer's Disease in the Dominantly Inherited Alzheimer Network (DIAN). Alzheimer's and Dementia, 2016, 12, P379.	0.8	1
252	Response to "Neuroacanthocytosis: A case with unusual clinical features and novel response to treatment―by Wu et al Journal of the Neurological Sciences, 2017, 373, 347.	0.6	1

#	Article	IF	Citations
253	Chorein Deficiency and Alzheimer Disease: An Intriguing, Yet Premature Speculation. Alzheimer Disease and Associated Disorders, 2017, 31, 80-81.	1.3	1
254	[O1–O2–O5]: GENOTYPIC VARIANCE MAY EXPLAIN THE BALANCE OF EARLY CORTICAL VERSUS STRIATAL AMYLOID DEPOSITION IN AUTOSOMAL DOMINANT AD. Alzheimer's and Dementia, 2017, 13, P187.	0.8	1
255	FDGâ€PET in a Case of Very Lateâ€onset Huntington's Disease. Movement Disorders Clinical Practice, 2018, 5, 227-228.	1.5	1
256	18 Fâ€Plâ€2620 tauâ€PET in corticobasal syndrome (ActiGliA cohort). Alzheimer's and Dementia, 2020, 16, e041469.	0.8	1
257	"Huntington disease-like phenotype in a patient with ANO3 mutation―Expert commentary. Parkinsonism and Related Disorders, 2021, 90, 123-124.	2.2	1
258	Practice effects in genetic frontotemporal dementia and at-risk individuals: a GENFI study. Journal of Neurology, Neurosurgery and Psychiatry, 2022, 93, 336-339.	1.9	1
259	Pick-Komplex und andere fokale Hirnatrophien. , 2009, , 123-139.		1
260	Eighth International Chorea–Acanthocytosis Symposium: Summary of Workshop Discussion and Action Points. Tremor and Other Hyperkinetic Movements, 2020, 7, 428.	2.0	1
261	Subtype and stage inference identifies distinct atrophy patterns in genetic frontotemporal dementia that MAP onto specific MAPT mutations. Alzheimer's and Dementia, 2020, 16, e042996.	0.8	1
262	Structural brain splitting is a hallmark of Granulin-related frontotemporal dementia. Neurobiology of Aging, 2022, , .	3.1	1
263	Anomia is present pre-symptomatically in frontotemporal dementia due to MAPT mutations. Journal of Neurology, 2022, 269, 4322-4332.	3.6	1
264	Obsessive-compulsive Disorders Due to Neuroacanthocytosis Treated with Citalopram. Pharmacopsychiatry, 2007, 40, 132-132.	3.3	0
265	Questions on the Movement of the Human Eye (Part II). Strabismus, 2007, 15, 221-225.	0.7	0
266	1.309 Clinical and molecular studies of patients screened for Huntington's disease in a movement disorders clinic from Brazil. Parkinsonism and Related Disorders, 2007, 13, S79.	2.2	0
267	Questions on the Movement of the Human Eye (Part I). Strabismus, 2007, 15, 119-125.	0.7	0
268	A PATIENT WITH CHOREA-ACANTHOCYTOSIS AND DILATED CARDIOMYOPATHY. Journal of Neurology, Neurosurgery and Psychiatry, 2012, 83, A14.4-A15.	1.9	0
269	Oligoclonal bands in hereditary diffuse leukencephalopathy with spheroids. European Journal of Neurology, 2015, 22, e48-e48.	3.3	0
270	O5-02-01: Longitudinal Clinical and Biomarker Changes in Dominantly Inherited Alzheimer's Disease: The Dominantly Inherited Alzheimer Network., 2016, 12, P378-P379.		0

#	Article	IF	CITATIONS
271	Heinrich Simon Frenkel (1860–1931). Journal of Neurology, 2017, 264, 1301-1303.	3.6	O
272	[P3–263]: MOTOR SYMPTOMS IN FAMILIAL ALZHEIMER's DISEASE: FREQUENCY, SEVERITY AND PREDICTIVE VALUE. Alzheimer's and Dementia, 2017, 13, P1043.	0.8	0
273	[O1–02–06]: ELEVATED CSF STREM2 IN AUTOSOMAL DOMINANTLY INHERITED ALZHEIMER's DISEASE ASSOCIATED WITH REGIONAL FIBER TRACT INJURY: RESULTS FROM THE DIAN STUDY. Alzheimer's and Dementia, 2017, 13, P188.	0.8	O
274	HyperCKemia instead of Hyperkalemia in Chorea-Acanthocytosis. Cellular Physiology and Biochemistry, 2017, 41, 1267-1268.	1.6	0
275	P1â€497: CLINICAL CORRELATES OF LEWY BODY PATHOLOGY IN AUTOSOMAL DOMINANT ALZHEIMER DISEASE. Alzheimer's and Dementia, 2018, 14, P520.	0.8	O
276	F20â€Life expectancy and mortality in neuroacanthocytosis. , 2018, , .		0
277	O4â€04â€05: FIBERâ€TRACTâ€"SPECIFIC DECLINE IN WHITEâ€MATTER INTEGRITY DURING THE ADULT LIFESPAN PRECLINICAL ALZHEIMER'S DISEASE: RESULTS FROM THE DALLAS LIFESPAN BRAIN STUDY AND DIAN. Alzheimer's and Dementia, 2018, 14, P1410.	AND 0.8	O
278	P2â€628: RELATIONSHIP BETWEEN PHYSICAL ACTIVITY, COGNITION AND ALZHEIMER PATHOLOGY IN AUTOSOMAL DOMINANT ALZHEIMER'S DISEASE. Alzheimer's and Dementia, 2018, 14, P982.	0.8	0
279	Acute frontal eye field infarction. Neurology, 2019, 92, 193-195.	1.1	O
280	Trajectory of apathy, cognition and neural correlates in the decades before symptoms in frontotemporal dementia. Alzheimer's and Dementia, 2020, 16, e041821.	0.8	0
281	Neuropathological characteristics associated with a recently identified rare PSEN1 deletion mutation (F175del). Alzheimer's and Dementia, 2020, 16, e045048.	0.8	O
282	The Free Cued Selective Reminding Test detects episodic memory impairment in the presymptomatic period of familial frontotemporal dementia within the GENFI cohort. Alzheimer's and Dementia, 2020, 16, e045768.	0.8	0
283	Living with global amnesia: self-established compensation strategies of a patient with severe memory impairment – a narrative report. Neurocase, 2021, 27, 287-296.	0.6	O
284	Comment on "A New Allelic Variant in the PANK2 Gene in a Patient with Incomplete HARP Syndrome― Journal of Movement Disorders, 2021, 14, 254-255.	1.3	0
285	F28â€Novel mutations and findings in a cohort of McLeod neuroacanthocytosis, an X-linked HD phenocopy. , 2021, , .		O
286	"Virtual Neuroacanthocytosis Institute― A Look Forward. , 2008, , 287-292.		0
287	Neuroacanthocytosis. Clinics, 2008, 63, 135.	1.5	O
288	Sleep Disorders in Neuroacanthocytosis. , 2008, , 249-253.		0

#	Article	IF	Citations
289	Neuropsychologische Befunde bei McLeod-Neuro-akanthozytose. Verhandlungen Der Deutschen Gesellschaft FÃ-¼r Neurologie, 1995, , 426-429.	0.0	0
290	Bilaterale bewegungs-korrelierte kortikale Potentiale bei erblichen spiegelbildlichen Mitbewegungen (Mirror-Movements). Verhandlungen Der Deutschen Gesellschaft FA½r Neurologie, 1995, , 146-149.	0.0	0
291	A longitudinal biomarker study of Patients with Corticobasal Syndrom: Activity of Cerebral Networks, Amyloid and Microglia in Aging and Alzheimer's disease (ActiGliA) – In vivo Imaging of Microglial Activation by TSPO PET. , 2019, 58, .		0
292	Combined in vivo PET imaging of astrogliosis and tau facilitates differential diagnosis of parkinsonian syndromes in correlation with the phenotype. , 2019, 58, .		0
293	Recurring Facial Nerve Palsy. Deutsches Ärzteblatt International, 2019, 116, 701.	0.9	0
294	$18\ \text{F-Pl}2620\ \text{Tau-PET}$ for Assessment of Heterogeneous Neuropathology in Corticobasal Syndrome. , 2020, $59,$ .		0
295	Value of FDG-PET as a Supporting Biomarker for Dementia with Lewy Bodies at Stages of Early Dopaminergic Loss., 2020, 59, .		0
296	Dual-phase $\hat{l}^2$ -Amyloid PET for Assessment of Neuronal Injury and Amyloidosis in Corticobasal Syndrome. , 2020, 59, .		0
297	Alzheimer: Wie man nach einer kausalen Therapie sucht. , 0, , .		0
298	Factors influencing atrophy progression in primary progressive aphasia. Alzheimer's and Dementia, 2021, 17, .	0.8	0
299	Detecting clinical progression from abnormal regional brain volumes at baseline in genetic frontotemporal dementia: A GENFI study. Alzheimer's and Dementia, 2021, 17, .	0.8	0
300	From brain volumes to subgroup classification in genetic mutation carriers for frontotemporal dementia: A cluster analysis in the GENFI study. Alzheimer's and Dementia, 2021, 17, .	0.8	0