

Martina Minnerop

List of Publications by Year in descending order

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47
papers

1,563
citations

331670

21
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315739

38
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all docs

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docs citations

49
times ranked

2563
citing authors

#	ARTICLE	IF	CITATIONS
1	Quantitative susceptibility mapping reveals alterations of dentate nuclei in common types of degenerative cerebellar ataxias. <i>Brain Communications</i> , 2022, 4, fcab306.	3.3	15
2	Regional changes of brain structure during progression of idiopathic Parkinson's disease – A longitudinal study using deformation based morphometry. <i>Cortex</i> , 2022, 151, 188-210.	2.4	11
3	The CCAS-scale in hereditary ataxias: helpful on the group level, particularly in SCA3, but limited in individual patients. <i>Journal of Neurology</i> , 2022, 269, 4363-4374.	3.6	13
4	System Comparison for Gait and Balance Monitoring Used for the Evaluation of a Home-Based Training. <i>Sensors</i> , 2022, 22, 4975.	3.8	4
5	Neurochemical Differences in Spinocerebellar Ataxia Type 14 and 1. <i>Cerebellum</i> , 2021, 20, 169-178.	2.5	0
6	Spinocerebellar ataxia type 14: refining clinicogenetic diagnosis in a rare adult-onset disorder. <i>Annals of Clinical and Translational Neurology</i> , 2021, 8, 774-789.	3.7	13
7	Reference values for the Cerebellar Cognitive Affective Syndrome Scale: age and education matter. <i>Brain</i> , 2021, 144, e20-e20.	7.6	14
8	Investigation of Visual System Involvement in Spinocerebellar Ataxia Type 14. <i>Cerebellum</i> , 2020, 19, 469-482.	2.5	3
9	Cerebellar Involvement in DYT-THAP1 Dystonia. <i>Cerebellum</i> , 2019, 18, 969-971.	2.5	2
10	Reply: Biallelic <i>POLR3A</i> variants confirmed as a frequent cause of hereditary ataxia and spastic paraparesis. <i>Brain</i> , 2019, 142, e13-e13.	7.6	4
11	Tracking the brain in myotonic dystrophies: A 5-year longitudinal follow-up study. <i>PLoS ONE</i> , 2019, 14, e0213381.	2.5	31
12	Reply: <i>POLR3A</i> variants in hereditary spastic paraplegia and ataxia. <i>Brain</i> , 2018, 141, e2-e2.	7.6	10
13	Derivation of Fiber Orientations From Oblique Views Through Human Brain Sections in 3D-Polarized Light Imaging. <i>Frontiers in Neuroanatomy</i> , 2018, 12, 75.	1.7	21
14	Current Progress in CNS Imaging of Myotonic Dystrophy. <i>Frontiers in Neurology</i> , 2018, 9, 646.	2.4	50
15	Expanded phenotype and hippocampal involvement in a novel compound heterozygosity of adult <i>PLA2G6</i> associated neurodegeneration (<i>PARK14</i>). <i>Parkinsonism and Related Disorders</i> , 2017, 37, 111-113.	2.2	7
16	Hypomorphic mutations in <i>POLR3A</i> are a frequent cause of sporadic and recessive spastic ataxia. <i>Brain</i> , 2017, 140, 1561-1578.	7.6	85
17	Accuracy and repeatability of two methods of gait analysis – GaitRite™ and Mobility Lab™ – in subjects with cerebellar ataxia. <i>Gait and Posture</i> , 2016, 48, 194-201.	1.4	59
18	SPECT and PET. , 2016, , 359-365.		0

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19	Autosomal Recessive Cerebellar Ataxia 3 Due to Homozygote c.132dupA Mutation Within the <i>ANO10</i> Gene. <i>JAMA Neurology</i> , 2015, 72, 238.	9.0	5
20	Cytoarchitectonic mapping of the human brain cerebellar nuclei in stereotaxic space and delineation of their co-activation patterns. <i>Frontiers in Neuroanatomy</i> , 2015, 09, 54.	1.7	35
21	Structural and functional MRI abnormalities of cerebellar cortex and nuclei in SCA3, SCA6 and Friedreich's ataxia. <i>Brain</i> , 2015, 138, 1182-1197.	7.6	106
22	Cerebellar neurochemical alterations in spinocerebellar ataxia type 14 appear to include glutathione deficiency. <i>Journal of Neurology</i> , 2015, 262, 1927-1935.	3.6	13
23	Studying variability in human brain aging in a population-based German cohort: rationale and design of 1000BRAINS. <i>Frontiers in Aging Neuroscience</i> , 2014, 6, 149.	3.4	97
24	Clinical and Neurophysiological Profile of Four German Families with Spinocerebellar Ataxia Type 14. <i>Cerebellum</i> , 2014, 13, 89-96.	2.5	42
25	Longitudinal changes in brains of patients with fluent primary progressive aphasia. <i>Brain and Language</i> , 2014, 131, 11-19.	1.6	13
26	Reply to: Cognitive dysfunction in spinocerebellar ataxia type 3: Variable topographies and patterns. <i>Movement Disorders</i> , 2014, 29, 157-158.	3.9	3
27	Progressive cognitive dysfunction in spinocerebellar ataxia type 3. <i>Movement Disorders</i> , 2013, 28, 1435-1438.	3.9	36
28	Never Neglect Inspecting the Leg in Movement Disorders. <i>Archives of Neurology</i> , 2012, 69, 782-3.	4.5	0
29	Alexithymia in healthy young men: A voxel-based morphometric study. <i>Journal of Affective Disorders</i> , 2012, 136, 1252-1256.	4.1	24
30	The time course of neurolinguistic and neuropsychological symptoms in three cases of logopenic primary progressive aphasia. <i>Neuropsychologia</i> , 2012, 50, 1708-1718.	1.6	33
31	The brain in myotonic dystrophy 1 and 2: evidence for a predominant white matter disease. <i>Brain</i> , 2011, 134, 3530-3546.	7.6	199
32	Stochastic resonance therapy in Parkinson's disease. <i>NeuroRehabilitation</i> , 2011, 28, 353-358.	1.3	33
33	Early signs of VCP-related frontotemporal dementia: a neuropsychological, FDG-PET and fMRI study. <i>Journal of Neurology</i> , 2011, 258, 515-518.	3.6	3
34	Neuropsychological Features of Patients with Spinocerebellar Ataxia (SCA) Types 1, 2, 3, and 6. <i>Cerebellum</i> , 2010, 9, 433-442.	2.5	125
35	Depression in Patients with Spinocerebellar Ataxia Type 3 (SCA3). <i>Cerebellum</i> , 2010, 9, 606-607.	2.5	1
36	Callosal tissue loss in multiple system atrophy: A one-year follow-up study. <i>Movement Disorders</i> , 2010, 25, 2613-2620.	3.9	24

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37	In vivo voxel-based relaxometry in amyotrophic lateral sclerosis. <i>Journal of Neurology</i> , 2009, 256, 28-34.	3.6	18
38	Exon deletions and intragenic insertions are not rare in ataxia with oculomotor apraxia 2. <i>BMC Medical Genetics</i> , 2009, 10, 87.	2.1	18
39	Bell's palsy. <i>Journal of Neurology</i> , 2008, 255, 1726-1730.	3.6	49
40	Grey and white matter loss along cerebral midline structures in myotonic dystrophy type 2. <i>Journal of Neurology</i> , 2008, 255, 1904-1909.	3.6	27
41	Smoking upregulates $\alpha 4 \beta 2$ nicotinic acetylcholine receptors in the human brain. <i>Neuroscience Letters</i> , 2008, 430, 34-37.	2.1	64
42	REM sleep behavioral disorder in pure autonomic failure (PAF). <i>Neurology</i> , 2006, 66, 608-609.	1.1	34
43	Dopamine Transporter Positron Emission Tomography in Spinocerebellar Ataxias Type 1, 2, 3, and 6. <i>Archives of Neurology</i> , 2005, 62, 1280.	4.5	89
44	Voxel-based analysis of multiple-system atrophy of cerebellar type: complementary results by combining voxel-based morphometry and voxel-based relaxometry. <i>NeuroImage</i> , 2005, 25, 287-293.	4.2	58
45	CNS infection with <i>C. pneumoniae</i> complicated by multiple strokes. <i>Journal of Neurology</i> , 2003, 250, 1128-1128.	3.6	0
46	In Vivo Voxel-Based Morphometry in Multiple System Atrophy of the Cerebellar Type. <i>Archives of Neurology</i> , 2003, 60, 1431.	4.5	66
47	CNS infection with <i>Chlamydia pneumoniae</i> complicated by multiple strokes. <i>Journal of Neurology</i> , 2002, 249, 1329-1331.	3.6	5