Jörg B. Schulz

List of Publications by Year in descending order

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391 papers

34,266 citations

90 h-index 4774 169 g-index

417 all docs

417 docs citations

417 times ranked

34627 citing authors

#	Article	IF	CITATIONS
1	Diagnosis and management of dementia with Lewy bodies. Neurology, 2005, 65, 1863-1872.	1.1	4,604
2	Glutathione, oxidative stress and neurodegeneration. FEBS Journal, 2000, 267, 4904-4911.	0.2	1,017
3	Modelling neural correlates of working memory: A coordinate-based meta-analysis. NeuroImage, 2012, 60, 830-846.	4.2	777
4	Superoxide Dismutase Activity, Oxidative Damage, and Mitochondrial Energy Metabolism in Familial and Sporadic Amyotrophic Lateral Sclerosis. Journal of Neurochemistry, 1993, 61, 2322-2325.	3.9	555
5	Early Outcome of Carotid Angioplasty and Stenting With and Without Cerebral Protection Devices. Stroke, 2003, 34, 813-819.	2.0	551
6	Loss of function mutations in the gene encoding Omi/HtrA2 in Parkinson's disease. Human Molecular Genetics, 2005, 14, 2099-2111.	2.9	514
7	Repetitive Bilateral Arm Training and Motor Cortex Activation in Chronic Stroke. JAMA - Journal of the American Medical Association, 2004, 292, 1853.	7.4	487
8	Senataxin, the ortholog of a yeast RNA helicase, is mutant in ataxia-ocular apraxia 2. Nature Genetics, 2004, 36, 225-227.	21.4	454
9	Pre-fibrillar \hat{l} ±-synuclein variants with impaired \hat{l}^2 -structure increase neurotoxicity in Parkinson's disease models. EMBO Journal, 2009, 28, 3256-3268.	7.8	411
10	Inhibition of Neuronal Nitric Oxide Synthase by 7â€Nitroindazole Protects Against MPTPâ€Induced Neurotoxicity in Mice. Journal of Neurochemistry, 1995, 64, 936-939.	3.9	377
11	Treatment with simvastatin in normocholesterolemic patients with Alzheimer's disease: A 26â€week randomized, placeboâ€controlled, doubleâ€blind trial. Annals of Neurology, 2002, 52, 346-350.	5. 3	372
12	Neuroprotective Role of the Reaper-Related Serine Protease HtrA2/Omi Revealed by Targeted Deletion in Mice. Molecular and Cellular Biology, 2004, 24, 9848-9862.	2.3	367
13	Protection by pioglitazone in the MPTP model of Parkinson's disease correlates with ll°Bl± induction and block of NFl°B and iNOS activation. Journal of Neurochemistry, 2004, 88, 494-501.	3.9	347
14	Potassium Deprivation-Induced Apoptosis of Cerebellar Granule Neurons: A Sequential Requirement for New mRNA and Protein Synthesis, ICE-Like Protease Activity, and Reactive Oxygen Species. Journal of Neuroscience, 1996, 16, 4696-4706.	3.6	330
15	Transgenic rat model of Huntington's disease. Human Molecular Genetics, 2003, 12, 617-624.	2.9	329
16	Cellular pathology of Parkinson?s disease: astrocytes, microglia and inflammation. Cell and Tissue Research, 2004, 318, 149-161.	2.9	327
17	Caspases as treatment targets in stroke and neurodegenerative diseases. Annals of Neurology, 1999, 45, 421-429.	5.3	315
18	Deficiency of Inducible Nitric Oxide Synthase Protects Against MPTP Toxicity In Vivo. Journal of Neurochemistry, 2008, 74, 2213-2216.	3.9	299

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19	Involvement of Free Radicals in Excitotoxicity In Vivo. Journal of Neurochemistry, 1995, 64, 2239-2247.	3.9	290
20	Elevated free nitrotyrosine levels, but not protein-bound nitrotyrosine or hydroxyl radicals, throughout amyotrophic lateral sclerosis (ALS)-like disease implicate tyrosine nitration as an aberrant in vivo property of one familial ALS-linked superoxide dismutase 1 mutant. Proceedings of the National Academy of Sciences of the United States of America, 1997, 94, 7606-7611.	7.1	279
21	Neuroprotection by Hypoxic Preconditioning Requires Sequential Activation of Vascular Endothelial Growth Factor Receptor and Akt. Journal of Neuroscience, 2002, 22, 6401-6407.	3.6	279
22	Current and experimental treatments of Parkinson disease: A guide for neuroscientists. Journal of Neurochemistry, 2016, 139, 325-337.	3.9	268
23	Chemoresistance of glioblastoma cancer stem cells - much more complex than expected. Molecular Cancer, 2011, 10, 128.	19.2	265
24	The natural history of spinocerebellar ataxia type 1, 2, 3, and 6. Neurology, 2011, 77, 1035-1041.	1.1	259
25	Update on the pathogenesis of Parkinson's disease. Journal of Neurology, 2008, 255, 3-7.	3.6	258
26	Magnetic resonance imaging-based volumetry differentiates idiopathic Parkinson's syndrome from multiple system atrophy and progressive supranuclear palsy. Annals of Neurology, 1999, 45, 65-74.	5.3	255
27	Efficient Inhibition of the Alzheimer's Disease Î ² -Secretase by Membrane Targeting. Science, 2008, 320, 520-523.	12.6	254
28	Loss of pain perception in diabetes is dependent on a receptor of the immunoglobulin superfamily. Journal of Clinical Investigation, 2004, 114, 1741-1751.	8.2	247
29	Increased 3â€nitrotyrosine and oxidative damage in mice with a human copper/zinc superoxide dismutase mutation. Annals of Neurology, 1997, 42, 326-334.	5.3	244
30	Deep brain stimulation. Cell and Tissue Research, 2004, 318, 275-288.	2.9	231
31	Diagnosis and treatment of Friedreich ataxia: a European perspective. Nature Reviews Neurology, 2009, 5, 222-234.	10.1	231
32	Long-term disease progression in spinocerebellar ataxia types 1, 2, 3, and 6: a longitudinal cohort study. Lancet Neurology, The, 2015, 14, 1101-1108.	10.2	213
33	Gene transfer of the JNK interacting protein-1 protects dopaminergic neurons in the MPTP model of Parkinson's disease. Proceedings of the National Academy of Sciences of the United States of America, 2001, 98, 10433-10438.	7.1	208
34	Neuroprotection by the Inhibition of Apoptosis. Brain Pathology, 2000, 10, 283-292.	4.1	203
35	Knockdown of transactive response DNA-binding protein (TDP-43) downregulates histone deacetylase 6. EMBO Journal, 2010, 29, 209-221.	7.8	200
36	Multiple system atrophy: natural history, MRI morphology, and dopamine receptor imaging with 123IBZM-SPECT Journal of Neurology, Neurosurgery and Psychiatry, 1994, 57, 1047-1056.	1.9	198

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37	Protection by Synergistic Effects of Adenovirus-Mediated X-Chromosome-Linked Inhibitor of Apoptosis and Glial Cell Line-Derived Neurotrophic Factor Gene Transfer in the 1-Methyl-4-Phenyl-1,2,3,6-Tetrahydropyridine Model of Parkinson's Disease. Journal of Neuroscience, 2000, 20, 9126-9134.	3.6	194
38	1-Methyl-4-phenyl-1,2,3,6-tetrahydropyride Neurotoxicity Is Attenuated in Mice Overexpressing Bcl-2. Journal of Neuroscience, 1998, 18, 8145-8152.	3.6	193
39	Patterns of Age-related Shrinkage in Cerebellum and Brainstem Observed In Vivo Using Three-dimensional MRI Volumetry. Cerebral Cortex, 1999, 9, 712-721.	2.9	192
40	Two molecular pathways initiate mitochondria-dependent dopaminergic neurodegeneration in experimental Parkinson's disease. Proceedings of the National Academy of Sciences of the United States of America, 2007, 104, 8161-8166.	7.1	190
41	Induction of Nitric Oxide Synthase and Nitric Oxideâ€Mediated Apoptosis in Neuronal PC12 Cells After Stimulation with Tumor Necrosis FActorâ€Î±/Lipopolysaccharide. Journal of Neurochemistry, 1998, 71, 88-94.	3.9	186
42	Systemic administration of rotenone produces selective damage in the striatum and globus pallidus, but not in the substantia nigra. Brain Research, 1997, 753, 157-162.	2.2	184
43	Coenzyme Q ₁₀ and nicotinamide block striatal lesions produced by the mitochondrial toxin malonate. Annals of Neurology, 1994, 36, 882-888.	5.3	183
44	PML in a Patient Treated with Fumaric Acid. New England Journal of Medicine, 2013, 368, 1657-1658.	27.0	176
45	Short and long-term motor skill learning in an accelerated rotarod training paradigm. Neurobiology of Learning and Memory, 2004, 81, 211-216.	1.9	172
46	Drosophila melanogaster as a model organism for Alzheimer's disease. Molecular Neurodegeneration, 2013, 8, 35.	10.8	171
47	Biological and clinical characteristics of individuals at risk for spinocerebellar ataxia types 1, 2, 3, and 6 in the longitudinal RISCA study: analysis of baseline data. Lancet Neurology, The, 2013, 12, 650-658.	10.2	167
48	Glutathione depletion and neuronal cell death: the role of reactive oxygen intermediates and mitochondrial function. Brain Research, 1999, 826, 53-62.	2.2	166
49	TDP-43-Mediated Neuron Loss In Vivo Requires RNA-Binding Activity. PLoS ONE, 2010, 5, e12247.	2.5	166
50	The multidrug resistance protein 1 (Mrp1), but not Mrp5, mediates export of glutathione and glutathione disulfide from brain astrocytes. Journal of Neurochemistry, 2006, 97, 373-384.	3.9	165
51	Neurodegeneration and Motor Dysfunction in a Conditional Model of Parkinson's Disease. Journal of Neuroscience, 2008, 28, 2471-2484.	3.6	164
52	Visualization, quantification and correlation of brain atrophy with clinical symptoms in spinocerebellar ataxia types 1, 3 and 6. NeuroImage, 2010, 49, 158-168.	4.2	162
53	Biological and clinical characteristics of the European Friedreich's Ataxia Consortium for Translational Studies (EFACTS) cohort: a cross-sectional analysis of baseline data. Lancet Neurology, The, 2015, 14, 174-182.	10.2	159
54	Autosomal dominant cerebellar ataxia type I. MRI-based volumetry of posterior fossa structures and basal ganglia in spinocerebellar ataxia types 1, 2 and 3. Brain, 1998, 121, 1687-1693.	7. 6	157

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55	Overexpression of the myelin proteolipid protein leads to accumulation of cholesterol and proteolipid protein in endosomes/lysosomes. Journal of Cell Biology, 2002, 157, 327-336.	5.2	154
56	The multidrug resistance protein MRP1 mediates the release of glutathione disulfide from rat astrocytes during oxidative stress. Journal of Neurochemistry, 2001, 76, 627-636.	3.9	153
57	Responsiveness of different rating instruments in spinocerebellar ataxia patients. Neurology, 2010, 74, 678-684.	1.1	153
58	Neuron-Specific Expression of Therapeutic Proteins: Evaluation of Different Cellular Promoters in Recombinant Adenoviral Vectors. Molecular and Cellular Neurosciences, 2001, 17, 78-96.	2.2	152
59	Magnetic resonance imaging–based volumetry differentiates idiopathic Parkinson's syndrome from multiple system atrophy and progressive supranuclear palsy. Annals of Neurology, 1999, 45, 65-74.	5. 3	152
60	Neuroprotective strategies for treatment of lesions produced by mitochondrial toxins: Implications for neurodegenerative diseases. Neuroscience, 1996, 71, 1043-1048.	2.3	150
61	Glutathione depletion potentiates MPTP and MPP+ toxicity in nigral dopaminergic neurones. NeuroReport, 1996, 7, 921-923.	1.2	149
62	Title is missing!. Molecular and Cellular Biochemistry, 1997, 174, 193-197.	3.1	145
63	Magnetic resonance imaging-based volumetry differentiates progressive supranuclear palsy from corticobasal degeneration. Neurolmage, 2004, 21, 714-724.	4.2	145
64	Mitochondrial Protein Quality Control by the Proteasome Involves Ubiquitination and the Protease Omi. Journal of Biological Chemistry, 2008, 283, 12681-12685.	3.4	145
65	Motor Skill Learning Depends on Protein Synthesis in Motor Cortex after Training. Journal of Neuroscience, 2004, 24, 6515-6520.	3.6	140
66	Systematic Review of Early Recurrent Stenosis After Carotid Angioplasty and Stenting. Stroke, 2005, 36, 367-373.	2.0	139
67	INVOLVEMENT OF OXIDATIVE STRESS IN 3-NITROPROPIONIC ACID NEUROTOXICITY. Neurochemistry International, 1996, 29, 167-171.	3.8	131
68	Comparison of three clinical rating scales in Friedreich ataxia (FRDA). Movement Disorders, 2009, 24, 1779-1784.	3.9	131
69	Lesion location alters brain activation in chronically impaired stroke survivors. NeuroImage, 2004, 21, 924-935.	4.2	130
70	The Montreal Cognitive Assessment (MoCA) - A Sensitive Screening Instrument for Detecting Cognitive Impairment in Chronic Hemodialysis Patients. PLoS ONE, 2014, 9, e106700.	2.5	130
71	Glutathione release from cultured brain cells: Multidrug resistance protein 1 mediates the release of GSH from rat astroglial cells. Journal of Neuroscience Research, 2002, 69, 318-326.	2.9	128
72	Genotype-specific patterns of atrophy progression are more sensitive than clinical decline in SCA1, SCA3 and SCA6. Brain, 2013, 136, 905-917.	7.6	128

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73	Sensitivity to MPTP is not increased in Parkinson's diseaseâ€associated mutant αâ€synuclein transgenic mice. Journal of Neurochemistry, 2001, 77, 1181-1184.	3.9	125
74	Identification and functional characterization of a novel R621C mutation in the synphilin-1 gene in Parkinson's disease. Human Molecular Genetics, 2003, 12, 1223-1231.	2.9	124
75	<scp>COVID</scp> â€19 Vaccineâ€Associated Cerebral Venous Thrombosis in Germany. Annals of Neurology, 2021, 90, 627-639.	5.3	122
76	The Mitochondrial Chaperone Protein TRAP1 Mitigates \hat{l}_{\pm} -Synuclein Toxicity. PLoS Genetics, 2012, 8, e1002488.	3.5	120
77	The Heart in Friedreich Ataxia. Circulation, 2012, 125, 1626-1634.	1.6	119
78	Novel homozygous p.E64D mutation in DJ1 in early onset Parkinson disease (PARK7). Human Mutation, 2004, 24, 321-329.	2.5	117
79	Progression characteristics of the European Friedreich's Ataxia Consortium for Translational Studies (EFACTS): a 2 year cohort study. Lancet Neurology, The, 2016, 15, 1346-1354.	10.2	117
80	Adenovirus-Mediated Gene Transfer of Inhibitors of Apoptosis Proteins Delays Apoptosis in Cerebellar Granule Neurons. Journal of Neurochemistry, 1999, 72, 292-301.	3.9	116
81	Granulocyteâ€colony stimulating factor is neuroprotective in a model of Parkinson's disease. Journal of Neurochemistry, 2006, 97, 675-686.	3.9	109
82	Apoptotic mechanisms and antiapoptotic therapy in the MPTP model of Parkinson's disease. Toxicology Letters, 2003, 139, 135-151.	0.8	102
83	Coenzyme Q10 and nicotinamide and a free radical spin trap protect against MPTP neurotoxicity. Experimental Neurology, 1995, 132, 279-283.	4.1	101
84	Relation between Regional Functional MRI Activation and Vascular Reactivity to Carbon Dioxide during Normal Aging. Journal of Cerebral Blood Flow and Metabolism, 2003, 23, 565-573.	4.3	100
85	NGF, BDNF and NT-5, but not NT-3 protect against MPP+ toxicity and oxidative stress in neonatal animals. Brain Research, 1996, 713, 178-185.	2.2	97
86	Coâ€enzyme Q ₁₀ and idebenone use in Friedreich's ataxia. Journal of Neurochemistry, 2013, 126, 125-141.	3.9	97
87	Role of nitric oxide in neurodegenerative diseases. Current Opinion in Neurology, 1995, 8, 480-486.	3.6	95
88	Exogenous Administration of Gangliosides Displaces GPI-anchored Proteins from Lipid Microdomains in Living Cells. Molecular Biology of the Cell, 1999, 10, 3187-3196.	2.1	95
89	Cooperative Interception of Neuronal Apoptosis by BCLâ€2 and BAGâ€1 Expression: Prevention of Caspase Activation and Reduced Production of Reactive Oxygen Species. Journal of Neurochemistry, 1997, 69, 2075-2086.	3.9	94
90	Extended therapeutic window for caspase inhibition and synergy with MK-801 in the treatment of cerebral histotoxic hypoxia. Cell Death and Differentiation, 1998, 5, 847-857.	11,2	93

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91	Alternate-Form Reliability of the Montreal Cognitive Assessment Screening Test in a Clinical Setting. Dementia and Geriatric Cognitive Disorders, 2012, 33, 379-384.	1.5	93
92	Feasibility of Prehospital Teleconsultation in Acute Stroke – A Pilot Study in Clinical Routine. PLoS ONE, 2012, 7, e36796.	2.5	91
93	Malonate produces striatal lesions by indirect NMDA receptor activation. Brain Research, 1994, 647, 161-166.	2.2	90
94	Rescue from death but not from functional impairment: caspase inhibition protects dopaminergic cells against 6-hydroxydopamine-induced apoptosis but not against the loss of their terminals. Journal of Neurochemistry, 2001, 77, 263-273.	3.9	89
95	Differentiated parietal connectivity of frontal regions for "what―and "where―memory. Brain Structure and Function, 2013, 218, 1551-1567.	2.3	86
96	<i>Drosophila</i> as a screening tool to study human neurodegenerative diseases. Journal of Neurochemistry, 2013, 127, 453-460.	3.9	86
97	Tatâ€Hsp70 protects dopaminergic neurons in midbrain cultures and in the substantia nigra in models of Parkinson's disease. Journal of Neurochemistry, 2008, 105, 853-864.	3.9	85
98	Basic Fibroblast Growth Factor Protects against Excitotoxicity and Chemical Hypoxia in Both Neonatal and Adult Rats. Journal of Cerebral Blood Flow and Metabolism, 1995, 15, 619-623.	4.3	83
99	Efficient Gene Therapy for Parkinson's Disease Using Astrocytes as Hosts for Localized Neurotrophic Factor Delivery. Molecular Therapy, 2012, 20, 534-543.	8.2	82
100	Altered restingâ€state connectivity in Huntington's Disease. Human Brain Mapping, 2014, 35, 2582-2593.	3.6	82
101	Statin Therapy at Carotid Angioplasty and Stent Placement: Effect on Procedure-related Stroke, Myocardial Infarction, and Death. Radiology, 2006, 240, 145-151.	7.3	81
102	TRAP1 rescues PINK1 loss-of-function phenotypes. Human Molecular Genetics, 2013, 22, 2829-2841.	2.9	81
103	Non-Invasive Neurochemical Analysis of Focal Excitotoxic Lesions in Models of Neurodegenerative Illness Using Spectroscopic Imaging. Journal of Cerebral Blood Flow and Metabolism, 1996, 16, 450-461.	4.3	80
104	Cholesterol depletion reduces aggregation of amyloid-beta peptide in hippocampal neurons. Neurobiology of Disease, 2006, 23, 573-577.	4.4	80
105	Silencing of the <i>Pink1</i> Gene Expression by Conditional RNAi Does Not Induce Dopaminergic Neuron Death in Mice. International Journal of Biological Sciences, 2007, 3, 242-250.	6.4	80
106	Neuronal pathology in Parkinson?s disease. Cell and Tissue Research, 2004, 318, 135-147.	2.9	79
107	The Cancer Stem Cell Subtype Determines Immune Infiltration of Glioblastoma. Stem Cells and Development, 2012, 21, 2753-2761.	2.1	79
108	A new semiautomated, three-dimensional technique allowing precise quantification of total and regional cerebellar volume using MRI. Magnetic Resonance in Medicine, 1998, 40, 143-151.	3.0	77

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109	Cascade of Caspase Activation in Potassium-Deprived Cerebellar Granule Neurons: Targets for Treatment with Peptide and Protein Inhibitors of Apoptosis. Molecular and Cellular Neurosciences, 2001, 17, 717-731.	2.2	77
110	Expanded phenotypic spectrum of the m.8344A>G "MERRF―mutation: data from the German mitoNET registry. Journal of Neurology, 2016, 263, 961-972.	3.6	77
111	Sporadic late-onset nemaline myopathy: clinico-pathological characteristics and review of 76 cases. Orphanet Journal of Rare Diseases, 2017, 12, 86.	2.7	77
112	Consensus clinical management guidelines for Friedreich ataxia. Orphanet Journal of Rare Diseases, 2014, 9, 184.	2.7	76
113	Subtypes of mild cognitive impairment in patients with Parkinson's disease: evidence from the LANDSCAPE study. Journal of Neurology, Neurosurgery and Psychiatry, 2016, 87, 1099-1105.	1.9	76
114	Brain imaging findings in idiopathic REM sleep behavior disorder (RBD) – A systematic review on potential biomarkers for neurodegeneration. Sleep Medicine Reviews, 2017, 34, 23-33.	8.5	76
115	Mitochondrial dysfunction in movement disorders. Current Opinion in Neurology, 1994, 7, 333-339.	3.6	7 5
116	Effectiveness of intravenous immunoglobin therapy in cerebellar ataxia associated with gluten sensitivity. Annals of Neurology, 2001, 50, 827-828.	5.3	74
117	Comparison of angioplasty and stenting with cerebral protection versus endarterectomy for treatment of internal carotid artery stenosis in elderly patients. Journal of Vascular Surgery, 2004, 40, 945-951.	1.1	74
118	Penguins and hummingbirds: Midbrain atrophy in progressive supranuclear palsy. Neurology, 2006, 66, 949-950.	1.1	74
119	Selfâ€rated health status in spinocerebellar ataxiaâ€"Results from a European multicenter study. Movement Disorders, 2010, 25, 587-595.	3.9	74
120	Spinocerebellar Ataxia Types 1, 2, 3 and 6: the Clinical Spectrum of Ataxia and Morphometric Brainstem and Cerebellar Findings. Cerebellum, 2012, 11, 155-166.	2.5	74
121	Clinical Predictors of Transient Ischemic Attack, Stroke, or Death Within 30 Days of Carotid Angioplasty and Stenting. Stroke, 2005, 36, 787-791.	2.0	73
122	Neurofilaments in spinocerebellar ataxia type 3: blood biomarkers at the preataxic and ataxic stage in humans and mice. EMBO Molecular Medicine, 2020, 12, e11803.	6.9	73
123	Striatal Malonate Lesions Are Attenuated in Neuronal Nitric Oxide Synthase Knockout Mice. Journal of Neurochemistry, 1996, 67, 430-433.	3.9	72
124	Investigating function and connectivity of morphometric findings $\hat{a} \in$ Exemplified on cerebellar atrophy in spinocerebellar ataxia 17 (SCA17). NeuroImage, 2012, 62, 1354-1366.	4.2	72
125	Characterization of motor skill and instrumental learning time scales in a skilled reaching task in rat. Behavioural Brain Research, 2004, 155, 249-256.	2.2	71
126	Diagnostic hallmarks and pitfalls in late-onset progressive transthyretin-related amyloid-neuropathy. Journal of Neurology, 2013, 260, 3093-3108.	3.6	71

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127	High level expression of expanded full-length ataxin-3 in vitro causes cell death and formation of intranuclear inclusions in neuronal cells. Human Molecular Genetics, 1999, 8, 1169-1176.	2.9	69
128	Differential effects of l-buthionine sulfoximine and ethacrynic acid on glutathione levels and mitochondrial function in PC12 cells. Neuroscience Letters, 1999, 264, 1-4.	2.1	69
129	Depression comorbidity in spinocerebellar ataxia. Movement Disorders, 2011, 26, 870-876.	3.9	69
130	Survival in patients with spinocerebellar ataxia types 1, 2, 3, and 6 (EUROSCA): a longitudinal cohort study. Lancet Neurology, The, 2018, 17, 327-334.	10.2	69
131	Parkinson's disease: one biochemical pathway to fit all genes?. Trends in Molecular Medicine, 2002, 8, 236-240.	6.7	68
132	Long-term EMG recordings differentiate between parkinsonian and essential tremor. Journal of Neurology, 2008, 255, 103-111.	3.6	68
133	Frequent genes in rare diseases: panelâ€based next generation sequencing to disclose causal mutations in hereditary neuropathies. Journal of Neurochemistry, 2017, 143, 507-522.	3.9	68
134	Cerebral changes improved by physical activity during cognitive decline: A systematic review on MRI studies. NeuroImage: Clinical, 2019, 23, 101933.	2.7	68
135	Effects of dopamine on the glutathione metabolism of cultured astroglial cells: implications for Parkinson's disease. Journal of Neurochemistry, 2002, 82, 458-467.	3.9	67
136	Magnetic resonance imaging in spinocerebellar ataxias. Cerebellum, 2008, 7, 204-214.	2.5	67
137	Increased brain tissue sodium concentration in Huntington's Disease â€" A sodium imaging study at 4T. Neurolmage, 2012, 63, 517-524.	4.2	67
138	Neuroanatomic changes and their association with cognitive decline in mild cognitive impairment: a meta-analysis. Brain Structure and Function, 2012, 217, 115-125.	2.3	67
139	Ret is essential to mediate GDNF's neuroprotective and neuroregenerative effect in a Parkinson disease mouse model. Cell Death and Disease, 2016, 7, e2359-e2359.	6.3	67
140	Long COVIDâ€19: Objectifying most selfâ€reported neurological symptoms. Annals of Clinical and Translational Neurology, 2022, 9, 141-154.	3.7	67
141	Accumulation and clearance of αâ€synuclein aggregates demonstrated by timeâ€lapse imaging. Journal of Neurochemistry, 2008, 106, 529-540.	3.9	66
142	Receptor for advanced glycation endproducts (RAGE) deficiency protects against MPTP toxicity. Neurobiology of Aging, 2012, 33, 2478-2490.	3.1	66
143	Loss of <scp>FBXO</scp> 7 (<scp>PARK</scp> 15) results in reduced proteasome activity and models a parkinsonismâ€ike phenotype in mice. EMBO Journal, 2016, 35, 2008-2025.	7.8	65
144	Consistent Neurodegeneration and Its Association with Clinical Progression in Huntington's Disease: A Coordinate-Based Meta-Analysis. Neurodegenerative Diseases, 2013, 12, 23-35.	1.4	64

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145	Rab7 induces clearance of αâ€synuclein aggregates. Journal of Neurochemistry, 2016, 138, 758-774.	3.9	63
146	UBE2E Ubiquitin-conjugating Enzymes and Ubiquitin Isopeptidase Y Regulate TDP-43 Protein Ubiquitination. Journal of Biological Chemistry, 2014, 289, 19164-19179.	3.4	62
147	Potential Synergistic Protection of Retinal Ganglion Cells from Axotomy-Induced Apoptosis by Adenoviral Administration of Clial Cell Line-Derived Neurotrophic Factor and X-Chromosome-Linked Inhibitor of Apoptosis. Neurobiology of Disease, 2002, 11, 123-133.	4.4	61
148	Lesion of the pedunculopontine nucleus reverses hyperactivity of the subthalamic nucleus and substantia nigra pars reticulata in a 6-hydroxydopamine rat model. European Journal of Neuroscience, 2006, 24, 2275-2282.	2.6	60
149	Cortical stimulation mapping using epidurally implanted thin-film microelectrode arrays. Journal of Neuroscience Methods, 2007, 161, 118-125.	2.5	60
150	Malonate-Induced Generation of Reactive Oxygen Species in Rat Strium Depends on Dopamine Release but Not on NMDA Receptor Activation. Journal of Neurochemistry, 2001, 73, 1329-1332.	3.9	58
151	Clinical experience with high-dose idebenone in Friedreich ataxia. Journal of Neurology, 2009, 256, 42-45.	3.6	58
152	Impaired retrograde transport by the Dynein/Dynactin complex contributes to Tau-induced toxicity. Human Molecular Genetics, 2015, 24, 3623-3637.	2.9	58
153	Transgenic rat model of Huntington's disease. Human Molecular Genetics, 2003, 12, 617-624.	2.9	58
154	Improved Therapeutic Window for Treatment of Histotoxic Hypoxia with a Free Radical Spin Trap. Journal of Cerebral Blood Flow and Metabolism, 1995, 15, 948-952.	4.3	57
155	Lactate as a diagnostic marker in transient loss of consciousness. Seizure: the Journal of the British Epilepsy Association, 2016, 40, 71-75.	2.0	56
156	Bloodâ€based neurochemical diagnosis of vascular dementia: a pilot study. Journal of Neurochemistry, 2007, 103, 467-474.	3.9	55
157	Chemotherapyâ€induced cell death in primary cerebellar granule neurons but not in astrocytes: <i>iin vitro</i>) paradigm of differential neurotoxicity. Journal of Neurochemistry, 2004, 91, 1067-1074.	3.9	54
158	Cognitive decline in Parkinson's disease: the impact of the motor phenotype on cognition. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 171-179.	1.9	54
159	Intracellular acidification by inhibition of the Na+/H+-exchanger leads to caspase-independent death of cerebellar granule neurons resembling paraptosis. Cell Death and Differentiation, 2004, 11, 760-770.	11.2	53
160	FasL (CD95L/APO-1L) Resistance of Neurons Mediated by Phosphatidylinositol 3-Kinase-Akt/Protein Kinase B-Dependent Expression of Lifeguard/Neuronal Membrane Protein 35. Journal of Neuroscience, 2005, 25, 6765-6774.	3.6	53
161	Visualization and quantification of disease progression in multiple system atrophy. Movement Disorders, 2006, 21, 1674-1681.	3.9	53
162	RET signaling does not modulate MPTP toxicity but is required for regeneration of dopaminergic axon terminals. Proceedings of the National Academy of Sciences of the United States of America, 2007, 104, 20049-20054.	7.1	53

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