Armin Giese

List of Publications by Year in descending order

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23567 19190 15,204 170 58 118 citations h-index g-index papers 178 178 178 14924 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Seizure prevalence in neurodegenerative diseases—a study of autopsy proven cases. European Journal of Neurology, 2022, 29, 12-18.	3.3	6
2	Blood \hat{l}^2 -Synuclein and Neurofilament Light Chain During the Course of Prion Disease. Neurology, 2022, , 10.1212/WNL.0000000000200002.	1.1	11
3	Tau deposition patterns are associated with functional connectivity in primary tauopathies. Nature Communications, 2022, 13, 1362.	12.8	34
4	Safety, tolerability and pharmacokinetics of the oligomer modulator anle138b with exposure levels sufficient for therapeutic efficacy in a murine Parkinson model: A randomised, double-blind, placebo-controlled phase 1a trial. EBioMedicine, 2022, 80, 104021.	6.1	26
5	Genetic determinants of survival in progressive supranuclear palsy: a genome-wide association study. Lancet Neurology, The, 2021, 20, 107-116.	10.2	62
6	Extended stereotactic brain biopsy in suspected primary central nervous system angiitis: good diagnostic accuracy and high safety. Journal of Neurology, 2021, 268, 367-376.	3.6	9
7	Mutations within FGFR1 are associated with superior outcome in a series of 83 diffuse midline gliomas with H3F3A K27M mutations. Acta Neuropathologica, 2021, 141, 323-325.	7.7	20
8	First symptom guides diagnosis and prognosis in neurodegenerative diseases—a retrospective study of autopsy proven cases. European Journal of Neurology, 2021, 28, 1801-1811.	3.3	11
9	Genotype–Phenotype Relations for the Atypical Parkinsonism Genes: MDSGene Systematic Review. Movement Disorders, 2021, 36, 1499-1510.	3.9	22
10	The number of methylated CpG sites within the MGMT promoter region linearly correlates with outcome in glioblastoma receiving alkylating agents. Acta Neuropathologica Communications, 2021, 9, 35.	5.2	11
11	Phenotypic diversity of genetic Creutzfeldt–Jakob disease: a histo-molecular-based classification. Acta Neuropathologica, 2021, 142, 707-728.	7.7	24
12	A Multi-Scale and Multi-Technique Approach for the Characterization of the Effects of Spatially Fractionated X-ray Radiation Therapies in a Preclinical Model. Cancers, 2021, 13, 4953.	3.7	4
13	[11C]MODAG-001â€"towards a PET tracer targeting α-synuclein aggregates. European Journal of Nuclear Medicine and Molecular Imaging, 2021, 48, 1759-1772.	6.4	50
14	Tau-induced mitochondrial membrane perturbation is dependent upon cardiolipin. Biochimica Et Biophysica Acta - Biomembranes, 2020, 1862, 183064.	2.6	38
15	Validation of the Movement Disorder Society Criteria for the Diagnosis of 4â€Repeat Tauopathies. Movement Disorders, 2020, 35, 171-176.	3.9	37
16	¹¹ C Radiolabeling of anle253b: a Putative PET Tracer for Parkinson's Disease That Binds to αâ€Synuclein Fibrils inâ€vitro and Crosses the Bloodâ€Brain Barrier. ChemMedChem, 2020, 15, 411-415.	3.2	35
17	Targeting α-synuclein by PD03 AFFITOPE® and Anle138b rescues neurodegenerative pathology in a model of multiple system atrophy: clinical relevance. Translational Neurodegeneration, 2020, 9, 38.	8.0	22
18	Clinical Conditions "Suggestive of Progressive Supranuclear Palsyâ€â€"Diagnostic Performance. Movement Disorders, 2020, 35, 2301-2313.	3.9	22

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19	Distribution patterns of tau pathology in progressive supranuclear palsy. Acta Neuropathologica, 2020, 140, 99-119.	7.7	210
20	Potential sources of interference with the highly sensitive detection and quantification of alphaâ€synuclein seeds by qRTâ€QuIC. FEBS Open Bio, 2020, 10, 883-893.	2.3	6
21	Copathology in Progressive Supranuclear Palsy: Does It Matter?. Movement Disorders, 2020, 35, 984-993.	3.9	48
22	Longitudinal correlation between neurofilament light chain and UMSARS in Multiple System Atrophy. Clinical Neurology and Neurosurgery, 2020, 195, 105924.	1.4	3
23	Travel-associated neurological disease terminated in a postmortem diagnosed atypical HSV-1 encephalitis after high-dose steroid therapy – a case report. BMC Infectious Diseases, 2020, 20, 150.	2.9	3
24	Binding of Metal-Ion-Induced Tau Oligomers to Lipid Surfaces Is Enhanced by GSK-3 \hat{l}^2 -Mediated Phosphorylation. ACS Chemical Neuroscience, 2020, 11, 880-887.	3.5	8
25	Investigation of potential adverse central nervous system effects after long term oral administration of gadolinium in mice. PLoS ONE, 2020, 15, e0231495.	2.5	11
26	Late-stage Anle138b treatment ameliorates tau pathology and metabolic decline in a mouse model of human Alzheimer's disease tau. Alzheimer's Research and Therapy, 2019, 11, 67.	6.2	28
27	Cardiolipin Promotes Pore-Forming Activity of Alpha-Synuclein Oligomers in Mitochondrial Membranes. ACS Chemical Neuroscience, 2019, 10, 3815-3829.	3.5	60
28	Safety and efficacy of epigallocatechin gallate in multiple system atrophy (PROMESA): a randomised, double-blind, placebo-controlled trial. Lancet Neurology, The, 2019, 18, 724-735.	10.2	79
29	Iron-mediated aggregation and toxicity in a novel neuronal cell culture model with inducible alpha-synuclein expression. Scientific Reports, 2019, 9, 9100.	3.3	8
30	Depopulation of dense α-synuclein aggregates is associated with rescue of dopamine neuron dysfunction and death in a new Parkinson's disease model. Acta Neuropathologica, 2019, 138, 575-595.	7.7	79
31	How to apply the movement disorder society criteria for diagnosis of progressive supranuclear palsy. Movement Disorders, 2019, 34, 1228-1232.	3.9	93
32	Protein-Drug Interactions in the Membrane: The Small Molecule Anle138b and its Binding to α-Synuclein Oligomers. Biophysical Journal, 2019, 116, 352a.	0.5	2
33	PATH-04. INFLUENCE OF INDIVIDUAL CpG METHYLATION STATUS OF THE MGMT PROMOTOR ON OUTCOME IN ADULT PATIENTS WITH GLIOBLASTOMA MULTIFORME RECEIVING ALKYLATING AGENT TREATMENT. Neuro-Oncology, 2019, 21, vi143-vi143.	1.2	0
34	The small molecule inhibitor anle 145c thermodynamically traps human islet amyloid peptide in the form of non-cytotoxic oligomers. Scientific Reports, 2019, 9, 19023.	3.3	16
35	Contrast enhancement is a prognostic factor in IDH1/2 mutant, but not in wild-type WHO grade II/III glioma as confirmed by machine learning. European Journal of Cancer, 2019, 107, 15-27.	2.8	30
36	Different Effects of α-Synuclein Mutants on Lipid Binding and Aggregation Detected by Single Molecule Fluorescence Spectroscopy and ThT Fluorescence-Based Measurements. ACS Chemical Neuroscience, 2019, 10, 1649-1659.	3.5	44

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37	Anle138b modulates αâ€synuclein oligomerization and prevents motor decline and neurodegeneration in a mouse model of multiple system atrophy. Movement Disorders, 2019, 34, 255-263.	3.9	72
38	Prion-related peripheral neuropathy in sporadic Creutzfeldt-Jakob disease. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 424-427.	1.9	31
39	Epilepsy surgery in the first months of life: a large type IIb focal cortical dysplasia causing neonatal drugâ€resistant epilepsy. Epileptic Disorders, 2019, 21, 122-127.	1.3	5
40	Regional pattern of microgliosis in sporadic Creutzfeldtâ€Jakob disease in relation to phenotypic variants and disease progression. Neuropathology and Applied Neurobiology, 2018, 44, 574-589.	3.2	17
41	Photophysics of diphenyl-pyrazole compounds in solutions and α-synuclein aggregates. Biochimica Et Biophysica Acta - General Subjects, 2018, 1862, 800-807.	2.4	10
42	Identification of time-to-peak on dynamic 18F-FET-PET as a prognostic marker specifically in IDH1/2 mutant diffuse astrocytoma. Neuro-Oncology, 2018, 20, 279-288.	1.2	71
43	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
44	The Small Molecule anle 138b Shows Interaction with $\hat{l}\pm$ -Synuclein Oligomers in Phospholipid Membranes. Biophysical Journal, 2018, 114, 560a.	0.5	2
45	Cell Type and Species-specific Patterns in Neuronal and Non-neuronal Methylomes of Human and Chimpanzee Cortices. Cerebral Cortex, 2018, 28, 3724-3739.	2.9	7
46	TERT promoter mutation is associated with worse prognosis in WHO grade II and III meningiomas. Journal of Neuro-Oncology, 2018, 139, 671-678.	2.9	51
47	DNA methylation analysis on purified neurons and glia dissects age and Alzheimer's disease-specific changes in the human cortex. Epigenetics and Chromatin, 2018, 11, 41.	3.9	173
48	MGMT promoter methylation is not correlated with integrin expression in malignant gliomas: clarifying recent clinical trial results. Medical Oncology, 2018, 35, 103.	2.5	2
49	The diphenylpyrazole compound anle 138b blocks $\hat{Al^2}$ channels and rescues disease phenotypes in a mouse model for amyloid pathology. EMBO Molecular Medicine, 2018, 10, 32-47.	6.9	63
50	Contrast enhancement as a prognostic factor in IDH1/2 mutant glioma Journal of Clinical Oncology, 2018, 36, 2029-2029.	1.6	0
51	Which ante mortem clinical features predict progressive supranuclear palsy pathology?. Movement Disorders, 2017, 32, 995-1005.	3.9	121
52	Clinical diagnosis of progressive supranuclear palsy: The movement disorder society criteria. Movement Disorders, 2017, 32, 853-864.	3.9	1,402
53	Treatment with diphenyl–pyrazole compound anle138b/c reveals that α-synuclein protects melanoma cells from autophagic cell death. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, E4971-E4977.	7.1	25
54	MR imaging differentiation of Fe2+ and Fe3+ based on relaxation and magnetic susceptibility properties. Neuroradiology, 2017, 59, 403-409.	2.2	36

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55	Seeding and transgenic overexpression of alphaâ€synuclein triggers dendritic spine pathology in the neocortex. EMBO Molecular Medicine, 2017, 9, 716-731.	6.9	58
56	K27M midline gliomas display malignant progression by imaging and histology. Neuropathology and Applied Neurobiology, 2017, 43, 458-462.	3.2	9
57	Immunoprofiling of glial tumours of the neurohypophysis suggests a common pituicytic origin of neoplastic cells. Pituitary, 2017, 20, 211-217.	2.9	26
58	Substitutions of PrP N-terminal histidine residues modulate scrapie disease pathogenesis and incubation time in transgenic mice. PLoS ONE, 2017, 12, e0188989.	2.5	11
59	The Differential Diagnosis and Treatment of Atypical Parkinsonism. Deutsches Ärzteblatt International, 2016, 113, 61-9.	0.9	135
60	Anle138b Partly Ameliorates Motor Deficits Despite Failure of Neuroprotection in a Model of Advanced Multiple System Atrophy. Frontiers in Neuroscience, 2016, 10, 99.	2.8	23
61	An autopsy-confirmed case of progressive supranuclear palsy with predominant postural instability. Acta Neuropathologica Communications, 2016, 4, 120.	5. 2	22
62	Neurofilaments in blood and CSF for diagnosis and prediction of onset in Creutzfeldt-Jakob disease. Scientific Reports, 2016, 6, 38737.	3.3	81
63	Effect of the Novel Amyloid Inhibitor "anle145c―on Aggregation of Islet Amyloid Polypeptide and how it is Modulated by Membranes. Biophysical Journal, 2016, 110, 420a.	0.5	0
64	Analysis of Conformational Stability of Abnormal Prion Protein Aggregates across the Spectrum of Creutzfeldt-Jakob Disease Prions. Journal of Virology, 2016, 90, 6244-6254.	3.4	29
65	Re-irradiation strategies in combination with bevacizumab for recurrent malignant glioma. Journal of Neuro-Oncology, 2016, 130, 591-599.	2.9	28
66	Parkinson's disease: SNCA-, PARK2-, and LRRK2- targeting microRNAs elevated in cingulate gyrus. Parkinsonism and Related Disorders, 2016, 33, 115-121.	2.2	72
67	Generation and deposition of Aβ43 by the virtually inactive presenilinâ€1 L435F mutant contradicts the presenilin lossâ€ofâ€function hypothesis of Alzheimer's disease. EMBO Molecular Medicine, 2016, 8, 458-465.	6.9	60
68	Intracellular formation of \hat{l} ±-synuclein oligomers and the effect of heat shock protein 70 characterized by confocal single particle spectroscopy. Biochemical and Biophysical Research Communications, 2016, 477, 76-82.	2.1	4
69	The PROMESA-protocol: progression rate of multiple system atrophy under EGCG supplementation as anti-aggregation-approach. Journal of Neural Transmission, 2016, 123, 439-445.	2.8	32
70	Spinal neuronal cannabinoid receptors mediate urodynamic effects of systemic fatty acid amide hydrolase (FAAH) inhibition in rats. Neurourology and Urodynamics, 2016, 35, 464-470.	1.5	12
71	Quantifying prion disease penetrance using large population control cohorts. Science Translational Medicine, 2016, 8, 322ra9.	12.4	289
72	Interaction of α-synuclein with biomembranes in Parkinson's disease â€"role of cardiolipin. Progress in Lipid Research, 2016, 61, 73-82.	11.6	73

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73	Distinct Histomorphology in Molecular Subgroups of Glioblastomas in Young Patients. Journal of Neuropathology and Experimental Neurology, 2016, 75, 408-414.	1.7	35
74	Ageâ€Dependent Levels of 5â€Methylâ€, 5â€Hydroxymethylâ€, and 5â€Formylcytosine in Human and Mouse Bra Tissues. Angewandte Chemie - International Edition, 2015, 54, 12511-12514.	ain 13.8	116
75	Histopathology and clinical course of MOG-antibody-associated encephalomyelitis. Annals of Clinical and Translational Neurology, 2015, 2, 295-301.	3.7	160
76	Dynamic ¹⁸ <scp>Fâ€FET PET</scp> in suspected <scp>WHO</scp> grade II gliomas defines distinct biological subgroups with different clinical courses. International Journal of Cancer, 2015, 136, 2132-2145.	5.1	68
77	The Novel Inhibitor "Anle145C―Efficiently Inhibits Fibril Formation of Islet Amyloid Polypeptide (IAPP) and uses Distinctly Different Modes of Action in the Absence and Presence of Membranes. Biophysical Journal, 2015, 108, 256a.	0.5	0
78	Anle138b and related compounds are aggregation specific fluorescence markers and reveal high affinity binding to α-synuclein aggregates. Biochimica Et Biophysica Acta - General Subjects, 2015, 1850, 1884-1890.	2.4	52
79	Quantitative Real-Time Quaking-Induced Conversion Allows Monitoring of Disease-Modifying Therapy in the Urine of Prion-Infected Mice. Journal of Neuropathology and Experimental Neurology, 2015, 74, 924-933.	1.7	10
80	Distribution of dipeptide repeat proteins in cellular models and C9orf72 mutation cases suggests link to transcriptional silencing. Acta Neuropathologica, 2015, 130, 537-555.	7.7	157
81	The association of aphasia and right-sided motor impairment in corticobasal syndrome. Journal of Neurology, 2015, 262, 2241-2246.	3.6	9
82	Cysteine-Sparing CADASIL Mutations in <i>NOTCH3</i> Show Proaggregatory Properties In Vitro. Stroke, 2015, 46, 786-792.	2.0	46
83	Extracellular vesicle sorting of \hat{l}_{\pm} -Synuclein is regulated by sumoylation. Acta Neuropathologica, 2015, 129, 695-713.	7.7	136
84	Reducing tau aggregates with anle138b delays disease progression in a mouse model of tauopathies. Acta Neuropathologica, 2015, 130, 619-631.	7.7	58
85	Correlation of dynamic $\langle \sup 18 \rangle 18 \rangle$ sup $\langle \sup FET$ -PET with IDH 1 mutation for prediction of outcome in anaplastic astrocytoma WHO° III independently from tumor vascularisation Journal of Clinical Oncology, 2015, 33, 2037-2037.	1.6	2
86	Modelling Ser129 Phosphorylation Inhibits Membrane Binding of Pore-Forming Alpha-Synuclein Oligomers. PLoS ONE, 2014, 9, e98906.	2.5	24
87	The phenotypic spectrum of progressive supranuclear palsy: A retrospective multicenter study of 100 definite cases. Movement Disorders, 2014, 29, 1758-1766.	3.9	286
88	Identification of tetrahydrocarbazoles as novel multifactorial drug candidates for treatment of Alzheimer's disease. Translational Psychiatry, 2014, 4, e489-e489.	4.8	15
89	Scanning for Intensely Fluorescent Targets (SIFT) in the Study of Protein Aggregation at the Single-Particle Level. , 2014, , 167-176.		0
90	Diffuse leukoencephalopathy with spheroids: Biopsy findings and a novel mutation. Clinical Neurology and Neurosurgery, 2014, 122, 113-115.	1.4	10

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91	Plasminogen activator inhibitor-1 influences cerebrovascular complications and death in pneumococcal meningitis. Acta Neuropathologica, 2014, 127, 553-564.	7.7	17
92	Molecular stereotactic biopsy technique improves diagnostic accuracy and enables personalized treatment strategies in glioma patients. Acta Neurochirurgica, 2014, 156, 1427-1440.	1.7	51
93	Sequestration of latent TGF- \hat{l}^2 binding protein 1 into CADASIL-related Notch3-ECD deposits. Acta Neuropathologica Communications, 2014, 2, 96.	5.2	54
94	The oligomer modulator anle 138b inhibits disease progression in a Parkinson mouse model even with treatment started after disease onset. Acta Neuropathologica, 2014, 127, 779-780.	7.7	103
95	Piperazine derivatives inhibit PrP/PrPres propagation in vitro and in vivo. Biochemical and Biophysical Research Communications, 2014, 445, 23-29.	2.1	9
96	Mitochondrial membrane permeabilisation by amyloid aggregates and protection by polyphenols. Biochimica Et Biophysica Acta - Biomembranes, 2013, 1828, 2532-2543.	2.6	120
97	Establishing quantitative real-time quaking-induced conversion (qRT-QuIC) for highly sensitive detection and quantification of PrPSc in prion-infected tissues. Acta Neuropathologica Communications, $2013, 1, 44$.	5.2	36
98	Presence of Phosphodiesterase Type 5 in the Spinal Cord and its Involvement in Bladder Outflow Obstruction Related Bladder Overactivity. Journal of Urology, 2013, 190, 1430-1435.	0.4	13
99	Anle138b: a novel oligomer modulator for disease-modifying therapy of neurodegenerative diseases such as prion and Parkinson's disease. Acta Neuropathologica, 2013, 125, 795-813.	7.7	327
100	Analyses of Protease Resistance and Aggregation State of Abnormal Prion Protein across the Spectrum of Human Prions. Journal of Biological Chemistry, 2013, 288, 27972-27985.	3.4	34
101	Adjunctive <i>N</i> -Acetyl- <scp>I</scp> -Cysteine in Treatment of Murine Pneumococcal Meningitis. Antimicrobial Agents and Chemotherapy, 2013, 57, 4825-4830.	3.2	9
102	Development and Implementation of a High-Throughput Compound Screening Assay for Targeting Disrupted ER Calcium Homeostasis in Alzheimer's Disease. PLoS ONE, 2013, 8, e80645.	2.5	18
103	An Unusual Case of Ectopic ACTH Syndrome. Experimental and Clinical Endocrinology and Diabetes, 2012, 120, 63-67.	1.2	24
104	Consensus classification of human prion disease histotypes allows reliable identification of molecular subtypes: an inter-rater study among surveillance centres in Europe and USA. Acta Neuropathologica, 2012, 124, 517-529.	7.7	184
105	Limited cleavage of tau with matrix-metalloproteinase MMP-9, but not MMP-3, enhances tau oligomer formation. Experimental Neurology, 2012, 237, 470-476.	4.1	41
106	Polyphenolic compounds are novel protective agents against lipid membrane damage by α-synuclein aggregates in vitro. Biochimica Et Biophysica Acta - Biomembranes, 2012, 1818, 2502-2510.	2.6	50
107	Synthesis of benzamide derivatives and their evaluation as antiprion agents. Bioorganic and Medicinal Chemistry, 2012, 20, 5001-5011.	3.0	10
108	Two Different Binding Modes of $\hat{l}\pm$ -Synuclein to Lipid Vesicles Depending on its Aggregation State. Biophysical Journal, 2012, 102, 1646-1655.	0.5	39

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109	Synergistic influence of phosphorylation and metal ions on tau oligomer formation and coaggregation with α-synuclein at the single molecule level. Molecular Neurodegeneration, 2012, 7, 35.	10.8	60
110	Single-Channel Electrophysiology Reveals a Distinct and Uniform Pore Complex Formed by \hat{l}_{\pm} -Synuclein Oligomers in Lipid Membranes. PLoS ONE, 2012, 7, e42545.	2.5	75
111	Identification of Polyphenolic Compounds and Black Tea Extract as Potent Inhibitors of Lipid Membrane Destabilization by AÎ ² 42 Aggregates. Journal of Alzheimer's Disease, 2011, 27, 767-779.	2.6	46
112	Single particle analysis of tau oligomer formation induced by metal ions and organic solvents. Biochemical and Biophysical Research Communications, 2011, 411, 190-196.	2.1	51
113	Comprehensive Neuropathologic Analysis of Genetic Prion Disease Associated With the E196K Mutation inPRNPReveals Phenotypic Heterogeneity. Journal of Neuropathology and Experimental Neurology, 2011, 70, 192-200.	1.7	8
114	\hat{l}_{\pm} -Internexin in the Diagnosis of Oligodendroglial Tumors and Association With $1p/19q$ Status. Journal of Neuropathology and Experimental Neurology, 2011, 70, 970-978.	1.7	19
115	Generation of Ferric Iron Links Oxidative Stress to $\hat{l}\pm$ -Synuclein Oligomer Formation. Journal of Parkinson's Disease, 2011, 1, 205-216.	2.8	58
116	AMPA-receptor-mediated excitatory synaptic transmission is enhanced by iron-induced \hat{l}_{\pm} -synuclein oligomers. Journal of Neurochemistry, 2011, 117, 868-878.	3.9	60
117	Inhibition and disaggregation of \hat{l}_{\pm} -synuclein oligomers by natural polyphenolic compounds. FEBS Letters, 2011, 585, 1113-1120.	2.8	240
118	Phenotypic variability of sporadic human prion disease and its molecular basis: past, present, and future. Acta Neuropathologica, 2011, 121, 91-112.	7.7	134
119	Next-Generation Sequencing Reveals Regional Differences of the $\hat{l}\pm$ -Synuclein Methylation State Independent of Lewy Body Disease. NeuroMolecular Medicine, 2011, 13, 310-320.	3.4	61
120	Reduced spiral ganglion neuronal loss by adjunctive neurotrophin-3 in experimental pneumococcal meningitis. Journal of Neuroinflammation, 2011, 8, 7.	7.2	26
121	From Highâ€Throughput Cell Culture Screening to Mouse Model: Identification of New Inhibitor Classes against Prion Disease. ChemMedChem, 2011, 6, 1928-1937.	3.2	21
122	Diphenylpyrazole-Derived Compounds Increase Survival Time of Mice after Prion Infection. Antimicrobial Agents and Chemotherapy, 2011, 55, 4774-4781.	3.2	12
123	Co-aggregate formation of CADASIL-mutant NOTCH3: a single-particle analysis. Human Molecular Genetics, 2011, 20, 3256-3265.	2.9	87
124	Divergent Molecular Effects of Desmin Mutations on Protein Assembly in Myofibrillar Myopathy. Journal of Neuropathology and Experimental Neurology, 2010, 69, 415-424.	1.7	13
125	Inhibition of mitochondrial fusion by α-synuclein is rescued by PINK1, Parkin and DJ-1. EMBO Journal, 2010, 29, 3571-3589.	7.8	431
126	Elevated Levels of Methylmalonate and Homocysteine in Parkinson's Disease, Progressive Supranuclear Palsy and Amyotrophic Lateral Sclerosis. Dementia and Geriatric Cognitive Disorders, 2010, 29, 553-559.	1.5	40

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127	Agent strain variation in human prion disease: insights from a molecular and pathological review of the National Institutes of Health series of experimentally transmitted disease. Brain, 2010, 133, 3030-3042.	7.6	69
128	Converse modulation of toxic α-synuclein oligomers in living cells by N′-benzylidene-benzohydrazide derivates and ferric iron. Biochemical and Biophysical Research Communications, 2010, 391, 461-466.	2.1	52
129	Design of anti- and pro-aggregation variants to assess the effects of methionine oxidation in human prion protein. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 7756-7761.	7.1	98
130	CADASIL mutations enhance spontaneous multimerization of NOTCH3. Human Molecular Genetics, 2009, 18, 2761-2767.	2.9	72
131	Incidence and spectrum of sporadic Creutzfeldt–Jakob disease variants with mixed phenotype and co-occurrence of PrPSc types: an updated classification. Acta Neuropathologica, 2009, 118, 659-671.	7.7	224
132	Increased \hat{l}_{\pm} -synuclein aggregation following limited cleavage by certain matrix metalloproteinases. Experimental Neurology, 2009, 215, 201-208.	4.1	80
133	Differential effects of prion particle size on infectivity in vivo and in vitro. Biochemical and Biophysical Research Communications, 2008, 369, 924-928.	2.1	11
134	Characterization of Truncated Forms of Abnormal Prion Protein in Creutzfeldt-Jakob Disease. Journal of Biological Chemistry, 2008, 283, 30557-30565.	3.4	75
135	Single Particle Characterization of Iron-induced Pore-forming α-Synuclein Oligomers. Journal of Biological Chemistry, 2008, 283, 10992-11003.	3.4	204
136	Different Species of \hat{l}_{\pm} -Synuclein Oligomers Induce Calcium Influx and Seeding. Journal of Neuroscience, 2007, 27, 9220-9232.	3.6	708
137	Identification of Anti-prion Compounds as Efficient Inhibitors of Polyglutamine Protein Aggregation in a Zebrafish Model. Journal of Biological Chemistry, 2007, 282, 9195-9203.	3.4	95
138	Repeated Peripheral Administrations of CpG Oligodeoxynucleotides Lead to Sustained CNS Immune Activation. Immunopharmacology and Immunotoxicology, 2007, 29, 413-424.	2.4	11
139	A refined method for molecular typing reveals that co-occurrence of PrPSc types in Creutzfeldt–Jakob disease is not the rule. Laboratory Investigation, 2007, 87, 1103-1112.	3.7	60
140	Generation of genuine prion infectivity by serial PMCA. Veterinary Microbiology, 2007, 123, 346-357.	1.9	51
141	Photo-induced crosslinking of prion protein oligomers and prions. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2006, 13 , 67 - 77 .	3.0	18
142	Chaperonin TRiC Promotes the Assembly of polyQ Expansion Proteins into Nontoxic Oligomers. Molecular Cell, 2006, 23, 887-897.	9.7	259
143	Unaltered prion protein cleavage in plasminogen-deficient mice. NeuroReport, 2006, 17, 527-530.	1.2	10
144	Cell-free formation of misfolded prion protein with authentic prion infectivity. Proceedings of the National Academy of Sciences of the United States of America, 2006, 103, 15818-15823.	7.1	50

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145	Mouse Brain Synaptosomes Accumulate Copper-67 Efficiently by Two Distinct Processes Independent of Cellular Prion Protein. Journal of Molecular Neuroscience, 2005, 27, 347-354.	2.3	27
146	Systematic Identification of Antiprion Drugs by High-Throughput Screening Based on Scanning for Intensely Fluorescent Targets. Journal of Virology, 2005, 79, 7785-7791.	3.4	64
147	Automated PrPres amplification using indirect sonication. Journal of Proteomics, 2005, 63, 213-221.	2.4	13
148	Breakage of PrP aggregates is essential for efficient autocatalytic propagation of misfolded prion protein. Biochemical and Biophysical Research Communications, 2005, 326, 339-343.	2.1	29
149	Single particle analysis of manganese-induced prion protein aggregates. Biochemical and Biophysical Research Communications, 2005, 329, 1200-1207.	2.1	45
150	Single particle detection and characterization of synuclein co-aggregation. Biochemical and Biophysical Research Communications, 2005, 333, 1202-1210.	2.1	47
151	Molecular Dynamics Simulations Indicate a Possible Role of Parallel β-Helices in Seeded Aggregation of Poly-Gln. Biophysical Journal, 2005, 88, 2442-2451.	0.5	66
152	Effects of Different Experimental Conditions on the PrPSc Core Generated by Protease Digestion. Journal of Biological Chemistry, 2004, 279, 16797-16804.	3.4	118
153	Autocatalytic self-propagation of misfolded prion protein. Proceedings of the National Academy of Sciences of the United States of America, 2004, 101, 12207-12211.	7.1	109
154	Effect of metal ions on de novo aggregation of full-length prion protein. Biochemical and Biophysical Research Communications, 2004, 320, 1240-1246.	2.1	92
155	Decreased prion protein expression in human peripheral blood leucocytes from patients with paroxysmal nocturnal haemoglobinuria. British Journal of Haematology, 2001, 112, 658-662.	2. 5	4
156	Prion Protein as Copper-Binding Protein at the Synapse., 2001, 59, 17-30.		3
157	Current clinical diagnosis in Creutzfeldt-Jakob disease: Identification of uncommon variants. Annals of Neurology, 2000, 48, 323-329.	5 . 3	159
158	Differential constitutive and activation-dependent expression of prion protein in human peripheral blood leucocytes. British Journal of Haematology, 2000, 108, 488-495.	2.5	75
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