Markus Glatzel

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/6090412/publications.pdf

Version: 2024-02-01

280 papers

17,684 citations

62 h-index

18482

120 g-index

307 all docs

307 docs citations

times ranked

307

24881 citing authors

#	Article	IF	Citations
1	Anchorless risk or released benefit? An updated view on the ADAM10-mediated shedding of the prion protein. Cell and Tissue Research, 2023, 392, 215-234.	2.9	4
2	Detection of SARS-CoV-2 genomic and subgenomic RNA in retina and optic nerve of patients with COVID-19. British Journal of Ophthalmology, 2022, 106, 1313-1317.	3.9	30
3	Dying of VOC-202012/01 — multimodal investigations in a death case of the SARS-CoV-2 variant. International Journal of Legal Medicine, 2022, 136, 193-202.	2.2	3
4	The blood-brain barrier is dysregulated in COVID-19 and serves as a CNS entry route for SARS-CoV-2. Stem Cell Reports, 2022, 17, 307-320.	4.8	138
5	Validation of Revised International Creutzfeldt-Jakob Disease Surveillance Network Diagnostic Criteria for Sporadic Creutzfeldt-Jakob Disease. JAMA Network Open, 2022, 5, e2146319.	5.9	28
6	Transcriptional Alterations in X-Linked Dystonia–Parkinsonism Caused by the SVA Retrotransposon. International Journal of Molecular Sciences, 2022, 23, 2231.	4.1	6
7	Meprin β knockout reduces brain Aβ levels and rescues learning and memory impairments in the APP/lon mouse model for Alzheimer's disease. Cellular and Molecular Life Sciences, 2022, 79, 168.	5.4	3
8	Comprehensive profiling of myxopapillary ependymomas identifies a distinct molecular subtype with relapsing disease. Neuro-Oncology, 2022, 24, 1689-1699.	1.2	11
9	Organ manifestations of COVID-19: what have we learned so far (not only) from autopsies?. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2022, 481, 139-159.	2.8	28
10	NeuroCOVID: Insights into Neuroinvasion and Pathophysiology. Clinical and Translational Neuroscience, 2022, 6, 10.	0.9	1
11	The prion protein and its ligands: Insights into structure-function relationships. Biochimica Et Biophysica Acta - Molecular Cell Research, 2022, 1869, 119240.	4.1	10
12	Paul Kleihues (1936–2022), neuropathology innovator and entrepreneur. Brain Pathology, 2022, 32, e13073.	4.1	0
13	Response to: SARS-CoV-2 and type I interferon signaling in brain endothelial cells: Blurring the lines between friend or foe. Stem Cell Reports, 2022, 17, 1014-1015.	4.8	5
14	Diagnostic potential of extracellular vesicles in meningioma patients. Neuro-Oncology, 2022, 24, 2078-2090.	1.2	6
15	EPEN-06. Comprehensive profiling of myxopapillary ependymomas identifies a distinct molecular subtype with relapsing disease. Neuro-Oncology, 2022, 24, i39-i39.	1.2	0
16	Distinct tau neuropathology and cellular profiles of an APOE3 Christchurch homozygote protected against autosomal dominant Alzheimer's dementia. Acta Neuropathologica, 2022, 144, 589-601.	7.7	32
17	Ependymoma relapse goes along with a relatively stable epigenome, but a severely altered tumor morphology. Brain Pathology, 2021, 31, 33-44.	4.1	8
18	Co-expression of intermediate filaments glial fibrillary acidic protein and cytokeratin in pituitary adenoma. Pituitary, 2021, 24, 62-67.	2.9	2

#	Article	IF	CITATIONS
19	A multifactorial model of pathology for age of onset heterogeneity in familial Alzheimer's disease. Acta Neuropathologica, 2021, 141, 217-233.	7.7	33
20	Disordered structure and flexible roles: using the prion protein N1 fragment for neuroprotective and regenerative therapy. Neural Regeneration Research, 2021, 16, 1431.	3.0	3
21	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
22	A tribute to our case of the month and a warm welcome to "under your microscope― Brain Pathology, 2021, 31, 3-3.	4.1	1
23	Assessment of small fiber neuropathy in patients carrying the nonâ€classical <scp>Fabry</scp> variant <scp>p.D313Y</scp> . Muscle and Nerve, 2021, 63, 745-750.	2.2	5
24	Neuroserpin Is Strongly Expressed in the Developing and Adult Mouse Neocortex but Its Absence Does Not Perturb Cortical Lamination and Synaptic Proteome. Frontiers in Neuroanatomy, 2021, 15, 627896.	1.7	6
25	Prion protein oligomers cause neuronal cytoskeletal damage in rapidly progressive Alzheimer's disease. Molecular Neurodegeneration, 2021, 16, 11.	10.8	15
26	Presence of SARS-CoV-2 RNA in the Cornea of Viremic Patients With COVID-19. JAMA Ophthalmology, 2021, 139, 383.	2.5	62
27	G392E neuroserpin causing the dementia FENIB is secreted from cells but is not synaptotoxic. Scientific Reports, 2021, 11, 8766.	3.3	7
28	Differential expression of stem cell markers in proliferating cells in glioma. Journal of Cancer Research and Clinical Oncology, 2021, 147, 2969-2982.	2.5	8
29	CD74 and CD44 Expression on CTCs in Cancer Patients with Brain Metastasis. International Journal of Molecular Sciences, 2021, 22, 6993.	4.1	26
30	Deep spatial profiling of human COVID-19 brains reveals neuroinflammation with distinct microanatomical microglia-T-cell interactions. Immunity, 2021, 54, 1594-1610.e11.	14.3	210
31	Targeting Runt-Related Transcription Factor 1 Prevents Pulmonary Fibrosis and Reduces Expression of Severe Acute Respiratory Syndrome Coronavirus 2 Host Mediators. American Journal of Pathology, 2021, 191, 1193-1208.	3.8	14
32	Prospective postmortem evaluation of 735 consecutive SARS-CoV-2-associated death cases. Scientific Reports, 2021, 11, 19342.	3.3	28
33	Neuropathology associated with SARS-CoV-2 infection. Lancet, The, 2021, 397, 276.	13.7	13
34	Genome-wide methylation profiling of glioblastoma cell-derived extracellular vesicle DNA allows tumor classification. Neuro-Oncology, 2021, 23, 1087-1099.	1.2	59
35	The SARS-CoV-2 main protease Mpro causes microvascular brain pathology by cleaving NEMO in brain endothelial cells. Nature Neuroscience, 2021, 24, 1522-1533.	14.8	164
36	Co-activation of Sonic hedgehog and Wnt signaling in murine retinal precursor cells drives ocular lesions with features of intraocular medulloepithelioma. Oncogenesis, 2021, 10, 78.	4.9	0

#	Article	IF	Citations
37	Ligands binding to the prion protein induce its proteolytic release with therapeutic potential in neurodegenerative proteinopathies. Science Advances, 2021, 7, eabj1826.	10.3	18
38	BIOM-39. METHYLATION AND MUTATION PROFILES IN MENINGIOMA CELL-DERIVED EXTRACELLULAR VESICLE DNA REFLECT EPIGENETIC AND GENOMIC ALTERATIONS IN ORIGINAL TUMORS. Neuro-Oncology, 2021, 23, vi19-vi19.	1.2	0
39	Reactive Astrocytes Contribute to Alzheimer's Disease-Related Neurotoxicity and Synaptotoxicity in a Neuron-Astrocyte Co-culture Assay. Frontiers in Cellular Neuroscience, 2021, 15, 739411.	3.7	7
40	Distinctive low epidermal nerve fiber density in schwannomatosis patients provides a major parameter for diagnosis and differential diagnosis. Brain Pathology, 2020, 30, 386-391.	4.1	4
41	Enzyme replacement therapy with recombinant pro-CTSD (cathepsin D) corrects defective proteolysis and autophagy in neuronal ceroid lipofuscinosis. Autophagy, 2020, 16, 811-825.	9.1	70
42	Molecular characterization of histopathological ependymoma variants. Acta Neuropathologica, 2020, 139, 305-318.	7.7	43
43	Shortening heparan sulfate chains prolongs survival and reduces parenchymal plaques in prion disease caused by mobile, ADAM10-cleaved prions. Acta Neuropathologica, 2020, 139, 527-546.	7.7	23
44	Deficits in developmental neurogenesis and dendritic spine maturation in mice lacking the serine protease inhibitor neuroserpin. Molecular and Cellular Neurosciences, 2020, 102, 103420.	2.2	25
45	The histone H2B ubiquitin ligase RNF40 is required for HER2-driven mammary tumorigenesis. Cell Death and Disease, 2020, 11, 873.	6.3	10
46	Neuropathology of patients with COVID-19 in Germany: a post-mortem case series. Lancet Neurology, The, 2020, 19, 919-929.	10.2	957
47	Germany's first COVID-19 deceased: a 59-year-old man presenting with diffuse alveolar damage due to SARS-CoV-2 infection. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2020, 477, 335-339.	2.8	29
48	Influence of Methanol on Prion Reduction during High Temperature and High Pressure Oleochemical Processes. European Journal of Lipid Science and Technology, 2020, 122, 2000136.	1.5	1
49	Decreased Deposition of Beta-Amyloid 1-38 and Increased Deposition of Beta-Amyloid 1-42 in Brain Tissue of Presenilin-1 E280A Familial Alzheimer's Disease Patients. Frontiers in Aging Neuroscience, 2020, 12, 220.	3.4	13
50	Characterization of brainâ€derived extracellular vesicles reveals changes in cellular origin after stroke and enrichment of the prion protein with a potential role in cellular uptake. Journal of Extracellular Vesicles, 2020, 9, 1809065.	12.2	47
51	<i>Brain Pathology</i> moves to open access. Brain Pathology, 2020, 30, 1011-1011.	4.1	1
52	Assessment of Peripheral Nervous System Alterations in Patients with the Fabry Related GLA-Variant p.A143T. Diagnostics, 2020, 10, 1027.	2.6	4
53	Multiorgan and Renal Tropism of SARS-CoV-2. New England Journal of Medicine, 2020, 383, 590-592.	27.0	1,523
54	Prion protein post-translational modifications modulate heparan sulfate binding and limit aggregate size in prion disease. Neurobiology of Disease, 2020, 142, 104955.	4.4	5

#	Article	IF	CITATIONS
55	Neuropathology of COVIDâ€19: where are the neuropathologists?. Brain Pathology, 2020, 30, 729-729.	4.1	11
56	Cerebellar heterotopia of infancy in sudden infant death syndrome: an observational neuropathological study of four cases. International Journal of Legal Medicine, 2020, 134, 2143-2147.	2.2	3
57	Transgenic Overexpression of the Disordered Prion Protein N1 Fragment in Mice Does Not Protect Against Neurodegenerative Diseases Due to Impaired ER Translocation. Molecular Neurobiology, 2020, 57, 2812-2829.	4.0	17
58	ALCAM contributes to brain metastasis formation in non-small-cell lung cancer through interaction with the vascular endothelium. Neuro-Oncology, 2020, 22, 955-966.	1.2	36
59	Susceptibility to cellular stress in PS1 mutant N2a cells is associated with mitochondrial defects and altered calcium homeostasis. Scientific Reports, 2020, 10, 6455.	3.3	6
60	Prion protein glycans reduce intracerebral fibril formation and spongiosis in prion disease. Journal of Clinical Investigation, 2020, 130, 1350-1362.	8.2	32
61	C-Fiber Loss as a Possible Cause of Neuropathic Pain in Schwannomatosis. International Journal of Molecular Sciences, 2020, 21, 3569.	4.1	5
62	Standardizing Brain Cancer Reporting. Brain Pathology, 2019, 29, 465-465.	4.1	0
63	Early-onset stroke in two siblings with Neurofibromatosis type 1. European Journal of Medical Genetics, 2019, 62, 103710.	1.3	4
64	Clonality of circulating tumor cells in breast cancer brain metastasis patients. Breast Cancer Research, 2019, 21, 101.	5.0	54
65	The cellular prion protein and its derived fragments in human prion diseases and their role as potential biomarkers. Expert Review of Molecular Diagnostics, 2019, 19, 1007-1018.	3.1	5
66	Severe meningo-/encephalitis after daclizumab therapy for multiple sclerosis. Multiple Sclerosis Journal, 2019, 25, 1618-1632.	3.0	32
67	Complement 3+-astrocytes are highly abundant in prion diseases, but their abolishment led to an accelerated disease course and early dysregulation of microglia. Acta Neuropathologica Communications, 2019, 7, 83.	5.2	84
68	Phagocytosis of Apoptotic Cells Is Specifically Upregulated in ApoE4 Expressing Microglia in vitro. Frontiers in Cellular Neuroscience, 2019, 13, 181.	3.7	26
69	Impact of USP8 Gene Mutations on Protein Deregulation in Cushing Disease. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 2535-2546.	3.6	29
70	EGFR and HER3 expression in circulating tumor cells and tumor tissue from non-small cell lung cancer patients. Scientific Reports, 2019, 9, 7406.	3.3	73
71	Dear Reader: Data citation in changing times. Brain Pathology, 2019, 29, 153-154.	4.1	1
72	Imaging flow cytometry facilitates multiparametric characterization of extracellular vesicles in malignant brain tumours. Journal of Extracellular Vesicles, 2019, 8, 1588555.	12.2	86

#	Article	IF	CITATIONS
73	The Colombian–German network for neurodegenerative research: UndoAD. Lancet Neurology, The, 2019, 18, 29.	10.2	1
74	Recent advances on the molecular pathogenesis of prion diseases. Brain Pathology, 2019, 29, 245-247.	4.1	0
75	GPI-anchor signal sequence influences PrPC sorting, shedding and signalling, and impacts on different pathomechanistic aspects of prion disease in mice. PLoS Pathogens, 2019, 15, e1007520.	4.7	34
76	Cellular and Molecular Mechanisms of Prion Disease. Annual Review of Pathology: Mechanisms of Disease, 2019, 14, 497-516.	22.4	83
77	Molecular Mechanisms of Prion Diseases. , 2019, , .		0
78	Abstract 1114: Upregulation of ALCAM is a marker for non-small-cell lung cancer brain metastases. , 2019, , .		0
79	Infectious prions do not induce $\hat{Al^2}$ deposition in an in vivo seeding model. Acta Neuropathologica, 2018, 135, 965-967.	7.7	8
80	Hemodynamic Forces Tune the Arrest, Adhesion, and Extravasation of Circulating Tumor Cells. Developmental Cell, 2018, 45, 33-52.e12.	7.0	219
81	No difference in the prevalence of Alzheimer-type neurodegenerative changes in the brains of suicides when compared with controls: an explorative neuropathologic study. European Archives of Psychiatry and Clinical Neuroscience, 2018, 268, 509-517.	3.2	4
82	Structural and mechanistic aspects influencing the ADAM10-mediated shedding of the prion protein. Molecular Neurodegeneration, 2018, 13, 18.	10.8	45
83	DNA methylation-based reclassification of olfactory neuroblastoma. Acta Neuropathologica, 2018, 136, 255-271.	7.7	59
84	In vivo regulation of the A disintegrin and metalloproteinase 10 (ADAM10) by the tetraspanin 15. Cellular and Molecular Life Sciences, 2018, 75, 3251-3267.	5.4	37
85	Diagnostic red flags: steroidâ€treated malignant CNS lymphoma mimicking autoimmune inflammatory demyelination. Brain Pathology, 2018, 28, 225-233.	4.1	28
86	IL-23 (Interleukin-23)–Producing Conventional Dendritic Cells Control the Detrimental IL-17 (Interleukin-17) Response in Stroke. Stroke, 2018, 49, 155-164.	2.0	81
87	Oncogenic Amplification of Zygotic Dux Factors in Regenerating p53-Deficient Muscle Stem Cells Defines a Molecular Cancer Subtype. Cell Stem Cell, 2018, 23, 794-805.e4.	11.1	21
88	Frequency of Circulating Tumor Cells (CTC) in Patients with Brain Metastases: Implications as a Risk Assessment Marker in Oligo-Metastatic Disease. Cancers, 2018, 10, 527.	3.7	45
89	Dear Reader: another changing of the guard. Brain Pathology, 2018, 28, 789-789.	4.1	0
90	Alterations in the brain interactome of the intrinsically disordered N-terminal domain of the cellular prion protein (PrPC) in Alzheimer's disease. PLoS ONE, 2018, 13, e0197659.	2.5	20

#	Article	IF	Citations
91	Distinct microglia profile in Creutzfeldt–Jakob disease and Alzheimer's disease is independent of disease kinetics. Neuropathology, 2018, 38, 591-600.	1.2	3
92	Muskelin Coordinates PrPC Lysosome versus Exosome Targeting and Impacts Prion Disease Progression. Neuron, 2018, 99, 1155-1169.e9.	8.1	39
93	Validation and utilization of amended diagnostic criteria in Creutzfeldt-Jakob disease surveillance. Neurology, 2018, 91, e331-e338.	1.1	84
94	The secreted glycolytic enzyme GPI/AMF stimulates glioblastoma cell migration and invasion in an autocrine fashion but can have anti-proliferative effects. Neuro-Oncology, 2018, 20, 1594-1605.	1.2	21
95	RNase H2 Loss in Murine Astrocytes Results in Cellular Defects Reminiscent of Nucleic Acid-Mediated Autoinflammation. Frontiers in Immunology, 2018, 9, 587.	4.8	14
96	Inverse Perfusion Requirements of Supra- and Infratentorial Brain Metastases Formation. Frontiers in Neurology, 2018, 9, 391.	2.4	5
97	N-Glycosylation of Extracellular Vesicles from HEK-293 and Glioma Cell Lines. Analytical Chemistry, 2018, 90, 7871-7879.	6.5	42
98	The TREM2-APOE Pathway Drives the Transcriptional Phenotype of Dysfunctional Microglia in Neurodegenerative Diseases. Immunity, 2017, 47, 566-581.e9.	14.3	1,741
99	The serine protease inhibitor neuroserpin is required for normal synaptic plasticity and regulates learning and social behavior. Learning and Memory, 2017, 24, 650-659.	1.3	24
100	Amyloid polymorphisms constitute distinct clouds of conformational variants in different etiological subtypes of Alzheimer's disease. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, 13018-13023.	7.1	170
101	Subtypes of primary angiitis of the CNS identified by MRI patterns reflect the size of affected vessels. Journal of Neurology, Neurosurgery and Psychiatry, 2017, 88, 749-755.	1.9	38
102	Diverse functions of the prion protein – Does proteolytic processing hold the key?. Biochimica Et Biophysica Acta - Molecular Cell Research, 2017, 1864, 2128-2137.	4.1	60
103	Tetraspanin 3: A central endocytic membrane component regulating the expression of ADAM10, presenilin and the amyloid precursor protein. Biochimica Et Biophysica Acta - Molecular Cell Research, 2017, 1864, 217-230.	4.1	26
104	IgG4â€related hypophysitis is highly prevalent among cases of histologically confirmed hypophysitis. Brain Pathology, 2017, 27, 839-845.	4.1	42
105	Immunoprofiling of glial tumours of the neurohypophysis suggests a common pituicytic origin of neoplastic cells. Pituitary, 2017, 20, 211-217.	2.9	26
106	Activation of microglia by retroviral infection correlates with transient clearance of prions from the brain but does not change incubation time. Brain Pathology, 2017, 27, 590-602.	4.1	19
107	Losing sleep over mitochondria: a new player in the pathophysiology of fatal familial insomnia. Brain Pathology, 2017, 27, 107-108.	4.1	4
108	Exosomes and the Prion Protein: More than One Truth. Frontiers in Neuroscience, 2017, 11, 194.	2.8	60

#	Article	IF	Citations
109	YKL-40 in the brain and cerebrospinal fluid of neurodegenerative dementias. Molecular Neurodegeneration, 2017, 12, 83.	10.8	140
110	PTEN mediates the cross talk between breast and glial cells in brain metastases leading to rapid disease progression. Oncotarget, 2017, 8, 6155-6168.	1.8	35
111	Functional characterization of the lysosomal membrane protein TMEM192 in mice. Oncotarget, 2017, 8, 43635-43652.	1.8	8
112	Exosomes in Prion Diseases. Neuromethods, 2017, , 197-207.	0.3	0
113	Aromatase Expression in the Hippocampus of AD Patients and 5xFAD Mice. Neural Plasticity, 2016, 2016, 1-11.	2.2	20
114	Evidence of a pathogenic role for CD8 ⁺ T cells in anti-GABA _B receptor limbic encephalitis. Neurology: Neuroimmunology and NeuroInflammation, 2016, 3, e232.	6.0	46
115	Exosomal cellular prion protein drives fibrillization of amyloid beta and counteracts amyloid betaâ€mediated neurotoxicity. Journal of Neurochemistry, 2016, 137, 88-100.	3.9	117
116	IL-17 production by CSF lymphocytes as a biomarker for cerebral vasculitis. Neurology: Neuroimmunology and NeuroInflammation, 2016, 3, e214.	6.0	29
117	Amyloid- \hat{l}^2 Precursor Protein Modulates the Sorting of Testican-1 and Contributes to Its Accumulation in Brain Tissue and Cerebrospinal Fluid from Patients with Alzheimer Disease. Journal of Neuropathology and Experimental Neurology, 2016, 75, 903-916.	1.7	18
118	Misfolding leads the way to unraveling signaling pathways in the pathophysiology of prion diseases. Prion, 2016, 10, 434-443.	1.8	2
119	Secretory pathway retention of mutant prion protein induces p38-MAPK activation and lethal disease in mice. Scientific Reports, 2016, 6, 24970.	3.3	22
120	Generation of aggregation prone N-terminally truncated amyloid \hat{l}^2 peptides by meprin \hat{l}^2 depends on the sequence specificity at the cleavage site. Molecular Neurodegeneration, 2016, 11, 19.	10.8	65
121	Mannose 6-phosphate-dependent targeting of lysosomal enzymes is required for normal craniofacial and dental development. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2016, 1862, 1570-1580.	3.8	15
122	Limited Unfolded Protein Response and Inflammation in Neuroserpinopathy. Journal of Neuropathology and Experimental Neurology, 2016, 75, 121-133.	1.7	8
123	Epidermal growth factor receptor overexpression is common and not correlated to gene copy number in ependymoma. Child's Nervous System, 2016, 32, 281-290.	1.1	7
124	Frontal lobe dementia syndrome as a first manifestation of primary angiitis of the central nervous system (PACNS). Clinical Neurology and Neurosurgery, 2016, 141, 92-94.	1.4	7
125	Abstract 1571: PTEN is an important mediator of tumor and glia cell crosstalk in breast cancer brain metastasis. Cancer Research, 2016, 76, 1571-1571.	0.9	0
126	Comment on "Primary Central Nervous System (CNS) Lymphoma B Cell Receptors Recognize CNS Proteins― Journal of Immunology, 2015, 195, 4549-4550.	0.8	5

#	Article	IF	CITATIONS
127	Shedding light on prion disease. Prion, 2015, 9, 244-256.	1.8	17
128	Encephalopathy and death in infants with abusive head trauma is due to hypoxic-ischemic injury following local brain trauma to vital brainstem centers. International Journal of Legal Medicine, 2015, 129, 105-114.	2.2	36
129	Upregulation of Shiga Toxin Receptor <scp>CD</scp> 77/ <scp>G</scp> b3 and Interleukinâ€1β Expression in the Brain of <scp>EHEC</scp> Patients with Hemolytic Uremic Syndrome and Neurologic Symptoms. Brain Pathology, 2015, 25, 146-156.	4.1	12
130	Deficiency of the miR-29a/b-1 cluster leads to ataxic features and cerebellar alterations in mice. Neurobiology of Disease, 2015, 73, 275-288.	4.4	46
131	Distribution and prognostic relevance of tumor-infiltrating lymphocytes (TILs) and PD-1/PD-L1 immune checkpoints in human brain metastases. Oncotarget, 2015, 6, 40836-40849.	1.8	106
132	The sheddase ADAM10 is a potent modulator of prion disease. ELife, 2015, 4, .	6.0	66
133	Abstract P6-16-09: SPARC expression in brain metastases of breast cancer patients., 2015,,.		0
134	Vessel and Mast Cell Densities in Sporadic and Syndrome-associated Peripheral Nerve Sheath Tumors. Anticancer Research, 2015, 35, 4713-22.	1.1	4
135	Vascular Innervation in Benign Neurofibromas of Patients with Neurofibromatosis Type 1. Anticancer Research, 2015, 35, 6509-16.	1.1	3
136	Immune Activation in Amyloid- \hat{l}^2 -Related Angiitis Correlates with Decreased Parenchymal Amyloid- \hat{l}^2 Plaque Load. Neurodegenerative Diseases, 2014, 13, 38-44.	1.4	26
137	The GPI-anchoring of PrP. Prion, 2014, 8, 11-18.	1.8	49
138	Creutzfeldt-Jakob disease mimicking autoimmune encephalitis with CASPR2 antibodies. BMC Neurology, 2014, 14, 227.	1.8	16
139	Dissemination of Orientia tsutsugamushi and Inflammatory Responses in a Murine Model of Scrub Typhus. PLoS Neglected Tropical Diseases, 2014, 8, e3064.	3.0	62
140	Transient Receptor Potential Melastatin Subfamily Member 2 Cation Channel Regulates Detrimental Immune Cell Invasion in Ischemic Stroke. Stroke, 2014, 45, 3395-3402.	2.0	85
141	Hematogenous dissemination of glioblastoma multiforme. Science Translational Medicine, 2014, 6, 247ra101.	12.4	264
142	The lectin OS-9 delivers mutant neuroserpin to endoplasmic reticulum associated degradation in familial encephalopathy with neuroserpin inclusion bodies. Neurobiology of Aging, 2014, 35, 2394-2403.	3.1	23
143	Human CLP1 Mutations Alter tRNA Biogenesis, Affecting Both Peripheral and Central Nervous System Function. Cell, 2014, 157, 636-650.	28.9	189
144	TRPM2 cation channel regulates detrimental immune cell invasion in ischemic stroke. Journal of Neuroimmunology, 2014, 275, 99.	2.3	0

#	Article	IF	Citations
145	LIMP-2 expression is critical for \hat{l}^2 -glucocerebrosidase activity and $\hat{l}\pm$ -synuclein clearance. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 15573-15578.	7.1	109
146	High molecular mass assemblies of amyloid-β oligomers bind prion protein in patients with Alzheimer's disease. Brain, 2014, 137, 873-886.	7.6	96
147	Familial Alzheimer's disease–associated presenilin-1 alters cerebellar activity and calcium homeostasis. Journal of Clinical Investigation, 2014, 124, 1552-1567.	8.2	104
148	B-Cell Receptors of Primary Central Nervous System Lymphoma Recognize Antigens in the Brain. Blood, 2014, 124, 3003-3003.	1.4	1
149	Loss of CADM1 expression is associated with poor prognosis and brain metastasis in breast cancer patients. Oncotarget, 2014, 5, 3076-3087.	1.8	35
150	Abstract 46: Loss of CADM1 expression is associated with poor prognosis and brain metastasis in breast cancer patients. , 2014 , , .		0
151	No reactivation of JCV and CMV infections in the temporal cortex and cerebellum of sporadic Creutzfeldt-Jakob disease patients. American Journal of Neurodegenerative Disease, 2014, 3, 152-7.	0.1	0
152	Postnatal Disruption of the Disintegrin/Metalloproteinase ADAM10 in Brain Causes Epileptic Seizures, Learning Deficits, Altered Spine Morphology, and Defective Synaptic Functions. Journal of Neuroscience, 2013, 33, 12915-12928.	3.6	107
153	Hypoxia and oxygenation induce a metabolic switch between pentose phosphate pathway and glycolysis in glioma stem-like cells. Acta Neuropathologica, 2013, 126, 763-780.	7.7	106
154	A Novel Interaction Between Aging and ER Overload in a Protein Conformational Dementia. Genetics, 2013, 193, 865-876.	2.9	21
155	STAT3 silencing inhibits glioma single cell infiltration and tumor growth. Neuro-Oncology, 2013, 15, 840-852.	1.2	57
156	Myositis facilitates preclinical accumulation of pathological prion protein in muscle. Acta Neuropathologica Communications, 2013, 1, 78.	5.2	1
157	Intravenous immunoglobulin suppresses NLRP1 and NLRP3 inflammasome-mediated neuronal death in ischemic stroke. Cell Death and Disease, 2013, 4, e790-e790.	6.3	331
158	Plasma levels of neuron specific enolase quantify the extent of neuronal injury in murine models of ischemic stroke and multiple sclerosis. Neurobiology of Disease, 2013, 59, 177-182.	4.4	16
159	Frequent Genetic Alterations in EGFR- and HER2-Driven Pathways in Breast Cancer Brain Metastases. American Journal of Pathology, 2013, 183, 83-95.	3.8	63
160	CLP1 links tRNA metabolism to progressive motor-neuron loss. Nature, 2013, 495, 474-480.	27.8	231
161	Active vaccination with ankyrin G reduces \hat{l}^2 -amyloid pathology in APP transgenic mice. Molecular Psychiatry, 2013, 18, 358-368.	7.9	23
162	Roles of endoproteolytic αâ€cleavage and shedding of the prion protein in neurodegeneration. FEBS Journal, 2013, 280, 4338-4347.	4.7	48

#	Article	IF	Citations
163	BSE-associated Prion-Amyloid Cardiomyopathy in Primates. Emerging Infectious Diseases, 2013, 19, 985-988.	4.3	10
164	Protease-sensitive prion species in neoplastic spleens of prion-infected mice with uncoupling of PrPSc and prion infectivity. Journal of General Virology, 2013, 94, 453-463.	2.9	13
165	Specific de-SUMOylation triggered by acquisition of spatial learning is related to epigenetic changes in the rat hippocampus. NeuroReport, 2013, 24, 976-981.	1.2	6
166	Deficiency in Serine Protease Inhibitor Neuroserpin Exacerbates Ischemic Brain Injury by Increased Postischemic Inflammation. PLoS ONE, 2013, 8, e63118.	2.5	37
167	Amyloid-Precursor-Protein-Lowering Small Molecules for Disease Modifying Therapy of Alzheimer's Disease. PLoS ONE, 2013, 8, e82255.	2.5	6
168	The neurological syndrome in adults during the 2011 northern German E. coli serotype O104:H4 outbreak. Brain, 2012, 135, 1850-1859.	7.6	105
169	Phenotypic Profile of Early-Onset Familial Alzheimer's Disease Caused by Presenilin-1 E280A Mutation. Journal of Alzheimer's Disease, 2012, 32, 1-12.	2.6	55
170	Neutralization of the IL-17 axis diminishes neutrophil invasion and protects from ischemic stroke. Blood, 2012, 120, 3793-3802.	1.4	374
171	Relevance of PTEN loss in brain metastasis formation in breast cancer patients. Breast Cancer Research, 2012, 14, R49.	5.0	93
172	Persistent retroviral infection with MoMuLV influences neuropathological signature and phenotype of prion disease. Acta Neuropathologica, 2012, 124, 111-126.	7.7	14
173	Hyaluronan in intraâ€operative edema of NF1â€associated neurofibromas. Neuropathology, 2012, 32, 406-414.	1.2	7
174	Opposite roles of <i>FOXA1</i> and <i>NKX2â€1</i> in lung cancer progression. Genes Chromosomes and Cancer, 2012, 51, 618-629.	2.8	27
175	Proteolytic processing of the prion protein in health and disease. American Journal of Neurodegenerative Disease, 2012, 1, 15-31.	0.1	58
176	Non-human primates in prion research. , 2012, 50, 57-67.		3
177	Deposition of Hyperphosphorylated Tau in Cerebellum of PS1 E280A Alzheimer's Disease. Brain Pathology, 2011, 21, 452-463.	4.1	65
178	Podoplanin and CD34 in peripheral nerve sheath tumours: focus on neurofibromatosis 1-associated atypical neurofibroma. Journal of Neuro-Oncology, 2011, 103, 239-245.	2.9	21
179	Lack of a-disintegrin-and-metalloproteinase ADAM10 leads to intracellular accumulation and loss of shedding of the cellular prion protein in vivo. Molecular Neurodegeneration, 2011, 6, 36.	10.8	93
180	Treatment of glioblastoma with poly(isohexyl cyanoacrylate) nanoparticles. International Journal of Pharmaceutics, 2011, 415, 244-251.	5.2	44

#	Article	IF	CITATIONS
181	Complementary and alternative medicine (CAM) use in terminally ill patients. Trace Elements and Electrolytes, 2011, 28, 49-51.	0.1	4
182	Selenium substitution during radiotherapy of solid tumors. Trace Elements and Electrolytes, 2011, 28, 101-104.	0.1	0
183	Efficient Chemotherapy of Rat Glioblastoma Using Doxorubicin-Loaded PLGA Nanoparticles with Different Stabilizers. PLoS ONE, 2011, 6, e19121.	2.5	138
184	N-Glycans and Glycosylphosphatidylinositol-Anchor Act on Polarized Sorting of Mouse PrPC in Madin-Darby Canine Kidney Cells. PLoS ONE, 2011, 6, e24624.	2.5	19
185	Preclinical Deposition of Pathological Prion Protein in Muscle of Experimentally Infected Primates. PLoS ONE, 2010, 5, e13906.	2.5	19
186	Chronic wasting disease prions are not transmissible to transgenic mice overexpressing human prion protein. Journal of General Virology, 2010, 91, 2651-2657.	2.9	106
187	Dietary Curcumin Attenuates Glioma Growth in a Syngeneic Mouse Model by Inhibition of the JAK1,2/STAT3 Signaling Pathway. Clinical Cancer Research, 2010, 16, 5781-5795.	7.0	120
188	The Disintegrin/Metalloproteinase ADAM10 Is Essential for the Establishment of the Brain Cortex. Journal of Neuroscience, 2010, 30, 4833-4844.	3.6	327
189	Embryonic stem cell-derived L1 overexpressing neural aggregates enhance recovery in Parkinsonian mice. Brain, 2010, 133, 189-204.	7.6	54
190	A Novel Single-Chain Antibody Fragment for Detection of Mannose 6-Phosphate-Containing Proteins. American Journal of Pathology, 2010, 177, 240-247.	3.8	33
191	Selenium substitution during radiotherapy in head and neck cancer. Trace Elements and Electrolytes, 2010, 27, 235-239.	0.1	9
192	Trace elements selenium and zinc as tumor markers in patients with advanced head and neck cancer. Trace Elements and Electrolytes, 2010, 27, 246-249.	0.1	3
193	Nonaccidental Head Injury Is the Most Common Cause of Subdural Bleeding in Infants & Dit; 1 Year of Age. Pediatrics, 2009, 124, 1587-1594.	2.1	50
194	Genomic Profiles Associated with Early Micrometastasis in Lung Cancer: Relevance of 4q Deletion. Clinical Cancer Research, 2009, 15, 1566-1574.	7.0	87
195	Shaken Baby Syndrome. Deutsches Ärzteblatt International, 2009, 106, 211-7.	0.9	60
196	A case-control study of sporadic Creutzfeldt-Jakob disease in Switzerland: analysis of potential risk factors with regard to an increased CJD incidence in the years 2001–2004. BMC Public Health, 2009, 9, 18.	2.9	30
197	Autoantibodies against βâ€amyloid are common in Alzheimer's disease and help control plaque burden. Annals of Neurology, 2009, 65, 24-31.	5. 3	119
198	Ocular pathology in shaken baby syndrome and other forms of infantile non-accidental head injury. International Journal of Legal Medicine, 2009, 123, 189-197.	2.2	50

#	Article	IF	Citations
199	Selenium Does Not Prevent Radiation-induced Toxicities in Radiotherapy (RT) for Head and Neck Cancer. International Journal of Radiation Oncology Biology Physics, 2009, 75, S519.	0.8	0
200	Mast Cell-Mediated Antigen Presentation Regulates CD8+ T Cell Effector Functions. Immunity, 2009, 31, 665-676.	14.3	145
201	Plasminogen Activator Inhibitor Type 1 Up-Regulation Is Associated with Skeletal Muscle Atrophy and Associated Fibrosis. American Journal of Pathology, 2009, 175, 763-771.	3.8	25
202	Increased Numbers of Injections of Doxorubicin Bound to Nanoparticles Lead to Enhanced Efficacy Against Rat Glioblastoma 101/8. Journal of Nanoneuroscience, 2009, 1, 144-151.	0.5	10
203	Efficient systemic therapy of rat glioblastoma by nanoparticle-bound doxorubicin is due to antiangiogenic effects., 2009, 28, 153-164.		53
204	Postoperative serum and whole blood selenium levels in patients with squamous cell and adenocarcinomas of the uterus after curative surgical treatment. Trace Elements and Electrolytes, 2009, 26, 78-82.	0.1	6
205	Heightened incidence of sporadic Creutzfeldt-Jakob disease is associated with a shift in clinicopathological profiles. Journal of Neurology, 2008, 255, 1464-1472.	3.6	9
206	Tenascin-R Promotes Neuronal Differentiation of Embryonic Stem Cells and Recruitment of Host-Derived Neural Precursor Cells After Excitotoxic Lesion of the Mouse Striatum. Stem Cells, 2008, 26, 1973-1984.	3.2	33
207	Accumulation of bis(monoacylglycero)phosphate and gangliosides in mouse models of neuronal ceroid lipofuscinosis. Journal of Neurochemistry, 2008, 106, 1415-1425.	3.9	46
208	Experimental Chronic Wasting Disease (CWD) in the Ferret. Journal of Comparative Pathology, 2008, 138, 189-196.	0.4	47
209	Association between Deposition of Beta-Amyloid and Pathological Prion Protein in Sporadic Creutzfeldt-Jakob Disease. Neurodegenerative Diseases, 2008, 5, 347-354.	1.4	50
210	Transport and distribution of 3-hydroxyglutaric acid before and during induced encephalopathic crises in a mouse model of glutaric aciduria type 1. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2008, 1782, 385-390.	3.8	29
211	Transmissible spongiform encephalopathies. Journal of the American Veterinary Medical Association, 2008, 233, 1705-1712.	0.5	9
212	Biomarker profile for the diagnosis of Creutzfeldt–Jakob disease. Nature Clinical Practice Neurology, 2008, 4, 70-71.	2.5	1
213	Multicenter, phase-III study comparing selenium supplementation with observation in gynecologic radiation oncology. Journal of Clinical Oncology, 2008, 26, 9539-9539.	1.6	2
214	Understanding the natural variability of prion diseases. Vaccine, 2007, 25, 5631-5636.	3.8	23
215	Accumulation of Mutant Neuroserpin Precedes Development of Clinical Symptoms in Familial Encephalopathy with Neuroserpin Inclusion Bodies. American Journal of Pathology, 2007, 170, 1305-1313.	3.8	33
216	3-Hydroxyglutaric acid is transported via the sodium-dependent dicarboxylate transporter NaDC3. Journal of Molecular Medicine, 2007, 85, 763-770.	3.9	33

#	Article	IF	Citations
217	Induced Prion Protein Controls Immune-Activated Retroviruses in the Mouse Spleen. PLoS ONE, 2007, 2, e1158.	2.5	29
218	The LPS Receptor, CD14 in Experimental Autoimmune Encephalomyelitis and Multiple Sclerosis. Cellular Physiology and Biochemistry, 2006, 17, 167-172.	1.6	33
219	Cerebrovascular P-glycoprotein expression is decreased in Creutzfeldt–Jakob disease. Acta Neuropathologica, 2006, 111, 436-443.	7.7	40
220	Human transmissible spongiform encephalopathies in eleven countries: diagnostic pattern across time, 1993–2002. BMC Public Health, 2006, 6, 278.	2.9	28
221	Sleep-wake disturbances in sporadic Creutzfeldt-Jakob disease. Neurology, 2006, 66, 1418-1424.	1.1	74
222	Strain Fidelity of Chronic Wasting Disease upon Murine Adaptation. Journal of Virology, 2006, 80, 12303-12311.	3.4	74
223	Determinants of diagnostic investigation sensitivities across the clinical spectrum of sporadic Creutzfeldt-Jakob disease. Brain, 2006, 129, 2278-2287.	7.6	283
224	Prion infections, blood and transfusions. Nature Clinical Practice Neurology, 2006, 2, 321-329.	2.5	36
225	Letters to the Editor. Nursing Science Quarterly, 2006, 19, 276-280.	0.8	2
226	Coexistence of multiple PrPSc types in individuals with Creutzfeldt-Jakob disease. Lancet Neurology, The, 2005, 4, 805-814.	10.2	192
227	Sporadic Creutzfeldt?Jakob disease. Journal of Neurology, 2005, 252, 338-342.	3.6	25
228	Genetic prion disease: the EUROCJD experience. Human Genetics, 2005, 118, 166-174.	3.8	391
229	Progressive scoliosis in central core disease. European Spine Journal, 2005, 14, 900-905.	2.2	11
230	Thoracolumbar intradural extramedullary bronchiogenic cyst. Acta Neurochirurgica, 2005, 147, 317-319.	1.7	23
231	Analysis of Prion Strains by PrPSc Profiling in Sporadic Creutzfeldt–Jakob Disease. PLoS Medicine, 2005, 3, e14.	8.4	90
232	No influence of amyloid- \hat{l}^2 -degrading neprilysin activity on prion pathogenesis. Journal of General Virology, 2005, 86, 1861-1867.	2.9	5
233	Mortality from Creutzfeldt–Jakob disease and related disorders in Europe, Australia, and Canada. Neurology, 2005, 64, 1586-1591.	1.1	306
234	Letters to the Editor. Veterinary Pathology, 2005, 42, 107-107.	1.7	2

#	Article	IF	Citations
235	Human Prion Diseases. Archives of Neurology, 2005, 62, 545.	4.5	113
236	Treatment of myositis with etanercept (Enbrel $\hat{A}^{@}$), a recombinant human soluble fusion protein of TNF- $\hat{l}\pm$ type II receptor and IgG1. Rheumatology, 2004, 43, 524-526.	1.9	83
237	Extraneural Pathologic Prion Protein. New England Journal of Medicine, 2004, 350, 732-733.	27.0	4
238	Mimicry of Variant Creutzfeldt-Jakob Disease by Sporadic Creutzfeldt-Jakob Disease: Importance of the Pulvinar Sign. Archives of Neurology, 2004, 61, 445.	4.5	11
239	Predictors of survival in sporadic Creutzfeldt-Jakob disease and other human transmissible spongiform encephalopathies. Brain, 2004, 127, 2348-2359.	7.6	244
240	Intrinsic Resistance of Oligodendrocytes to Prion Infection. Journal of Neuroscience, 2004, 24, 5974-5981.	3.6	46
241	Creutzfeldt–Jakob disease and inclusion body myositis: Abundant diseaseâ€essociated prion protein in muscle. Annals of Neurology, 2004, 55, 121-125.	5.3	47
242	Variant Creutzfeldt–Jakob disease: between lymphoid organs and brain. Trends in Microbiology, 2004, 12, 51-53.	7.7	16
243	Serial EEG findings in sporadic and iatrogenic Creutzfeldt–Jakob disease. Clinical Neurophysiology, 2004, 115, 2467-2478.	1.5	66
244	vCJD tissue distribution and transmission by transfusion—a worst-case scenario coming true?. Lancet, The, 2004, 363, 411-412.	13.7	36
245	Testing for prions: a novel protocol for vCJD prevalence studies. Lancet, The, 2004, 364, 1196-1197.	13.7	3
246	The Peripheral Nervous System and the Pathogenesis of Prion Diseases. Current Molecular Medicine, 2004, 4, 355-359.	1.3	15
247	Prion Diseases. Journal of NeuroVirology, 2003, 9, 183-193.	2.1	66
248	Clinical and radiological mimicry of vCJD in a valine homozygous PrP Sc type 1 sCJD patient. Journal of Neurology, 2003, 250, 491-493.	3.6	17
249	Human prion diseases: epidemiology and integrated risk assessment. Lancet Neurology, The, 2003, 2, 757-763.	10.2	62
250	Positioning of follicular dendritic cells within the spleen controls prion neuroinvasion. Nature, 2003, 425, 957-962.	27.8	195
251	Immunochemical detection of prion protein on dipsticks prepared with crystalline bacterial cell-surface layers. Transfusion, 2003, 43, 1677-1682.	1.6	15
252	Oral Prion Infection Requires Normal Numbers of Peyer's Patches but Not of Enteric Lymphocytes. American Journal of Pathology, 2003, 162, 1103-1111.	3.8	125

#	Article	IF	Citations
253	Immune system and peripheral nerves in propagation of prions to CNS. British Medical Bulletin, 2003, 66, 141-159.	6.9	51
254	Extraneural Pathologic Prion Protein in Sporadic Creutzfeldt–Jakob Disease. New England Journal of Medicine, 2003, 349, 1812-1820.	27.0	299
255	Analysis of the Prion Protein in Primates Reveals a New Polymorphism in Codon 226 (Y226F). Biological Chemistry, 2002, 383, 1021-5.	2.5	10
256	Incidence of Creutzfeldt-Jakob disease in Switzerland. Lancet, The, 2002, 360, 139-141.	13.7	84
257	Unhampered Prion Neuroinvasion despite Impaired Fast Axonal Transport in Transgenic Mice Overexpressing Four-Repeat Tau. Journal of Neuroscience, 2002, 22, 7471-7477.	3.6	61
258	The shifting biology of prions. Brain Research Reviews, 2001, 36, 241-248.	9.0	25
259	Prionsâ€"Role of the Peripheral Nervous System. Virus Research, 2001, 82, 53.	2.2	3
260	Sympathetic Innervation of Lymphoreticular Organs Is Rate Limiting for Prion Neuroinvasion. Neuron, 2001, 31, 25-34.	8.1	223
261	Plasminogen binds to disease-associated prion protein of multiple species. Lancet, The, 2001, 357, 2026-2028.	13.7	79
262	Sympathetic Prions. Scientific World Journal, The, 2001, 1, 555-556.	2.1	5
263	Spongiform encephalopathies: Insights from transgenic models. Advances in Virus Research, 2001, 56, 313-352.	2.1	15
264	Interventional strategies against prion diseases. Nature Reviews Neuroscience, 2001, 2, 745-749.	10.2	76
265	Prions: From Neurografts to Neuroinvasion. , 2001, 59, 129-137.		1
266	Peripheral pathogenesis of prion diseases. Microbes and Infection, 2000, 2, 613-619.	1.9	35
267	Neuroinvasion of Prions: Insights from Mouse Models. Experimental Physiology, 2000, 85, 705-712.	2.0	18
268	Deletions in the spinal muscular atrophy gene region in a newborn with neuropathy and extreme generalized muscular weakness. European Journal of Paediatric Neurology, 2000, 4, 35-38.	1.6	14
269	Neuroinvasion of prions: insights from mouse models. Experimental Physiology, 2000, 85, 705-712.	2.0	2
270	Creutzfeldt-Jakob Disease Presenting as Isolated Dysarthria and Dysphagia due to Pseudobulbar Palsy. European Neurology, 2000, 44, 126-127.	1.4	14

#	Article	IF	CITATIONS
271	Adenoviral and adeno-associated viral transfer of genes to the peripheral nervous system. Proceedings of the National Academy of Sciences of the United States of America, 2000, 97, 442-447.	7.1	120
272	Impaired Prion Replication in Spleens of Mice Lacking Functional Follicular Dendritic Cells. Science, 2000, 288, 1257-1259.	12.6	341
273	Prions: Pathogenesis and Reverse Genetics. Annals of the New York Academy of Sciences, 2000, 920, 140-157.	3.8	15
274	PrPC expression in the peripheral nervous system is a determinant of prion neuroinvasion. Journal of General Virology, 2000, 81, 2813-2821.	2.9	121
275	No Complementation Between TP53 or RBâ€1 and vâ€ <i>src</i> in Astrocytomas of GFAPâ€vâ€ <i>src</i> Transgenic Mice. Brain Pathology, 1999, 9, 627-637.	4.1	15
276	Antimicroglia antibodies in sera of Alzheimer's disease patients. Biological Psychiatry, 1999, 45, 508-511.	1.3	7
277	The Blood-Brain Barrier is Dysregulated in COVID-19 and Serves as a CNS Entry Route for SARS-CoV-2. SSRN Electronic Journal, 0, , .	0.4	3
278	Transmissible Spongiform Encephalopathies., 0,, 1859-1866.		2
279	Young COVID-19 Patients Show a Higher Degree of Microglial Activation When Compared to Controls. Frontiers in Neurology, 0, 13, .	2.4	7
280	Evidence of beta amyloid independent small vessel disease in familial Alzheimer's disease. Brain Pathology, 0, , .	4.1	4