Herbert Budka

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/3321300/publications.pdf

Version: 2024-02-01

		4370	8138
486	29,003	86	148
papers	citations	h-index	g-index

539	539	539	21220
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Coâ€incidental C9orf72 expansion mutationâ€related frontotemporal lobar degeneration pathology and sporadic Creutzfeldtâ"Jakob disease. European Journal of Neurology, 2021, 28, 1009-1015.	1.7	2
2	Histotype-Dependent Oligodendroglial PrP Pathology in Sporadic CJD: A Frequent Feature of the M2C â€œStrainâ€• Viruses, 2021, 13, 1796.	1.5	1
3	Identification of novel risk loci and causal insights for sporadic Creutzfeldt-Jakob disease: a genome-wide association study. Lancet Neurology, The, 2020, 19, 840-848.	4.9	42
4	The autophagic marker p62 highlights Alzheimer type II astrocytes in metabolic/hepatic encephalopathy. Neuropathology, 2020, 40, 358-366.	0.7	4
5	Phenotypic and functional complexity of brain-infiltrating T cells in Rasmussen encephalitis. Neurology: Neuroimmunology and NeuroInflammation, 2018, 5, e419.	3.1	34
6	Distinctive cerebral neuropathology in an adult case of sensory ataxic neuropathy with dysarthria and ophthalmoplegia (SANDO) syndrome. Neuropathology and Applied Neurobiology, 2018, 44, 639-642.	1.8	3
7	K27/G34 versus K28/G35 in histone H3-mutant gliomas: A note of caution. Acta Neuropathologica, 2018, 136, 175-176.	3.9	12
8	Iatrogenic Creutzfeldtâ€"Jakob disease. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2018, 153, 207-218.	1.0	20
9	SAMP8 mice as a neuropathological model of accelerated brain aging and dementia: Toshio Takeda's legacy and future directions. Neuropathology, 2017, 37, 293-305.	0.7	127
10	Multisite Assessment of Aging-Related Tau Astrogliopathy (ARTAG). Journal of Neuropathology and Experimental Neurology, 2017, 76, 605-619.	0.9	38
11	Tau pathology in Creutzfeldtâ€Jakob disease revisited. Brain Pathology, 2017, 27, 332-344.	2.1	61
12	Comorbidity and prognosis in disproportionately enlarged subarachnoid space hydrocephalus (DESH-iNPH): Japanese and Austrian cohort studies. Journal of the Neurological Sciences, 2017, 381, 314.	0.3	0
13	Cystatin F is a biomarker of prion pathogenesis in mice. PLoS ONE, 2017, 12, e0171923.	1.1	20
14	Mechanisms of immune escape in central nervous system infection with neurotropic <scp>JC</scp> virus variant. Annals of Neurology, 2016, 79, 404-418.	2.8	40
15	Neuropathological criteria of anti-IgLON5-related tauopathy. Acta Neuropathologica, 2016, 132, 531-543.	3.9	173
16	Dura mater is a potential source of AÎ² seeds. Acta Neuropathologica, 2016, 131, 911-923.	3.9	85
17	Fatal cerebral haemorrhage in a hypertensive seven-year-old boy. JRSM Open, 2016, 7, 205427041664928.	0.2	0
18	Aging-related tau astrogliopathy (ARTAG): harmonized evaluation strategy. Acta Neuropathologica, 2016, 131, 87-102.	3.9	380

#	Article	IF	Citations
19	Amyloid-Î² pathology and cerebral amyloid angiopathy are frequent in iatrogenic Creutzfeldt-Jakob disease after dural grafting. Swiss Medical Weekly, 2016, 146, w14287.	0.8	89
20	A Genome Wide Association Study Links Glutamate Receptor Pathway to Sporadic Creutzfeldt-Jakob Disease Risk. PLoS ONE, 2015, 10, e0123654.	1.1	28
21	Iatrogenic and sporadic Creutzfeldt-Jakob disease in 2 sisters without mutation in the prion protein gene. Prion, 2015, 9, 444-448.	0.9	4
22	A case of variably protease-sensitive prionopathy treated with doxycyclin. Journal of Neurology, Neurosurgery and Psychiatry, 2015, 86, 816-818.	0.9	23
23	Structure-based drug design identifies polythiophenes as antiprion compounds. Science Translational Medicine, 2015, 7, 299ra123.	5.8	130
24	Reduction in Serum Aquaporin-4 Antibody Titers During Development of a Tumor-Like Brain Lesion in a Patient With Neuromyelitis Optica: A Serum Antibodyâ€“Consuming Effect?. Journal of Neuropathology and Experimental Neurology, 2015, 74, 194-197.	0.9	6
25	The end of the BSE saga: do we still need surveillance for human prion diseases?. Swiss Medical Weekly, 2015, 145, w14212.	0.8	12
26	The Role of the NADPH Oxidase NOX2 in Prion Pathogenesis. PLoS Pathogens, 2014, 10, e1004531.	2.1	57
27	Disproportionate subarachnoid space hydrocephalusâ€"outcome and perivascular space. Annals of Clinical and Translational Neurology, 2014, 1, 562-569.	1.7	18
28	Intracellular processing of disease-associated Î±-synuclein in the human brain suggests prion-like cell-to-cell spread. Neurobiology of Disease, 2014, 69, 76-92.	2.1	110
29	MRI features of Binswanger's disease predict prognosis and associated pathology. Annals of Clinical and Translational Neurology, 2014, 1, 813-821.	1.7	13
30	ISN Information Days at the University of Witwatersrand Medical School, Johannesburg (South) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 30	2.1	0
31	Non-Alzheimer neurodegenerative pathologies and their combinations are more frequent than commonly believed in the elderly brain: a community-based autopsy series. Acta Neuropathologica, 2013, 126, 365-384.	3.9	264
32	Cerebellar dysfunction in a family harboring the PSEN1 mutation co-segregating with a Cathepsin D variant p.A58V. Journal of the Neurological Sciences, 2013, 326, 75-82.	0.3	18
33	Globular glial tauopathies (GGT): consensus recommendations. Acta Neuropathologica, 2013, 126, 537-544.	3.9	168
34	Rapidly progressive dementia with thalamic degeneration and peculiar cortical prion protein immunoreactivity, but absence of proteinase K resistant PrP: a new disease entity?. Acta Neuropathologica Communications, 2013, 1, 72.	2.4	12
35	JC virus granule cell neuronopathy and GCNâ€“IRIS under natalizumab treatment. Annals of Neurology, 2013, 74, 622-626.	2.8	41
36	Expression of myogenic regulatory factors and myo-endothelial remodeling in sporadic inclusion body myositis. Neuromuscular Disorders, 2013, 23, 75-83.	0.3	32

#	Article	IF	Citations
37	Imported human rabies in Switzerland, 2012: A diagnostic conundrum. Journal of Clinical Virology, 2013, 57, 178-181.	1.6	3
38	Neuropathology of the hippocampus in FTLDâ€Tau with Pick bodies: a study of the BrainNet Europe Consortium. Neuropathology and Applied Neurobiology, 2013, 39, 166-178.	1.8	54
39	Intensity of human prion disease surveillance predicts observed disease incidence. Journal of Neurology, Neurosurgery and Psychiatry, 2013, 84, 1372-1377.	0.9	49
40	Prion diseases: an introduction. Food Safety Assurance and Veterinary Public Health, 2013, , 231-254.	0.4	0
41	Gerstmannâ€“StrÃ¤usslerâ€“Scheinker Disease. , 2013, , 327-329.		0
42	The Spectrum of Tau Pathology in Human Prion Disease. , 2013, , 103-119.		6
43	External granular cell layer bobbling: a distinct histomorphological feature of the developing human cerebellum. , 2013, 32, 42-50.		4
44	Stabilization of a Prion Strain of Synthetic Origin Requires Multiple Serial Passages. Journal of Biological Chemistry, 2012, 287, 30205-30214.	1.6	51
45	Intraneuronal Immunoreactivity for the Prion Protein Distinguishes a Subset of E200K Genetic From Sporadic Creutzfeldt-Jakob Disease. Journal of Neuropathology and Experimental Neurology, 2012, 71, 223-232.	0.9	18
46	Antibody 9D5 Recognizes Oligomeric Pyroglutamate Amyloid-Î² in a Fraction of Amyloid-Î² Deposits in Alzheimer's Disease without Cross-Reactivity with other Protein Aggregates. Journal of Alzheimer's Disease, 2012, 29, 361-371.	1.2	17
47	Scientific Opinion on Norovirus (NoV) in oysters: methods, limits and control options. EFSA Journal, 2012, 10, 2500.	0.9	82
48	Scientific Opinion on the minimum hygiene criteria to be applied to clean seawater and on the public health risks and hygiene criteria for bottled seawater intended for domestic use. EFSA Journal, 2012, 10, 2613.	0.9	8
49	Scientific Opinion on the evaluation of new TSE rapid tests submitted in the framework of the Commission Call for expression of interest 2007/S204-247339. EFSA Journal, 2012, 10, 2660.	0.9	3
50	Scientific Opinion on the development of a risk ranking framework on biological hazards. EFSA Journal, 2012, 10, 2724.	0.9	36
51	Scientific Opinion on Reflecting on the experiences and lessons learnt from modelling on biological hazards. EFSA Journal, 2012, 10, 2725.	0.9	6
52	Risk assessment of biological hazards for consumer protection. EFSA Journal, 2012, 10, s1003.	0.9	4
53	Novel crystalloid oligodendrogliopathy in hereditary spastic paraplegia. Acta Neuropathologica, 2012, 124, 583-591.	3.9	8
54	The need to unify neuropathological assessments of vascular alterations in the ageing brain. Experimental Gerontology, 2012, 47, 825-833.	1.2	57

#	Article	IF	Citations
55	A New Mechanism for Transmissible Prion Diseases. Journal of Neuroscience, 2012, 32, 7345-7355.	1.7	72
56	Targeting of prion-infected lymphoid cells to the central nervous system accelerates prion infection. Journal of Neuroinflammation, 2012, 9, 58.	3.1	3
57	Scientific Opinion on Public health risks represented by certain composite products containing food of animal origin. EFSA Journal, 2012, 10, 2662.	0.9	25
58	Scientific Opinion on a review on the European Union Summary Reports on trends and sources zoonoses, zoonotic agents and foodâ€borne outbreaks in 2009 and 2010 â€“ specifically for the data on Salmonella, Campylobacter, verotoxigenic Escherichia coli, Listeria monocytogenes and foodborne outbreaks. EFSA Journal, 2012, 10, 2726.	0.9	10
59	An antibody with high reactivity for disease-associated Î±-synuclein reveals extensive brain pathology. Acta Neuropathologica, 2012, 124, 37-50.	3.9	133
60	Advocacy for Neuropathology in practice: ISN Information Days. Brain Pathology, 2012, 22, i-i.	2.1	0
61	Asymmetry of neurodegenerative disease-related pathologies: a cautionary note. Acta Neuropathologica, 2012, 123, 449-452.	3.9	35
62	Creutzfeldt-Jakob disease with unusually extensive neuropathology in a child treated with native human growth hormone. , 2012, 31, 127-134.		2
63	Prion protein (PrP) deposits in the tectum of experimental Gerstmann-StrÃussler-Scheinker disease following intraocular inoculation. , 2012, 50, 85-8.		8
64	Long-term clinical improvement of progressive multifocal leukoencephalopathy associated with prominent inflammatory response and follicular lymphoma. Leukemia and Lymphoma, 2011, 52, 2190-2192.	0.6	0
65	Identification of common variants influencing risk of the tauopathy progressive supranuclear palsy. Nature Genetics, 2011, 43, 699-705.	9.4	502
66	Scientific Opinion on the evaluation of a new processing method for ABP Category 2 materials of fish origin. EFSA Journal, 2011, 9, 2389.	0.9	1
67	Scientific Opinion on Hatchery Waste as animal byâ€products. EFSA Journal, 2011, 9, 2321.	0.9	3
68	Scientific Opinion on the revision of the quantitative risk assessment (QRA) of the BSE risk posed by processed animal proteins (PAPs). EFSA Journal, 2011, 9, 1947.	0.9	36
69	Scientific Opinion on the efficacy and microbiological safety of irradiation of food. EFSA Journal, 2011, 9, 2103.	0.9	29
70	Prediction of Preadolescent Depressive Symptoms From Child Temperament, Maternal Distress, and Gender: Results of a Prospective, Longitudinal Study. Journal of Developmental and Behavioral Pediatrics, 2011, 32, 18-26.	0.6	20
71	Joint Scientific Opinion on any possible epidemiological or molecular association between TSEs in animals and humans. EFSA Journal, 2011, 9, 1945.	0.9	33
72	Scientific Opinion on a review of the BSEâ€related risk in bovine intestines. EFSA Journal, 2011, 9, 2104.	0.9	1

#	Article	IF	Citations
73	Scientific Opinion on a quantitative estimation of the public health impact of setting a new target for the reduction of Salmonella in broilers. EFSA Journal, 2011, 9, 2106.	0.9	19
74	Scientific Opinion on risk based control of biogenic amine formation in fermented foods. EFSA Journal, 2011, 9, 2393.	0.9	602
75	Editorial: The European Response to BSE: A Success Story. EFSA Journal, 2011, 9, e991.	0.9	5
76	Scientific Opinion on the public health risks of bacterial strains producing extended-spectrum Î²-lactamases and/or AmpC Î²-lactamases in food and food-producing animals. EFSA Journal, 2011, 9, 2322.	0.9	235
77	Scientific Opinion on the maintenance of the list of QPS biological agents intentionally added to food and feed (2011 update). EFSA Journal, 2011, 9, 2497.	0.9	44
78	Scientific Opinion onCampylobacterin broiler meat production: control options and performance objectives and/or targets at different stages of the food chain. EFSA Journal, 2011, 9, 2105.	0.9	379
79	Scientific Opinion on assessment of epidemiological data in relation to the health risks resulting from the presence of parasites in wild caught fish from fishing grounds in the Baltic Sea. EFSA Journal, 2011, 9, 2320.	0.9	4
80	Scientific Opinion on the public health hazards to be covered by inspection of meat (swine). EFSA Journal, 2011, 9, 2351.	0.9	154
81	Post-Salzburg Blues vs. Rise to New Horizons?. Brain Pathology, 2011, 21, i-i.	2.1	0
82	Compassion for our colleagues and friends in Japan. Brain Pathology, 2011, 21, i-i.	2.1	0
83	Advocacy for Neuropathology. Brain Pathology, 2011, 21, iii-iii.	2.1	1
84	New lexicon and criteria for the diagnosis of Alzheimer's disease. Lancet Neurology, The, 2011, 10, 298-299.	4.9	26
85	Genetic Creutzfeldt-Jakob disease associated with the E200K mutation: characterization of a complex proteinopathy. Acta Neuropathologica, 2011, 121, 39-57.	3.9	105
86	A peculiar constellation of tau pathology defines a subset of dementia in the elderly. Acta Neuropathologica, 2011, 122, 205-222.	3.9	80
87	Peroxisomal alterations in Alzheimerâ€™s disease. Acta Neuropathologica, 2011, 122, 271-283.	3.9	176
88	Complex tauopathies vs. tangle predominant dementia. Acta Neuropathologica, 2011, 122, 517-517.	3.9	0
89	Fibulin-5 mutations link inherited neuropathies, age-related macular degeneration and hyperelastic skin. Brain, 2011, 134, 1839-1852.	3.7	64
90	Fatal Prion Disease in a Mouse Model of Genetic E200K Creutzfeldt-Jakob Disease. PLoS Pathogens, 2011, 7, e1002350.	2.1	68

#	Article	IF	Citations
91	Genesis of Mammalian Prions: From Non-infectious Amyloid Fibrils to a Transmissible Prion Disease. PLoS Pathogens, 2011, 7, e1002419.	2.1	98
92	Unclassifiable tauopathy associated with an A152T variation in MAPT exon 7. , 2011, 30, 3-10.		35
93	Multiple intracranial cavernomas with focal amyloid deposition â€“ diagnostic pitfalls. , 2011, 30, 324-7.		0
94	Cathepsin D (C224T) Polymorphism in Sporadic and Genetic Creutzfeldt-Jakob Disease. Alzheimer Disease and Associated Disorders, 2010, 24, 104-107.	0.6	12
95	Statement on technical assistance on the format for applications for new alternative methods for animal byâ€products. EFSA Journal, 2010, 8, .	0.9	4
96	Scientific Opinion on Quantification of the risk posed by broiler meat to human campylobacteriosis in the EU. EFSA Journal, 2010, 8, 1437.	0.9	181
97	Scientific Opinion on risk assessment of parasites in fishery products. EFSA Journal, 2010, 8, 1543.	0.9	214
98	Scientific Opinion on the link between Salmonella criteria at different stages of the poultry production chain. EFSA Journal, 2010, 8, 1545.	0.9	7
99	Scientific Opinion on a quantitative estimation of the public health impact of setting a new target for the reduction of Salmonella in laying hens. EFSA Journal, 2010, 8, 1546.	0.9	34
100	Scientific Opinion on a Quantitative Microbiological Risk Assessment of Salmonella in slaughter and breeder pigs. EFSA Journal, 2010, 8, 1547.	0.9	68
101	Scientific Opinion on Analytical sensitivity of approved TSE rapid tests â€“ new data for assessment of two rapid tests. EFSA Journal, 2010, 8, 1591.	0.9	3
102	Scientific Opinion on Q fever. EFSA Journal, 2010, 8, 1595.	0.9	129
103	Statement on Food safety considerations of novel H1N1 influenza virus infections in humans. EFSA Journal, 2010, 8, 1629.	0.9	4
104	Scientific Opinion on Lime Treatment of Solid Pig and Poultry Manure. EFSA Journal, 2010, 8, 1681.	0.9	4
105	Scientific Opinion on the Neste Oil Application for a new alternative method of disposal or use of Animal By-Products. EFSA Journal, 2010, 8, 1825.	0.9	3
106	Scientific Opinion on monitoring and assessment of the public health risk of â€œSalmonellaTyphimurium-likeâ€•strains. EFSA Journal, 2010, 8, 1826.	0.9	122
107	Scientific Opinion on the safety and efficacy of using recycled hot water as a decontamination technique for meat carcasses. EFSA Journal, 2010, 8, 1827.	0.9	8
108	Scientific Opinion on the results of the EU survey for Chronic Wasting Disease (CWD) in cervids. EFSA Journal, 2010, 8, 1861.	0.9	16

#	Article	IF	Citations
109	Scientific Opinion on Fish Oil for Human Consumption. Food Hygiene, including Rancidity. EFSA Journal, 2010, 8, 1874.	0.9	62
110	Scientific Opinion on a second update on the risk for human and animal health related to the revision of the BSE monitoring regime in some Member States. EFSA Journal, 2010, 8, 1946.	0.9	6
111	Recombinant prion protein induces a new transmissible prion disease in wild-type animals. Acta Neuropathologica, 2010, 119, 177-187.	3.9	256
112	Status of ICN 2010. Acta Neuropathologica, 2010, 119, 381-382.	3.9	0
113	Protein coding of neurodegenerative dementias: the neuropathological basis of biomarker diagnostics. Acta Neuropathologica, 2010, 119, 389-408.	3.9	98
114	Welcome to ICN 2010. Acta Neuropathologica, 2010, 120, 275-276.	3.9	0
115	Distribution of apoptosis-related proteins in sporadic Creutzfeldtâ€“Jakob disease. Brain Research, 2010, 1323, 192-199.	1.1	17
116	Tubulin polymerization promoting protein (TPPP/p25) as a marker for oligodendroglial changes in multiple sclerosis. Glia, 2010, 58, 1847-1857.	2.5	61
117	Viral meningoencephalitis: a review of diagnostic methods and guidelines for management. European Journal of Neurology, 2010, 17, 999.	1.7	176
118	Scientific Opinion on BSE/TSE infectivity in small ruminant tissues. EFSA Journal, 2010, 8, 1875.	0.9	15
119	Scientific Opinion on the maintenance of the list of QPS biological agents intentionally added to food and feed (2010 update). EFSA Journal, 2010, 8, 1944.	0.9	55
120	Ultrastructural Characteristics (or Evaluation) of Creutzfeldt-Jakob Disease and Other Human Transmissible Spongiform Encephalopathies or Prion Diseases. Ultrastructural Pathology, 2010, 34, 351-361.	0.4	30
121	Peroxisomal Localization of the Proopiomelanocortin-Derived Peptides Î²-Lipotropin and Î²-Endorphin. Endocrinology, 2010, 151, 4801-4810.	1.4	9
122	Increased neuronal Rab5 immunoreactive endosomes do not colocalize with TDP-43 in motor neuron disease. Experimental Neurology, 2010, 225, 133-139.	2.0	10
123	Current concepts of neuropathological diagnostics in practice: neurodegenerative diseases. , 2010, 29, 271-288.		68
124	Molecular Pathology of Human Prion Diseases. International Journal of Molecular Sciences, 2009, 10, 976-999.	1.8	73
125	Genetic Creutzfeldt-Jakob disease mimicking variant Creutzfeldt-Jakob disease. Journal of Neurology, Neurosurgery and Psychiatry, 2009, 80, 1410-1411.	0.9	5
126	TARDBP variation associated with frontotemporal dementia, supranuclear gaze palsy, and chorea. Movement Disorders, 2009, 24, 1842-1847.	2.2	182

#	Article	IF	Citations
127	No evidence for cognitive dysfunction or depression in patients with mild restless legs syndrome. Movement Disorders, 2009, 24, 1843-1847.	2.2	64
128	Ultrastructural study of florid plaques in variant Creutzfeldtâ€“Jakob disease: a comparison with amyloid plaques in kuru, sporadic Creutzfeldtâ€“Jakob disease and Gerstmannâ€“StrÃ¤uslerâ€“Scheinker disease. Neuropathology and Applied Neurobiology, 2009, 35, 46-59.	1.8	62
129	In vivo expression of proinflammatory cytokines in HIV encephalitis: an analysis of 11 autopsy cases. Neuropathology, 2009, 29, 433-442.	0.7	46
130	Interâ€Laboratory Assessment of PrP^{Sc} Typing in Creutzfeldtâ€“Jakob Disease: A Western Blot Study within the NeuroPrion Consortium. Brain Pathology, 2009, 19, 384-391.	2.1	50
131	Danon disease: Case report and detection of new mutation. Journal of Inherited Metabolic Disease, 2009, 32, 115-122.	1.7	23
132	Reduced Expression of Excitatory Amino Acid Transporter 2 and Diffuse Microglial Activation in the Cerebral Cortex in AIDS Cases With or Without HIV Encephalitis. Journal of Neuropathology and Experimental Neurology, 2009, 68, 199-209.	0.9	27
133	Protein-Based Neuropathology and Molecular Classification of Human Neurodegenerative Diseases. Focus on Structural Biology, 2009, , 251-272.	0.1	5
134	Inter-laboratory comparison of neuropathological assessments of Î²-amyloid protein: a study of the BrainNet Europe consortium. Acta Neuropathologica, 2008, 115, 533-546.	3.9	86
135	Management of a twenty-first century brain bank: experience in the BrainNet Europe consortium. Acta Neuropathologica, 2008, 115, 497-507.	3.9	101
136	Intrathecal anti-Î±B-crystallin IgG antibody responses: Potential inflammatory markers in Guillain-BarrÃ© syndrome. Journal of Neurology, 2008, 255, 917-924.	1.8	17
137	Nigral burden of Î±â€synuclein correlates with striatal dopamine deficit. Movement Disorders, 2008, 23, 1608-1612.	2.2	53
138	The role of parvalbumin and calbindin D28k in experimental scrapie. Neuropathology and Applied Neurobiology, 2008, 34, 435-445.	1.8	11
139	Neocortical neurones may be targeted by immune attack in antiâ€Yo paraneoplastic syndrome. Neuropathology and Applied Neurobiology, 2008, 34, 248-252.	1.8	21
140	Franz Seitelberger (4 December 1916 to 2 November 2007). Neuropathology and Applied Neurobiology, 2008, 34, 128-128.	1.8	0
141	Tubulovesicular structures are a consistent (and unexplained) finding in the brains of humans with prion diseases. Virus Research, 2008, 132, 226-228.	1.1	10
142	BSE and TSEs: Past, present and future. Trends in Food Science and Technology, 2008, 19, S34-S39.	7.8	8
143	Prion Diseases: From Protein to Cell Pathology. American Journal of Pathology, 2008, 172, 555-565.	1.9	126
144	Mixed Brain Pathologies in Dementia: The BrainNet Europe Consortium Experience. Dementia and Geriatric Cognitive Disorders, 2008, 26, 343-350.	0.7	148

#	Article	IF	Citations
145	Creutzfeldt-Jakob Disease in Austria: An Autopsy-Controlled Study. Neuroepidemiology, 2008, 30, 215-221.	1.1	31
146	Iatrogenic Creutzfeldt Jakob disease 22 years after human growth hormone therapy: clinical and radiological features. Journal of Neurology, Neurosurgery and Psychiatry, 2008, 79, 229-231.	0.9	13
147	A quantitative microbiological risk assessment on Salmonella in meat : Source attribution for human salmonellosis from meat - Scientific Opinion of the Panel on Biological Hazards. EFSA Journal, 2008, 6, 625.	0.9	8
148	Assessment of the possible effect of the four antimicrobial treatment substances on the emergence of antimicrobial resistance - Scientific Opinion of the Panel on Biological Hazards. EFSA Journal, 2008, 6, 659.	0.9	10
149	Further consideration of ageâ€related parameters on the Risk for Human and Animal Health related to the revision of the BSE Monitoring regime in some Member States. EFSA Journal, 2008, 6, 763.	0.9	0
150	Overview of methods for source attribution for human illness from foodâ€borne microbiological hazards â€•Scientific Opinion of the Panel on Biological Hazards. EFSA Journal, 2008, 6, 764.	0.9	8
151	The maintenance of the list of QPS microorganisms intentionally added to food or feed - Scientific Opinion of the Panel on Biological Hazards. EFSA Journal, 2008, 6, 923.	0.9	45
152	Assessment of Î±-Synuclein Pathology: A Study of the BrainNet Europe Consortium. Journal of Neuropathology and Experimental Neurology, 2008, 67, 125-143.	0.9	73
153	Endothelial and Myogenic Differentiation of Hematopoietic Progenitor Cells in Inflammatory Myopathies. Journal of Neuropathology and Experimental Neurology, 2008, 67, 711-719.	0.9	23
154	Accumulation of HtrA2/Omi in Neuronal and Glial Inclusions in Brains With Î±-Synucleinopathies. Journal of Neuropathology and Experimental Neurology, 2008, 67, 984-993.	0.9	44
155	White Matter Tauopathy With Globular Glial Inclusions: A Distinct Sporadic Frontotemporal Lobar Degeneration. Journal of Neuropathology and Experimental Neurology, 2008, 67, 963-975.	0.9	111
156	Excretion of Transmissible Spongiform Encephalopathy Infectivity in Urine. Emerging Infectious Diseases, 2008, 14, 1406-1412.	2.0	46
157	Scientific and technical clarification in the interpretation and consideration of some facets of the conclusions of its Opinion of 8 March 2007 on certain aspects related to the risk of Transmissible Spongiform Encephalopathies (TSEs) in ovine and caprine animals â€•Scientific Report of the Scientific Panel on Biological Hazards. EFSA Journal, 2008, 6, 626.	0.9	0
158	Risk for Human and Animal Health related to the revision of the BSE Monitoring regime in some Member States â€•Scientific Opinion. EFSA Journal, 2008, 6, 762.	0.9	0
159	Food Safety considerations of animal welfare aspects of husbandry systems for farmed fish - Scientific opinion of the Panel on Biological Hazards. EFSA Journal, 2008, 6, 867.	0.9	1
160	Fulminant central nervous system demyelination associated with interferon-Î± therapy and hepatitis C virus infection. Multiple Sclerosis Journal, 2007, 13, 1100-1106.	1.4	21
161	Fatal Neurological Disease in Scrapie-Infected Mice Induced for Experimental Autoimmune Encephalomyelitis. Journal of Virology, 2007, 81, 9942-9949.	1.5	15
162	Surveillance and monitoring of Toxoplasma in humans, food and animals - Scientific Opinion of the Panel on Biological Hazards. EFSA Journal, 2007, 5, 583.	0.9	13

#	Article	IF	Citations
163	Monitoring and identification of human enteropathogenic Yersinia spp. - Scientific Opinion of the Panel on Biological Hazards. EFSA Journal, 2007, 5, 595.	0.9	13
164	Involvement of the Endosomal-Lysosomal System Correlates With Regional Pathology in Creutzfeldt-Jakob Disease. Journal of Neuropathology and Experimental Neurology, 2007, 66, 628-636.	0.9	68
165	Brain Protein Preservation Largely Depends on the Postmortem Storage Temperature. Journal of Neuropathology and Experimental Neurology, 2007, 66, 35-46.	0.9	151
166	Verbal perseveration as the initial symptom in a case of Creutzfeldtâ€Jakob disease. Aphasiology, 2007, 21, 1079-1113.	1.4	2
167	Tickborne Encephalitis in Naturally Exposed Monkey (Macaca sylvanus). Emerging Infectious Diseases, 2007, 13, 905-907.	2.0	40
168	Opinion of the Scientific Panel on biological hazards (BIOHAZ) on the assessment of the health risks of feeding of ruminants with fishmeal in relation to the risk of TSE. EFSA Journal, 2007, 5, 443.	0.9	3
169	Immunohistochemical detection of class III ?-tubulin in primary brain tumours: variable expression in most tumour types limits utility as a differential diagnostic marker. Histopathology, 2007, 50, 949-952.	1.6	11
170	Effects of Formalin Fixation, Paraffin Embedding, and Time of Storage on DNA Preservation in Brain Tissue: A BrainNet Europe Study. Brain Pathology, 2007, 17, 297-303.	2.1	127
171	Bilateral striopallidodentate calcification (Fahr's syndrome) and multiple system atrophy in a patient with longstanding hypoparathyroidism. Neuropathology, 2007, 27, 453-456.	0.7	25
172	Papillary glioneuronal tumor. Neuropathology, 2007, 27, 468-473.	0.7	33
173	OLIG2 is a useful immunohistochemical marker in differential diagnosis of clear cell primary CNS neoplasms. Histopathology, 2007, 50, 365-370.	1.6	44
174	Synucleinopathy with features of both multiple system atrophy and dementia with Lewy bodies. Neuropathology and Applied Neurobiology, 2007, 33, 126-9.	1.8	17
175	Malignant predominantly minigemistocytic glioma in two infants: a distinctive glioma variant?. Neuropathology and Applied Neurobiology, 2007, 33, 169-178.	1.8	3
176	Secretagogin expression in tumours of the human brain and its coverings. Apmis, 2007, 115, 319-326.	0.9	7
177	Accumulation of Hsc70 and Hsp70 in glial cytoplasmic inclusions in patients with multiple system atrophy. Brain Research, 2007, 1136, 219-227.	1.1	34
178	Distribution and cellular localization of adrenoleukodystrophy protein in human tissues: Implications for X-linked adrenoleukodystrophy. Neurobiology of Disease, 2007, 28, 165-174.	2.1	47
179	How a neuropsychiatric brain bank should be run: a consensus paper of Brainnet Europe II. Journal of Neural Transmission, 2007, 114, 527-537.	1.4	49
180	The brain-specific protein TPPP/p25 in pathological protein deposits of neurodegenerative diseases. Acta Neuropathologica, 2007, 113, 153-161.	3.9	65

#	Article	IF	Citations
181	TPPP/p25 in brain tumours: expression in non-neoplastic oligodendrocytes but not in oligodendroglioma cells. Acta Neuropathologica, 2007, 113, 213-215.	3.9	28
182	Autopsy at 2 months after death: Brain is satisfactorily preserved for neuropathology. Forensic Science International, 2007, 168, 177-182.	1.3	10
183	Opinion of the Scientific Panel on biological hazards (BIOHAZ) on the request for review of the opinion on microbiological risks in infant formulae and follow-on formulae with regard to Enterobacteriaceae as indicators. EFSA Journal, 2007, 5, 444.	0.9	8
184	Histopathological prognostic factors in medulloblastoma: High expression of survivin is related to unfavourable outcome. European Journal of Cancer, 2006, 42, 2996-3003.	1.3	35
185	Opinion of the Scientific Panel on Biological Hazards (BIOHAZ) and of the Scientific Panel on Animal Health and Welfare (AHAW) on â€œReview of the Community Summary Report on Trends and Sources of Zoonoses, Zoonotic Agents and Antimicrobial Resistance in th. EFSA Journal, 2006, 4, 403.	0.9	5
186	Opinion of the Scientific Panel on Animal Health and Welfare (AHAW) on a request from the Commission related with the welfare aspects of the main systems of stunning and killing applied to commercially farmed deer, goats, rabbits, ostriches, ducks, geese. EFSA Journal, 2006, 4, 326.	0.9	12
187	Opinion of the Scientific Panel on biological hazards (BIOHAZ) related to the public health risks of feeding farmed animals with readyâ€toâ€use dairy products without further treatment. EFSA Journal, 2006, 4, 340.	0.9	0
188	Opinion of the Scientific Panel on biological hazards (BIOHAZ) on the evaluation of the efficacy of L (+) Lactic acid for carcass decontamination. EFSA Journal, 2006, 4, 342.	0.9	3
189	Opinion of the Scientific Panel on biological hazards (BIOHAZ) related to the evaluation of the efficacy of SANâ€PELÂ® for use as an antimicrobial substance applied on carcasses of chickens, turkeys, quails, pigs, beef, sheep, goats and game and in washing the shells of eggs. EFSA Journal, 2006, 4, 352.	0.9	0
190	Opinion of the Scientific Panel on biological hazards (BIOHAZ) on the Breeding programme for TSE resistance in sheep. EFSA Journal, 2006, 4, 382.	0.9	2
191	Guidance document on the submission of data for the evaluation of the safety and the efficacy of substances for the removal of microbial surface contamination of foods of animal origin from the Scientific Panels AFC/BIOHAZ. EFSA Journal, 2006, 4, 388.	0.9	0
192	Interlaboratory Comparison of Assessments of Alzheimer Disease-Related Lesions: A Study of the BrainNet Europe Consortium. Journal of Neuropathology and Experimental Neurology, 2006, 65, 740-757.	0.9	95
193	Endoplasmic Reticulum Stress Features Are Prominent in Alzheimer Disease but Not in Prion Diseases In Vivo. Journal of Neuropathology and Experimental Neurology, 2006, 65, 348-357.	0.9	196
194	Immunohistochemical Expression of Prion Protein (PrPC) in the Human Forebrain During Development. Journal of Neuropathology and Experimental Neurology, 2006, 65, 698-706.	0.9	40
195	Immunohistochemical Analysis of INI1 Protein in Malignant Pediatric CNS Tumors: Lack of INI1 in Atypical Teratoid/Rhabdoid Tumors and in a Fraction of Primitive Neuroectodermal Tumors without Rhabdoid Phenotype. American Journal of Surgical Pathology, 2006, 30, 1462-1468.	2.1	166
196	Prominent corticosteroid disturbance in experimental prion disease. European Journal of Neuroscience, 2006, 23, 2723-2730.	1.2	23
197	Introduction. Brain Pathology, 2006, 8, 553-553.	2.1	3
198	The Austrian FFI Cases. Brain Pathology, 2006, 8, 554-554.	2.1	3

#	Article	IF	Citations
199	Comparative analysis of NeuN immunoreactivity in primary brain tumours: conclusions for rational use in diagnostic histopathology. Histopathology, 2006, 48, 438-444.	1.6	62
200	Inflammatory response in human tick-borne encephalitis: analysis of postmortem brain tissue. Journal of NeuroVirology, 2006, 12, 322-327.	1.0	121
201	Immunohistochemical Analysis of Platelet-derived Growth Factor Receptor-Î±, -Î², c-kit, c-abl, and Arg Proteins in Glioblastoma: Possible Implications for Patient Selection for Imatinib Mesylate Therapy. Journal of Neuro-Oncology, 2006, 76, 105-109.	1.4	47
202	Human transmissible spongiform encephalopathies in eleven countries: diagnostic pattern across time, 1993â€“2002. BMC Public Health, 2006, 6, 278.	1.2	28
203	Acute motor and sensory axonal neuropathy in Burkitt-like lymphoma. Muscle and Nerve, 2006, 34, 494-498.	1.0	22
204	Upregulated Expression of 14-3-3 Proteins in Astrocytes From Human Cerebrovascular Ischemic Lesions. Stroke, 2006, 37, 830-835.	1.0	36
205	Phenotypic heterogeneity in inherited prion disease (P102L) is associated with differential propagation of protease-resistant wild-type and mutant prion protein. Brain, 2006, 129, 1557-1569.	3.7	91
206	Determinants of diagnostic investigation sensitivities across the clinical spectrum of sporadic Creutzfeldt-Jakob disease. Brain, 2006, 129, 2278-2287.	3.7	283
207	UPDATE ON TRANSMISSIBLE SPONGIFORM ENCEPHALOPATHIES. , 2006, , .		0
208	Central Pathogenesis of Prion Diseases. , 2005, , 49-82.		0
209	Opinion of the Scientific Panel on biological hazards (BIOHAZ) on the risk assessment of a revised inspection of slaughter animals in areas with low prevalence of Cysticercus. EFSA Journal, 2005, 3, 176.	0.9	4
210	Opinion of the Scientific Panel on biological hazards (BIOHAZ) on the on the â€œAssessment of the human and animal BSE risk posed by tallow with respect to residual BSE riskâ€• EFSA Journal, 2005, 3, 221.	0.9	4
211	Opinion of the Scientific Panel on biological hazards (BIOHAZ) on the â€œQuantitative risk assessment of the animal BSE risk posed by meat and bone meal with respect to the residual BSE riskâ€• EFSA Journal, 2005, 3, 257.	0.9	2
212	Opinion of the Scientific Panel on biological hazards (BIOHAZ) on the biological safety of heat treatment of manure. EFSA Journal, 2005, 3, 265.	0.9	2
213	Opinion of the Scientific Panel on biological hazards (BIOHAZ) on the â€œRequest for an opinion on the feasibility of establishing Trichinella-free areas, and if feasible on the risk increase to public health of not examining pigs from those areas for Trich. EFSA Journal, 2005, 3, 277.	0.9	3
214	Imaging ex vivo healthy and pathological human brain tissue with ultra-high-resolution optical coherence tomography. Journal of Biomedical Optics, 2005, 10, 011006.	1.4	82
215	Viral encephalitis: a review of diagnostic methods and guidelines for management. European Journal of Neurology, 2005, 12, 331-343.	1.7	256
216	Pleomorphic xanthoastrocytoma with anaplastic features presenting without GFAP immunoreactivity: Implications for differential diagnosis. Neuropathology, 2005, 25, 241-246.	0.7	23

#	Article	IF	Citations
217	Vascularization and expression of hypoxia-related tissue factors in intracranial ependymoma and their impact on patient survival. Acta Neuropathologica, 2005, 109, 211-216.	3.9	34
218	Pathogenesis of prion diseases. Acta Neuropathologica, 2005, 109, 32-48.	3.9	51
219	No prognostic impact of survivin expression in glioblastoma. Acta Neuropathologica, 2005, 109, 534-538.	3.9	26
220	Redox metals and oxidative abnormalities in human prion diseases. Acta Neuropathologica, 2005, 110, 232-238.	3.9	52
221	Prion disease with a 144 base pair insertion: unusual cerebellar prion protein immunoreactivity. Acta Neuropathologica, 2005, 110, 513-519.	3.9	23
222	Genetic prion disease: the EUROCJD experience. Human Genetics, 2005, 118, 166-174.	1.8	391
223	Opinion of the Scientific Panel on biological hazards (BIOHAZ) on vis-Ã-vis biological risks of biogas and compost treatment standards of animal by-products (ABP). EFSA Journal, 2005, 3, 264.	0.9	3
224	Survivin Expression in Intracranial Ependymomas and Its Correlation With Tumor Cell Proliferation and Patient Outcome. American Journal of Clinical Pathology, 2005, 124, 543-549.	0.4	32
225	Mortality from Creutzfeldt-Jakob disease and related disorders in Europe, Australia, and Canada. Neurology, 2005, 64, 1586-1591.	1.5	306
226	Megadolichobasilar anomaly with thrombosis in a family with Fabry's disease and a novel mutation in the Î±-galactosidase A gene. Brain, 2005, 128, 2078-2083.	3.7	47
227	Visualization of Central European Tick-Borne Encephalitis Infection in Fatal Human Cases. Journal of Neuropathology and Experimental Neurology, 2005, 64, 506-512.	0.9	164
228	Alzheimer-type neuropathology in a 28 year old patient with iatrogenic Creutzfeldt-Jakob disease after dural grafting. Journal of Neurology, Neurosurgery and Psychiatry, 2005, 77, 413-416.	0.9	42
229	The ubiquitinâ€“proteasome system in Creutzfeldtâ€“Jakob and Alzheimer disease: Intracellular redistribution of components correlates with neuronal vulnerability. Neurobiology of Disease, 2005, 19, 427-435.	2.1	20
230	Subcellular Localization of Disease-Associated Prion Protein in the Human Brain. American Journal of Pathology, 2005, 166, 287-294.	1.9	77
231	DEC1 expression in 1p-aberrant oligodendroglial neoplasms. Histology and Histopathology, 2005, 20, 1173-7.	0.5	12
232	UPDATE ON THE PATHOGENESIS OF TRANSMISSIBLE SPONGIFORM ENCEPHALOPATHIES. , 2005, , .		0
233	Rationale for diagnosing human prion disease. World Journal of Biological Psychiatry, 2004, 5, 83-91.	1.3	10
234	Predictors of survival in sporadic Creutzfeldt-Jakob disease and other human transmissible spongiform encephalopathies. Brain, 2004, 127, 2348-2359.	3.7	244

#	Article	IF	Citations
235	Standards for the assay of Creutzfeldtâ€“Jakob disease specimens. Journal of General Virology, 2004, 85, 1777-1784.	1.3	46
236	Meeting report. Neuropathology and Applied Neurobiology, 2004, 30, 308-310.	1.8	1
237	Presence of D110 antigen expressing immunocompetent cells in glioblastoma associates with prolonged survival. Neuropathology and Applied Neurobiology, 2004, 30, 608-614.	1.8	2
238	Antigenic profile of human recombinant PrP: generation and characterization of a versatile polyclonal antiserum. Journal of Neuroimmunology, 2004, 146, 22-32.	1.1	13
239	Increased 14-3-3 immunoreactivity in glial elements in patients with multiple sclerosis. Acta Neuropathologica, 2004, 107, 137-143.	3.9	19
240	Topographical and cytopathological lesion analysis of the white matter in Binswanger?s disease brains. Acta Neuropathologica, 2004, 107, 563-570.	3.9	21
241	Value and limits of immunohistochemistry in differential diagnosis of clear cell primary brain tumors. Acta Neuropathologica, 2004, 108, 24-30.	3.9	38
242	Enhanced expression of 14-3-3 proteins in reactive astrocytes in Creutzfeldt-Jakob disease brains. Acta Neuropathologica, 2004, 108, 302-308.	3.9	10
243	14-3-3 proteins in Lewy body-like hyaline inclusions in patients with sporadic amyotrophic lateral sclerosis. Acta Neuropathologica, 2004, 108, 531-537.	3.9	21
244	Concern about Mad Cow Disease: End of the beginning, or beginning of the end?. Wiener Klinische Wochenschrift, 2004, 116, 505-507.	1.0	0
245	The prion protein in human neuromuscular diseases. Journal of Pathology, 2004, 204, 241-247.	2.1	19
246	Creutzfeldt-Jakob disease and inclusion body myositis: Abundant disease-associated prion protein in muscle. Annals of Neurology, 2004, 55, 121-125.	2.8	47
247	Advances in broad bandwidth light sources for ultrahigh resolution optical coherence tomography. Physics in Medicine and Biology, 2004, 49, 1235-1246.	1.6	104
248	Neuropathology of white matter disease in Leber's hereditary optic neuropathy. Brain, 2004, 128, 35-41.	3.7	96
249	Complement activation in human prion disease. Neurobiology of Disease, 2004, 15, 21-28.	2.1	68
250	Natively unfolded tubulin polymerization promoting protein TPPP/p25 is a common marker of alpha-synucleinopathies. Neurobiology of Disease, 2004, 17, 155-162.	2.1	140
251	Ki-67 Immunolabeling Index Is an Accurate Predictor of Outcome in Patients With Intracranial Ependymoma. American Journal of Surgical Pathology, 2004, 28, 914-920.	2.1	78
252	Contribution of neuropathology to the understanding of human prion disease. Folia Neuropathologica, 2004, 42 Suppl A, 69-76.	0.5	3

#	Article	IF	Citations
253	Echigo-1: a panencephalopathic strain of creutzfeldt-jakob disease. I. neuropathological and immunohistochemical studies. Folia Neuropathologica, 2004, 42 Suppl B, 161-6.	0.5	0
254	Echigo-1: a panencephalopathic strain of Creutzfeldt-Jakob disease. II. Ultrastructural studies in hamsters. Folia Neuropathologica, 2004, 42 Suppl B, 167-75.	0.5	1
255	Severe depletion of mitochondrial DNA in spinal muscular atrophy. Acta Neuropathologica, 2003, 105, 245-251.	3.9	72
256	Distinctive cerebellar immunoreactivity for the prion protein in familial (E200K) Creutzfeldt-Jakob disease. Acta Neuropathologica, 2003, 105, 449-454.	3.9	25
257	Rosette-forming glioneuronal tumor of the fourth ventricle. Acta Neuropathologica, 2003, 106, 506-508.	3.9	63
258	Immunostaining for ubiquitin: efficient pretreatment. Neuropathology and Applied Neurobiology, 2003, 29, 174-177.	1.8	3
259	Fluorescent In Situ Hybridization on Isolated Tumor Cell Nuclei: A Sensitive Method for 1p and 19q Deletion Analysis in Paraffin-Embedded Oligodendroglial Tumor Specimens. Modern Pathology, 2003, 16, 708-715.	2.9	53
260	D18G Transthyretin Is Monomeric, Aggregation Prone, and Not Detectable in Plasma and Cerebrospinal Fluid:Â A Prescription for Central Nervous System Amyloidosis?â€. Biochemistry, 2003, 42, 6656-6663.	1.2	117
261	Neuropathology of prion diseases. British Medical Bulletin, 2003, 66, 121-130.	2.7	136
262	Distinct time pattern of complement activation and cytotoxic T cell response in Guillain-Barre syndrome. Brain, 2003, 126, 2034-2042.	3.7	94
263	Molecular classification of sporadic Creutzfeldtâ€“Jakob disease. Brain, 2003, 126, 1333-1346.	3.7	301
264	Nucleus-specific alteration of raphe neurons in human neurodegenerative disorders. NeuroReport, 2003, 14, 73-76.	0.6	69
265	Imaging brain morphology with ultrahigh-resolution optical coherence tomography. , 2003, , .		2
266	Vascular Patterns in Glioblastoma Influence Clinical Outcome and Associate with Variable Expression of Angiogenic Proteins: Evidence for Distinct Angiogenic Subtypes. Brain Pathology, 2003, 13, 133-143.	2.1	132
267	The cerebral cortex in Fetal Down Syndrome. Journal of Neural Transmission Supplementum, 2003, , 159-163.	0.5	8
268	14-3-3 Proteins in Lewy Bodies in Parkinson Disease and Diffuse Lewy Body Disease Brains. Journal of Neuropathology and Experimental Neurology, 2002, 61, 245-253.	0.9	145
269	Disease-Associated Prion Protein in Vessel Walls. American Journal of Pathology, 2002, 161, 1979-1984.	1.9	47
270	Oxidative Damage to Nucleic Acids in Human Prion Disease. Neurobiology of Disease, 2002, 9, 275-281.	2.1	68

#	Article	IF	Citations
271	The prion protein in human neurodegenerative disorders. Neuroscience Letters, 2002, 329, 269-272.	1.0	44
272	DNA topoisomerase IIÎ± expression in optic pathway gliomas of childhood. European Journal of Cancer, 2002, 38, 393-400.	1.3	17
273	High expression of DNA topoisomerase IIÎ± and Ki-67 antigen is associated with prolonged survival in glioblastoma patients. European Journal of Cancer, 2002, 38, 1343-1347.	1.3	34
274	Aging, the brain and human prion disease. Experimental Gerontology, 2002, 37, 603-605.	1.2	12
275	Spontaneous mutations in the prion protein gene causing transmissible spongiform encephalopathy. Annals of Neurology, 2002, 52, 355-359.	2.8	37
276	Accumulation of 14-3-3 proteins in glial cytoplasmic inclusions in multiple system atrophy. Annals of Neurology, 2002, 52, 722-731.	2.8	53
277	Author reply to previously published correspondence. Cancer, 2002, 94, 2316-2317.	2.0	0
278	BSE and variant Creutzfeldt-Jakob disease: never say never. Acta Neuropathologica, 2002, 103, 627-628.	3.9	5
279	Distribution of intraneuronal immunoreactivity for the prion protein in human prion diseases. Acta Neuropathologica, 2002, 104, 320-326.	3.9	19
280	A comparative study on the expression of cyclooxygenase and 5-lipoxygenase during cerebral ischemia in humans. Acta Neuropathologica, 2002, 104, 601-607.	3.9	84
281	Mutations of the Prion Protein Gene. Journal of Neurology, 2002, 249, 1567-1582.	1.8	251
282	Fibrous meningeal tumours with extensive non-calcifying collagenous whorls and glial fibrillary acidic protein expression: the whorling-sclerosing variant of meningioma. Neuropathology and Applied Neurobiology, 2002, 28, 42-47.	1.8	30
283	Proliferative activity as measured by MIB-1 labeling index and long-term outcome of cerebellar juvenile pilocytic astrocytomas. Journal of Neuro-Oncology, 2002, 58, 141-146.	1.4	17
284	Immunohistochemistry for the Prion Protein: Comparison of Different Monoclonal Antibodies in Human Prion Disease Subtypes. Brain Pathology, 2002, 12, 1-11.	2.1	96
285	Prognostic relevance of p53 protein expression in glioblastoma. Oncology Reports, 2002, 9, 703-7.	1.2	23
286	Prominent Stress Response of Purkinje Cells in Creutzfeldtâ€“Jakob Disease. Neurobiology of Disease, 2001, 8, 881-889.	2.1	29
287	Immunohistochemical Detection of Cell Growth Fraction in Formalin-Fixed and Paraffin-Embedded Murine Tissue. American Journal of Pathology, 2001, 158, 1991-1996.	1.9	42
288	Dysembryoplastic Neuroectodermal Tumor: An Ultrastructural Study of Six Cases. Ultrastructural Pathology, 2001, 25, 455-467.	0.4	7

#	Article	IF	Citations
289	Serotonergic nuclei of the raphe are not affected in human ageing. NeuroReport, 2001, 12, 669-671.	0.6	18
290	Fibroblasts Can Express Glial Fibrillary Acidic protein (GFAP) In Vivo. Journal of Neuropathology and Experimental Neurology, 2001, 60, 449-461.	0.9	65
291	Autoradiography with [3H]PK11195 of spinal tract degeneration in amyotrophic lateral sclerosis. Acta Neuropathologica, 2001, 101, 75-78.	3.9	26
292	Marked increase of neuronal prion protein immunoreactivity in Alzheimer's disease and human prion diseases. Acta Neuropathologica, 2001, 101, 417-423.	3.9	70
293	Expression of hypoxia-inducible factor-1Î± in oligodendrogliomas. Cancer, 2001, 92, 165-171.	2.0	171
294	Increased incidence of sporadic Creutzfeldt-Jakob disease on the island of Crete associated with a high rate ofPRNP 129-methionine homozygosity in the local population. Annals of Neurology, 2001, 50, 227-233.	2.8	23
295	Deposition patterns of disease-associated prion protein in captive mule deer brains with chronic wasting disease. Acta Neuropathologica, 2001, 102, 496-500.	3.9	49
296	Inherited prion disease with A117V mutation of the prion protein gene: a novel Hungarian family. Journal of Neurology, Neurosurgery and Psychiatry, 2001, 70, 802-805.	0.9	16
297	Nail-Patella Syndrome Associated with Respiratory Chain Disorder. European Neurology, 2001, 46, 92-95.	0.6	18
298	Increased incidence of sporadic Creutzfeldt-Jakob disease on the island of Crete associated with a high rate of PRNP 129-methionine homozygosity in the local population. , 2001, 50, 227.		2
299	Vascular Cell Components of the Medullary Arteries in Binswangerâ€™s Disease Brains. Stroke, 2000, 31, 1838-1842.	1.0	38
300	An autosomal dominant early adult-onset distal muscular dystrophy. Muscle and Nerve, 2000, 23, 1876-1879.	1.0	25
301	No tissue damage by chronic deep brain stimulation in Parkinson's disease. Annals of Neurology, 2000, 48, 372-376.	2.8	210
302	Alteration of the serotonergic nervous system in fatal familial insomnia. Annals of Neurology, 2000, 48, 788-791.	2.8	35
303	Neuropathological relationships between Austria and Japan. Neuropathology, 2000, 20, 124-126.	0.7	11
304	Cyclooxygenase-2 is induced in microglia during chronic cerebral ischemia in humans. Acta Neuropathologica, 2000, 99, 26-30.	3.9	83
305	Kynurenic acid metabolism in the brain of HIV-1 infected patients. Journal of Neural Transmission, 2000, 107, 1127-1138.	1.4	37
306	Binswanger's encephalopathy: serial sections and morphometry of the cerebral arteries. Acta Neuropathologica, 2000, 100, 347-355.	3.9	46

#	Article	IF	Citations
307	Analysis of EEG and CSF 14-3-3 proteins as aids to the diagnosis of Creutzfeldtâ€“Jakob disease. Neurology, 2000, 55, 811-815.	1.5	432
308	Evidence for Oxidative Stress in Experimental Prion Disease. Neurobiology of Disease, 2000, 7, 270-273.	2.1	128
309	No tissue damage by chronic deep brain stimulation in Parkinson's disease. , 2000, 48, 372.		1
310	No tissue damage by chronic deep brain stimulation in Parkinson's disease. , 2000, 48, 372.		5
311	Alteration of the serotonergic nervous system in fatal familial insomnia. Annals of Neurology, 2000, 48, 788-791.	2.8	2
312	Histopathology and immunohistochemistry of human transmissible spongiform encephalopathies (TSEs). , 2000, , 135-142.		12
313	Prions and transfusion medicine. Vox Sanguinis, 2000, 78 Suppl 2, 231-8.	0.7	3
314	No tissue damage by chronic deep brain stimulation in Parkinson's disease. Annals of Neurology, 2000, 48, 372-6.	2.8	68
315	Fatal familial insomnia: a new Austrian family. Brain, 1999, 122, 5-16.	3.7	74
316	Cytopathological alterations and therapeutic approaches in Binswanger's disease. Neuropathology, 1999, 19, 119-128.	0.7	6
317	Proliferative activity as measured by MIB-1 labeling index and long-term outcome of visual pathway astrocytomas in children. Journal of Neuro-Oncology, 1999, 42, 143-150.	1.4	13
318	Neuroaxonal pathology in Creutzfeldt-Jakob disease. Acta Neuropathologica, 1999, 97, 329-334.	3.9	46
319	Vascular changes in white matter lesions of Alzheimer's disease. Acta Neuropathologica, 1999, 97, 629-634.	3.9	30
320	Deposition of disease-associated prion protein involves the peripheral nervous system in experimental scrapie. Acta Neuropathologica, 1999, 98, 453-457.	3.9	65
321	Disease associated prion protein may deposit in the peripheral nervous system in human transmissible spongiform encephalopathies. Acta Neuropathologica, 1999, 98, 458-460.	3.9	54
322	Deposition of the prion protein (PrP) during the evolution of experimental Creutzfeldt-Jakob disease. Acta Neuropathologica, 1999, 98, 597-602.	3.9	24
323	A novel phenotype in familial Creutzfeldt-Jakob disease: Prion protein gene E200K mutation coupled with valine at codon 129 and type 2 protease-resistant prion protein. Annals of Neurology, 1999, 45, 812-816.	2.8	55
324	Classification of sporadic Creutzfeldt-Jakob disease based on molecular and phenotypic analysis of 300 subjects. Annals of Neurology, 1999, 46, 224-233.	2.8	1,314

#	Article	IF	Citations
325	Ancestral Origins and Worldwide Distribution of the PRNP 200K Mutation Causing Familial Creutzfeldt-Jakob Disease. American Journal of Human Genetics, 1999, 64, 1063-1070.	2.6	85
326	Selective Neuronal Vulnerability in Human Prion Diseases. American Journal of Pathology, 1999, 155, 1453-1457.	1.9	66
327	Early Destruction of the Extracellular Matrix around Parvalbumin-Immunoreactive Interneurons in Creutzfeldt-Jakob Disease. Neurobiology of Disease, 1999, 6, 269-279.	2.1	68
328	IMMUNOCYTOCHEMICAL DETECTION OF DISEASE-ASSOCIATED PRION PROTEIN IN THE PERIPHERAL NERVOUS SYSTEM OF PRION DISEASE. Journal of Neuropathology and Experimental Neurology, 1999, 58, 551.	0.9	0
329	A novel phenotype in familial Creutzfeldtâ€Jakob disease: Prion protein gene E200K mutation coupled with valine at codon 129 and type 2 proteaseâ€resistant prion protein. Annals of Neurology, 1999, 45, 812-816.	2.8	2
330	Classification of sporadic Creutzfeldt-Jakob disease based on molecular and phenotypic analysis of 300 subjects. , 1999, 46, 224.		9
331	A novel phenotype in familial Creutzfeldt-Jakob disease: prion protein gene E200K mutation coupled with valine at codon 129 and type 2 protease-resistant prion protein. Annals of Neurology, 1999, 45, 812-6.	2.8	21
332	Classification of sporadic Creutzfeldt-Jakob disease based on molecular and phenotypic analysis of 300 subjects. Annals of Neurology, 1999, 46, 224-33.	2.8	469
333	Prion diseases--light at the end of the tunnel?. Clinical and Experimental Pathology, 1999, 47, 123-4.	0.0	0
334	Severe, Early and Selective Loss of a Subpopulation of GABAergic Inhibitory Neurons in Experimental Transmissible Spongiform Encephalopathies. Brain Pathology, 1998, 8, 615-623.	2.1	78
335	Virus production, cell proliferation and cell death in progressive multifocal leukoencephalopathy. Neuropathology, 1998, 18, 206-210.	0.7	0
336	Development of HIV encephalitis in AIDS and TNF-Î± regulatory elements. Journal of Neuroimmunology, 1998, 91, 89-92.	1.1	21
337	Blood-brain barrier dysfunction in Binswanger's disease; an immunohistochemical study. Acta Neuropathologica, 1998, 95, 78-84.	3.9	93
338	Coexistence of Alzheimer-type neuropathology in Creutzfeldt-Jakob disease. Acta Neuropathologica, 1998, 96, 116-122.	3.9	136
339	Malnutrition-Induced Hypokalemic Myopathy in Chronic Alcoholism. Journal of Toxicology: Clinical Toxicology, 1998, 36, 369-373.	1.5	19
340	SELECTIVE, SEVERE AND EARLY LOSS OF A SUBSET OF INHIBITORY NEURONS IN EXPERIMENTAL TRANSMISSIBLE SPONGIFORM ENCEPHALOPATHIES. Journal of Neuropathology and Experimental Neurology, 1998, 57, 518.	0.9	2
341	NEURONAL APOPTOSIS IN HUMAN PRION DISEASES. Journal of Neuropathology and Experimental Neurology, 1998, 57, 493.	0.9	0
342	Different patterns of truncated prion protein fragments correlate with distinct phenotypes in P102L Gerstmann-Straussler-Scheinker disease. Proceedings of the National Academy of Sciences of the United States of America, 1998, 95, 8322-8327.	3.3	206

#	Article	IF	Citations
343	Neuronal Apoptosis in Fatal Familial Insomnia. Brain Pathology, 1998, 8, 531-537.	2.1	73
344	Primary Neuroendocrine (Merkel Cell) Carcinoma of the Anterior Skull Base. Skull Base, 1997, 7, 151-158.	0.4	6
345	Naturally Occurring Herpes Simplex Encephalitis in a Domestic Rabbit (Oryctolagus cuniculus). Veterinary Pathology, 1997, 34, 44-47.	0.8	34
346	Ganglionitis in paraneoplastic subacute sensory neuronopathy: A morphologic study. Neurology, 1997, 49, 1156-1159.	1.5	76
347	Encephalitogenic peptide (EP) in human cerebrovascular white matter lesions. NeuroReport, 1997, 8, 3727-3730.	0.6	5
348	Distribution of Parvalbumin-Immunoreactive Neurons in Brain Correlates with Hippocampal and Temporal Cortical Pathology in Creutzfeldt-Jakob Disease. Journal of Neuropathology and Experimental Neurology, 1997, 56, 1119-1124.	0.9	74
349	Central nervous system pathology in patients with the Guillain-Barre syndrome. Brain, 1997, 120, 451-464.	3.7	59
350	Nonâ€HTLVâ€I associated pleomorphic Tâ€cell lymphoma of the brain mimicking postâ€vaccinal acute inflammatory demyelination. Neuropathology and Applied Neurobiology, 1997, 23, 43-49.	1.8	7
351	The human prion diseases: from neuropathology to pathobiology and molecular genetics. Neuropathology and Applied Neurobiology, 1997, 23, 416-422.	1.8	6
352	Proliferation and Dna fragmentation in meningioma subtypes. Neuropathology and Applied Neurobiology, 1997, 23, 496-506.	1.8	46
353	Pathology and Immunocytochemistry of a Kuru Brain. Brain Pathology, 1997, 7, 547-553.	2.1	64
354	Absence of measles virus receptor (CD46) in lesions of subacute sclerosing panencephalitis brains. Acta Neuropathologica, 1997, 94, 444-449.	3.9	33
355	Metallothionein overexpression in human brain tumours. Acta Neuropathologica, 1997, 94, 599-604.	3.9	29
356	Progressive multifocal leukoencephalopathy in AIDS: initial and follow-up CT and MRI. Neuroradiology, 1997, 39, 611-618.	1.1	52
357	Proliferation and Dna fragmentation in meningioma subtypes. Neuropathology and Applied Neurobiology, 1997, 23, 496-506.	1.8	11
358	Alterations in Glia and Axons in the Brains of Binswangerâ€™s Disease Patients. Stroke, 1997, 28, 1423-1429.	1.0	108
359	The human prion diseases: from neuropathology to pathobiology and molecular genetics. Final report of an EU concerted action. Neuropathology and Applied Neurobiology, 1997, 23, 416-22.	1.8	1
360	Testing for prion protein does not confirm previously reported conjugal CJD. Lancet, The, 1996, 347, 616-617.	6.3	12

#	Article	IF	Citations
361	NEUROPATHOLOGICALLY CONFIRMED SPORADIC CREUTZFELDT-JAKOB DISEASE. Journal of Neuropathology and Experimental Neurology, 1996, 55, 661.	0.9	1
362	Familial meningocerebrovascular amyloidosis, Hungarian type, with mutant transthyretin (TTR) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 702	1.5	85
363	HIV and brain pathology: where do we go from here?. Brain Pathology, 1996, 6, 15-15.	2.1	1
364	T Cell-mediated ganglionitis associated with acute sensory neuronopathy. Annals of Neurology, 1996, 39, 543-547.	2.8	26
365	Creutzfeldt-Jakob disease in Austria.. Journal of Neurology, Neurosurgery and Psychiatry, 1996, 61, 139-142.	0.9	28
366	Subacute measles virus encephalitis. Neurology, 1996, 46, 586-587.	1.5	43
367	Meningocerebrovascular amyloidosis associated with a novel transthyretin mis-sense mutation at codon 18 (TTRD 18G). American Journal of Pathology, 1996, 148, 361-6.	1.9	71
368	Repermeation of partially embolized cerebral arteriovenous malformations: a clinical, radiologic, and histologic study. American Journal of Neuroradiology, 1996, 17, 1323-31.	1.2	48
369	Tissue Handling in Suspected Creutzfeldtâ€Jakob Disease (CJD) and Other Human Spongiform Encephalopathies (Prion Diseases). Brain Pathology, 1995, 5, 319-322.	2.1	103
370	The Original Gerstmannâ€StrÃ¤usslerâ€Scheinker Family of Austria: Divergent Clinicopathological Phenotypes but Constant PrP Genotype. Brain Pathology, 1995, 5, 201-211.	2.1	141
371	The significance of nucleolar organizer region (AgNOR) score in predicting meningioma recurrence. Cancer, 1995, 75, 130-131.	2.0	1
372	Diagnostic value of stereotactic biopsy of cerebral lesions in patients with AIDS. Acta Neurochirurgica, 1995, 134, 214-219.	0.9	21
373	Diffuse Lewy body disease as substrate of primary lateral sclerosis. Journal of Neurology, 1995, 242, 59-63.	1.8	24
374	Immunohistochemical study of apolipoprotein E in human cerebrovascular white matter lesions. Acta Neuropathologica, 1995, 90, 608-614.	3.9	11
375	Suprasellar meningioma with expression of glial fibrillary acidic protein: a peculiar variant. Acta Neuropathologica, 1995, 90, 539-544.	3.9	64
376	Ultra Structural Pathology of Gerstmann-StrÃ¤ussler-Scheinker Disease. Ultrastructural Pathology, 1995, 19, 23-36.	0.4	18
377	Neuropathological Diagnostic Criteria for Creutzfeldtâ€Jakob Disease (CJD) and Other Human Spongiform Encephalopathies (Prion Diseases). Brain Pathology, 1995, 5, 459-466.	2.1	378
378	Neuroaxonal dystrophy in experimental creutzfeldt-jakob disease: Electron microscopical and immunohistochemical demonstration of neurofilament accumulations within affected neurites. Journal of Comparative Pathology, 1995, 112, 243-255.	0.1	13

#	Article	IF	Citations
379	Suprasellar meningioma with expression of glial fibrillary acidic protein: a peculiar variant. Acta Neuropathologica, 1995, 90, 539-544.	3.9	4
380	Immunohistochemical study of apolipoprotein E in human cerebrovascular white matter lesions. Acta Neuropathologica, 1995, 90, 608-614.	3.9	2
381	Pathology of the peripheral nervous system in unselected AIDS autopsies. , 1995, , 159-172.		0
382	Trilobar holoprosencephaly ("triprosencephaly"): a unique type of cerebral malformation. Acta Neuropathologica, 1995, 89, 567-570.	3.9	0
383	Patterns of oligodendroglia pathology in multiple sclerosis. Brain, 1994, 117, 1311-1322.	3.7	381
384	Symposia on Neurodegenerative Disorders: Common Molecular Mechanisms, Ocho Rios, Jamaica, February 1993 * and April 1994. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 1994, 1, 206-210.	1.4	1
385	Tubulovesicular structures in Gerstmann-Strï¿½ussler-Scheinker disease. Acta Neuropathologica, 1994, 88, 491-492.	3.9	10
386	Lipomatous medulloblastoma in adults: A new tumor type with possible favorable prognosis. Human Pathology, 1994, 25, 730-731.	1.1	30
387	Tumoren des Zentralnervensystems (ZNS) und seiner Umgebungsstrukturen. , 1994, , 171-182.		0
388	Neocortical changes in Parkinson's disease, revisited. , 1994, 13, 55-9.		5
389	Neuronal damage in the cerebral cortex of AIDS brains: a morphometric study. Acta Neuropathologica, 1993, 85, 185-9.	3.9	86
390	Uncommon types of polyglucosan bodies in the human brain: distribution and relation to disease. Acta Neuropathologica, 1993, 86, 484-90.	3.9	23
391	Immunostaining for proliferating cell nuclear antigen: its role in determination of proliferation in routinely processed human brain tumor specimens. Acta Neuropathologica, 1993, 86, 582-589.	3.9	47
392	Microglia is a component of the prion protein amyloid plaque in the Gerstmann-Strï¿½ussler-Scheinker syndrome. Acta Neuropathologica, 1993, 85, 623-627.	3.9	51
393	Astroglial changes in the cerebral cortex of AIDS brains: a morphometric and immunohistochemical investigation. Neuropathology and Applied Neurobiology, 1993, 19, 329-335.	1.8	44
394	Clonal Expansion of Hypermutated Measles Virus in a SSPE Brain. Virology, 1993, 197, 188-195.	1.1	119
395	Fatal Encephalitis Caused by Concomitant Infection with Tick-Borne Encephalitis Virus and Borrelia burgdorferi. Clinical Infectious Diseases, 1993, 16, 392-396.	2.9	57
396	HIV-Related Dementia: Pathology and Possible Pathogenesis. , 1993, , 171-185.		6

#	Article	IF	Citations
397	PRESENCE, DISTRIBUTION AND SPREAD OF PRODUCTIVE VARICELLA ZOSTER VIRUS INFECTION IN NERVOUS TISSUES. Brain, 1992, 115, 383-398.	3.7	96
398	Devic's neuromyelitis optica and Schilder's myelinoclastic diffuse sclerosis. Journal of Neurology, Neurosurgery and Psychiatry, 1992, 55, 1194-1196.	0.9	13
399	Classic, atypical, and anaplastic meningioma: three histopathological subtypes of clinical relevance. Journal of Neurosurgery, 1992, 77, 616-623.	0.9	247
400	Tubulovesicular structures in Creutzfeldt-Jakob disease. Acta Neuropathologica, 1992, 84, 238-43.	3.9	29
401	Subacute diencephalic angioencephalopathy: an entity similar to angiodysgenetic necrotizing encephalopathy and Foix-Alajouanine disease. Journal of Neurology, 1992, 239, 379-381.	1.8	7
402	Morphological spectrum, distribution and clinical correlation of white matter lesions in AIDS brains. Neuropathology and Applied Neurobiology, 1992, 18, 489-501.	1.8	32
403	HIVâ€Associated Disease of the Nervous System: Review of Nomenclature and Proposal for Neuropathologyâ€Based Terminology. Brain Pathology, 1991, 1, 143-152.	2.1	323
404	Prion protein mutation in family first reported by Gerstmann, StrÃussler, and Scheinker. Lancet, The, 1991, 337, 1160.	6.3	108
405	HIVâ€Induced CNS Lesions. Brain Pathology, 1991, 1, 153-154.	2.1	7
406	Neuropathology of Human Immunodeficiency Virus Infection. Brain Pathology, 1991, 1, 163-175.	2.1	410
407	The Definition of HIVâ€specific Neuropathology. Pathology International, 1991, 41, 182-191.	0.6	14
408	Lipomatous differentiation in a medulloblastoma. Acta Neuropathologica, 1991, 81, 471-473.	3.9	52
409	Prominent cortical atrophy with neuronal loss as correlate of human immunodeficiency virus encephalopathy. Acta Neuropathologica, 1991, 82, 229-233.	3.9	70
410	Tubulovesicular structures in human and experimental Creutzfeldt-Jakob disease. European Journal of Epidemiology, 1991, 7, 551-555.	2.5	3
411	Determination of proliferative activities in human brain tumor specimens: a comparison of three methods. Journal of Neuro-Oncology, 1991, 10, 1-11.	1.4	34
412	Multifocal vacuolar leucoencephalopathy: a distinct HIVâ€associated lesion of the brain**. Neuropathology and Applied Neurobiology, 1990, 16, 437-443.	1.8	27
413	Loss of neurons in the frontal cortex in AIDS brains. Acta Neuropathologica, 1990, 80, 92-94.	3.9	355
414	Human immunodeficiency virus (HIV) envelope and core proteins in CNS tissues of patients with the acquired immune deficiency syndrome (AIDS). Acta Neuropathologica, 1990, 79, 611-619.	3.9	90

#	Article	IF	Citations
415	Monocyte subpopulations in human gliomas: expression of Fc and complement receptors and correlation with tumor proliferation. Acta Neuropathologica, 1990, 80, 287-294.	3.9	119
416	Progressive multifocal leukoencephalopathy (PML) in AIDS and in the pre-AIDS era. Acta Neuropathologica, 1990, 80, 375-380.	3.9	63
417	Argyrophilic nucleolar organizer region proteins (Ag-NORs) in human brain tumors: relations with grade of malignany and proliferation indices. Acta Neuropathologica, 1990, 80, 156-162.	3.9	52
418	Neuropathology of AIDS dementia. A review after 205 post mortem examinations. Acta Neurologica, 1990, 12, 32-5.	0.1	2
419	Fatal encephalitis in a patient with chronic graft-versus-host disease. Bone Marrow Transplantation, 1990, 6, 53-7.	1.3	35
420	In situ analysis of cell kinetics in human brain tumors. Acta Neuropathologica, 1989, 77, 276-282.	3.9	83
421	Vacuolar myelopathy with multinucleated giant cells in the acquired immune deficiency syndrome (AIDS). Acta Neuropathologica, 1989, 78, 497-503.	3.9	64
422	Cytomegalovirus (CMV) disease of the brain in AIDS and connatal infection: a comparative study by histology, immunocytochemistry and in situ DNA hybridization. Acta Neuropathologica, 1989, 79, 286-293.	3.9	61
423	Accumulation of abnormally phosphorylated Ï„ precedes the formation of neurofibrillary tangles in Alzheimer's disease. Brain Research, 1989, 477, 90-99.	1.1	790
424	Human immunodeficiency virus (HIV)-induced disease of the central nervous system: pathology and implications for pathogenesis. Acta Neuropathologica, 1989, 77, 225-236.	3.9	146
425	An Antigenic Profile of Lewy Bodies: Immunocytochemical Indication for Protein Phosphorylation and Ubiquitination. Journal of Neuropathology and Experimental Neurology, 1989, 48, 81-93.	0.9	137
426	Herpes Simplex Virus (HSV) DNA in Microglial Nodular Brainstem Encephalitis. Journal of Neuropathology and Experimental Neurology, 1989, 48, 645-652.	0.9	30
427	CNS DISTRIBUTION OF HUMAN IMMUNODEFICIENCY VIRUS (HIV) ANTIGENS IN AIDS. Journal of Neuropathology and Experimental Neurology, 1989, 48, 384.	0.9	3
428	Tau and ubiquitin immunoreactivity at different stages of formation of Alzheimer neurofibrillary tangles. Progress in Clinical and Biological Research, 1989, 317, 837-48.	0.2	34
429	Comparison of in situ DNA hybridization (ISH) and immunocytochemistry for diagnosis of herpes simplex virus (HSV) encephalitis in tissue. Virchows Archiv A, Pathological Anatomy and Histopathology, 1988, 414, 39-43.	1.4	13
430	Pathogenesis of Human Immunodeficiency Virus (HIV)-Associated Brain Lesions A Neuropathologic Evaluation. Annals of the New York Academy of Sciences, 1988, 540, 630-633.	1.8	5
431	Neuropathy Associated with Acrodermatitis Chronica Atrophicans Clinical and Morphological Features. Annals of the New York Academy of Sciences, 1988, 539, 35-45.	1.8	80
432	Human Immunodeficiency Virus in Vacuolar Myelopathy of the Acquired Immunodeficiency Syndrome. New England Journal of Medicine, 1988, 319, 1667-1668.	13.9	31

#	Article	IF	Citations
433	Human Immunodeficiency Virus in Glial Cells?. Journal of Infectious Diseases, 1988, 157, 203-203.	1.9	17
434	NIMODIPINE DOES NOT REDUCE BRAIN DAMAGE AFTER VENTRICULAR FIBRILLATION IN AN ACUTE PIG MODEL. Critical Care Medicine, 1988, 16, 386.	0.4	0
435	Very Long Chain Fatty Acids and Phytanic Acid in Genetic Peroxisomal Diseases. , 1988, , 423-428.		1
436	Glioblastoma developing at the site of a cerebellar medulloblastoma treated 6 years earlier. Journal of Neurosurgery, 1987, 67, 915-918.	0.9	36
437	Pathogenesis of HIV-associated brain lesions: A neuropathological evaluation. Journal of Neuroimmunology, 1987, 16, 24-25.	1.1	1
438	Brain pathology induced by infection with the human immunodeficiency virus (HIV). Acta Neuropathologica, 1987, 75, 185-198.	3.9	311
439	Neurofibrillary tangles in Alzheimer's disease and progressive supranuclear palsy: antigenic similarities and differences. Acta Neuropathologica, 1987, 74, 39-46.	3.9	124
440	Contribution of histiocytic cells to sarcomatous development of the gliosarcoma. Acta Neuropathologica, 1987, 73, 124-130.	3.9	31
441	Das morphologische Korrelat der HIV-Infektion des Gehirns. , 1987, , 117-132.		5
442	Non-glial specificities of immunocytochemistry for the glial fibrillary acidic protein (GFAP). Acta Neuropathologica, 1986, 72, 43-54.	3.9	75
443	Development of stroma in malignant lymphomas of the brain compared with epidural lymphomas. Acta Neuropathologica, 1986, 71, 125-129.	3.9	7
444	Multinucleated giant cells in brain: A hallmark of the acquired immune deficiency syndrome (AIDS). Acta Neuropathologica, 1986, 69, 253-258.	3.9	218
445	Glial fibrillary acidic protein and S-100 protein in human hepatic encephalopathy: Immunocytochemical demonstration of dissociation of two glia-associated proteins. Acta Neuropathologica, 1986, 70, 17-21.	3.9	62
446	Restricted Expression of Measles Virus Proteins in Brains from Cases of Subacute Sclerosing Panencephalitis. Journal of General Virology, 1986, 67, 2435-2444.	1.3	105
447	Expression of defective measles virus genes in brain tissues of patients with subacute sclerosing panencephalitis. Journal of Virology, 1986, 59, 472-478.	1.5	152
448	Unilateral Creutzfeldtâ€Jakob disease. Neurology, 1986, 36, 1517-1517.	1.5	39
449	An immunocytochemical comparison of the glia-associated proteins glial fibrillary acidic protein (GFAP) and S-100 protein (S100P) in human brain tumors. , 1986, 5, 21-7.		23
450	Occurrence of IgA Subclasses (IgA1 and IgA2) in the Human Nervous System. International Archives of Allergy and Immunology, 1985, 76, 107-115.	0.9	23

#	Article	IF	Citations
451	Shared antigenic determinants between human hemopoietic cells and nervous tissues and tumors. Acta Neuropathologica, 1985, 67, 58-66.	3.9	29
452	Progressive diffuse leukoencephalopathy in patients with acquired immune deficiency syndrome (AIDS). Acta Neuropathologica, 1985, 68, 333-339.	3.9	139
453	Cross-reactivity between human hemopoietic cells and brain tumors as defined by monoclonal antibodies. Journal of Neuro-Oncology, 1985, 3, 173-179.	1.4	9
454	Cross-reactivity of human brain tumors with lymphoid- and hemapoietic-associated monoclonal antibodies. Journal of Neuroimmunology, 1985, 10, 177.	1.1	0
455	Pathology of Midline Brain Tumors. Acta Neurochirurgica Supplementum, 1985, , 23-30.	0.5	6
456	Primitive neuroectodermal tumors including the medulloblastoma: glial differentiation signaled by immunoreactivity for GFAP is restricted to the pure desmoplastic medulloblastoma ("arachnoidal) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50		
457	Glial bundles in spinal nerve roots. Acta Neuropathologica, 1984, 65, 46-52.	3.9	8
458	Glial fibrillary acidic protein (GFAP) in oligodendroglial tumors: Gliofibrillary oligodendroglioma and transitional oligoastrocytoma as subtypes of oligodendroglioma. Acta Neuropathologica, 1984, 64, 265-272.	3.9	126
459	Production of glial fibrillary acidic protein (GFAP) by neoplastic cells: Adaptation to the microenvironment. Acta Neuropathologica, 1984, 64, 333-338.	3.9	29
460	23 MONOCLONAL ANTIBODIES RAISED AGAINST HEMOPOIETIC CELLS. Journal of Neuropathology and Experimental Neurology, 1984, 43, 300.	0.9	1
461	Tumoren des Zentralnervensystems. , 1984, , 140-146.		6
462	Diffuse type of Lewy body disease: progressive dementia with abundant cortical Lewy bodies and senile changes of varying degree--a new disease?. , 1984, 3, 185-92.		112
463	Supratentorial lobar ependymomas: Reports on the grading and survival periods in 80 cases, including 46 recurrences. Acta Neurochirurgica, 1983, 69, 243-251.	0.9	61
464	Brain tissue immunoglobulins in adrenoleukodystrophy: A comparison with multiple sclerosis and systemic lupus erythematosus. Acta Neuropathologica, 1983, 59, 95-102.	3.9	53
465	Immunohistological Demonstration of Serum Proteins and Structural and Viral Antigens in Paraffin Sections of Nervous Tissues. Annals of the New York Academy of Sciences, 1983, 420, 176-184.	1.8	14
466	Central Nervous System Immunoglobulins (Ig) in Adrenoleukodystrophy (ALD). Protides of the Biological Fluids; Proceedings of the Colloquium, 1983, 30, 171-173.	0.1	0
467	Papovavirus antigens in paraffin sections of PML brains. Progress in Clinical and Biological Research, 1983, 105, 299-309.	0.2	17
468	PAPOVA VIRAL ANTIGENS IN PML BRAINS. Journal of Neuropathology and Experimental Neurology, 1982, 41, 366.	0.9	1

#	Article	IF	Citations
469	Measles virus antigen in panencephalitis. Acta Neuropathologica, 1982, 56, 52-62.	3.9	27
470	Hyaline inclusions (pseudopsammoma bodies) in meningiomas: Immunocytochemical demonstration of epithel-like secretion of secretory component and immunoglobulins A and M. Acta Neuropathologica, 1982, 56, 294-298.	3.9	37
471	Accumulation of very long chain fatty acids is common to 3 variants of adrenoleukodystrophy (ALD). Journal of the Neurological Sciences, 1981, 51, 301-310.	0.3	44
472	Rabies and Herpes simplex virus encephalitis. Virchows Archiv A, Pathological Anatomy and Histology, 1981, 390, 353-364.	1.3	18
473	Inflammatory Demyelinating Polyradiculitis in a Patient With Multiple Sclerosis. Archives of Neurology, 1981, 38, 99-102.	4.9	84
474	Brain Pathology in the Collagen Vascular Diseases. Angiology, 1981, 32, 365-372.	0.8	9
475	Immunohistological Studies in Viral Encephalitis. Acta Neuropathologica Supplementum, 1981, 7, 142-144.	0.8	5
476	Benign mixed glial-mesenchymal tumour (â€œglio-fibromaâ€) of the spinal cord. Acta Neurochirurgica, 1980, 55, 141-145.	0.9	22
477	Diagnostic and pathomorphological aspects of glioma multiplicity. Neurosurgical Review, 1980, 3, 233-241.	1.2	35
478	Hereditary cerebellar atrophy (Holmes type) with optic atrophy. Archiv Fur Psychiatrie Und Nervenkrankheiten, 1979, 226, 311-318.	0.6	2
479	Intermittent meningitic reaction with severe basophilia and eosinophilia in CNS leukaemia. Journal of the Neurological Sciences, 1976, 28, 459-468.	0.3	15
480	Spastic paraplegia associated with addison's disease: Adult variant of adreno-leukodystrophy. Journal of Neurology, 1976, 213, 237-250.	1.8	97
481	Partially resected und irradiated cerebellar astrocytoma of childhood: Malignant evolution after 28 years. Acta Neurochirurgica, 1975, 32, 139-146.	0.9	71
482	Primary leptomeningeal sarcomatosis. Journal of Neurology, 1975, 211, 77-93.	1.8	23
483	SSPE-Like Inclusion Body Disorder in Treated Childhood Leukemia. , 1975, Suppl 6, 267-272.		13
484	Primary glioblastoma of the cerebellum. Acta Neurochirurgica, 1974, 31, 115-121.	0.9	19
485	Intracranial lipomatous hamartomas (Intracranial ?lipomas?). Acta Neuropathologica, 1974, 28, 205-222.	3.9	146
486	Prognostic relevance of p53 protein expression in glioblastoma. Oncology Reports, 0, , .	1.2	9