Herbert Budka

List of Publications by Year
in descending order

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501 papers	29,003 citations	4388 86 h-index	8167 148 g-index
539 all docs	539 docs citations	539 times ranked	21220 citing authors

#	Article	IF	Citations
1	Classification of sporadic Creutzfeldt-Jakob disease based on molecular and phenotypic analysis of 300 subjects. Annals of Neurology, 1999, 46, 224-233.	5.3	1,314
2	Accumulation of abnormally phosphorylated Ï„ precedes the formation of neurofibrillary tangles in Alzheimer's disease. Brain Research, 1989, 477, 90-99.	2.2	790
3	Scientific Opinion on risk based control of biogenic amine formation in fermented foods. EFSA Journal, 2011, 9, 2393.	1.8	602
4	Identification of common variants influencing risk of the tauopathy progressive supranuclear palsy. Nature Genetics, 2011, 43, 699-705.	21.4	502
5	Classification of sporadic Creutzfeldt-Jakob disease based on molecular and phenotypic analysis of 300 subjects. Annals of Neurology, 1999, 46, 224-33.	5.3	469
6	Analysis of EEG and CSF 14-3-3 proteins as aids to the diagnosis of Creutzfeldtâ€“Jakob disease. Neurology, 2000, 55, 811-815.	1.1	432
7	Neuropathology of Human Immunodeficiency Virus Infection. Brain Pathology, 1991, 1, 163-175.	4.1	410
8	Genetic prion disease: the EUROCJD experience. Human Genetics, 2005, 118, 166-174.	3.8	391
9	Patterns of oligodendroglia pathology in multiple sclerosis. Brain, 1994, 117, 1311-1322.	7.6	381
10	Aging-related tau astrogliopathy (ARTAG): harmonized evaluation strategy. Acta Neuropathologica, 2016, 131, 87-102.	7.7	380
11	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.	1.8	379
12	Neuropathological Diagnostic Criteria for Creutzfeldtâ€Jakob Disease (CJD) and Other Human Spongiform Encephalopathies (Prion Diseases). Brain Pathology, 1995, 5, 459-466.	4.1	378
13	Loss of neurons in the frontal cortex in AIDS brains. Acta Neuropathologica, 1990, 80, 92-94.	7.7	355
14	HIVâ€Associated Disease of the Nervous System: Review of Nomenclature and Proposal for Neuropathologyâ€Based Terminology. Brain Pathology, 1991, 1, 143-152.	4.1	323
15	Brain pathology induced by infection with the human immunodeficiency virus (HIV). Acta Neuropathologica, 1987, 75, 185-198.	7.7	311
16	Mortality from Creutzfeldtâ€“Jakob disease and related disorders in Europe, Australia, and Canada. Neurology, 2005, 64, 1586-1591.	1.1	306
17	Molecular classification of sporadic Creutzfeldtâ€“Jakob disease. Brain, 2003, 126, 1333-1346.	7.6	301
18	Determinants of diagnostic investigation sensitivities across the clinical spectrum of sporadic Creutzfeldt-Jakob disease. Brain, 2006, 129, 2278-2287.	7.6	283

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19	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.	7.7	264
20	Viral encephalitis: a review of diagnostic methods and guidelines for management. European Journal of Neurology, 2005, 12, 331-343.	3.3	256
21	Recombinant prion protein induces a new transmissible prion disease in wild-type animals. Acta Neuropathologica, 2010, 119, 177-187.	7.7	256
22	Mutations of the Prion Protein Gene. Journal of Neurology, 2002, 249, 1567-1582.	3.6	251
23	Classic, atypical, and anaplastic meningioma: three histopathological subtypes of clinical relevance. Journal of Neurosurgery, 1992, 77, 616-623.	1.6	247
24	Predictors of survival in sporadic Creutzfeldt-Jakob disease and other human transmissible spongiform encephalopathies. Brain, 2004, 127, 2348-2359.	7.6	244
25	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.	1.8	235
26	Multinucleated giant cells in brain: A hallmark of the acquired immune deficiency syndrome (AIDS). Acta Neuropathologica, 1986, 69, 253-258.	7.7	218
27	Scientific Opinion on risk assessment of parasites in fishery products. EFSA Journal, 2010, 8, 1543.	1.8	214
28	No tissue damage by chronic deep brain stimulation in Parkinson's disease. Annals of Neurology, 2000, 48, 372-376.	5.3	210
29	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.	7.1	206
30	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.	1.7	196
31	TARDBP variation associated with frontotemporal dementia, supranuclear gaze palsy, and chorea. Movement Disorders, 2009, 24, 1842-1847.	3.9	182
32	Scientific Opinion on Quantification of the risk posed by broiler meat to human campylobacteriosis in the EU. EFSA Journal, 2010, 8, 1437.	1.8	181
33	Viral meningoencephalitis: a review of diagnostic methods and guidelines for management. European Journal of Neurology, 2010, 17, 999.	3.3	176
34	Peroxisomal alterations in Alzheimerâ€™s disease. Acta Neuropathologica, 2011, 122, 271-283.	7.7	176
35	Neuropathological criteria of anti-IgLON5-related tauopathy. Acta Neuropathologica, 2016, 132, 531-543.	7.7	173
36	Expression of hypoxia-inducible factor-1Î± in oligodendrogliomas. Cancer, 2001, 92, 165-171.	4.1	171

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37	Globular glial tauopathies (GGT): consensus recommendations. Acta Neuropathologica, 2013, 126, 537-544.	7.7	168
38	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.	3.7	166
39	Visualization of Central European Tick-Borne Encephalitis Infection in Fatal Human Cases. Journal of Neuropathology and Experimental Neurology, 2005, 64, 506-512.	1.7	164
40	Scientific Opinion on the public health hazards to be covered by inspection of meat (swine). EFSA Journal, 2011, 9, 2351.	1.8	154
41	Expression of defective measles virus genes in brain tissues of patients with subacute sclerosing panencephalitis. Journal of Virology, 1986, 59, 472-478.	3.4	152
42	Brain Protein Preservation Largely Depends on the Postmortem Storage Temperature. Journal of Neuropathology and Experimental Neurology, 2007, 66, 35-46.	1.7	151
43	Mixed Brain Pathologies in Dementia: The BrainNet Europe Consortium Experience. Dementia and Geriatric Cognitive Disorders, 2008, 26, 343-350.	1.5	148
44	Intracranial lipomatous hamartomas (Intracranial ?lipomas?). Acta Neuropathologica, 1974, 28, 205-222.	7.7	146
45	Human immunodeficiency virus (HIV)-induced disease of the central nervous system: pathology and implications for pathogenesis. Acta Neuropathologica, 1989, 77, 225-236.	7.7	146
46	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.	1.7	145
47	The Original Gerstmannâ€šStrÃusslerâ€šScheinker Family of Austria: Divergent Clinicopathological Phenotypes but Constant PrP Genotype. Brain Pathology, 1995, 5, 201-211.	4.1	141
48	Natively unfolded tubulin polymerization promoting protein TPPP/p25 is a common marker of alpha-synucleinopathies. Neurobiology of Disease, 2004, 17, 155-162.	4.4	140
49	Progressive diffuse leukoencephalopathy in patients with acquired immune deficiency syndrome (AIDS). Acta Neuropathologica, 1985, 68, 333-339.	7.7	139
50	An Antigenic Profile of Lewy Bodies: Immunocytochemical Indication for Protein Phosphorylation and Ubiquitination. Journal of Neuropathology and Experimental Neurology, 1989, 48, 81-93.	1.7	137
51	Coexistence of Alzheimer-type neuropathology in Creutzfeldt-Jakob disease. Acta Neuropathologica, 1998, 96, 116-122.	7.7	136
52	Neuropathology of prion diseases. British Medical Bulletin, 2003, 66, 121-130.	6.9	136
53	An antibody with high reactivity for disease-associated Î±-synuclein reveals extensive brain pathology. Acta Neuropathologica, 2012, 124, 37-50.	7.7	133
54	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.	4.1	132

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55	Structure-based drug design identifies polythiophenes as antiprion compounds. Science Translational Medicine, 2015, 7, 299ra123.	12.4	130
56	Scientific Opinion on Q fever. EFSA Journal, 2010, 8, 1595.	1.8	129
57	Evidence for Oxidative Stress in Experimental Prion Disease. Neurobiology of Disease, 2000, 7, 270-273.	4.4	128
58	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.	4.1	127
59	SAMP8 mice as a neuropathological model of accelerated brain aging and dementia: Toshio Takeda's legacy and future directions. Neuropathology, 2017, 37, 293-305.	1.2	127
60	Glial fibrillary acidic protein (GFAP) in oligodendroglial tumors: Gliofibrillary oligodendroglioma and transitional oligoastrocytoma as subtypes of oligodendroglioma. Acta Neuropathologica, 1984, 64, 265-272.	7.7	126
61	Prion Diseases: From Protein to Cell Pathology. American Journal of Pathology, 2008, 172, 555-565.	3.8	126
62	Neurofibrillary tangles in Alzheimer's disease and progressive supranuclear palsy: antigenic similarities and differences. Acta Neuropathologica, 1987, 74, 39-46.	7.7	124
63	Scientific Opinion on monitoring and assessment of the public health risk of â€œSalmonellaTyphimurium-likeâ€•strains. EFSA Journal, 2010, 8, 1826.	1.8	122
64	Inflammatory response in human tick-borne encephalitis: analysis of postmortem brain tissue. Journal of NeuroVirology, 2006, 12, 322-327.	2.1	121
65	Monocyte subpopulations in human gliomas: expression of Fc and complement receptors and correlation with tumor proliferation. Acta Neuropathologica, 1990, 80, 287-294.	7.7	119
66	Clonal Expansion of Hypermutated Measles Virus in a SSPE Brain. Virology, 1993, 197, 188-195.	2.4	119
67	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.	2.5	117
68	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
69	White Matter Tauopathy With Globular Glial Inclusions: A Distinct Sporadic Frontotemporal Lobar Degeneration. Journal of Neuropathology and Experimental Neurology, 2008, 67, 963-975.	1.7	111
70	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.	4.4	110
71	Prion protein mutation in family first reported by Gerstmann, StrÃussler, and Scheinker. Lancet, The, 1991, 337, 1160.	13.7	108
72	Alterations in Glia and Axons in the Brains of Binswangerâ€™s Disease Patients. Stroke, 1997, 28, 1423-1429.	2.0	108

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73	Restricted Expression of Measles Virus Proteins in Brains from Cases of Subacute Sclerosing Panencephalitis. Journal of General Virology, 1986, 67, 2435-2444.	2.9	105
74	Genetic Creutzfeldt-Jakob disease associated with the E200K mutation: characterization of a complex proteinopathy. Acta Neuropathologica, 2011, 121, 39-57.	7.7	105
75	Advances in broad bandwidth light sources for ultrahigh resolution optical coherence tomography. Physics in Medicine and Biology, 2004, 49, 1235-1246.	3.0	104
76	Tissue Handling in Suspected Creutzfeldtâ€Jakob Disease (CJD) and Other Human Spongiform Encephalopathies (Prion Diseases). Brain Pathology, 1995, 5, 319-322.	4.1	103
77	Management of a twenty-first century brain bank: experience in the BrainNet Europe consortium. Acta Neuropathologica, 2008, 115, 497-507.	7.7	101
78	Protein coding of neurodegenerative dementias: the neuropathological basis of biomarker diagnostics. Acta Neuropathologica, 2010, 119, 389-408.	7.7	98
79	Genesis of Mammalian Prions: From Non-infectious Amyloid Fibrils to a Transmissible Prion Disease. PLoS Pathogens, 2011, 7, e1002419.	4.7	98
80	Spastic paraplegia associated with addison's disease: Adult variant of adreno-leukodystrophy. Journal of Neurology, 1976, 213, 237-250.	3.6	97
81	PRESENCE, DISTRIBUTION AND SPREAD OF PRODUCTIVE VARICELLA ZOSTER VIRUS INFECTION IN NERVOUS TISSUES. Brain, 1992, 115, 383-398.	7.6	96
82	Neuropathology of white matter disease in Leber's hereditary optic neuropathy. Brain, 2004, 128, 35-41.	7.6	96
83	Immunohistochemistry for the Prion Protein: Comparison of Different Monoclonal Antibodies in Human Prion Disease Subtypes. Brain Pathology, 2002, 12, 1-11.	4.1	96
84	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.	1.7	95
85	Distinct time pattern of complement activation and cytotoxic T cell response in Guillain-Barre syndrome. Brain, 2003, 126, 2034-2042.	7.6	94
86	Blood-brain barrier dysfunction in Binswanger's disease; an immunohistochemical study. Acta Neuropathologica, 1998, 95, 78-84.	7.7	93
87	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.	7.6	91
88	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.	7.7	90
89	Amyloid-Î² pathology and cerebral amyloid angiopathy are frequent in iatrogenic Creutzfeldt-Jakob disease after dural grafting. Swiss Medical Weekly, 2016, 146, w14287.	1.6	89
90	Neuronal damage in the cerebral cortex of AIDS brains: a morphometric study. Acta Neuropathologica, 1993, 85, 185-9.	7.7	86

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91	Inter-laboratory comparison of neuropathological assessments of Î²-amyloid protein: a study of the BrainNet Europe consortium. Acta Neuropathologica, 2008, 115, 533-546.	7.7	86
92	Familial meningocerebrovascular amyloidosis, Hungarian type, with mutant transthyretin (TTR) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 702	1.1	85
93	Ancestral Origins and Worldwide Distribution of the PRNP 200K Mutation Causing Familial Creutzfeldt-Jakob Disease. American Journal of Human Genetics, 1999, 64, 1063-1070.	6.2	85
94	Dura mater is a potential source of AÎ² seeds. Acta Neuropathologica, 2016, 131, 911-923.	7.7	85
95	Inflammatory Demyelinating Polyradiculitis in a Patient With Multiple Sclerosis. Archives of Neurology, 1981, 38, 99-102.	4.5	84
96	A comparative study on the expression of cyclooxygenase and 5-lipoxygenase during cerebral ischemia in humans. Acta Neuropathologica, 2002, 104, 601-607.	7.7	84
97	In situ analysis of cell kinetics in human brain tumors. Acta Neuropathologica, 1989, 77, 276-282.	7.7	83
98	Cyclooxygenase-2 is induced in microglia during chronic cerebral ischemia in humans. Acta Neuropathologica, 2000, 99, 26-30.	7.7	83
99	Imaging ex vivo healthy and pathological human brain tissue with ultra-high-resolution optical coherence tomography. Journal of Biomedical Optics, 2005, 10, 011006.	2.6	82
100	Scientific Opinion on Norovirus (NoV) in oysters: methods, limits and control options. EFSA Journal, 2012, 10, 2500.	1.8	82
101	Neuropathy Associated with Acrodermatitis Chronica Atrophicans Clinical and Morphological Features. Annals of the New York Academy of Sciences, 1988, 539, 35-45.	3.8	80
102	A peculiar constellation of tau pathology defines a subset of dementia in the elderly. Acta Neuropathologica, 2011, 122, 205-222.	7.7	80
103	Severe, Early and Selective Loss of a Subpopulation of GABAergic Inhibitory Neurons in Experimental Transmissible Spongiform Encephalopathies. Brain Pathology, 1998, 8, 615-623.	4.1	78
104	Ki-67 Immunolabeling Index Is an Accurate Predictor of Outcome in Patients With Intracranial Ependymoma. American Journal of Surgical Pathology, 2004, 28, 914-920.	3.7	78
105	Subcellular Localization of Disease-Associated Prion Protein in the Human Brain. American Journal of Pathology, 2005, 166, 287-294.	3.8	77
106	Ganglionitis in paraneoplastic subacute sensory neuronopathy: A morphologic study. Neurology, 1997, 49, 1156-1159.	1.1	76
107	Non-glial specificities of immunocytochemistry for the glial fibrillary acidic protein (GFAP). Acta Neuropathologica, 1986, 72, 43-54.	7.7	75
108	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.	1.7	74

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109	Fatal familial insomnia: a new Austrian family. Brain, 1999, 122, 5-16.	7.6	74
110	Neuronal Apoptosis in Fatal Familial Insomnia. Brain Pathology, 1998, 8, 531-537.	4.1	73
111	Assessment of Î±-Synuclein Pathology: A Study of the BrainNet Europe Consortium. Journal of Neuropathology and Experimental Neurology, 2008, 67, 125-143.	1.7	73
112	Molecular Pathology of Human Prion Diseases. International Journal of Molecular Sciences, 2009, 10, 976-999.	4.1	73
113	Severe depletion of mitochondrial DNA in spinal muscular atrophy. Acta Neuropathologica, 2003, 105, 245-251.	7.7	72
114	A New Mechanism for Transmissible Prion Diseases. Journal of Neuroscience, 2012, 32, 7345-7355.	3.6	72
115	Partially resected und irradiated cerebellar astrocytoma of childhood: Malignant evolution after 28 years. Acta Neurochirurgica, 1975, 32, 139-146.	1.7	71
116	Meningocerebrovascular amyloidosis associated with a novel transthyretin mis-sense mutation at codon 18 (TTRD 18G). American Journal of Pathology, 1996, 148, 361-6.	3.8	71
117	Prominent cortical atrophy with neuronal loss as correlate of human immunodeficiency virus encephalopathy. Acta Neuropathologica, 1991, 82, 229-233.	7.7	70
118	Marked increase of neuronal prion protein immunoreactivity in Alzheimer's disease and human prion diseases. Acta Neuropathologica, 2001, 101, 417-423.	7.7	70
119	Nucleus-specific alteration of raphe neurons in human neurodegenerative disorders. NeuroReport, 2003, 14, 73-76.	1.2	69
120	Early Destruction of the Extracellular Matrix around Parvalbumin-Immunoreactive Interneurons in Creutzfeldt-Jakob Disease. Neurobiology of Disease, 1999, 6, 269-279.	4.4	68
121	Oxidative Damage to Nucleic Acids in Human Prion Disease. Neurobiology of Disease, 2002, 9, 275-281.	4.4	68
122	Complement activation in human prion disease. Neurobiology of Disease, 2004, 15, 21-28.	4.4	68
123	Involvement of the Endosomal-Lysosomal System Correlates With Regional Pathology in Creutzfeldt-Jakob Disease. Journal of Neuropathology and Experimental Neurology, 2007, 66, 628-636.	1.7	68
124	Scientific Opinion on a Quantitative Microbiological Risk Assessment of Salmonella in slaughter and breeder pigs. EFSA Journal, 2010, 8, 1547.	1.8	68
125	Fatal Prion Disease in a Mouse Model of Genetic E200K Creutzfeldt-Jakob Disease. PLoS Pathogens, 2011, 7, e1002350.	4.7	68
126	Current concepts of neuropathological diagnostics in practice: neurodegenerative diseases. , 2010, 29, 271-288.		68

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127	No tissue damage by chronic deep brain stimulation in Parkinson's disease. Annals of Neurology, 2000, 48, 372-6.	5.3	68
128	Selective Neuronal Vulnerability in Human Prion Diseases. American Journal of Pathology, 1999, 155, 1453-1457.	3.8	66
129	Deposition of disease-associated prion protein involves the peripheral nervous system in experimental scrapie. Acta Neuropathologica, 1999, 98, 453-457.	7.7	65
130	Fibroblasts Can Express Glial Fibrillary Acidic protein (GFAP) In Vivo. Journal of Neuropathology and Experimental Neurology, 2001, 60, 449-461.	1.7	65
131	The brain-specific protein TPPP/p25 in pathological protein deposits of neurodegenerative diseases. Acta Neuropathologica, 2007, 113, 153-161.	7.7	65
132	Vacuolar myelopathy with multinucleated giant cells in the acquired immune deficiency syndrome (AIDS). Acta Neuropathologica, 1989, 78, 497-503.	7.7	64
133	Suprasellar meningioma with expression of glial fibrillary acidic protein: a peculiar variant. Acta Neuropathologica, 1995, 90, 539-544.	7.7	64
134	Pathology and Immunocytochemistry of a Kuru Brain. Brain Pathology, 1997, 7, 547-553.	4.1	64
135	No evidence for cognitive dysfunction or depression in patients with mild restless legs syndrome. Movement Disorders, 2009, 24, 1843-1847.	3.9	64
136	Fibulin-5 mutations link inherited neuropathies, age-related macular degeneration and hyperelastic skin. Brain, 2011, 134, 1839-1852.	7.6	64
137	Progressive multifocal leukoencephalopathy (PML) in AIDS and in the pre-AIDS era. Acta Neuropathologica, 1990, 80, 375-380.	7.7	63
138	Rosette-forming glioneuronal tumor of the fourth ventricle. Acta Neuropathologica, 2003, 106, 506-508.	7.7	63
139	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.	7.7	62
140	Comparative analysis of NeuN immunoreactivity in primary brain tumours: conclusions for rational use in diagnostic histopathology. Histopathology, 2006, 48, 438-444.	2.9	62
141	Ultrastructural study of florid plaques in variant Creutzfeldtâ€“Jakob disease: a comparison with amyloid plaques in kuru, sporadic Creutzfeldtâ€“Jakob disease and Gerstmannâ€“StrÃusslerâ€“Scheinker disease. Neuropathology and Applied Neurobiology, 2009, 35, 46-59.	3.2	62
142	Scientific Opinion on Fish Oil for Human Consumption. Food Hygiene, including Rancidity. EFSA Journal, 2010, 8, 1874.	1.8	62
143	Supratentorial lobar ependymomas: Reports on the grading and survival periods in 80 cases, including 46 recurrences. Acta Neurochirurgica, 1983, 69, 243-251.	1.7	61
144	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.	7.7	61

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145	Tubulin polymerization promoting protein (TPPP/p25) as a marker for oligodendroglial changes in multiple sclerosis. Glia, 2010, 58, 1847-1857.	4.9	61
146	Tau pathology in Creutzfeldtâ€Jakob disease revisited. Brain Pathology, 2017, 27, 332-344.	4.1	61
147	Central nervous system pathology in patients with the Guillain-Barre syndrome. Brain, 1997, 120, 451-464.	7.6	59
148	Fatal Encephalitis Caused by Concomitant Infection with Tick-Borne Encephalitis Virus and Borrelia burgdorferi. Clinical Infectious Diseases, 1993, 16, 392-396.	5.8	57
149	The need to unify neuropathological assessments of vascular alterations in the ageing brain. Experimental Gerontology, 2012, 47, 825-833.	2.8	57
150	The Role of the NADPH Oxidase NOX2 in Prion Pathogenesis. PLoS Pathogens, 2014, 10, e1004531.	4.7	57
151	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.	5.3	55
152	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.	1.8	55
153	Disease associated prion protein may deposit in the peripheral nervous system in human transmissible spongiform encephalopathies. Acta Neuropathologica, 1999, 98, 458-460.	7.7	54
154	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.	3.2	54
155	Brain tissue immunoglobulins in adrenoleukodystrophy: A comparison with multiple sclerosis and systemic lupus erythematosus. Acta Neuropathologica, 1983, 59, 95-102.	7.7	53
156	Accumulation of 14-3-3 proteins in glial cytoplasmic inclusions in multiple system atrophy. Annals of Neurology, 2002, 52, 722-731.	5.3	53
157	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.	5.5	53
158	Nigral burden of Î±â€synuclein correlates with striatal dopamine deficit. Movement Disorders, 2008, 23, 1608-1612.	3.9	53
159	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.	7.7	52
160	Lipomatous differentiation in a medulloblastoma. Acta Neuropathologica, 1991, 81, 471-473.	7.7	52
161	Progressive multifocal leukoencephalopathy in AIDS: initial and follow-up CT and MRI. Neuroradiology, 1997, 39, 611-618.	2.2	52
162	Redox metals and oxidative abnormalities in human prion diseases. Acta Neuropathologica, 2005, 110, 232-238.	7.7	52

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163	Microglia is a component of the prion protein amyloid plaque in the Gerstmann-Strï¿½ussler-Scheinker syndrome. Acta Neuropathologica, 1993, 85, 623-627.	7.7	51
164	Pathogenesis of prion diseases. Acta Neuropathologica, 2005, 109, 32-48.	7.7	51
165	Stabilization of a Prion Strain of Synthetic Origin Requires Multiple Serial Passages. Journal of Biological Chemistry, 2012, 287, 30205-30214.	3.4	51
166	Interâ€Łaboratory Assessment of PrP^{Sc} Typing in Creutzfeldtâ€“Jakob Disease: A Western Blot Study within the NeuroPrion Consortium. Brain Pathology, 2009, 19, 384-391.	4.1	50
167	Deposition patterns of disease-associated prion protein in captive mule deer brains with chronic wasting disease. Acta Neuropathologica, 2001, 102, 496-500.	7.7	49
168	How a neuropsychiatric brain bank should be run: a consensus paper of Brainnet Europe II. Journal of Neural Transmission, 2007, 114, 527-537.	2.8	49
169	Intensity of human prion disease surveillance predicts observed disease incidence. Journal of Neurology, Neurosurgery and Psychiatry, 2013, 84, 1372-1377.	1.9	49
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