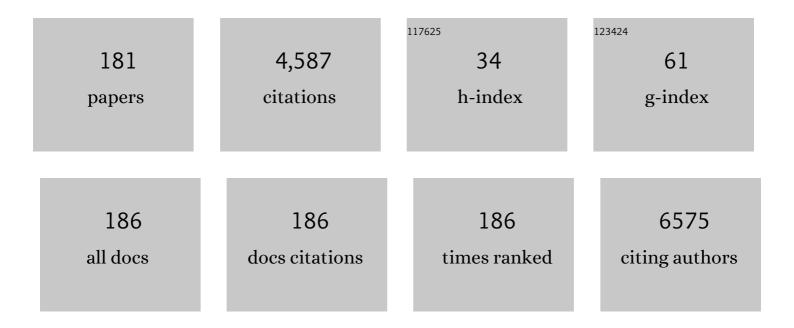
List of Publications by Year in descending order

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ΤοριιΙνναι

#	Article	IF	CITATIONS
1	Insulin resistance is associated with the pathology of Alzheimer disease. Neurology, 2010, 75, 764-770.	1.1	382
2	Altered Expression of Diabetes-Related Genes in Alzheimer's Disease Brains: The Hisayama Study. Cerebral Cortex, 2014, 24, 2476-2488.	2.9	294
3	Trends in dementia prevalence, incidence, and survival rate in a Japanese community. Neurology, 2017, 88, 1925-1932.	1.1	154
4	Comparative profiling of cortical gene expression in Alzheimer's disease patients and mouse models demonstrates a link between amyloidosis and neuroinflammation. Scientific Reports, 2017, 7, 17762.	3.3	138
5	Trends in prevalence of Alzheimer's disease and vascular dementia in a Japanese community: the Hisayama Study. Acta Psychiatrica Scandinavica, 2010, 122, 319-325.	4.5	123
6	Expression of 8-oxoguanine DNA glycosylase is reduced and associated with neurofibrillary tangles in Alzheimer's disease brain. Acta Neuropathologica, 2002, 103, 20-25.	7.7	122
7	Hakata Antigen, a New Member of the Ficolin/Opsonin p35 Family, Is a Novel Human Lectin Secreted into Bronchus/Alveolus and Bile. Journal of Histochemistry and Cytochemistry, 1999, 47, 777-785.	2.5	121
8	Differentiation of high-grade and low-grade diffuse gliomas by intravoxel incoherent motion MR imaging. Neuro-Oncology, 2016, 18, 132-141.	1.2	109
9	Molecular pathophysiology of impaired glucose metabolism, mitochondrial dysfunction, and oxidative DNA damage in Alzheimer's disease brain. Mechanisms of Ageing and Development, 2017, 161, 95-104.	4.6	105
10	Sense and antisense modification of glial alpha B-crystallin production results in alterations of stress fiber formation and thermoresistance Journal of Cell Biology, 1994, 125, 1385-1393.	5.2	92
11	Association of Alzheimer disease pathology with abnormal lipid metabolism. Neurology, 2011, 77, 1068-1075.	1.1	92
12	Multiple system degeneration with basophilic inclusions in Japanese ALS patients with FUS mutation. Acta Neuropathologica, 2010, 119, 355-364.	7.7	90
13	Grading diffuse gliomas without intense contrast enhancement by amide proton transfer MR imaging: comparisons with diffusion- and perfusion-weighted imaging. European Radiology, 2017, 27, 578-588.	4.5	90
14	Connexin 43 Astrocytopathy Linked to Rapidly Progressive Multiple Sclerosis and Neuromyelitis Optica. PLoS ONE, 2013, 8, e72919.	2.5	89
15	Clusterin/apolipoprotein J is associated with cortical Lewy bodies: immunohistochemical study in cases with 1±-synucleinopathies. Acta Neuropathologica, 2002, 104, 225-230.	7.7	75
16	Epithelial properties of pleomorphic xanthoastrocytomas determined in ultrastructural and immunohistochemical studies. Acta Neuropathologica, 1987, 74, 142-150.	7.7	74
17	Preferential expression of αB-crystallin in astrocytic elements of neuroectodermal tumors. Cancer, 1991, 68, 2230-2240.	4.1	69
18	α-Synuclein is expressed in a variety of brain tumors showing neuronal differentiation. Acta Neuropathologica, 2000, 99, 154-160.	7.7	69

#	Article	IF	CITATIONS
19	Prevalence and clinicopathological features of H3.3 G34-mutant high-grade gliomas: a retrospective study of 411 consecutive glioma cases in a single institution. Brain Tumor Pathology, 2017, 34, 103-112.	1.7	69

Distinctive immunohistochemical profiles of small heat shock proteins (Heat shock protein 27 and) Tj ETQq0 0 0 rg $_{4.1}^{BT}$ /Overlock 10 Tf 50

21	A comparative immunohistochemical study of Kuru and senile plaques with a special reference to glial reactions at various stages of amyloid plaque formation. American Journal of Pathology, 1991, 139, 589-98.	3.8	62
22	Amyloid imaging probes are useful for detection of prion plaques and treatment of transmissible spongiform encephalopathies. Journal of General Virology, 2004, 85, 1785-1790.	2.9	58
23	Extensive loss of connexins in Baló's disease: evidence for an auto-antibody-independent astrocytopathy via impaired astrocyte–oligodendrocyte/myelin interaction. Acta Neuropathologica, 2012, 123, 887-900.	7.7	57
24	Midlife and Lateâ€Life Smoking and Risk of Dementia in the Community: The Hisayama Study. Journal of the American Geriatrics Society, 2015, 63, 2332-2339.	2.6	56
25	Chordoma in Early Childhood: A Clinicopathological Study. Neurosurgery, 1991, 29, 442-446.	1.1	55
26	Aquaporin-4 astrocytopathy in Baló's disease. Acta Neuropathologica, 2010, 120, 651-660.	7.7	53
27	Expression of hMTH1 in the hippocampi of control and Alzheimer's disease. NeuroReport, 2001, 12, 2895-2899.	1.2	49
28	Amplification and Overexpression of mdm2 Gene in Ependymomas. Modern Pathology, 2000, 13, 548-553.	5.5	43
29	Differential Expression of Metallothioneins in Human Prion Diseases. Dementia and Geriatric Cognitive Disorders, 2000, 11, 251-262.	1.5	41
30	Reappraisal of Aquaporinâ€4 Astrocytopathy in Asian Neuromyelitis Optica and Multiple Sclerosis Patients. Brain Pathology, 2011, 21, 516-532.	4.1	41
31	Loss of <scp>hnRNPA1</scp> in <scp>ALS</scp> spinal cord motor neurons with <scp>TDP</scp> â€43â€positive inclusions. Neuropathology, 2015, 35, 37-43.	1.2	41
32	Clinicopathological review of solitary fibrous tumors: dedifferentiation is a major cause of patient death. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2019, 475, 467-477.	2.8	40
33	Cognitive dysfunction in patients with amyotrophic lateral sclerosis is associated with spherical or crescent-shaped ubiquitinated intraneuronal inclusions in the parahippocampal gyrus and amygdala, but not in the neostriatum. Acta Neuropathologica, 2001, 102, 467-472.	7.7	35
34	Diagnostic utility of intravoxel incoherent motion mr imaging in differentiating primary central nervous system lymphoma from glioblastoma multiforme. Journal of Magnetic Resonance Imaging, 2016, 44, 1256-1261.	3.4	35
35	Skein-like inclusions in the neostriatum from a case of amyotrophic lateral sclerosis with dementia. Acta Neuropathologica, 1998, 96, 541-545.	7.7	34
36	An atypical case of sporadic Creutzfeldt-Jakob disease with Parkinson's disease. Neuropathology, 2001, 21, 294-297.	1.2	33

#	Article	IF	CITATIONS
37	A comprehensive analysis identifies <i>BRAF</i> hotspot mutations associated with gliomas with peculiar epithelial morphology. Neuropathology, 2017, 37, 191-199.	1.2	33
38	Widespread distribution of tau in the astrocytic elements of glial tumors. Acta Neuropathologica, 1993, 86, 236-241.	7.7	32
39	Immunohistochemical demonstration of alphaB-crystallin in hamartomas of tuberous sclerosis. American Journal of Pathology, 1991, 139, 1303-8.	3.8	32
40	Allelic Losses of Chromosome 10 in Glioma Tissues Detected by Quantitative Single-Strand Conformation Polymorphism Analysis. Clinical Chemistry, 2006, 52, 370-378.	3.2	31
41	Discrepancy in Programmed Cell Death-Ligand 1 Between Primary and Metastatic Non-small Cell Lung Cancer. Anticancer Research, 2017, 37, 4223-4228.	1.1	30
42	Immunohistochemical analysis of spinal cord lesions in amyotrophic lateral sclerosis using microtubule-associated protein 2 (MAP2) antibodies. Acta Neuropathologica, 1999, 97, 13-21.	7.7	29
43	Prostaglandin D Synthase (β-Trace) in Meningeal Hemangiopericytoma. Modern Pathology, 2001, 14, 197-201.	5.5	29
44	Association of adipocyte enhancerâ€binding protein 1 with <scp>A</scp> lzheimer's disease pathology in human hippocampi. Brain Pathology, 2018, 28, 58-71.	4.1	28
45	Detection of SV40 T antigen genome in human gliomas. Brain Tumor Pathology, 1997, 14, 125-129.	1.7	27
46	A Simple VNTR-PCR Method for Detecting Maternal Cell Contamination in Prenatal Diagnosis. Genetic Testing and Molecular Biomarkers, 1998, 2, 347-350.	1.7	27
47	Molecular diagnosis of diffuse glioma using a chip-based digital PCR system to analyze IDH, TERT, and H3 mutations in the cerebrospinal fluid. Journal of Neuro-Oncology, 2021, 152, 47-54.	2.9	27
48	Early and extensive spinal white matter involvement in neuromyelitis optica. Brain Pathology, 2017, 27, 249-265.	4.1	26
49	Advanced glycosylation endâ€products and heat shock proteins accumulate in the basophilic degeneration of the myocardium and the corpora amylacea of the glia. Pathology International, 1996, 46, 757-763.	1.3	25
50	Autopsy case of autosomal recessive hereditary spastic paraplegia with reference to the muscular pathology. Neuropathology, 2001, 21, 212-217.	1.2	25
51	Correlation between arterial spin-labeling perfusion and histopathological vascular density of pediatric intracranial tumors. Journal of Neuro-Oncology, 2017, 135, 561-569.	2.9	25
52	Measurement of the perfusion fraction in brain tumors with intravoxel incoherent motion MR imaging: validation with histopathological vascular density in meningiomas. British Journal of Radiology, 2018, 91, 20170912.	2.2	25
53	Expression of the lysosome-associated membrane proteins in myopathies with rimmed vacuoles. Acta Neuropathologica, 2001, 101, 579-584.	7.7	24
54	Defense mechanism to oxidative DNA damage in glial cells. Neuropathology, 2004, 24, 125-130.	1.2	23

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55	Preferential neurodegeneration in the cervical spinal cord of progressive supranuclear palsy. Acta Neuropathologica, 1999, 97, 577-584.	7.7	22
56	An autopsied case of sporadic adult-onset amyotrophic lateral sclerosis with FUS-positive basophilic inclusions. Neuropathology, 2011, 31, 71-76.	1.2	22
57	Downâ€regulation of <scp>MET</scp> in hippocampal neurons of <scp>A</scp> lzheimer's disease brains. Neuropathology, 2014, 34, 284-290.	1.2	22
58	Quantitative digital assessment of MGMT immunohistochemical expression in glioblastoma tissue. Brain Tumor Pathology, 2011, 28, 25-31.	1.7	21
59	Trends in autopsyâ€verified dementia prevalence over 29Âyears of the Hisayama study. Neuropathology, 2016, 36, 383-387.	1.2	21
60	Clinical significance of <i>CDKN2A</i> homozygous deletion in combination with methylated <i>MGMT</i> status for <i>IDH</i> â€wildtype glioblastoma. Cancer Medicine, 2021, 10, 3177-3187.	2.8	21
61	Predicting TERT promoter mutation using MR images in patients with wild-type IDH1 glioblastoma. Diagnostic and Interventional Imaging, 2019, 100, 411-419.	3.2	20
62	Reclassification of 400 consecutive glioma cases based on the revised 2016WHO classification. Brain Tumor Pathology, 2018, 35, 81-89.	1.7	19
63	Clinical Significance of PD-L1 Expression in Brain Metastases from Non-small Cell Lung Cancer. Anticancer Research, 2018, 38, 553-557.	1.1	19
64	An immunohistochemical study of tissue transglutaminase in gliomas with reference to their cell dying processes. American Journal of Pathology, 1994, 145, 776-81.	3.8	19
65	Cell kinetics of the malignant evolution of meningothelial meningioma. Acta Neuropathologica, 1987, 74, 243-247.	7.7	18
66	Extensive distribution of glial cytoplasmic inclusions in an autopsied case of multiple system atrophy with a prolonged 18â€year clinical course. Neuropathology, 2012, 32, 69-76.	1.2	18
67	DCTN1 F52L mutation case of Perry syndrome with progressive supranuclear palsy-like tauopathy. Parkinsonism and Related Disorders, 2018, 51, 105-110.	2.2	18
68	Highâ€resolution melting and immunohistochemical analysis efficiently detects mutually exclusive genetic alterations of adamantinomatous and papillary craniopharyngiomas. Neuropathology, 2018, 38, 3-10.	1.2	18
69	Diagnostic accuracy for the epileptogenic zone detection in focal epilepsy could be higher in FDG-PET/MRI than in FDG-PET/CT. European Radiology, 2021, 31, 2915-2922.	4.5	18
70	Immunohistochemistry of chondromodulin-I in the human intervertebral discs with special reference to the degenerative changes. The Histochemical Journal, 2000, 32, 545-550.	0.6	17
71	MUTYH Actively Contributes to Microglial Activation and Impaired Neurogenesis in the Pathogenesis of Alzheimer's Disease. Oxidative Medicine and Cellular Longevity, 2021, 2021, 1-30.	4.0	17
72	Preferential involvement of the short arm in chromosome 8-derived supernumerary markers and ring as identified by chromosome arm painting. American Journal of Medical Genetics Part A, 2000, 90, 276-282.	2.4	16

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73	Intravoxel Incoherent Motion MR Imaging of Pediatric Intracranial Tumors: Correlation with Histology and Diagnostic Utility. American Journal of Neuroradiology, 2019, 40, 878-884.	2.4	16
74	MOG antibody disease manifesting as progressive cognitive deterioration and behavioral changes with primary central nervous system vasculitis. Multiple Sclerosis and Related Disorders, 2019, 30, 48-50.	2.0	16
75	Ubiquitin-immunoreactive skein-like inclusions in the neostriatum are not restricted to amyotrophic lateral sclerosis, but are rather aging-related structures. Acta Neuropathologica, 2000, 100, 43-49.	7.7	15
76	Increased asymmetric pulvinar magnetic resonance imaging signals in Creutzfeldt-Jakob disease with florid plaques following a cadaveric dura mater graft. Neuropathology, 2006, 26, 82-88.	1.2	15
77	Tauopathy in basal ganglia involvement is exacerbated in a subset of patients with Alzheimer's disease: The Hisayama study. Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring, 2019, 11, 415-423.	2.4	15
78	Expanded polyglutamine impairs normal nuclear distribution of fused in sarcoma and poly (rC)â€binding protein 1 in Huntington's disease. Neuropathology, 2019, 39, 358-367.	1.2	15
79	Case of lipoblastoma with two derivative chromosomes 8 containing homogeneously staining-like regions and a review of the literature:. Cancer Genetics and Cytogenetics, 2001, 125, 10-13.	1.0	14
80	Radiological Features of Brain Metastases from Non-small Cell Lung Cancer Harboring <i>EGFR</i> Mutation. Anticancer Research, 2018, 38, 3731-3734.	1.1	14
81	Establishment and Characterization of Choroid Plexus Carcinoma Cell Lines: Connection between Choroid Plexus and Immune Systems. Japanese Journal of Cancer Research, 1996, 87, 893-899.	1.7	13
82	EWS/FLI-1 fusion signal inserted into chromosome 11 in one patient with morphologic features of Ewing sarcoma, but lacking t(11;22). Cancer Genetics and Cytogenetics, 2002, 133, 72-75.	1.0	13
83	An astroblastoma case associated with loss of heterozygosity on chromosome 9p. Journal of Neuro-Oncology, 2006, 80, 69-73.	2.9	13
84	Sporadic <scp>C</scp> reutzfeldt– <scp>J</scp> akob Disease <scp>MM1+2C</scp> and <scp>MM</scp> 1 are Identical in Transmission Properties. Brain Pathology, 2016, 26, 95-101.	4.1	13
85	CD206 Expression in Induced Microglia-Like Cells From Peripheral Blood as a Surrogate Biomarker for the Specific Immune Microenvironment of Neurosurgical Diseases Including Glioma. Frontiers in Immunology, 2021, 12, 670131.	4.8	13
86	A case of intracranial solitary fibrous tumor/hemangiopericytoma with dedifferentiated component. Neuropathology, 2015, 35, 260-265.	1.2	12
87	Recent Increases in Hippocampal Tau Pathology in the Aging Japanese Population: The Hisayama Study. Journal of Alzheimer's Disease, 2016, 55, 613-624.	2.6	12
88	Insular primary glioblastomas with <i>IDH</i> mutations: Clinical and biological specificities. Neuropathology, 2017, 37, 200-206.	1.2	12
89	Pediatric ganglioglioma with an H3 K27M mutation arising from the cervical spinal cord. Neuropathology, 2018, 38, 422-427.	1.2	12
90	Dynactin is involved in Lewy body pathology. Neuropathology, 2018, 38, 583-590.	1.2	12

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91	Differentiation of high-grade from low-grade diffuse gliomas using diffusion-weighted imaging: a comparative study of mono-, bi-, and stretched-exponential diffusion models. Neuroradiology, 2020, 62, 815-823.	2.2	12
92	Hemangiopericytoma of the meninges: a clinicopathologic and immunohistochemical study. , 1988, 7, 93-9.		12
93	IgH Intronic Enhancer Element HE2 (μB) Functions as a cis-Activator in Choroid Plexus Cells at the Cellular Level as well as in Transgenic Mice. Journal of Neurochemistry, 2002, 64, 961-966.	3.9	11
94	Deferred radiotherapy and upfront procarbazine–ACNU–vincristine administration for 1p19q codeleted oligodendroglial tumors are associated with favorable outcome without compromising patient performance, regardless of WHO grade. OncoTargets and Therapy, 2016, Volume 9, 7123-7131.	2.0	11
95	â€~PrP systemic deposition disease': clinical and pathological characteristics of novel familial prion disease with 2â€bp deletion in codon 178. European Journal of Neurology, 2016, 23, 196-200.	3.3	11
96	Distinct microglial and macrophage distribution patterns in the concentric and lamellar lesions in BalÃ3's disease and neuromyelitis optica spectrum disorders. Brain Pathology, 2020, 30, 1144-1157.	4.1	11
97	C-Terminal-Deleted Prion Protein Fragment Is a Major Accumulated Component of Systemic PrP Deposits in Hereditary Prion Disease With a 2-Bp (CT) Deletion in <i>PRNP</i> Codon 178. Journal of Neuropathology and Experimental Neurology, 2016, 75, 1008-1019.	1.7	10
98	Arterial spin-labeling is useful for the diagnosis of residual or recurrent meningiomas. European Radiology, 2018, 28, 4334-4342.	4.5	10
99	Accumulation of Astrocytic Aquaporin 4 and Aquaporin 1 in Prion Protein Plaques. Journal of Neuropathology and Experimental Neurology, 2020, 79, 419-429.	1.7	10
100	Abnormal prion protein deposits with high seeding activities in the skeletal muscle, femoral nerve, and scalp of an autopsied case of sporadic Creutzfeldt–Jakob disease. Neuropathology, 2021, 41, 152-158.	1.2	10
101	Microsphere formation in a subtype of <scp>C</scp> reutzfeldt– <scp>J</scp> akob disease with a <scp>V180I</scp> mutation and codon 129 <scp>MM</scp> polymorphism. Neuropathology and Applied Neurobiology, 2013, 39, 844-848.	3.2	9
102	Upregulation of Annexin A1 in Reactive Astrocytes and Its Subtle Induction in Microglia at the Boundaries of Human Brain Infarcts. Journal of Neuropathology and Experimental Neurology, 2019, 78, 961-970.	1.7	9
103	Transactivation response DNAâ€binding protein of 43 kDa proteinopathy and lysosomal abnormalities in spastic paraplegia type 11. Neuropathology, 2021, 41, 253-265.	1.2	9
104	Leser-Trélat sign with anaplastic ependymoma - an autopsy case. Acta Neuropathologica, 1997, 93, 97-100.	7.7	8
105	Forced retraction of spinal root injury enhances activation of p38 MAPK cascade in infiltrating macrophages. Neuropathology, 2005, 25, 37-47.	1.2	8
106	ABL1 gene involvement within a complex three-way translocation (2;9;4) in perineurioma characterized by molecular cytogenetic methods. Cancer Genetics, 2014, 207, 263-267.	0.4	8
107	Dura mater graftâ€associated Creutzfeldtâ€Jakob disease with 30â€year incubation period. Neuropathology, 2017, 37, 275-281.	1.2	8
108	A Novel Combination of Prion Strain Co-Occurrence in Patients with Sporadic Creutzfeldt-Jakob Disease. American Journal of Pathology, 2019, 189, 1276-1283.	3.8	8

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109	Co-existence of alternative forms of 8q gain in cytogenetic clones of three patients with acute myeloid leukemia, pointing to 8q22â^1⁄48qter as a region of biologic significance. Cancer Genetics and Cytogenetics, 2001, 126, 20-25.	1.0	7
110	Sporadic hereditary diffuse leukoencephalopathy with axonal spheroids showing numerous lesions with restricted diffusivity caused by a novel splice site mutation in the <i><scp>CSF</scp>1R</i> gene. Clinical and Experimental Neuroimmunology, 2013, 4, 76-81.	1.0	7
111	Expression of CRYM in different rat organs during development and its decreased expression in degenerating pyramidal tracts in amyotrophic lateral sclerosis. Neuropathology, 2018, 38, 247-259.	1.2	7
112	Relevance of calcification and contrast enhancement pattern for molecular diagnosis and survival prediction of gliomas based on the 2016 World Health Organization Classification. Clinical Neurology and Neurosurgery, 2019, 187, 105556.	1.4	7
113	Toxic myopathy with multiple deletions in mitochondrial DNA associated with longâ€ŧerm use of oral antiâ€viral drugs for hepatitis B: A case study. Neuropathology, 2019, 39, 162-167.	1.2	6
114	PCBP2 Is Downregulated in Degenerating Neurons and Rarely Observed in TDP-43-Positive Inclusions in Sporadic Amyotrophic Lateral Sclerosis. Journal of Neuropathology and Experimental Neurology, 2021, 80, 220-228.	1.7	6
115	Symmetrical glial hyperplasia in the brainstem of fibrodysplasia ossificans progressiva. Neuropathology, 2021, 41, 146-151.	1.2	6
116	Clinical implications of molecular analysis in diffuse glioma stratification. Brain Tumor Pathology, 2021, 38, 210-217.	1.7	6
117	Concurrent cardiac transthyretin and brain \hat{l}^2 amyloid accumulation among the older adults: The Hisayama study. Brain Pathology, 2021, , e13014.	4.1	6
118	Different responses of benign and atypical meningiomas to gamma-knife radiosurgery: report of two cases with immunohistochemical analysis. Brain Tumor Pathology, 2001, 18, 61-66.	1.7	5
119	Unusual aberration involving the short arm of chromosome 11 in an 8-month-old patient with a supratentorial primitive neuroectodermal tumor. Cancer Genetics and Cytogenetics, 2003, 141, 143-147.	1.0	5
120	Coexistence of neocentromeric marker 3q and trisomy 3 in two different tissues in a 3-year-old boy with peripheral T-cell lymphoma: support for a gene dosage effect hypothesis. Cancer Genetics and Cytogenetics, 2006, 170, 152-157.	1.0	5
121	Accumulation of class I mutant p53 and apoptosis induced by carboplatin in a human glioma cell line. Brain Tumor Pathology, 1998, 15, 77-82.	1.7	4
122	Different Complicated Brain Pathologies in Monozygotic Twins With Gerstmann–StrÃ ¤ ssler–Scheinker Disease. Journal of Neuropathology and Experimental Neurology, 2017, 76, 854-863.	1.7	4
123	Fourâ€repeat tau dominant pathology in a congenital myotonic dystrophy type 1 patient with mental retardation. Brain Pathology, 2018, 28, 431-433.	4.1	4
124	Mitochondrial dysfunction and altered ribostasis in hippocampal neurons with cytoplasmic inclusions of multiple system atrophy. Neuropathology, 2018, 38, 361-371.	1.2	4
125	Differences between primary central nervous system lymphoma and glioblastoma: topographic analysis using voxel-based morphometry. Clinical Radiology, 2019, 74, 816.e1-816.e8.	1.1	4
126	Frequent Detection of Pituitary-Derived PrPres in Human Prion Diseases. Journal of Neuropathology and Experimental Neurology, 2019, 78, 922-929.	1.7	4

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127	A case of overlapping adultâ€onset linear scleroderma and Parryâ€Romberg syndrome presenting with widespread ipsilateral neurogenic involvement. Neuropathology, 2020, 40, 109-115.	1.2	4
128	Alectinibâ€responsive infantile anaplastic ganglioglioma with a novel <i>VCL–ALK</i> gene fusion. Pediatric Blood and Cancer, 2021, 68, e29122.	1.5	4
129	Histological background of dedifferentiated solitary fibrous tumour. Journal of Clinical Pathology, 2022, 75, 397-403.	2.0	4
130	Endonasal endoscopic surgery for temporal lobe epilepsy associated with sphenoidal encephalocele. , 2021, 12, 379.		4
131	A case of ganglioglioma grade 3 with <scp>H3 K27M</scp> mutation arising in the medial temporal lobe in an elderly patient. Neuropathology, 2022, , .	1.2	4
132	Quantitative relaxometry using synthetic MRI could be better than T2-FLAIR mismatch sign for differentiation of IDH-mutant gliomas: a pilot study. Scientific Reports, 2022, 12, .	3.3	4
133	A comparative immunohistochemical study of tissue transglutaminase and factor XIIIa in hemangioblastoma. Neuropathology, 1998, 18, 199-205.	1.2	3
134	An elderly case of malignant small cell glioma with hemorrhage coexistent with a calcified pilocytic astrocytoma component in the cerebellar hemisphere. Neuropathology, 2018, 38, 493-497.	1.2	3
135	A juvenile case of epilepsyâ€associated, isocitrate dehydrogenase wildâ€type/histone 3 wildâ€type diffuse glioma with a rare BRAF A598T mutation. Neuropathology, 2020, 40, 646-650.	1.2	3
136	Immunotherapyâ€refractory vacuolar myopathy with mucin deposition in scleromyxedema: A possible role of fibroblast growth factor 2. Neuropathology, 2020, 40, 492-495.	1.2	3
137	Primary Pineal Yolk Sac Tumor: Ultrastructural and Immunohistochemical Features. Neurologia Medico-Chirurgica, 1986, 26, 564-570.	2.2	2
138	Hypothalamic mass in a 28â€yearâ€old man with diabetes insipidus ataxia, nystagmus and dysarthria. Neuropathology, 2001, 21, 99-100.	1.2	2
139	An intragenic deletion of the gene <i>MNAT1</i> in a family with pectus deformities. American Journal of Medical Genetics, Part A, 2014, 164, 1293-1297.	1.2	2
140	Predictors of recurrence and postoperative outcomes in patients with non-skull base meningiomas based on modern neurosurgical standards. Interdisciplinary Neurosurgery: Advanced Techniques and Case Management, 2019, 15, 30-37.	0.3	2
141	Intraventricular mucinâ€producing glioblastoma arising in the septum pellucidum at the frontal horn of the lateral ventricle: A case report. Neuropathology, 2021, 41, 381-386.	1.2	2
142	Distinctive immunohistochemical profiles of small heat shock proteins (Heat shock protein 27 and) Tj ETQq0 0 () rgBT /Ove	erlock 10 Tf 5
143	Cytoplasmic inclusions of astrocytic elements of glial tumors: special reference to round granulated body and eosinophilic hyaline droplets. Acta Neuropathologica, 1994, 88, 501-510.	7.7	2

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#	Article	IF	CITATIONS
145	Prion Gene PRNP Y162X Truncation Mutation Can Induce a Refractory Esophageal Achalasia. American Journal of Gastroenterology, 2021, 116, 1350-1351.	0.4	2
146	Changes in the Relapse Pattern and Prognosis of Glioblastoma After Approval of First-Line Bevacizumab: A Single-Center Retrospective Study. World Neurosurgery, 2022, 159, e479-e487.	1.3	2
147	The 92nd Neuropathological Meeting of Kyushu District 22 July 2006. Neuropathology, 2006, 26, 597-597.	1.2	1
148	The 97th Neuropathological Meeting of Kyushu DistrictÃ⁻Âį¼26 December 2008. Neuropathology, 2009, 29, 207-207.	1.2	1
149	Masked hypodiploidy in anaplastic meningiomas by duplication of the original clone found in atypical meningiomas: Illustration of the evolution of genetic alterations. Neuropathology, 2014, 34, 353-359.	1.2	1
150	Spindle cell/sclerosing rhabdomyosarcoma with intracranial invasion without destroying the bone of the skull base: a case report and literature review. Acta Radiologica Open, 2017, 6, 205846011772731.	0.6	1
151	Detection of cutaneous prion protein deposits could help diagnose GPIâ€anchorless prion disease with neuropathy. European Journal of Neurology, 2021, 28, 2133-2137.	3.3	1
152	Acute aortic dissection associated with wildâ€ŧype transthyretin amyloid. Pathology International, 2021, 71, 556-558.	1.3	1
153	Papillary craniopharyngioma coexisting with an intratumoral abscess in a pediatric patient: A case report and review of the literature. Acta Radiologica Open, 2021, 10, 205846012110306.	0.6	1
154	Optic nerve atrophy and visual disturbance following PRNP Y162X truncation mutation. Journal of the Neurological Sciences, 2021, 428, 117614.	0.6	1
155	The Combined Use of Composite Ceramic Granules and Fibrin Glue for Cranioplasties : Results of a Rat Model Study and Clinical Findings with Regard to Biocompatibility. Japanese Journal of Neurosurgery, 1995, 4, 543-547.	0.0	1
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