

Toru Iwaki

List of Publications by Year in descending order

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Version: 2024-02-01

181
papers

4,587
citations

117625

34
h-index

123424

61
g-index

186
all docs

186
docs citations

186
times ranked

6575
citing authors

#	ARTICLE	IF	CITATIONS
1	Histological background of dedifferentiated solitary fibrous tumour. <i>Journal of Clinical Pathology</i> , 2022, 75, 397-403.	2.0	4
2	A Case of Cerebral Tuberculoma with Mild Posterior Cervical Pain as the Main Symptom Despite Extensive Brain Lesions. <i>Internal Medicine</i> , 2022, , .	0.7	0
3	A case of ganglioglioma grade 3 with <sc>H3 K27M</sc> mutation arising in the medial temporal lobe in an elderly patient. <i>Neuropathology</i> , 2022, , .	1.2	4
4	Changes in the Relapse Pattern and Prognosis of Glioblastoma After Approval of First-Line Bevacizumab: A Single-Center Retrospective Study. <i>World Neurosurgery</i> , 2022, 159, e479-e487.	1.3	2
5	A Comparative Study of Site-Specific Distribution of Aging-Related Tau Astroglialopathy and Its Risk Factors Between Alzheimer Disease and Cognitive Healthy Brains: The Hisayama Study. <i>Journal of Neuropathology and Experimental Neurology</i> , 2022, 81, 106-116.	1.7	1
6	Quantitative relaxometry using synthetic MRI could be better than T2-FLAIR mismatch sign for differentiation of IDH-mutant gliomas: a pilot study. <i>Scientific Reports</i> , 2022, 12, .	3.3	4
7	Diagnostic accuracy for the epileptogenic zone detection in focal epilepsy could be higher in FDG-PET/MRI than in FDG-PET/CT. <i>European Radiology</i> , 2021, 31, 2915-2922.	4.5	18
8	PCBP2 Is Downregulated in Degenerating Neurons and Rarely Observed in TDP-43-Positive Inclusions in Sporadic Amyotrophic Lateral Sclerosis. <i>Journal of Neuropathology and Experimental Neurology</i> , 2021, 80, 220-228.	1.7	6
9	Ruptured anterior communicating artery aneurysms associated with left common carotid artery occlusion due to Takayasu arteritis: an autopsy case report. <i>Nosotchu</i> , 2021, , .	0.1	0
10	Molecular diagnosis of diffuse glioma using a chip-based digital PCR system to analyze IDH, TERT, and H3 mutations in the cerebrospinal fluid. <i>Journal of Neuro-Oncology</i> , 2021, 152, 47-54.	2.9	27
11	Symmetrical glial hyperplasia in the brainstem of fibrodysplasia ossificans progressiva. <i>Neuropathology</i> , 2021, 41, 146-151.	1.2	6
12	Detection of cutaneous prion protein deposits could help diagnose GPIâ€ anchorless prion disease with neuropathy. <i>European Journal of Neurology</i> , 2021, 28, 2133-2137.	3.3	1
13	Abnormal prion protein deposits with high seeding activities in the skeletal muscle, femoral nerve, and scalp of an autopsied case of sporadic Creutzfeldtâ€ Jakob disease. <i>Neuropathology</i> , 2021, 41, 152-158.	1.2	10
14	Clinical significance of <i>CDKN2A</i> homozygous deletion in combination with methylated <i>MGMT</i> status for <i>IDH</i>â€wildtype glioblastoma. <i>Cancer Medicine</i> , 2021, 10, 3177-3187.	2.8	21
15	Alectinibâ€responsive infantile anaplastic ganglioglioma with a novel <i>VCLâ€ALK</i> gene fusion. <i>Pediatric Blood and Cancer</i> , 2021, 68, e29122.	1.5	4
16	Transactivation response DNAâ€binding protein of 43 kDa proteinopathy and lysosomal abnormalities in spastic paraplegia type 11. <i>Neuropathology</i> , 2021, 41, 253-265.	1.2	9
17	Acute aortic dissection associated with wildâ€type transthyretin amyloid. <i>Pathology International</i> , 2021, 71, 556-558.	1.3	1
18	CD206 Expression in Induced Microglia-Like Cells From Peripheral Blood as a Surrogate Biomarker for the Specific Immune Microenvironment of Neurosurgical Diseases Including Glioma. <i>Frontiers in Immunology</i> , 2021, 12, 670131.	4.8	13

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19	Hemorrhagic stroke due to leukostasis in pediatric mixed-phenotype acute leukemia. <i>Blood Research</i> , 2021, 56, 60-60.	1.3	0
20	Clinical implications of molecular analysis in diffuse glioma stratification. <i>Brain Tumor Pathology</i> , 2021, 38, 210-217.	1.7	6
21	Endonasal endoscopic surgery for temporal lobe epilepsy associated with sphenoidal encephalocele. , 2021, 12, 379.		4
22	Papillary craniopharyngioma coexisting with an intratumoral abscess in a pediatric patient: A case report and review of the literature. <i>Acta Radiologica Open</i> , 2021, 10, 205846012110306.	0.6	1
23	Intraventricular mucin-producing glioblastoma arising in the septum pellucidum at the frontal horn of the lateral ventricle: A case report. <i>Neuropathology</i> , 2021, 41, 381-386.	1.2	2
24	Concurrent cardiac transthyretin and brain β^2 amyloid accumulation among the older adults: The Hisayama study. <i>Brain Pathology</i> , 2021, , e13014.	4.1	6
25	Optic nerve atrophy and visual disturbance following PRNP Y162X truncation mutation. <i>Journal of the Neurological Sciences</i> , 2021, 428, 117614.	0.6	1
26	Acute-phase electroencephalography for an infantile atypical teratoid/rhabdoid tumor. <i>Clinical Neurology and Neurosurgery</i> , 2021, 209, 106922.	1.4	0
27	Prion Gene PRNP Y162X Truncation Mutation Can Induce a Refractory Esophageal Achalasia. <i>American Journal of Gastroenterology</i> , 2021, 116, 1350-1351.	0.4	2
28	MUTYH Actively Contributes to Microglial Activation and Impaired Neurogenesis in the Pathogenesis of Alzheimer's Disease. <i>Oxidative Medicine and Cellular Longevity</i> , 2021, 2021, 1-30.	4.0	17
29	A case of overlapping adult-onset linear scleroderma and Parry-Romberg syndrome presenting with widespread ipsilateral neurogenic involvement. <i>Neuropathology</i> , 2020, 40, 109-115.	1.2	4
30	A juvenile case of epilepsy-associated, isocitrate dehydrogenase wild-type/histone 3 wild-type diffuse glioma with a rare BRAF A598T mutation. <i>Neuropathology</i> , 2020, 40, 646-650.	1.2	3
31	Distinct microglial and macrophage distribution patterns in the concentric and lamellar lesions in Baló's disease and neuromyelitis optica spectrum disorders. <i>Brain Pathology</i> , 2020, 30, 1144-1157.	4.1	11
32	Differentiation of high-grade from low-grade diffuse gliomas using diffusion-weighted imaging: a comparative study of mono-, bi-, and stretched-exponential diffusion models. <i>Neuroradiology</i> , 2020, 62, 815-823.	2.2	12
33	Immunotherapy-refractory vacuolar myopathy with mucin deposition in scleromyxedema: A possible role of fibroblast growth factor 2. <i>Neuropathology</i> , 2020, 40, 492-495.	1.2	3
34	Intronic variant in IQGAP3 associated with hereditary neuropathy with proximal lower dominance, urinary disturbance, and paroxysmal dry cough. <i>Journal of Human Genetics</i> , 2020, 65, 717-725.	2.3	0
35	Accumulation of Astrocytic Aquaporin 4 and Aquaporin 1 in Prion Protein Plaques. <i>Journal of Neuropathology and Experimental Neurology</i> , 2020, 79, 419-429.	1.7	10
36	HGG-24. HIGH-GRADE GLIOMA WITH A NOVEL FUSION GENE OF VCL-ALK. <i>Neuro-Oncology</i> , 2020, 22, iii348-iii348.	1.2	2

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37	Clinicopathological review of solitary fibrous tumors: dedifferentiation is a major cause of patient death. <i>Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin</i> , 2019, 475, 467-477.	2.8	40
38	Differences between primary central nervous system lymphoma and glioblastoma: topographic analysis using voxel-based morphometry. <i>Clinical Radiology</i> , 2019, 74, 816.e1-816.e8.	1.1	4
39	Tauopathy in basal ganglia involvement is exacerbated in a subset of patients with Alzheimer's disease: The Hisayama study. <i>Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring</i> , 2019, 11, 415-423.	2.4	15
40	Relevance of calcification and contrast enhancement pattern for molecular diagnosis and survival prediction of gliomas based on the 2016 World Health Organization Classification. <i>Clinical Neurology and Neurosurgery</i> , 2019, 187, 105556.	1.4	7
41	Expanded polyglutamine impairs normal nuclear distribution of fused in sarcoma and poly (rC)â€binding protein 1 in Huntington's disease. <i>Neuropathology</i> , 2019, 39, 358-367.	1.2	15
42	Frequent Detection of Pituitary-Derived PrPres in Human Prion Diseases. <i>Journal of Neuropathology and Experimental Neurology</i> , 2019, 78, 922-929.	1.7	4
43	Upregulation of Annexin A1 in Reactive Astrocytes and Its Subtle Induction in Microglia at the Boundaries of Human Brain Infarcts. <i>Journal of Neuropathology and Experimental Neurology</i> , 2019, 78, 961-970.	1.7	9
44	Multiple mtDNA deletions due to mitochondrion toxicity of antiâ€hepadnaviral drugs: Comments to the letter from J. Finsterer. <i>Neuropathology</i> , 2019, 39, 326-327.	1.2	0
45	Intravoxel Incoherent Motion MR Imaging of Pediatric Intracranial Tumors: Correlation with Histology and Diagnostic Utility. <i>American Journal of Neuroradiology</i> , 2019, 40, 878-884.	2.4	16
46	Toxic myopathy with multiple deletions in mitochondrial DNA associated with longâ€term use of oral antiâ€viral drugs for hepatitis B: A case study. <i>Neuropathology</i> , 2019, 39, 162-167.	1.2	6
47	A Novel Combination of Prion Strain Co-Occurrence in Patients with Sporadic Creutzfeldt-Jakob Disease. <i>American Journal of Pathology</i> , 2019, 189, 1276-1283.	3.8	8
48	Predicting TERT promoter mutation using MR images in patients with wild-type IDH1 glioblastoma. <i>Diagnostic and Interventional Imaging</i> , 2019, 100, 411-419.	3.2	20
49	MOG antibody disease manifesting as progressive cognitive deterioration and behavioral changes with primary central nervous system vasculitis. <i>Multiple Sclerosis and Related Disorders</i> , 2019, 30, 48-50.	2.0	16
50	Predictors of recurrence and postoperative outcomes in patients with non-skull base meningiomas based on modern neurosurgical standards. <i>Interdisciplinary Neurosurgery: Advanced Techniques and Case Management</i> , 2019, 15, 30-37.	0.3	2
51	DCTN1 F52L mutation case of Perry syndrome with progressive supranuclear palsy-like tauopathy. <i>Parkinsonism and Related Disorders</i> , 2018, 51, 105-110.	2.2	18
52	Expression of CRYM in different rat organs during development and its decreased expression in degenerating pyramidal tracts in amyotrophic lateral sclerosis. <i>Neuropathology</i> , 2018, 38, 247-259.	1.2	7
53	Pediatric ganglioglioma with an H3 K27M mutation arising from the cervical spinal cord. <i>Neuropathology</i> , 2018, 38, 422-427.	1.2	12
54	Arterial spin-labeling is useful for the diagnosis of residual or recurrent meningiomas. <i>European Radiology</i> , 2018, 28, 4334-4342.	4.5	10

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55	Le hasard ou la nécessité? Comorbid pathologies of neurodegenerative diseases. <i>Neuropathology</i> , 2018, 38, 62-63.	1.2	0
56	Measurement of the perfusion fraction in brain tumors with intravoxel incoherent motion MR imaging: validation with histopathological vascular density in meningiomas. <i>British Journal of Radiology</i> , 2018, 91, 20170912.	2.2	25
57	Reclassification of 400 consecutive glioma cases based on the revised 2016WHO classification. <i>Brain Tumor Pathology</i> , 2018, 35, 81-89.	1.7	19
58	Association of adipocyte enhancer-binding protein 1 with Alzheimer's disease pathology in human hippocampi. <i>Brain Pathology</i> , 2018, 28, 58-71.	4.1	28
59	High-resolution melting and immunohistochemical analysis efficiently detects mutually exclusive genetic alterations of adamantinomatous and papillary craniopharyngiomas. <i>Neuropathology</i> , 2018, 38, 3-10.	1.2	18
60	Dynactin is involved in Lewy body pathology. <i>Neuropathology</i> , 2018, 38, 583-590.	1.2	12
61	An elderly case of malignant small cell glioma with hemorrhage coexistent with a calcified pilocytic astrocytoma component in the cerebellar hemisphere. <i>Neuropathology</i> , 2018, 38, 493-497.	1.2	3
62	Four-repeat tau dominant pathology in a congenital myotonic dystrophy type 1 patient with mental retardation. <i>Brain Pathology</i> , 2018, 28, 431-433.	4.1	4
63	Mitochondrial dysfunction and altered ribostasis in hippocampal neurons with cytoplasmic inclusions of multiple system atrophy. <i>Neuropathology</i> , 2018, 38, 361-371.	1.2	4
64	Radiological Features of Brain Metastases from Non-small Cell Lung Cancer Harboring EGFR Mutation. <i>Anticancer Research</i> , 2018, 38, 3731-3734.	1.1	14
65	Clinical Significance of PD-L1 Expression in Brain Metastases from Non-small Cell Lung Cancer. <i>Anticancer Research</i> , 2018, 38, 553-557.	1.1	19
66	Grading diffuse gliomas without intense contrast enhancement by amide proton transfer MR imaging: comparisons with diffusion- and perfusion-weighted imaging. <i>European Radiology</i> , 2017, 27, 578-588.	4.5	90
67	Molecular pathophysiology of impaired glucose metabolism, mitochondrial dysfunction, and oxidative DNA damage in Alzheimer's disease brain. <i>Mechanisms of Ageing and Development</i> , 2017, 161, 95-104.	4.6	105
68	Early and extensive spinal white matter involvement in neuromyelitis optica. <i>Brain Pathology</i> , 2017, 27, 249-265.	4.1	26
69	Insular primary glioblastomas with IDH mutations: Clinical and biological specificities. <i>Neuropathology</i> , 2017, 37, 200-206.	1.2	12
70	Trends in dementia prevalence, incidence, and survival rate in a Japanese community. <i>Neurology</i> , 2017, 88, 1925-1932.	1.1	154
71	Prevalence and clinicopathological features of H3.3 G34-mutant high-grade gliomas: a retrospective study of 411 consecutive glioma cases in a single institution. <i>Brain Tumor Pathology</i> , 2017, 34, 103-112.	1.7	69
72	Dura mater graft-associated Creutzfeldt-Jakob disease with 30-year incubation period. <i>Neuropathology</i> , 2017, 37, 275-281.	1.2	8

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73	Correlation between arterial spin-labeling perfusion and histopathological vascular density of pediatric intracranial tumors. <i>Journal of Neuro-Oncology</i> , 2017, 135, 561-569.	2.9	25
74	Spindle cell/sclerosing rhabdomyosarcoma with intracranial invasion without destroying the bone of the skull base: a case report and literature review. <i>Acta Radiologica Open</i> , 2017, 6, 205846011772731.	0.6	1
75	Different Complicated Brain Pathologies in Monozygotic Twins With Gerstmannâ€“Strâ€“usslerâ€“Scheinker Disease. <i>Journal of Neuropathology and Experimental Neurology</i> , 2017, 76, 854-863.	1.7	4
76	A comprehensive analysis identifies <i>BRAF</i> hotspot mutations associated with gliomas with peculiar epithelial morphology. <i>Neuropathology</i> , 2017, 37, 191-199.	1.2	33
77	Comparative profiling of cortical gene expression in Alzheimerâ€™s disease patients and mouse models demonstrates a link between amyloidosis and neuroinflammation. <i>Scientific Reports</i> , 2017, 7, 17762.	3.3	138
78	A Case of Suprasellar Ganglioglioma arising from the Genu-Rostrum of the Corpus Callosum composed of Tumor Cells in Various Stages of Neuronal Differentiation. <i>Japanese Journal of Neurosurgery</i> , 2017, 26, 362-368.	0.0	0
79	Discrepancy in Programmed Cell Death-Ligand 1 Between Primary and Metastatic Non-small Cell Lung Cancer. <i>Anticancer Research</i> , 2017, 37, 4223-4228.	1.1	30
80	Recent Increases in Hippocampal Tau Pathology in the Aging Japanese Population: The Hisayama Study. <i>Journal of Alzheimer's Disease</i> , 2016, 55, 613-624.	2.6	12
81	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. <i>OncoTargets and Therapy</i> , 2016, Volume 9, 7123-7131.	2.0	11
82	Diagnostic utility of intravoxel incoherent motion mr imaging in differentiating primary central nervous system lymphoma from glioblastoma multiforme. <i>Journal of Magnetic Resonance Imaging</i> , 2016, 44, 1256-1261.	3.4	35
83	Trends in autopsyâ€“verified dementia prevalence over 29Â“years of the Hisayama study. <i>Neuropathology</i> , 2016, 36, 383-387.	1.2	21
84	Sporadic <i>C</i> reutzfeldtâ€“akob Disease <i>MM1+2C</i> and <i>MM</i> 1 are Identical in Transmission Properties. <i>Brain Pathology</i> , 2016, 26, 95-101.	4.1	13
85	â€“PrP systemic deposition diseaseâ€™: clinical and pathological characteristics of novel familial prion disease with 2â€“bp deletion in codon 178. <i>European Journal of Neurology</i> , 2016, 23, 196-200.	3.3	11
86	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. <i>Journal of Neuropathology and Experimental Neurology</i> , 2016, 75, 1008-1019.	1.7	10
87	Differentiation of high-grade and low-grade diffuse gliomas by intravoxel incoherent motion MR imaging. <i>Neuro-Oncology</i> , 2016, 18, 132-141.	1.2	109
88	Midlife and Lateâ€“Life Smoking and Risk of Dementia in the Community: The Hisayama Study. <i>Journal of the American Geriatrics Society</i> , 2015, 63, 2332-2339.	2.6	56
89	Multiphasic acute disseminated encephalomyelitis associated with atypical rubella virus infection â€“ response to the letter from Wu et al. <i>Multiple Sclerosis Journal</i> , 2015, 21, 1089-1089.	3.0	0
90	A case of intracranial solitary fibrous tumor/hemangiopericytoma with dedifferentiated component. <i>Neuropathology</i> , 2015, 35, 260-265.	1.2	12

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91	Loss of <i>hnRNPA1</i> in <i>ALS</i> spinal cord motor neurons with <i>TDP-43</i> positive inclusions. <i>Neuropathology</i> , 2015, 35, 37-43.	1.2	41
92	Cryptic insertion of <i>FOXO1</i> into inverted chromosome arm 2q in the presence of two normal chromosome 13s and 13 small interstitial duplications in a patient with alveolar rhabdomyosarcoma. <i>Cancer Genetics</i> , 2015, 208, 428-433.	0.4	0
93	Masked hypodiploidy in anaplastic meningiomas by duplication of the original clone found in atypical meningiomas: Illustration of the evolution of genetic alterations. <i>Neuropathology</i> , 2014, 34, 353-359.	1.2	1
94	Down-regulation of <i>MET</i> in hippocampal neurons of <i>AD</i> brains. <i>Neuropathology</i> , 2014, 34, 284-290.	1.2	22
95	An intragenic deletion of the gene <i>MNAT1</i> in a family with pectus deformities. <i>American Journal of Medical Genetics, Part A</i> , 2014, 164, 1293-1297.	1.2	2
96	Altered Expression of Diabetes-Related Genes in Alzheimer's Disease Brains: The Hisayama Study. <i>Cerebral Cortex</i> , 2014, 24, 2476-2488.	2.9	294
97	<i>ABL1</i> gene involvement within a complex three-way translocation (2;9;4) in perineurioma characterized by molecular cytogenetic methods. <i>Cancer Genetics</i> , 2014, 207, 263-267.	0.4	8
98	Detection of prion protein oligomers by single molecule fluorescence imaging. <i>Neuropathology</i> , 2013, 33, 1-6.	1.2	0
99	The 102nd Neuropathological Meeting of Kyushu District 3 November 2012. <i>Neuropathology</i> , 2013, 33, 219-219.	1.2	0
100	Sporadic hereditary diffuse leukoencephalopathy with axonal spheroids showing numerous lesions with restricted diffusivity caused by a novel splice site mutation in the <i>CSF1R</i> gene. <i>Clinical and Experimental Neuroimmunology</i> , 2013, 4, 76-81.	1.0	7
101	Microsphere formation in a subtype of <i>Creutzfeldt-Jakob</i> disease with a <i>V180I</i> mutation and codon 129 <i>MM</i> polymorphism. <i>Neuropathology and Applied Neurobiology</i> , 2013, 39, 844-848.	3.2	9
102	Connexin 43 Astrocytopathy Linked to Rapidly Progressive Multiple Sclerosis and Neuromyelitis Optica. <i>PLoS ONE</i> , 2013, 8, e72919.	2.5	89
103	Extensive loss of connexins in <i>Balb/c</i> disease: evidence for an auto-antibody-independent astrocytopathy via impaired astrocyte-oligodendrocyte/myelin interaction. <i>Acta Neuropathologica</i> , 2012, 123, 887-900.	7.7	57
104	Extensive distribution of glial cytoplasmic inclusions in an autopsied case of multiple system atrophy with a prolonged 18-year clinical course. <i>Neuropathology</i> , 2012, 32, 69-76.	1.2	18
105	Association of Alzheimer disease pathology with abnormal lipid metabolism. <i>Neurology</i> , 2011, 77, 1068-1075.	1.1	92
106	An autopsied case of sporadic adult-onset amyotrophic lateral sclerosis with FUS-positive basophilic inclusions. <i>Neuropathology</i> , 2011, 31, 71-76.	1.2	22
107	The 99th Neuropathological Meeting of Kyushu District 4 December 2010. <i>Neuropathology</i> , 2011, 31, 199-199.	1.2	0
108	The 100th Neuropathological Meeting of Kyushu District 9 July 2011. <i>Neuropathology</i> , 2011, 31, 560-560.	1.2	0

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109	Reappraisal of Aquaporin-4 Astrocytopathy in Asian Neuromyelitis Optica and Multiple Sclerosis Patients. <i>Brain Pathology</i> , 2011, 21, 516-532.	4.1	41
110	Quantitative digital assessment of MGMT immunohistochemical expression in glioblastoma tissue. <i>Brain Tumor Pathology</i> , 2011, 28, 25-31.	1.7	21
111	Multiple system degeneration with basophilic inclusions in Japanese ALS patients with FUS mutation. <i>Acta Neuropathologica</i> , 2010, 119, 355-364.	7.7	90
112	Aquaporin-4 astrocytopathy in Baló's disease. <i>Acta Neuropathologica</i> , 2010, 120, 651-660.	7.7	53
113	Trends in prevalence of Alzheimer's disease and vascular dementia in a Japanese community: the Hisayama Study. <i>Acta Psychiatrica Scandinavica</i> , 2010, 122, 319-325.	4.5	123
114	Insulin resistance is associated with the pathology of Alzheimer disease. <i>Neurology</i> , 2010, 75, 764-770.	1.1	382
115	The 97th Neuropathological Meeting of Kyushu District 26 December 2008. <i>Neuropathology</i> , 2009, 29, 207-207.	1.2	1
116	The 95th Neuropathological Meeting of Kyushu District 15 December 2007. <i>Neuropathology</i> , 2008, 28, 346-346.	1.2	0
117	The 96th Neuropathological Meeting of Kyushu District 5 July 2008. <i>Neuropathology</i> , 2008, 28, 667-667.	1.2	0
118	The 94th Neuropathological Meeting of Kyushu District 30 June 2007. <i>Neuropathology</i> , 2007, 27, 508-508.	1.2	0
119	The 93rd Neuropathological Meeting of Kyushu District 2 December 2006. <i>Neuropathology</i> , 2007, 27, 150-150.	1.2	0
120	Increased asymmetric pulvinar magnetic resonance imaging signals in Creutzfeldt-Jakob disease with florid plaques following a cadaveric dura mater graft. <i>Neuropathology</i> , 2006, 26, 82-88.	1.2	15
121	The 91st Neuropathological Meeting of Kyushu District 17 December 2005. <i>Neuropathology</i> , 2006, 26, 275-275.	1.2	0
122	The 92nd Neuropathological Meeting of Kyushu District 22 July 2006. <i>Neuropathology</i> , 2006, 26, 597-597.	1.2	1
123	An astroblastoma case associated with loss of heterozygosity on chromosome 9p. <i>Journal of Neuro-Oncology</i> , 2006, 80, 69-73.	2.9	13
124	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. <i>Cancer Genetics and Cytogenetics</i> , 2006, 170, 152-157.	1.0	5
125	Allelic Losses of Chromosome 10 in Glioma Tissues Detected by Quantitative Single-Strand Conformation Polymorphism Analysis. <i>Clinical Chemistry</i> , 2006, 52, 370-378.	3.2	31
126	Forced retraction of spinal root injury enhances activation of p38 MAPK cascade in infiltrating macrophages. <i>Neuropathology</i> , 2005, 25, 37-47.	1.2	8

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127	The 90th Neuropathological Meeting of Kyushu District 25 June 2005. <i>Neuropathology</i> , 2005, 25, 284-284.	1.2	0
128	Amyloid imaging probes are useful for detection of prion plaques and treatment of transmissible spongiform encephalopathies. <i>Journal of General Virology</i> , 2004, 85, 1785-1790.	2.9	58
129	Defense mechanism to oxidative DNA damage in glial cells. <i>Neuropathology</i> , 2004, 24, 125-130.	1.2	23
130	Unusual aberration involving the short arm of chromosome 11 in an 8-month-old patient with a supratentorial primitive neuroectodermal tumor. <i>Cancer Genetics and Cytogenetics</i> , 2003, 141, 143-147.	1.0	5
131	The 85th Neuropathological Meeting of Kyushu District 30 November 2002. <i>Neuropathology</i> , 2003, 23, 109-109.	1.2	0
132	EWS/FLI-1 fusion signal inserted into chromosome 11 in one patient with morphologic features of Ewing sarcoma, but lacking t(11;22). <i>Cancer Genetics and Cytogenetics</i> , 2002, 133, 72-75.	1.0	13
133	Clusterin/apolipoprotein J is associated with cortical Lewy bodies: immunohistochemical study in cases with α -synucleinopathies. <i>Acta Neuropathologica</i> , 2002, 104, 225-230.	7.7	75
134	Expression of 8-oxoguanine DNA glycosylase is reduced and associated with neurofibrillary tangles in Alzheimer's disease brain. <i>Acta Neuropathologica</i> , 2002, 103, 20-25.	7.7	122
135	The 84th Neuropathological Meeting of Kyushu District 1 June 2002. <i>Neuropathology</i> , 2002, 22, 218-218.	1.2	0
136	Sixty-one-year-old woman presenting with deep coma during treatment for adrenal insufficiency. <i>Neuropathology</i> , 2002, 22, 353-355.	1.2	0
137	IgH Intronic Enhancer Element HE2 ($\frac{1}{4}$ B) Functions as a cis-Activator in Choroid Plexus Cells at the Cellular Level as well as in Transgenic Mice. <i>Journal of Neurochemistry</i> , 2002, 64, 961-966.	3.9	11
138	Expression of hMTH1 in the hippocampi of control and Alzheimer's disease. <i>NeuroReport</i> , 2001, 12, 2895-2899.	1.2	49
139	Expression of the lysosome-associated membrane proteins in myopathies with rimmed vacuoles. <i>Acta Neuropathologica</i> , 2001, 101, 579-584.	7.7	24
140	Different responses of benign and atypical meningiomas to gamma-knife radiosurgery: report of two cases with immunohistochemical analysis. <i>Brain Tumor Pathology</i> , 2001, 18, 61-66.	1.7	5
141	Autopsy case of autosomal recessive hereditary spastic paraplegia with reference to the muscular pathology. <i>Neuropathology</i> , 2001, 21, 212-217.	1.2	25
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143	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. <i>Acta Neuropathologica</i> , 2001, 102, 467-472.	7.7	35
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