Hiroyuki Honda

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	The changing prevalence and incidence of dementia over time — current evidence. Nature Reviews Neurology, 2017, 13, 327-339.	4.9	503
2	Altered Expression of Diabetes-Related Genes in Alzheimer's Disease Brains: The Hisayama Study. Cerebral Cortex, 2014, 24, 2476-2488.	1.6	294
3	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.	0.7	41
4	Proteaseâ€resistant PrP and PrP oligomers in the brain in human prion diseases after intraventricular pentosan polysulfate infusion. Neuropathology, 2012, 32, 124-132.	0.7	30
5	Association of adipocyte enhancerâ€binding protein 1 with <scp>A</scp> lzheimer's disease pathology in human hippocampi. Brain Pathology, 2018, 28, 58-71.	2.1	28
6	Elevated expression of fatty acid synthase and nuclear localization of carnitine palmitoyltransferase <scp>1C</scp> are common among human gliomas. Neuropathology, 2014, 34, 465-474.	0.7	26
7	An autopsied case of sporadic adult-onset amyotrophic lateral sclerosis with FUS-positive basophilic inclusions. Neuropathology, 2011, 31, 71-76.	0.7	22
8	Downâ€regulation of <scp>MET</scp> in hippocampal neurons of <scp>A</scp> lzheimer's disease brains. Neuropathology, 2014, 34, 284-290.	0.7	22
9	Trends in autopsyâ€verified dementia prevalence over 29Âyears of the Hisayama study. Neuropathology, 2016, 36, 383-387.	0.7	21
10	Prion protein oligomers in Creutzfeldtâ€ j akob disease detected by gelâ€ f iltration centrifuge columns. Neuropathology, 2009, 29, 536-542.	0.7	18
11	DCTN1 F52L mutation case of Perry syndrome with progressive supranuclear palsy-like tauopathy. Parkinsonism and Related Disorders, 2018, 51, 105-110.	1.1	18
12	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.	1.2	15
13	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.	0.7	15
14	Recent Increases in Hippocampal Tau Pathology in the Aging Japanese Population: The Hisayama Study. Journal of Alzheimer's Disease, 2016, 55, 613-624.	1.2	12
15	Dynactin is involved in Lewy body pathology. Neuropathology, 2018, 38, 583-590.	0.7	12
16	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.	0.9	10
17	Accumulation of Astrocytic Aquaporin 4 and Aquaporin 1 in Prion Protein Plaques. Journal of Neuropathology and Experimental Neurology, 2020, 79, 419-429.	0.9	10
18	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.	0.7	10

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19	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.	0.9	9
20	Transactivation response DNAâ€binding protein of 43 kDa proteinopathy and lysosomal abnormalities in spastic paraplegia type 11. Neuropathology, 2021, 41, 253-265.	0.7	9
21	Dura mater graftâ€associated Creutzfeldtâ€Jakob disease with 30â€year incubation period. Neuropathology, 2017, 37, 275-281.	0.7	8
22	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.	0.7	7
23	Toxic myopathy with multiple deletions in mitochondrial DNA associated with longâ€term use of oral antiâ€viral drugs for hepatitis B: A case study. Neuropathology, 2019, 39, 162-167.	0.7	6
24	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.	0.9	6
25	Symmetrical glial hyperplasia in the brainstem of fibrodysplasia ossificans progressiva. Neuropathology, 2021, 41, 146-151.	0.7	6
26	Concurrent cardiac transthyretin and brain \hat{l}^2 amyloid accumulation among the older adults: The Hisayama study. Brain Pathology, 2021, , e13014.	2.1	6
27	Proposal of new diagnostic criteria for fatal familial insomnia. Journal of Neurology, 2022, 269, 4909-4919.	1.8	5
28	Different Complicated Brain Pathologies in Monozygotic Twins With Gerstmann–StrÃ u ssler–Scheinker Disease. Journal of Neuropathology and Experimental Neurology, 2017, 76, 854-863.	0.9	4
29	Fourâ€repeat tau dominant pathology in a congenital myotonic dystrophy type 1 patient with mental retardation. Brain Pathology, 2018, 28, 431-433.	2.1	4
30	Mitochondrial dysfunction and altered ribostasis in hippocampal neurons with cytoplasmic inclusions of multiple system atrophy. Neuropathology, 2018, 38, 361-371.	0.7	4
31	Frequent Detection of Pituitary-Derived PrPres in Human Prion Diseases. Journal of Neuropathology and Experimental Neurology, 2019, 78, 922-929.	0.9	4
32	Immunotherapyâ€refractory vacuolar myopathy with mucin deposition in scleromyxedema: A possible role of fibroblast growth factor 2. Neuropathology, 2020, 40, 492-495.	0.7	3
33	Prion Gene PRNP Y162X Truncation Mutation Can Induce a Refractory Esophageal Achalasia. American Journal of Gastroenterology, 2021, 116, 1350-1351.	0.2	2
34	Detection of cutaneous prion protein deposits could help diagnose GPIâ€anchorless prion disease with neuropathy. European Journal of Neurology, 2021, 28, 2133-2137.	1.7	1
35	Optic nerve atrophy and visual disturbance following PRNP Y162X truncation mutation. Journal of the Neurological Sciences, 2021, 428, 117614.	0.3	1
36	A Comparative Study of Site-Specific Distribution of Aging-Related Tau Astrogliopathy and Its Risk Factors Between Alzheimer Disease and Cognitive Healthy Brains: The Hisayama Study. Journal of Neuropathology and Experimental Neurology, 2022, 81, 106-116.	0.9	1

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37	Multiple mtDNA deletions due to mitochondrion toxicity of antiâ€hepadnaviral drugs: Comments to the letter from J. Finsterer. Neuropathology, 2019, 39, 326-327.	0.7	0