

# Hiroyuki Honda

## List of Publications by Year in descending order

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37  
papers

1,193  
citations

758635

12  
h-index

395343

33  
g-index

37  
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37  
docs citations

37  
times ranked

2419  
citing authors

#	ARTICLE	IF	CITATIONS
1	A Comparative Study of Site-Specific Distribution of Aging-Related Tau Astroglipopathy 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.	0.9	1
2	Proposal of new diagnostic criteria for fatal familial insomnia. <i>Journal of Neurology</i> , 2022, 269, 4909-4919.	1.8	5
3	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.	0.9	6
4	Symmetrical glial hyperplasia in the brainstem of fibrodysplasia ossificans progressiva. <i>Neuropathology</i> , 2021, 41, 146-151.	0.7	6
5	Detection of cutaneous prion protein deposits could help diagnose GPI anchorless prion disease with neuropathy. <i>European Journal of Neurology</i> , 2021, 28, 2133-2137.	1.7	1
6	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. <i>Neuropathology</i> , 2021, 41, 152-158.	0.7	10
7	Transactivation response DNA-binding protein of 43 kDa proteinopathy and lysosomal abnormalities in spastic paraplegia type 11. <i>Neuropathology</i> , 2021, 41, 253-265.	0.7	9
8	Concurrent cardiac transthyretin and brain $\beta^2$ amyloid accumulation among the older adults: The Hisayama study. <i>Brain Pathology</i> , 2021, , e13014.	2.1	6
9	Optic nerve atrophy and visual disturbance following PRNP Y162X truncation mutation. <i>Journal of the Neurological Sciences</i> , 2021, 428, 117614.	0.3	1
10	Prion Gene PRNP Y162X Truncation Mutation Can Induce a Refractory Esophageal Achalasia. <i>American Journal of Gastroenterology</i> , 2021, 116, 1350-1351.	0.2	2
11	Immunotherapy refractory vacuolar myopathy with mucin deposition in scleromyxedema: A possible role of fibroblast growth factor 2. <i>Neuropathology</i> , 2020, 40, 492-495.	0.7	3
12	Accumulation of Astrocytic Aquaporin 4 and Aquaporin 1 in Prion Protein Plaques. <i>Journal of Neuropathology and Experimental Neurology</i> , 2020, 79, 419-429.	0.9	10
13	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.	1.2	15
14	Expanded polyglutamine impairs normal nuclear distribution of fused in sarcoma and poly (rC) binding protein 1 in Huntington's disease. <i>Neuropathology</i> , 2019, 39, 358-367.	0.7	15
15	Frequent Detection of Pituitary-Derived PrPres in Human Prion Diseases. <i>Journal of Neuropathology and Experimental Neurology</i> , 2019, 78, 922-929.	0.9	4
16	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.	0.9	9
17	Multiple mtDNA deletions due to mitochondrion toxicity of anti-hepadnaviral drugs: Comments to the letter from J. Finsterer. <i>Neuropathology</i> , 2019, 39, 326-327.	0.7	0
18	Toxic myopathy with multiple deletions in mitochondrial DNA associated with long-term use of oral anti-viral drugs for hepatitis B: A case study. <i>Neuropathology</i> , 2019, 39, 162-167.	0.7	6

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19	DCTN1 F52L mutation case of Perry syndrome with progressive supranuclear palsy-like tauopathy. <i>Parkinsonism and Related Disorders</i> , 2018, 51, 105-110.	1.1	18
20	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.	0.7	7
21	Association of adipocyte enhancer-binding protein 1 with Alzheimer's disease pathology in human hippocampi. <i>Brain Pathology</i> , 2018, 28, 58-71.	2.1	28
22	Dynactin is involved in Lewy body pathology. <i>Neuropathology</i> , 2018, 38, 583-590.	0.7	12
23	Four-repeat tau dominant pathology in a congenital myotonic dystrophy type 1 patient with mental retardation. <i>Brain Pathology</i> , 2018, 28, 431-433.	2.1	4
24	Mitochondrial dysfunction and altered ribostasis in hippocampal neurons with cytoplasmic inclusions of multiple system atrophy. <i>Neuropathology</i> , 2018, 38, 361-371.	0.7	4
25	The changing prevalence and incidence of dementia over time – current evidence. <i>Nature Reviews Neurology</i> , 2017, 13, 327-339.	4.9	503
26	Dura mater graft-associated Creutzfeldt-Jakob disease with 30-year incubation period. <i>Neuropathology</i> , 2017, 37, 275-281.	0.7	8
27	Different Complicated Brain Pathologies in Monozygotic Twins With Gerstmann-Sträussler-Scheinker Disease. <i>Journal of Neuropathology and Experimental Neurology</i> , 2017, 76, 854-863.	0.9	4
28	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.	1.2	12
29	Trends in autopsy-verified dementia prevalence over 29 years of the Hisayama study. <i>Neuropathology</i> , 2016, 36, 383-387.	0.7	21
30	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.	0.9	10
31	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.	0.7	41
32	Elevated expression of fatty acid synthase and nuclear localization of carnitine palmitoyltransferase <i>1C</i> are common among human gliomas. <i>Neuropathology</i> , 2014, 34, 465-474.	0.7	26
33	Down-regulation of <i>MET</i> in hippocampal neurons of Alzheimer's disease brains. <i>Neuropathology</i> , 2014, 34, 284-290.	0.7	22
34	Altered Expression of Diabetes-Related Genes in Alzheimer's Disease Brains: The Hisayama Study. <i>Cerebral Cortex</i> , 2014, 24, 2476-2488.	1.6	294
35	Protease-resistant PrP and PrP oligomers in the brain in human prion diseases after intraventricular pentosan polysulfate infusion. <i>Neuropathology</i> , 2012, 32, 124-132.	0.7	30
36	An autopsied case of sporadic adult-onset amyotrophic lateral sclerosis with FUS-positive basophilic inclusions. <i>Neuropathology</i> , 2011, 31, 71-76.	0.7	22

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37	Prion protein oligomers in Creutzfeldtâ€­Jakob disease detected by gelâ€­filtration centrifuge columns. <i>Neuropathology</i> , 2009, 29, 536-542.	0.7	18