

# Fumiaki Mori

## List of Publications by Year in descending order

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Version: 2024-02-01

63  
papers

3,086  
citations

159585

30  
h-index

161849

54  
g-index

63  
all docs

63  
docs citations

63  
times ranked

3809  
citing authors

#	ARTICLE	IF	CITATIONS
1	Neuropathology of Multiple System Atrophy, a Glioneuronal Degenerative Disease. <i>Cerebellum</i> , 2024, 23, 2-12.	2.5	3
2	Cabergoline, a long-acting dopamine agonist, attenuates L-dopa-induced dyskinesia without L-dopa sparing in a rat model of Parkinson's disease. <i>Neuroscience Research</i> , 2022, 178, 93-97.	1.9	2
3	The clinical and neuropathological picture of adult neuronal intranuclear inclusion disease with no radiological abnormality. <i>Neuropathology</i> , 2022, 42, 204-211.	1.2	5
4	Accumulation of Nonfibrillar TDP-43 in the Rough Endoplasmic Reticulum Is the Early-Stage Pathology in Amyotrophic Lateral Sclerosis. <i>Journal of Neuropathology and Experimental Neurology</i> , 2022, 81, 271-281.	1.7	5
5	Effects of Aging on Levo-Dihydroxyphenylalanine- Induced Dyskinesia in a Rat Model of Parkinson's Disease. <i>Frontiers in Aging Neuroscience</i> , 2021, 13, 650350.	3.4	2
6	Role of VAPB and vesicular profiles in $\alpha$ -synuclein aggregates in multiple system atrophy. <i>Brain Pathology</i> , 2021, 31, e13001.	4.1	5
7	GABA storage and release in the medial globus pallidus in L-DOPA-induced dyskinesia priming. <i>Neurobiology of Disease</i> , 2020, 143, 104979.	4.4	14
8	Phosphorylated TDP-43 aggregates in skeletal and cardiac muscle are a marker of myogenic degeneration in amyotrophic lateral sclerosis and various conditions. <i>Acta Neuropathologica Communications</i> , 2019, 7, 165.	5.2	35
9	Autophagy Is a Common Degradation Pathway for Bunina Bodies and TDP-43 Inclusions in Amyotrophic Lateral Sclerosis. <i>Journal of Neuropathology and Experimental Neurology</i> , 2019, 78, 910-921.	1.7	7
10	MicroRNA expression profiles of neuron-derived extracellular vesicles in plasma from patients with amyotrophic lateral sclerosis. <i>Neuroscience Letters</i> , 2019, 708, 134176.	2.1	66
11	A mouse model of adult-onset multiple system atrophy. <i>Neurobiology of Disease</i> , 2019, 127, 339-349.	4.4	14
12	Alteration of autophagy-related proteins in peripheral blood mononuclear cells of patients with Parkinson's disease. <i>Neurobiology of Aging</i> , 2018, 63, 33-43.	3.1	54
13	Immunohistochemical localization of exoribonucleases (DIS3L2 and XRN1) in intranuclear inclusion body disease. <i>Neuroscience Letters</i> , 2018, 662, 389-394.	2.1	11
14	Colocalization of Bunina bodies and TDP-43 inclusions in a case of sporadic amyotrophic lateral sclerosis with Lewy body-like hyaline inclusions. <i>Neuropathology</i> , 2018, 38, 521-528.	1.2	11
15	Autophagy mediators (FOXO1, SESN3 and TSC2) in Lewy body disease and aging. <i>Neuroscience Letters</i> , 2018, 684, 35-41.	2.1	19
16	PLA2G6 accumulates in Lewy bodies in PARK14 and idiopathic Parkinson's disease. <i>Neuroscience Letters</i> , 2017, 645, 40-45.	2.1	34
17	Alteration of Upstream Autophagy-Related Proteins (ULK1, ULK2) in Multiple System Atrophy. <i>Journal of Neuropathology</i> , 2016, 26, 359-370.	4.1	40
18	Accumulation of phosphorylated $\alpha$ -synuclein in subpial and periventricular astrocytes in multiple system atrophy of long duration. <i>Neuropathology</i> , 2016, 36, 157-167.	1.2	38

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19	Î±-Synuclein pathology in the cranial and spinal nerves in Lewy body disease. <i>Neuropathology</i> , 2016, 36, 262-269.	1.2	17
20	Novel eosinophilic neuronal cytoplasmic inclusions in the external cuneate nucleus of humans. <i>Neuropathology</i> , 2016, 36, 441-447.	1.2	1
21	G protein-coupled receptor 26 immunoreactivity in intranuclear inclusions associated with polyglutamine and intranuclear inclusion body diseases. <i>Neuropathology</i> , 2016, 36, 50-55.	1.2	11
22	Isopentenyl diphosphate isomerase, a cholesterol synthesizing enzyme, is localized in Lewy bodies. <i>Neuropathology</i> , 2015, 35, 432-440.	1.2	31
23	Spontaneous epileptic seizures in transgenic rats harboring a human ADNFLE missense mutation in the Î²2-subunit of the nicotinic acetylcholine receptor. <i>Neuroscience Research</i> , 2015, 100, 46-54.	1.9	17
24	Filamentous aggregations of phosphorylated Î±-synuclein in Schwann cells (Schwann cell cytoplasmic) Tj ETQq0 0 Q rgBT /Overlock 10 T	5.2	49
25	Sortilin-related receptor CNS expressed 2 (SorCS2) is localized to Bunina bodies in amyotrophic lateral sclerosis. <i>Neuroscience Letters</i> , 2015, 608, 6-11.	2.1	8
26	Analysis of microRNA from archived formalin-fixed paraffin-embedded specimens of amyotrophic lateral sclerosis. <i>Acta Neuropathologica Communications</i> , 2014, 2, 173.	5.2	33
27	Accumulation of the sigma-1 receptor is common to neuronal nuclear inclusions in various neurodegenerative diseases. <i>Neuropathology</i> , 2014, 34, 148-158.	1.2	52
28	ALS-associated protein FIG4 is localized in Pick and Lewy bodies, and also neuronal nuclear inclusions, in polyglutamine and intranuclear inclusion body diseases. <i>Neuropathology</i> , 2014, 34, 19-26.	1.2	27
29	Ubiquitin-negative, eosinophilic neuronal cytoplasmic inclusions associated with stress granules and autophagy: An immunohistochemical investigation of two cases. <i>Neuropathology</i> , 2014, 34, 140-147.	1.2	2
30	Valosin-containing protein immunoreactivity in tauopathies, synucleinopathies, polyglutamine diseases and intranuclear inclusion body disease. <i>Neuropathology</i> , 2013, 33, 637-644.	1.2	20
31	A novel prophylactic effect of furosemide treatment on autosomal dominant nocturnal frontal lobe epilepsy (ADNFLE). <i>Epilepsy Research</i> , 2013, 107, 127-137.	1.6	20
32	The Lewy Body in Parkinson's Disease and Related Neurodegenerative Disorders. <i>Molecular Neurobiology</i> , 2013, 47, 495-508.	4.0	323
33	An autopsy case of preclinical multiple system atrophy (MSA-C). <i>Neuropathology</i> , 2013, 33, 667-672.	1.2	27
34	Endosomal sorting related protein CHMP2B is localized in Lewy bodies and glial cytoplasmic inclusions in Î±-synucleinopathy. <i>Neuroscience Letters</i> , 2012, 527, 16-21.	2.1	24
35	Optineurin immunoreactivity in neuronal nuclear inclusions of polyglutamine diseases (Huntington's,) Tj ETQq1,1 0.784314 rgBT 10	7.7	32
36	Ubiquilin immunoreactivity in cytoplasmic and nuclear inclusions in synucleinopathies, polyglutamine diseases and intranuclear inclusion body disease. <i>Acta Neuropathologica</i> , 2012, 124, 149-151.	7.7	41

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37	Immunohistochemical analysis of Marinesco bodies, using antibodies against proteins implicated in the ubiquitin-proteasome system, autophagy and aggresome formation. <i>Neuropathology</i> , 2012, 32, 261-266.	1.2	30
38	Ubiquitin-related proteins in neuronal and glial intranuclear inclusions in intranuclear inclusion body disease. <i>Pathology International</i> , 2012, 62, 407-411.	1.3	17
39	Incipient intranuclear inclusion body disease in a 78-year-old woman. <i>Neuropathology</i> , 2011, 31, 188-193.	1.2	23
40	Enhancement of native and phosphorylated TDP-43 immunoreactivity by proteinase K treatment following autoclave heating. <i>Neuropathology</i> , 2011, 31, 401-404.	1.2	6
41	Alteration of autophagosomal proteins (LC3, GABARAP and GATE-16) in Lewy body disease. <i>Neurobiology of Disease</i> , 2011, 43, 690-697.	4.4	102
42	Proteinase K-resistant $\alpha$ -synuclein is deposited in presynapses in human Lewy body disease and A53T $\alpha$ -synuclein transgenic mice. <i>Acta Neuropathologica</i> , 2010, 120, 145-154.	7.7	87
43	Involvement of the peripheral nervous system in synucleinopathies, tauopathies and other neurodegenerative proteinopathies of the brain. <i>Acta Neuropathologica</i> , 2010, 120, 1-12.	7.7	131
44	Widespread occurrence of eosinophilic neuronal cytoplasmic inclusions in an asymptomatic adult: A novel ubiquitin-negative filamentous inclusion. <i>Neuropathology</i> , 2010, 30, 648-653.	1.2	2
45	Decreased Cystatin C Immunoreactivity in Spinal Motor Neurons and Astrocytes in Amyotrophic Lateral Sclerosis. <i>Journal of Neuropathology and Experimental Neurology</i> , 2009, 68, 1200-1206.	1.7	24
46	$\alpha$ -Synuclein pathology in the neostriatum in Parkinson's disease. <i>Acta Neuropathologica</i> , 2008, 115, 453-459.	7.7	52
47	Maturation process of TDP-43-positive neuronal cytoplasmic inclusions in amyotrophic lateral sclerosis with and without dementia. <i>Acta Neuropathologica</i> , 2008, 116, 193-203.	7.7	111
48	Epitope mapping of 2E2-D3, a monoclonal antibody directed against human TDP-43. <i>Neuroscience Letters</i> , 2008, 434, 170-174.	2.1	35
49	Axonal $\alpha$ -synuclein aggregates herald centripetal degeneration of cardiac sympathetic nerve in Parkinson's disease. <i>Brain</i> , 2008, 131, 642-650.	7.6	416
50	Rats Harboring S284L <i>Chrna4</i> Mutation Show Attenuation of Synaptic and Extrasynaptic GABAergic Transmission and Exhibit the Nocturnal Frontal Lobe Epilepsy Phenotype. <i>Journal of Neuroscience</i> , 2008, 28, 12465-12476.	3.6	62
51	Immunohistochemical localization of NUB1, a synphilin-1-binding protein, in neurodegenerative disorders. <i>Acta Neuropathologica</i> , 2007, 114, 365-371.	7.7	23
52	NUB1 Suppresses the Formation of Lewy Body-Like Inclusions by Proteasomal Degradation of Synphilin-1. <i>American Journal of Pathology</i> , 2006, 169, 553-565.	3.8	56
53	Effects of interleukin-1 $\beta$ on hippocampal glutamate and GABA releases associated with Ca <sup>2+</sup> -induced Ca <sup>2+</sup> releasing systems. <i>Epilepsy Research</i> , 2006, 71, 107-116.	1.6	66
54	Degeneration of cardiac sympathetic nerve can occur in multiple system atrophy. <i>Acta Neuropathologica</i> , 2006, 113, 81-86.	7.7	115

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55	An autopsy case of early (â€œminimal changeâ€œ) olivopontocerebellar atrophy (multiple system) Tj ETQq1 1 0.784314 rgBT /Overloc	7.7	73
56	Hypertrophy of medial globus pallidus and substantia nigra reticulata in 6-hydroxydopamine-lesioned rats treated with L-DOPA: Implication for L-DOPA-induced dyskinesia in Parkinson's disease. <i>Neuropathology</i> , 2004, 24, 290-295.	1.2	19
57	Accumulation of phosphorylated a-synuclein in the brain and peripheral ganglia of patients with multiple system atrophy. <i>Acta Neuropathologica</i> , 2004, 107, 292-298.	7.7	108
58	Î±-Synuclein pathology affecting Bergmann glia of the cerebellum in patients with Î±-synucleinopathies. <i>Acta Neuropathologica</i> , 2003, 105, 403-409.	7.7	54
59	Î±-Synuclein Accumulates in Purkinje Cells in Lewy Body Disease but not in Multiple System Atrophy. <i>Journal of Neuropathology and Experimental Neurology</i> , 2003, 62, 812-819.	1.7	66
60	Demonstration of Î±-Synuclein Immunoreactivity in Neuronal and Glial Cytoplasm in Normal Human Brain Tissue Using Proteinase K and Formic Acid Pretreatment. <i>Experimental Neurology</i> , 2002, 176, 98-104.	4.1	128
61	?-Synuclein immunoreactivity in normal and neoplastic Schwann cells. <i>Acta Neuropathologica</i> , 2002, 103, 145-151.	7.7	41
62	Immunocytochemical localization of synphilin-1, an Î±-synuclein-associated protein, in neurodegenerative disorders. <i>Acta Neuropathologica</i> , 2002, 103, 209-214.	7.7	76
63	Expression of ??-synuclein in a human glioma cell line and its up-regulation by interleukin-1??. <i>NeuroReport</i> , 2001, 12, 1909-1912.	1.2	63