

Mari Tada

List of Publications by Year in descending order

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Version: 2024-02-01

38
papers

1,230
citations

394421

19
h-index

377865

34
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41
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41
docs citations

41
times ranked

2082
citing authors

#	ARTICLE	IF	CITATIONS
1	Brain TDP-43 pathology in corticobasal degeneration: Topographical correlation with neuronal loss. <i>Neuropathology and Applied Neurobiology</i> , 2022, 48, .	3.2	8
2	Endogenous human retrovirus-K is not increased in the affected tissues of Japanese ALS patients. <i>Neuroscience Research</i> , 2022, 178, 78-82.	1.9	1
3	Praja1 <sc>RING</sc>-finger <sc>E3</sc> ubiquitin ligase is a common suppressor of neurodegenerative disease-associated protein aggregation. <i>Neuropathology</i> , 2022, 42, 488-504.	1.2	5
4	A novel splicing variant of ANXA11 in a patient with amyotrophic lateral sclerosis: histologic and biochemical features. <i>Acta Neuropathologica Communications</i> , 2021, 9, 106.	5.2	10
5	Age-related demethylation of the TDP-43 autoregulatory region in the human motor cortex. <i>Communications Biology</i> , 2021, 4, 1107.	4.4	10
6	Alzheimer's A β assembly binds sodium pump and blocks endothelial NOS activity via ROS-PKC pathway in brain vascular endothelial cells. <i>IScience</i> , 2021, 24, 102936.	4.1	7
7	Parkinson's disease and parkinsonism: Clinicopathological discrepancies on diagnosis in three patients. <i>Neuropathology</i> , 2021, 41, 450-456.	1.2	1
8	Toward allele-specific targeting therapy and pharmacodynamic marker for spinocerebellar ataxia type 3. <i>Science Translational Medicine</i> , 2020, 12, .	12.4	32
9	Praja1 <sc>RING</sc>-finger <sc>E3</sc> ubiquitin ligase suppresses neuronal cytoplasmic <sc>TDP</sc>-43 aggregate formation. <i>Neuropathology</i> , 2020, 40, 570-586.	1.2	23
10	Novel CHP1 mutation in autosomal-recessive cerebellar ataxia: autopsy features of two siblings. <i>Acta Neuropathologica Communications</i> , 2020, 8, 134.	5.2	1
11	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
12	Vertical Gaze Palsy Caused by Selective Unilateral Rostral Midbrain Infarction. <i>Neuro-Ophthalmology</i> , 2018, 42, 309-311.	1.0	6
13	Loss of Motor Neurons Innervating Cervical Muscles in Patients With Multiple System Atrophy and Dropped Head. <i>Journal of Neuropathology and Experimental Neurology</i> , 2018, 77, 317-324.	1.7	2
14	Neuronal intranuclear inclusion disease showing intranuclear inclusions in renal biopsy 12 years earlier. <i>Neurology</i> , 2018, 91, 884-886.	1.1	24
15	Clinicopathologic Features of Two Patients With Sporadic Amyotrophic Lateral Sclerosis Who Maintained Communication Ability for Over 30 Years. <i>Journal of Neuropathology and Experimental Neurology</i> , 2018, 77, 981-986.	1.7	1
16	Expression of Aquaporin 1 and Aquaporin 4 in the Temporal Neocortex of Patients with Parkinson's Disease. <i>Brain Pathology</i> , 2017, 27, 160-168.	4.1	57
17	Heterogeneity of cerebral TDP-43 pathology in sporadic amyotrophic lateral sclerosis: Evidence for clinico-pathologic subtypes. <i>Acta Neuropathologica Communications</i> , 2016, 4, 61.	5.2	38
18	Pathological and Clinical Spectrum of Progressive Supranuclear Palsy: With Special Reference to Astrocytic Tau Pathology. <i>Brain Pathology</i> , 2016, 26, 155-166.	4.1	28

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19	Characteristic microglial features in patients with hereditary diffuse leukoencephalopathy with spheroids. <i>Annals of Neurology</i> , 2016, 80, 554-565.	5.3	43
20	Globular Glial Mixed Four Repeat Tau and τ 43 Proteinopathy with Motor Neuron Disease and Frontotemporal Dementia. <i>Brain Pathology</i> , 2016, 26, 82-94.	4.1	25
21	Increased neuronal and astroglial aquaporin-1 immunoreactivity in rat striatum by chemical preconditioning with 3-nitropropionic acid. <i>Neuroscience Letters</i> , 2016, 626, 48-53.	2.1	5
22	Na, K-ATPase β 3 is a death target of Alzheimer patient amyloid- β 2 assembly. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2015, 112, E4465-74.	7.1	112
23	Progressive myoclonus epilepsy: extraneuronal brown pigment deposition and system neurodegeneration in the brains of Japanese patients with novel SCARB2 mutations. <i>Neuropathology and Applied Neurobiology</i> , 2014, 40, 551-563.	3.2	14
24	Pathology and sensitivity of current clinical criteria in corticobasal syndrome. <i>Movement Disorders</i> , 2014, 29, 238-244.	3.9	47
25	Haploinsufficiency of CSF-1R and clinicopathologic characterization in patients with HDLS. <i>Neurology</i> , 2014, 82, 139-148.	1.1	103
26	C9ORF72 repeat-associated non-ATG translated polypeptides are distributed independently of τ 43 in a Japanese patient with c9ALS. <i>Neuropathology and Applied Neurobiology</i> , 2014, 40, 783-788.	3.2	4
27	A fatal neuromuscular disease in an adult patient after poliomyelitis in early childhood: Consideration of the pathology of post-polio syndrome. <i>Neuropathology</i> , 2013, 33, 93-101.	1.2	9
28	Early clinical features of patients with progressive supranuclear palsy with predominant cerebellar ataxia. <i>Parkinsonism and Related Disorders</i> , 2013, 19, 1149-1151.	2.2	43
29	What is the key player in τ 43 pathology in ALS: Disappearance from the nucleus or inclusion formation in the cytoplasm?. <i>Neurology and Clinical Neuroscience</i> , 2013, 1, 11-17.	0.4	9
30	Difference in MSA Phenotype Distribution between Populations: Genetics or Environment?. <i>Journal of Parkinson's Disease</i> , 2012, 2, 7-18.	2.8	33
31	Coexistence of Huntington's disease and amyotrophic lateral sclerosis: a clinicopathologic study. <i>Acta Neuropathologica</i> , 2012, 124, 749-760.	7.7	48
32	Isolation and Characterization of Patient-derived, Toxic, High Mass Amyloid- β 2-Protein ($A\beta$) Assembly from Alzheimer Disease Brains. <i>Journal of Biological Chemistry</i> , 2009, 284, 32895-32905.	3.4	162
33	Depletion of medullary serotonergic neurons in patients with multiple system atrophy who succumbed to sudden death. <i>Brain</i> , 2009, 132, 1810-1819.	7.6	98
34	Marinesco-Björk syndrome with atrophy of the brain stem tegmentum and dysplastic cytoarchitecture in the cerebral cortex. <i>Neuropathology</i> , 2008, 28, 541-546.	1.2	12
35	Early Development of Autonomic Dysfunction May Predict Poor Prognosis in Patients With Multiple System Atrophy. <i>Archives of Neurology</i> , 2007, 64, 256.	4.5	130
36	Long-term therapeutic efficacy and safety of low-dose tacrolimus (FK506) for myasthenia gravis. <i>Journal of the Neurological Sciences</i> , 2006, 247, 17-20.	0.6	32

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37	Multiple Gas-Forming Brain Microabscesses Due to <i>Klebsiella pneumoniae</i> . <i>Archives of Neurology</i> , 2006, 63, 608.	4.5	3
38	Babinski-Nageotte Syndrome With Ipsilateral Hemiparesis. <i>Archives of Neurology</i> , 2005, 62, 676.	4.5	8