

Masato Hasegawa

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

102
papers

11,168
citations

44
h-index

105
g-index

111
ext. papers

13,355
ext. citations

8.5
avg, IF

6.2
L-index

#	Paper	IF	Citations
102	TDP-43 is a component of ubiquitin-positive tau-negative inclusions in frontotemporal lobar degeneration and amyotrophic lateral sclerosis. <i>Biochemical and Biophysical Research Communications</i> , 2006 , 351, 602-11	3.4	1771
101	alpha-Synuclein is phosphorylated in synucleinopathy lesions. <i>Nature Cell Biology</i> , 2002 , 4, 160-4	23.4	1385
100	Prion-like spreading of pathological β -synuclein in brain. <i>Brain</i> , 2013 , 136, 1128-38	11.2	551
99	Phosphorylated TDP-43 in frontotemporal lobar degeneration and amyotrophic lateral sclerosis. <i>Annals of Neurology</i> , 2008 , 64, 60-70	9.4	484
98	Inhibition of heparin-induced tau filament formation by phenothiazines, polyphenols, and porphyrins. <i>Journal of Biological Chemistry</i> , 2005 , 280, 7614-23	5.4	409
97	Tau proteins with FTDP-17 mutations have a reduced ability to promote microtubule assembly. <i>FEBS Letters</i> , 1998 , 437, 207-10	3.8	367
96	Reconsideration of Amyloid Hypothesis and Tau Hypothesis in Alzheimer's Disease. <i>Frontiers in Neuroscience</i> , 2018 , 12, 25	5.1	366
95	Prion-like properties of pathological TDP-43 aggregates from diseased brains. <i>Cell Reports</i> , 2013 , 4, 124-34.6	34.6	322
94	Phosphorylated alpha-synuclein is ubiquitinated in alpha-synucleinopathy lesions. <i>Journal of Biological Chemistry</i> , 2002 , 277, 49071-6	5.4	315
93	Frontotemporal dementia and corticobasal degeneration in a family with a P301S mutation in tau. <i>Journal of Neuropathology and Experimental Neurology</i> , 1999 , 58, 667-77	3.1	301
92	Accumulation of phosphorylated alpha-synuclein in aging human brain. <i>Journal of Neuropathology and Experimental Neurology</i> , 2003 , 62, 644-54	3.1	273
91	Misfolded proteinase K-resistant hyperphosphorylated β -synuclein in aged transgenic mice with locomotor deterioration and in human β -synucleinopathies. <i>Journal of Clinical Investigation</i> , 2002 , 110, 1429-1439	15.9	253
90	Phosphorylated TDP-43 in Alzheimer's disease and dementia with Lewy bodies. <i>Acta Neuropathologica</i> , 2009 , 117, 125-36	14.3	248
89	Seeded aggregation and toxicity of {alpha}-synuclein and tau: cellular models of neurodegenerative diseases. <i>Journal of Biological Chemistry</i> , 2010 , 285, 34885-98	5.4	235
88	Structures of β -synuclein filaments from multiple system atrophy. <i>Nature</i> , 2020 , 585, 464-469	50.4	195
87	Novel tau filament fold in corticobasal degeneration. <i>Nature</i> , 2020 , 580, 283-287	50.4	188
86	Hyperphosphorylation of tau in PHF. <i>Neurobiology of Aging</i> , 1995 , 16, 365-71; discussion 371-80	5.6	180

85	TDP-43 is deposited in the Guam parkinsonism-dementia complex brains. <i>Brain</i> , 2007 , 130, 1386-94	11.2	177
84	Misfolded proteinase K-resistant hyperphosphorylated alpha-synuclein in aged transgenic mice with locomotor deterioration and in human alpha-synucleinopathies. <i>Journal of Clinical Investigation</i> , 2002 , 110, 1429-39	15.9	156
83	Pathological alpha-synuclein propagates through neural networks. <i>Acta Neuropathologica Communications</i> , 2014 , 2, 88	7.3	155
82	Alzheimer-like changes in microtubule-associated protein Tau induced by sulfated glycosaminoglycans. Inhibition of microtubule binding, stimulation of phosphorylation, and filament assembly depend on the degree of sulfation. <i>Journal of Biological Chemistry</i> , 1997 , 272, 33118-24	5.4	148
81	Identification of amino-terminally cleaved tau fragments that distinguish progressive supranuclear palsy from corticobasal degeneration. <i>Annals of Neurology</i> , 2004 , 55, 72-9	9.4	146
80	Regulation of mitochondrial transport and inter-microtubule spacing by tau phosphorylation at the sites hyperphosphorylated in Alzheimer's disease. <i>Journal of Neuroscience</i> , 2012 , 32, 2430-41	6.6	123
79	Biochemical classification of tauopathies by immunoblot, protein sequence and mass spectrometric analyses of sarkosyl-insoluble and trypsin-resistant tau. <i>Acta Neuropathologica</i> , 2016 , 131, 267-280	14.3	122
78	Propagation of pathological synuclein in marmoset brain. <i>Acta Neuropathologica Communications</i> , 2017 , 5, 12	7.3	115
77	Ubiquitination of alpha-synuclein. <i>Biochemistry</i> , 2005 , 44, 361-8	3.2	106
76	Phosphorylated and cleaved TDP-43 in ALS, FTLN and other neurodegenerative disorders and in cellular models of TDP-43 proteinopathy. <i>Neuropathology</i> , 2010 , 30, 170-81	2	87
75	Progranulin regulates lysosomal function and biogenesis through acidification of lysosomes. <i>Human Molecular Genetics</i> , 2017 , 26, 969-988	5.6	84
74	Mass spectrometric analysis of accumulated TDP-43 in amyotrophic lateral sclerosis brains. <i>Scientific Reports</i> , 2016 , 6, 23281	4.9	83
73	Accumulation of phosphorylated TDP-43 in brains of patients with argyrophilic grain disease. <i>Acta Neuropathologica</i> , 2009 , 117, 151-8	14.3	80
72	Conversion of wild-type alpha-synuclein into mutant-type fibrils and its propagation in the presence of A30P mutant. <i>Journal of Biological Chemistry</i> , 2009 , 284, 7940-50	5.4	77
71	Phosphorylated TDP-43 pathology and hippocampal sclerosis in progressive supranuclear palsy. <i>Acta Neuropathologica</i> , 2010 , 120, 55-66	14.3	77
70	Molecular analysis and biochemical classification of TDP-43 proteinopathy. <i>Brain</i> , 2012 , 135, 3380-91	11.2	75
69	Neuronal and glial inclusions in frontotemporal dementia with or without motor neuron disease are immunopositive for p62. <i>Neuroscience Letters</i> , 2003 , 342, 41-4	3.3	73
68	Plasma phosphorylated-TDP-43 protein levels correlate with brain pathology in frontotemporal lobar degeneration. <i>Acta Neuropathologica</i> , 2009 , 118, 647-58	14.3	65

67	Methylene blue reduced abnormal tau accumulation in P301L tau transgenic mice. <i>PLoS ONE</i> , 2012 , 7, e52389	3.7	64
66	Templated Aggregation of TAR DNA-binding Protein of 43 kDa (TDP-43) by Seeding with TDP-43 Peptide Fibrils. <i>Journal of Biological Chemistry</i> , 2016 , 291, 8896-907	5.4	62
65	Cysteine misincorporation in bacterially expressed human alpha-synuclein. <i>FEBS Letters</i> , 2006 , 580, 1775-398	3.8	61
64	Structure-based classification of tauopathies. <i>Nature</i> , 2021 , 598, 359-363	50.4	59
63	The Effect of Fragmented Pathogenic β -Synuclein Seeds on Prion-like Propagation. <i>Journal of Biological Chemistry</i> , 2016 , 291, 18675-88	5.4	58
62	Fibrillogenic nuclei composed of P301L mutant tau induce elongation of P301L tau but not wild-type tau. <i>Journal of Biological Chemistry</i> , 2007 , 282, 20309-18	5.4	57
61	β -Synuclein Fibrils Exhibit Gain of Toxic Function, Promoting Tau Aggregation and Inhibiting Microtubule Assembly. <i>Journal of Biological Chemistry</i> , 2016 , 291, 15046-56	5.4	51
60	Extracellular association of APP and tau fibrils induces intracellular aggregate formation of tau. <i>Acta Neuropathologica</i> , 2015 , 129, 895-907	14.3	50
59	Extensive deamidation at asparagine residue 279 accounts for weak immunoreactivity of tau with RD4 antibody in Alzheimer's disease brain. <i>Acta Neuropathologica Communications</i> , 2013 , 1, 54	7.3	47
58	Prion-like mechanisms and potential therapeutic targets in neurodegenerative disorders. <i>Pharmacology & Therapeutics</i> , 2017 , 172, 22-33	13.9	43
57	Potent prion-like behaviors of pathogenic β -synuclein and evaluation of inactivation methods. <i>Acta Neuropathologica Communications</i> , 2018 , 6, 29	7.3	43
56	Molecular Mechanisms in the Pathogenesis of Alzheimer's disease and Tauopathies-Prion-Like Seeded Aggregation and Phosphorylation. <i>Biomolecules</i> , 2016 , 6,	5.9	43
55	Pathological Endogenous β -Synuclein Accumulation in Oligodendrocyte Precursor Cells Potentially Induces Inclusions in Multiple System Atrophy. <i>Stem Cell Reports</i> , 2018 , 10, 356-365	8	40
54	Isomerase Pin1 stimulates dephosphorylation of tau protein at cyclin-dependent kinase (Cdk5)-dependent Alzheimer phosphorylation sites. <i>Journal of Biological Chemistry</i> , 2013 , 288, 7968-7977	5.4	40
53	Molecular dissection of TDP-43 proteinopathies. <i>Journal of Molecular Neuroscience</i> , 2011 , 45, 480-5	3.3	40
52	Seeded assembly in vitro does not replicate the structures of β -synuclein filaments from multiple system atrophy. <i>FEBS Open Bio</i> , 2021 , 11, 999-1013	2.7	36
51	Progranulin reduction is associated with increased tau phosphorylation in P301L tau transgenic mice. <i>Journal of Neuropathology and Experimental Neurology</i> , 2015 , 74, 158-65	3.1	35
50	The twenty-four kDa C-terminal tau fragment increases with aging in tauopathy mice: implications of prion-like properties. <i>Human Molecular Genetics</i> , 2015 , 24, 6403-16	5.6	35

49	The basis of clinicopathological heterogeneity in TDP-43 proteinopathy. <i>Acta Neuropathologica</i> , 2019 , 138, 751-770	14.3	34
48	Wild-Type Monomeric β Synuclein Can Impair Vesicle Endocytosis and Synaptic Fidelity via Tubulin Polymerization at the Calyx of Held. <i>Journal of Neuroscience</i> , 2017 , 37, 6043-6052	6.6	33
47	Isoform-independent and -dependent phosphorylation of microtubule-associated protein tau in mouse brain during postnatal development. <i>Journal of Biological Chemistry</i> , 2018 , 293, 1781-1793	5.4	28
46	Pathological tau deposition in Motor Neurone Disease and frontotemporal lobar degeneration associated with TDP-43 proteinopathy. <i>Acta Neuropathologica Communications</i> , 2016 , 4, 33	7.3	26
45	Accumulation of multiple neurodegenerative disease-related proteins in familial frontotemporal lobar degeneration associated with granulin mutation. <i>Scientific Reports</i> , 2017 , 7, 1513	4.9	25
44	LATE to the PART-y. <i>Brain</i> , 2019 , 142, e47	11.2	25
43	β Synuclein strains that cause distinct pathologies differentially inhibit proteasome. <i>ELife</i> , 2020 , 9,	8.9	23
42	β Synuclein: Experimental Pathology. <i>Cold Spring Harbor Perspectives in Medicine</i> , 2016 , 6,	5.4	23
41	Mutations in CHCHD2 cause β Synuclein aggregation. <i>Human Molecular Genetics</i> , 2019 , 28, 3895-3911	5.6	22
40	TDP-43 pathology in familial British dementia. <i>Acta Neuropathologica</i> , 2009 , 118, 303-11	14.3	22
39	TDP-43 Prions. <i>Cold Spring Harbor Perspectives in Medicine</i> , 2018 , 8,	5.4	21
38	Comparison of Common and Disease-Specific Post-translational Modifications of Pathological Tau Associated With a Wide Range of Tauopathies. <i>Frontiers in Neuroscience</i> , 2020 , 14, 581936	5.1	18
37	Epitope mapping of antibodies against TDP-43 and detection of protease-resistant fragments of pathological TDP-43 in amyotrophic lateral sclerosis and frontotemporal lobar degeneration. <i>Biochemical and Biophysical Research Communications</i> , 2012 , 417, 116-21	3.4	17
36	Structure of pathological TDP-43 filaments from ALS with FTLD. <i>Nature</i> , 2021 ,	50.4	16
35	Quantitative and combinatorial determination of in situ phosphorylation of tau and its FTDP-17 mutants. <i>Scientific Reports</i> , 2016 , 6, 33479	4.9	15
34	The Abundance of Nonphosphorylated Tau in Mouse and Human Tauopathy Brains Revealed by the Use of Phos-Tag Method. <i>American Journal of Pathology</i> , 2016 , 186, 398-409	5.8	14
33	Prion-like propagation of β Synuclein in neurodegenerative diseases. <i>Progress in Molecular Biology and Translational Science</i> , 2019 , 168, 323-348	4	13
32	Effect of L-DOPA/Benserazide on Propagation of Pathological β Synuclein. <i>Frontiers in Neuroscience</i> , 2019 , 13, 595	5.1	11

31	Human NPCs can degrade β syn fibrils and transfer them preferentially in a cell contact-dependent manner possibly through TNT-like structures. <i>Neurobiology of Disease</i> , 2019 , 132, 104609	7.5	11
30	β Synuclein fibrils subvert lysosome structure and function for the propagation of protein misfolding between cells through tunneling nanotubes. <i>PLoS Biology</i> , 2021 , 19, e3001287	9.7	11
29	Tau isoform expression and phosphorylation in marmoset brains. <i>Journal of Biological Chemistry</i> , 2019 , 294, 11433-11444	5.4	10
28	Structurally distinct β Synuclein fibrils induce robust parkinsonian pathology. <i>Movement Disorders</i> , 2020 , 35, 256-267	7	10
27	-Methyl-D-Aspartate Receptor Link to the MAP Kinase Pathway in Cortical and Hippocampal Neurons and Microglia Is Dependent on Calcium Sensors and Is Blocked by β Synuclein, Tau, and Phospho-Tau in Non-transgenic and Transgenic APP Mice. <i>Frontiers in Molecular Neuroscience</i> , 2018 , 11, 273	6.1	10
26	An autopsy case of globular glial tauopathy presenting with clinical features of motor neuron disease with dementia and iron deposition in the motor cortex. <i>Neuropathology</i> , 2018 , 38, 372	2	9
25	Tau progression in single severe frontal traumatic brain injury in human brains. <i>Journal of the Neurological Sciences</i> , 2019 , 407, 116495	3.2	7
24	Factors associated with development and distribution of granular/fuzzy astrocytes in neurodegenerative diseases. <i>Brain Pathology</i> , 2020 , 30, 811-830	6	7
23	Age-dependent formation of TMEM106B amyloid filaments in human brains.. <i>Nature</i> , 2022 ,	50.4	6
22	Globular glial tauopathy Type I presenting with behavioral variant frontotemporal dementia. <i>Neuropathology</i> , 2020 , 40, 515-525	2	5
21	Asparagine residue 368 is involved in Alzheimer's disease tau strain-specific aggregation. <i>Journal of Biological Chemistry</i> , 2020 , 295, 13996-14014	5.4	5
20	Tau Protein and Frontotemporal Dementias. <i>Advances in Experimental Medicine and Biology</i> , 2021 , 1281, 177-199	3.6	5
19	Human tauopathy-derived tau strains determine the substrates recruited for templated amplification. <i>Brain</i> , 2021 , 144, 2333-2348	11.2	4
18	Distinct phosphorylation profiles of tau in brains of patients with different tauopathies. <i>Neurobiology of Aging</i> , 2021 , 108, 72-79	5.6	4
17	Dextran sulphate-induced tau assemblies cause endogenous tau aggregation and propagation in wild-type mice. <i>Brain Communications</i> , 2020 , 2, fcaa091	4.5	3
16	Development of a novel tau propagation mouse model endogenously expressing 3 and 4 repeat tau isoforms. <i>Brain</i> , 2021 ,	11.2	3
15	Experimental models of prion-like protein propagation. <i>Neuropathology</i> , 2020 , 40, 460-466	2	2
14	Phosphorylation of endogenous β synuclein induced by extracellular seeds initiates at the pre-synaptic region and spreads to the cell body.. <i>Scientific Reports</i> , 2022 , 12, 1163	4.9	2

13	In vivo visualization of propagating β synuclein pathologies in mouse and marmoset models by a bimodal imaging probe, C05-05		2
12	Progression of phosphorylated β synuclein in <i>Macaca fuscata</i> . <i>Brain Pathology</i> , 2021 , 31, e12952	6	2
11	Seeded assembly in vitro does not replicate the structures of β synuclein filaments from multiple system atrophy		1
10	Amyotrophic lateral sclerosis with speech apraxia, predominant upper motor neuron signs, and prominent iron accumulation in the frontal operculum and precentral gyrus. <i>Neuropathology</i> , 2021 , 41, 324-331	2	1
9	Electron Microscopic Analysis of β synuclein Fibrils. <i>Methods in Molecular Biology</i> , 2021 , 2322, 17-25	1.4	1
8	An autopsy case of corticobasal syndrome due to asymmetric degeneration of the motor cortex and substantia nigra with TDP-43 proteinopathy, associated with Alzheimer's disease pathology. <i>Neuropathology</i> , 2021 , 41, 214-225	2	1
7	Ultrastructural and biochemical classification of pathogenic tau, β synuclein and TDP-43.. <i>Acta Neuropathologica</i> , 2022 , 143, 613-640	14.3	1
6	The hot cross bun sign in corticobasal degeneration. <i>Neuropathology</i> , 2021 , 41, 376-380	2	0
5	Common Marmoset Model of β synuclein Propagation. <i>Methods in Molecular Biology</i> , 2021 , 2322, 131-139	1.4	0
4	Independent distribution between tauopathy secondary to subacute sclerotic panencephalitis and measles virus: An immunohistochemical analysis in autopsy cases including cases treated with aggressive antiviral therapies.. <i>Brain Pathology</i> , 2022 , e13069	6	0
3	Structures of tau and β synuclein filaments from brains of patients with neurodegenerative diseases. <i>Neurochemistry International</i> , 2022 , 105362	4.4	0
2	An autopsy case of Alzheimer's disease with amygdala-predominant Lewy pathology presenting with frontotemporal dementia-like psychiatric symptoms.. <i>Neuropathology</i> , 2022 ,	2	
1	An autopsy case of corticobasal degeneration with inferior olivary hypertrophy. <i>Neuropathology</i> , 2021 , 41, 226-235	2	