

# Wado Akamatsu

## List of Publications by Year in descending order

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Version: 2024-02-01

56  
papers

3,775  
citations

117625

34  
h-index

168389

53  
g-index

57  
all docs

57  
docs citations

57  
times ranked

6576  
citing authors

#	ARTICLE	IF	CITATIONS
1	Modeling familial Alzheimer's disease with induced pluripotent stem cells. <i>Human Molecular Genetics</i> , 2011, 20, 4530-4539.	2.9	527
2	Mitochondrial dysfunction associated with increased oxidative stress and $\beta$ -synuclein accumulation in PARK2 iPSC-derived neurons and postmortem brain tissue. <i>Molecular Brain</i> , 2012, 5, 35.	2.6	333
3	Nestin-EGFP Transgenic Mice: Visualization of the Self-Renewal and Multipotency of CNS Stem Cells. <i>Molecular and Cellular Neurosciences</i> , 2001, 17, 259-273.	2.2	298
4	Increased L1 Retrotransposition in the Neuronal Genome in Schizophrenia. <i>Neuron</i> , 2014, 81, 306-313.	8.1	277
5	The RNA-binding protein HuD regulates neuronal cell identity and maturation. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2005, 102, 4625-4630.	7.1	196
6	A human Dravet syndrome model from patient induced pluripotent stem cells. <i>Molecular Brain</i> , 2013, 6, 19.	2.6	111
7	Soluble epoxide hydrolase plays a key role in the pathogenesis of Parkinson's disease. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018, 115, E5815-E5823.	7.1	104
8	Generation of Human Melanocytes from Induced Pluripotent Stem Cells. <i>PLoS ONE</i> , 2011, 6, e16182.	2.5	102
9	Involvement of ER Stress in Dysmyelination of Pelizaeus-Merzbacher Disease with PLP1 Missense Mutations Shown by iPSC-Derived Oligodendrocytes. <i>Stem Cell Reports</i> , 2014, 2, 648-661.	4.8	100
10	RNA-Binding Protein HuD Controls Insulin Translation. <i>Molecular Cell</i> , 2012, 45, 826-835.	9.7	92
11	Neural Stem Cells Directly Differentiated from Partially Reprogrammed Fibroblasts Rapidly Acquire Gliogenic Competency. <i>Stem Cells</i> , 2012, 30, 1109-1119.	3.2	84
12	Controlling the Regional Identity of hPSC-Derived Neurons to Uncover Neuronal Subtype Specificity of Neurological Disease Phenotypes. <i>Stem Cell Reports</i> , 2015, 5, 1010-1022.	4.8	84
13	Differentiation of multipotent neural stem cells derived from Rett syndrome patients is biased toward the astrocytic lineage. <i>Molecular Brain</i> , 2015, 8, 31.	2.6	77
14	Establishment of In Vitro FUS-Associated Familial Amyotrophic Lateral Sclerosis Model Using Human Induced Pluripotent Stem Cells. <i>Stem Cell Reports</i> , 2016, 6, 496-510.	4.8	74
15	Establishment of Induced Pluripotent Stem Cells from Centenarians for Neurodegenerative Disease Research. <i>PLoS ONE</i> , 2012, 7, e41572.	2.5	72
16	Human Induced Pluripotent Stem Cell-Derived Ectodermal Precursor Cells Contribute to Hair Follicle Morphogenesis In Vivo. <i>Journal of Investigative Dermatology</i> , 2013, 133, 1479-1488.	0.7	72
17	Variants in saposin D domain of prosaposin gene linked to Parkinson's disease. <i>Brain</i> , 2020, 143, 1190-1205.	7.6	72
18	Suppression of Oct4 by Germ Cell Nuclear Factor Restricts Pluripotency and Promotes Neural Stem Cell Development in the Early Neural Lineage. <i>Journal of Neuroscience</i> , 2009, 29, 2113-2124.	3.6	64

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19	Prenatal Deletion of the RNA-Binding Protein HuD Disrupts Postnatal Cortical Circuit Maturation and Behavior. <i>Journal of Neuroscience</i> , 2014, 34, 3674-3686.	3.6	62
20	Astrocytes Protect Human Dopaminergic Neurons from $\alpha$ -Synuclein Accumulation and Propagation. <i>Journal of Neuroscience</i> , 2020, 40, 8618-8628.	3.6	57
21	I2020T mutant LRRK2 iPSC-derived neurons in the Sagamihara family exhibit increased Tau phosphorylation through the AKT/GSK-3 $\beta$ signaling pathway. <i>Human Molecular Genetics</i> , 2015, 24, 4879-4900.	2.9	56
22	Functional Neurons Generated from T Cell-Derived Induced Pluripotent Stem Cells for Neurological Disease Modeling. <i>Stem Cell Reports</i> , 2016, 6, 422-435.	4.8	56
23	Comparison of Genomic and Epigenomic Expression in Monozygotic Twins Discordant for Rett Syndrome. <i>PLoS ONE</i> , 2013, 8, e66729.	2.5	56
24	Efficient induction of dopaminergic neuron differentiation from induced pluripotent stem cells reveals impaired mitophagy in PARK2 neurons. <i>Biochemical and Biophysical Research Communications</i> , 2017, 483, 88-93.	2.1	55
25	Escape from Pluripotency via Inhibition of TGF- $\beta$ /BMP and Activation of Wnt Signaling Accelerates Differentiation and Aging in hPSC Progeny Cells. <i>Stem Cell Reports</i> , 2017, 9, 1675-1691.	4.8	54
26	CHARGE syndrome modeling using patient-iPSCs reveals defective migration of neural crest cells harboring CHD7 mutations. <i>ELife</i> , 2017, 6, .	6.0	52
27	Mutations in CHCHD2 cause $\alpha$ -synuclein aggregation. <i>Human Molecular Genetics</i> , 2019, 28, 3895-3911.	2.9	48
28	Differential gene expression profiles in neurons generated from lymphoblastoid B-cell line-derived iPSC cells from monozygotic twin cases with treatment-resistant schizophrenia and discordant responses to clozapine. <i>Schizophrenia Research</i> , 2017, 181, 75-82.	2.0	47
29	Enhanced Aggregation of Androgen Receptor in Induced Pluripotent Stem Cell-derived Neurons from Spinal and Bulbar Muscular Atrophy. <i>Journal of Biological Chemistry</i> , 2013, 288, 8043-8052.	3.4	45
30	Utility of Scalp Hair Follicles as a Novel Source of Biomarker Genes for Psychiatric Illnesses. <i>Biological Psychiatry</i> , 2015, 78, 116-125.	1.3	43
31	Down-regulation of ghrelin receptors on dopaminergic neurons in the substantia nigra contributes to Parkinson's disease-like motor dysfunction. <i>Molecular Brain</i> , 2018, 11, 6.	2.6	43
32	Identifying Therapeutic Agents for Amelioration of Mitochondrial Clearance Disorder in Neurons of Familial Parkinson Disease. <i>Stem Cell Reports</i> , 2020, 14, 1060-1075.	4.8	43
33	Evidence that phosphorylated ubiquitin signaling is involved in the etiology of Parkinson's disease. <i>Human Molecular Genetics</i> , 2017, 26, 3172-3185.	2.9	42
34	Translational derepression of Elavl4 isoforms at their alternative 5' UTRs determines neuronal development. <i>Nature Communications</i> , 2020, 11, 1674.	12.8	40
35	The RNA-binding Protein HuD Regulates Autophagosome Formation in Pancreatic $\beta$ Cells by Promoting Autophagy-related Gene 5 Expression. <i>Journal of Biological Chemistry</i> , 2014, 289, 112-121.	3.4	37
36	Regeneration of the damaged central nervous system through reprogramming technology: Basic concepts and potential application for cell replacement therapy. <i>Experimental Neurology</i> , 2014, 260, 12-18.	4.1	30

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37	Reprogramming non-human primate somatic cells into functional neuronal cells by defined factors. <i>Molecular Brain</i> , 2014, 7, 24.	2.6	26
38	Modeling neurological diseases with induced pluripotent cells reprogrammed from immortalized lymphoblastoid cell lines. <i>Molecular Brain</i> , 2016, 9, 88.	2.6	21
39	Generation of Human Melanocytes from Induced Pluripotent Stem Cells. <i>Methods in Molecular Biology</i> , 2013, 989, 193-215.	0.9	20
40	Cell-specific overexpression of COMT in dopaminergic neurons of Parkinson's disease. <i>Brain</i> , 2019, 142, 1675-1689.	7.6	13
41	Generation of neural cells using iPSCs from sleep bruxism patients with 5-HT2A polymorphism. <i>Journal of Prosthodontic Research</i> , 2017, 61, 242-250.	2.8	12
42	Brief exposure to small molecules allows induction of mouse embryonic fibroblasts into neural crest-like precursors. <i>FEBS Letters</i> , 2017, 591, 590-602.	2.8	11
43	Rostrocaudal Areal Patterning of Human PSC-Derived Cortical Neurons by FGF8 Signaling. <i>ENeuro</i> , 2018, 5, ENEURO.0368-17.2018.	1.9	11
44	Induced Pluripotent Stem Cells Reprogrammed with Three Inhibitors Show Accelerated Differentiation Potentials with High Levels of 2-Cell Stage Marker Expression. <i>Stem Cell Reports</i> , 2019, 12, 305-318.	4.8	10
45	BRIP1, an intracellular bilirubin modulator, exerts neuroprotective activity in a cellular Parkinson's disease model. <i>Journal of Neurochemistry</i> , 2020, 155, 81-97.	3.9	10
46	Developmental dysregulation of excitatory-to-inhibitory GABA-polarity switch may underlie schizophrenia pathology: A monozygotic-twin discordant case analysis in human iPS cell-derived neurons. <i>Neurochemistry International</i> , 2021, 150, 105179.	3.8	9
47	Establishment of an in vitro model for analyzing mitochondrial ultrastructure in PRKN-mutated patient iPSC-derived dopaminergic neurons. <i>Molecular Brain</i> , 2021, 14, 58.	2.6	8
48	Differential X Chromosome Inactivation Patterns during the Propagation of Human Induced Pluripotent Stem Cells. <i>Keio Journal of Medicine</i> , 2016, 66, 1-8.	1.1	6
49	Assessment of Mitophagy in iPS Cell-Derived Neurons. <i>Methods in Molecular Biology</i> , 2017, 1759, 59-67.	0.9	5
50	Differentiation of Midbrain from Human iPS Cells. <i>Methods in Molecular Biology</i> , 2021, 2322, 73-80.	0.9	3
51	Methods to Induce Small-Scale Differentiation of iPS Cells into Dopaminergic Neurons and to Detect Disease Phenotypes. <i>Methods in Molecular Biology</i> , 2021, , 271-279.	0.9	2
52	In vitro monitoring of HTR2A-positive neurons derived from human-induced pluripotent stem cells. <i>Scientific Reports</i> , 2021, 11, 15437.	3.3	2
53	Direct Induction of Neural Stem Cells from Somatic Cells. , 2015, , 103-106.		0
54	Induced Pluripotent Stem Cell Technology in Regenerative Medicine and Disease Modeling. <i>Juntendo Medical Journal</i> , 2017, 63, 37-41.	0.1	0

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55	Modeling Neurological Diseases with Induced Pluripotent Stem Cells. Juntendo Medical Journal, 2018, 64, 450-453.	0.1	0
56	Direct induction of neural cells from somatic cells. , 2020, , 179-185.		0