

Tatsuo Miyamoto

List of Publications by Year in Descending Order

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Version: 2024-04-27

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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.

35
papers

1,256
citations

16
h-index

35
g-index

38
ext. papers

1,544
ext. citations

6.6
avg, IF

3.64
L-index

#	Paper	IF	Citations
35	Imaging of the Ciliary Cholesterol Underlying the Sonic Hedgehog Signal Transduction. <i>Methods in Molecular Biology</i> , 2022 , 2374, 49-57	1.4	0
34	iPSC reprogramming-mediated aneuploidy correction in autosomal trisomy syndromes.. <i>PLoS ONE</i> , 2022 , 17, e0264965	3.7	0
33	Sulforaphane suppresses metastasis of triple-negative breast cancer cells by targeting the RAF/MEK/ERK pathway.. <i>Npj Breast Cancer</i> , 2022 , 8, 40	7.8	2
32	NBS1 I171V variant underlies individual differences in chromosomal radiosensitivity within human populations. <i>Scientific Reports</i> , 2021 , 11, 19661	4.9	0
31	Ciliary GPCR-based transcriptome as a key regulator of cilia length control. <i>FASEB BioAdvances</i> , 2021 , 3, 744-767	2.8	4
30	Premature aging syndrome showing random chromosome number instabilities with CDC20 mutation. <i>Aging Cell</i> , 2020 , 19, e13251	9.9	4
29	A novel CDK-independent function of p27 in preciliary vesicle trafficking during ciliogenesis. <i>Biochemical and Biophysical Research Communications</i> , 2020 , 527, 716-722	3.4	2
28	Genome-wide association meta-analysis identifies GP2 gene risk variants for pancreatic cancer. <i>Nature Communications</i> , 2020 , 11, 3175	17.4	14
27	Applications of Genome Editing Technology in Research on Chromosome Aneuploidy Disorders. <i>Cells</i> , 2020 , 9,	7.9	3
26	Insufficiency of ciliary cholesterol in hereditary Zellweger syndrome. <i>EMBO Journal</i> , 2020 , 39, e103499	13	17
25	Space Radiation Biology for "Living in Space". <i>BioMed Research International</i> , 2020 , 2020, 4703286	3	40
24	Characterization of Functional Primary Cilia in Human Induced Pluripotent Stem Cell-Derived Neurons. <i>Neurochemical Research</i> , 2019 , 44, 1736-1744	4.6	11
23	Coordinated regulation of the dorsal-ventral and anterior-posterior patterning of <i>Xenopus</i> embryos by the BTB/POZ zinc finger protein Zbtb14. <i>Development Growth and Differentiation</i> , 2018 , 60, 158-173	3	7
22	Exploration of genetic basis underlying individual differences in radiosensitivity within human populations using genome editing technology. <i>Journal of Radiation Research</i> , 2018 , 59, ii75-ii82	2.4	7
21	Updated summary of genome editing technology in human cultured cells linked to human genetics studies. <i>Journal of Human Genetics</i> , 2018 , 63, 133-143	4.3	3
20	Albatross/FBF1 contributes to both centriole duplication and centrosome separation. <i>Genes To Cells</i> , 2018 , 23, 1023-1042	2.3	3
19	PLK1-mediated phosphorylation of WDR62/MCPH2 ensures proper mitotic spindle orientation. <i>Human Molecular Genetics</i> , 2017 , 26, 4429-4440	5.6	26

18	Evaluation of ATM heterozygous mutations underlying individual differences in radiosensitivity using genome editing in human cultured cells. <i>Scientific Reports</i> , 2017 , 7, 5996	4.9	14
17	Cytoskeleton-related regulation of primary cilia shortening mediated by melanin-concentrating hormone receptor 1. <i>General and Comparative Endocrinology</i> , 2017 , 253, 44-52	3	19
16	Cilia play a role in breaking left-right symmetry of the sea urchin embryo. <i>Genes To Cells</i> , 2016 , 21, 568-78.3	6	
15	Functional consequence of fibulin-4 missense mutations associated with vascular and skeletal abnormalities and cutis laxa. <i>Matrix Biology</i> , 2016 , 56, 132-149	11.4	17
14	Analysis of individual differences in radiosensitivity using genome editing. <i>Annals of the ICRP</i> , 2016 , 45, 290-6	2.4	6
13	YAP is essential for tissue tension to ensure vertebrate 3D body shape. <i>Nature</i> , 2015 , 521, 217-221	50.4	154
12	The Microtubule-Depolymerizing Activity of a Mitotic Kinesin Protein KIF2A Drives Primary Cilia Disassembly Coupled with Cell Proliferation. <i>Cell Reports</i> , 2015 , 10, 664-673	10.6	78
11	TALEN-mediated single-base-pair editing identification of an intergenic mutation upstream of BUB1B as causative of PCS (MVA) syndrome. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014 , 111, 1461-6	11.5	43
10	Efficient TALEN construction and evaluation methods for human cell and animal applications. <i>Genes To Cells</i> , 2013 , 18, 315-26	2.3	171
9	Repeating pattern of non-RVD variations in DNA-binding modules enhances TALEN activity. <i>Scientific Reports</i> , 2013 , 3, 3379	4.9	165
8	Zinc-finger nuclease-mediated targeted insertion of reporter genes for quantitative imaging of gene expression in sea urchin embryos. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012 , 109, 10915-20	11.5	37
7	Two unrelated patients with MRE11A mutations and Nijmegen breakage syndrome-like severe microcephaly. <i>DNA Repair</i> , 2011 , 10, 314-21	4.3	46
6	Insufficiency of BUBR1, a mitotic spindle checkpoint regulator, causes impaired ciliogenesis in vertebrates. <i>Human Molecular Genetics</i> , 2011 , 20, 2058-70	5.6	41
5	In vivo imaging of tight junctions using claudin-EGFP transgenic medaka. <i>Methods in Molecular Biology</i> , 2011 , 762, 171-8	1.4	2
4	HpSulf, a heparan sulfate 6-O-endosulfatase, is involved in the regulation of VEGF signaling during sea urchin development. <i>Mechanisms of Development</i> , 2010 , 127, 235-45	1.7	30
3	Generation of transgenic medaka expressing claudin7-EGFP for imaging of tight junctions in living medaka embryos. <i>Cell and Tissue Research</i> , 2009 , 335, 465-71	4.2	6
2	Tight junctions in Schwann cells of peripheral myelinated axons: a lesson from claudin-19-deficient mice. <i>Journal of Cell Biology</i> , 2005 , 169, 527-38	7.3	141
1	Compartmentalization established by claudin-11-based tight junctions in stria vascularis is required for hearing through generation of endocochlear potential. <i>Journal of Cell Science</i> , 2004 , 117, 5087-96	5.3	137

