

# Tatsuo Miyamoto

## 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.

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	Efficient TALEN construction and evaluation methods for human cell and animal applications. <i>Genes To Cells</i> , <b>2013</b> , 18, 315-26	2.3	171
34	Repeating pattern of non-RVD variations in DNA-binding modules enhances TALEN activity. <i>Scientific Reports</i> , <b>2013</b> , 3, 3379	4.9	165
33	YAP is essential for tissue tension to ensure vertebrate 3D body shape. <i>Nature</i> , <b>2015</b> , 521, 217-221	50.4	154
32	Tight junctions in Schwann cells of peripheral myelinated axons: a lesson from claudin-19-deficient mice. <i>Journal of Cell Biology</i> , <b>2005</b> , 169, 527-38	7.3	141
31	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> , <b>2004</b> , 117, 5087-96	5.3	137
30	The Microtubule-Depolymerizing Activity of a Mitotic Kinesin Protein KIF2A Drives Primary Cilia Disassembly Coupled with Cell Proliferation. <i>Cell Reports</i> , <b>2015</b> , 10, 664-673	10.6	78
29	Two unrelated patients with MRE11A mutations and Nijmegen breakage syndrome-like severe microcephaly. <i>DNA Repair</i> , <b>2011</b> , 10, 314-21	4.3	46
28	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> , <b>2014</b> , 111, 1461-6	11.5	43
27	Insufficiency of BUBR1, a mitotic spindle checkpoint regulator, causes impaired ciliogenesis in vertebrates. <i>Human Molecular Genetics</i> , <b>2011</b> , 20, 2058-70	5.6	41
26	Space Radiation Biology for "Living in Space". <i>BioMed Research International</i> , <b>2020</b> , 2020, 4703286	3	40
25	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> , <b>2012</b> , 109, 10915-20	11.5	37
24	HpSulf, a heparan sulfate 6-O-endosulfatase, is involved in the regulation of VEGF signaling during sea urchin development. <i>Mechanisms of Development</i> , <b>2010</b> , 127, 235-45	1.7	30
23	PLK1-mediated phosphorylation of WDR62/MCPH2 ensures proper mitotic spindle orientation. <i>Human Molecular Genetics</i> , <b>2017</b> , 26, 4429-4440	5.6	26
22	Cytoskeleton-related regulation of primary cilia shortening mediated by melanin-concentrating hormone receptor 1. <i>General and Comparative Endocrinology</i> , <b>2017</b> , 253, 44-52	3	19
21	Functional consequence of fibulin-4 missense mutations associated with vascular and skeletal abnormalities and cutis laxa. <i>Matrix Biology</i> , <b>2016</b> , 56, 132-149	11.4	17
20	Insufficiency of ciliary cholesterol in hereditary Zellweger syndrome. <i>EMBO Journal</i> , <b>2020</b> , 39, e103499	13	17
19	Genome-wide association meta-analysis identifies GP2 gene risk variants for pancreatic cancer. <i>Nature Communications</i> , <b>2020</b> , 11, 3175	17.4	14

18	Evaluation of ATM heterozygous mutations underlying individual differences in radiosensitivity using genome editing in human cultured cells. <i>Scientific Reports</i> , <b>2017</b> , 7, 5996	4.9	14
17	Characterization of Functional Primary Cilia in Human Induced Pluripotent Stem Cell-Derived Neurons. <i>Neurochemical Research</i> , <b>2019</b> , 44, 1736-1744	4.6	11
16	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> , <b>2018</b> , 60, 158-173	3	7
15	Exploration of genetic basis underlying individual differences in radiosensitivity within human populations using genome editing technology. <i>Journal of Radiation Research</i> , <b>2018</b> , 59, ii75-ii82	2.4	7
14	Cilia play a role in breaking left-right symmetry of the sea urchin embryo. <i>Genes To Cells</i> , <b>2016</b> , 21, 568-78.3		6
13	Generation of transgenic medaka expressing claudin7-EGFP for imaging of tight junctions in living medaka embryos. <i>Cell and Tissue Research</i> , <b>2009</b> , 335, 465-71	4.2	6
12	Analysis of individual differences in radiosensitivity using genome editing. <i>Annals of the ICRP</i> , <b>2016</b> , 45, 290-6	2.4	6
11	Premature aging syndrome showing random chromosome number instabilities with CDC20 mutation. <i>Aging Cell</i> , <b>2020</b> , 19, e13251	9.9	4
10	Ciliary GPCR-based transcriptome as a key regulator of cilia length control. <i>FASEB BioAdvances</i> , <b>2021</b> , 3, 744-767	2.8	4
9	Applications of Genome Editing Technology in Research on Chromosome Aneuploidy Disorders. <i>Cells</i> , <b>2020</b> , 9,	7.9	3
8	Updated summary of genome editing technology in human cultured cells linked to human genetics studies. <i>Journal of Human Genetics</i> , <b>2018</b> , 63, 133-143	4.3	3
7	Albatross/FBF1 contributes to both centriole duplication and centrosome separation. <i>Genes To Cells</i> , <b>2018</b> , 23, 1023-1042	2.3	3
6	A novel CDK-independent function of p27 in preciliary vesicle trafficking during ciliogenesis. <i>Biochemical and Biophysical Research Communications</i> , <b>2020</b> , 527, 716-722	3.4	2
5	In vivo imaging of tight junctions using claudin-EGFP transgenic medaka. <i>Methods in Molecular Biology</i> , <b>2011</b> , 762, 171-8	1.4	2
4	Sulforaphane suppresses metastasis of triple-negative breast cancer cells by targeting the RAF/MEK/ERK pathway.. <i>Npj Breast Cancer</i> , <b>2022</b> , 8, 40	7.8	2
3	NBS1 I171V variant underlies individual differences in chromosomal radiosensitivity within human populations. <i>Scientific Reports</i> , <b>2021</b> , 11, 19661	4.9	0
2	Imaging of the Ciliary Cholesterol Underlying the Sonic Hedgehog Signal Transduction. <i>Methods in Molecular Biology</i> , <b>2022</b> , 2374, 49-57	1.4	0
1	iPSC reprogramming-mediated aneuploidy correction in autosomal trisomy syndromes.. <i>PLoS ONE</i> , <b>2022</b> , 17, e0264965	3.7	0

