

# Ryuji Morizane

## List of Publications by Year in descending order

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Version: 2024-02-01

37  
papers

3,630  
citations

331670

21  
h-index

377865

34  
g-index

38  
all docs

38  
docs citations

38  
times ranked

4538  
citing authors

#	ARTICLE	IF	CITATIONS
1	Nephron organoids derived from human pluripotent stem cells model kidney development and injury. <i>Nature Biotechnology</i> , 2015, 33, 1193-1200.	17.5	694
2	Modelling kidney disease with CRISPR-mutant kidney organoids derived from human pluripotent epiblast spheroids. <i>Nature Communications</i> , 2015, 6, 8715.	12.8	571
3	Flow-enhanced vascularization and maturation of kidney organoids in vitro. <i>Nature Methods</i> , 2019, 16, 255-262.	19.0	559
4	Prediction of DNA Repair Inhibitor Response in Short-Term Patient-Derived Ovarian Cancer Organoids. <i>Cancer Discovery</i> , 2018, 8, 1404-1421.	9.4	311
5	Rapid and Efficient Differentiation of Human Pluripotent Stem Cells into Intermediate Mesoderm That Forms Tubules Expressing Kidney Proximal Tubular Markers. <i>Journal of the American Society of Nephrology: JASN</i> , 2014, 25, 1211-1225.	6.1	271
6	Generation of nephron progenitor cells and kidney organoids from human pluripotent stem cells. <i>Nature Protocols</i> , 2017, 12, 195-207.	12.0	160
7	Interleukin-1 $\beta$ Activates a MYC-Dependent Metabolic Switch in Kidney Stromal Cells Necessary for Progressive Tubulointerstitial Fibrosis. <i>Journal of the American Society of Nephrology: JASN</i> , 2018, 29, 1690-1705.	6.1	152
8	Kidney Organoids: A Translational Journey. <i>Trends in Molecular Medicine</i> , 2017, 23, 246-263.	6.7	114
9	Differentiation of murine embryonic stem and induced pluripotent stem cells to renal lineage in vitro. <i>Biochemical and Biophysical Research Communications</i> , 2009, 390, 1334-1339.	2.1	99
10	The NIH Somatic Cell Genome Editing program. <i>Nature</i> , 2021, 592, 195-204.	27.8	84
11	Proximal tubule ATR regulates DNA repair to prevent maladaptive renal injury responses. <i>Journal of Clinical Investigation</i> , 2019, 129, 4797-4816.	8.2	73
12	The role of microRNA-145 in human embryonic stem cell differentiation into vascular cells. <i>Atherosclerosis</i> , 2011, 219, 468-474.	0.8	57
13	miR-34c attenuates epithelial-mesenchymal transition and kidney fibrosis with ureteral obstruction. <i>Scientific Reports</i> , 2014, 4, 4578.	3.3	54
14	Modeling injury and repair in kidney organoids reveals that homologous recombination governs tubular intrinsic repair. <i>Science Translational Medicine</i> , 2022, 14, eabj4772.	12.4	50
15	Kidney Specific Protein-Positive Cells Derived from Embryonic Stem Cells Reproduce Tubular Structures In Vitro and Differentiate into Renal Tubular Cells. <i>PLoS ONE</i> , 2013, 8, e64843.	2.5	42
16	Induction of human pluripotent stem cells into kidney tissues by synthetic mRNAs encoding transcription factors. <i>Scientific Reports</i> , 2019, 9, 913.	3.3	40
17	Generation of kidney tubular organoids from human pluripotent stem cells. <i>Scientific Reports</i> , 2016, 6, 38353.	3.3	36
18	Concise Review: Kidney Generation with Human Pluripotent Stem Cells. <i>Stem Cells</i> , 2017, 35, 2209-2217.	3.2	35

#	ARTICLE	IF	CITATIONS
19	Kidney organoids in translational medicine: Disease modeling and regenerative medicine. <i>Developmental Dynamics</i> , 2020, 249, 34-45.	1.8	33
20	Meclizine Preconditioning Protects the Kidney Against Ischemiaâ€œReperfusion Injury. <i>EBioMedicine</i> , 2015, 2, 1090-1101.	6.1	32
21	Selective depletion of mouse kidney proximal straight tubule cells causes acute kidney injury. <i>Transgenic Research</i> , 2012, 21, 51-62.	2.4	24
22	Epigenetic transcriptional reprogramming by WT1 mediates a repair response during podocyte injury. <i>Science Advances</i> , 2020, 6, eabb5460.	10.3	19
23	3D kidney organoids for bench-to-bedside translation. <i>Journal of Molecular Medicine</i> , 2021, 99, 477-487.	3.9	19
24	A case of atypical POEMS syndrome without polyneuropathy. <i>European Journal of Haematology</i> , 2008, 80, 452-455.	2.2	13
25	Kidney organoids: a pioneering model for kidney diseases. <i>Translational Research</i> , 2022, 250, 1-17.	5.0	12
26	Renal amyloidosis caused by apolipoprotein A-II without a genetic mutation in the coding sequence. <i>Clinical and Experimental Nephrology</i> , 2011, 15, 774-779.	1.6	11
27	CRISPR/Cas9â€œbased Targeted Genome Editing for the Development of Monogenic Diseases Models with Human Pluripotent Stem Cells. <i>Current Protocols in Stem Cell Biology</i> , 2018, 45, e50.	3.0	11
28	Directed Differentiation of Pluripotent Stem Cells into Kidney. <i>Biomarker Insights</i> , 2015, 10s1, BMI.S20055.	2.5	10
29	miR-363 induces transdifferentiation of human kidney tubular cells to mesenchymal phenotype. <i>Clinical and Experimental Nephrology</i> , 2016, 20, 394-401.	1.6	9
30	MPO-ANCA associated crescentic glomerulonephritis with numerous immune complexes: case report. <i>BMC Nephrology</i> , 2012, 13, 32.	1.8	8
31	Bioengineered Kidney Models: Methods and Functional Assessments. <i>Function</i> , 2021, 2, zqab026.	2.3	8
32	Kidney development to kidney organoids and back again. <i>Seminars in Cell and Developmental Biology</i> , 2022, 127, 68-76.	5.0	6
33	Modelling diabetic vasculopathy with human vessel organoids. <i>Nature Reviews Nephrology</i> , 2019, 15, 258-260.	9.6	5
34	Regenerative Medicine, Disease Modeling, and Drug Discovery in Human Pluripotent Stem Cell-derived Kidney Tissue. <i>European Medical Journal Reproductive Health</i> , 2017, 3, 57-67.	1.0	4
35	Organoids for modeling kidney disease. , 2018, , 227-245.		2
36	The application of iPSC-derived kidney organoids and genome editing in kidney disease modeling. , 2022, , 111-136.		2

#	ARTICLE	IF	CITATIONS
37	Revealing potential cardiac manifestation of ADPKD using iPS cell-derived cardiomyocytes. EBioMedicine, 2019, 40, 19-20.	6.1	0