

Minoru Takasato

List of Publications by Year in descending order

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Version: 2024-02-01

40
papers

3,791
citations

430874

18
h-index

395702

33
g-index

48
all docs

48
docs citations

48
times ranked

3972
citing authors

#	ARTICLE	IF	CITATIONS
1	Inducing human retinal pigment epithelium-like cells from somatic tissue. <i>Stem Cell Reports</i> , 2022, 17, 289-306.	4.8	3
2	On-Chip Compartmentalized Vascular Bed Preserves Kidney Organoid Differentiation. , 2022, , .		0
3	Kidney organoid research: current status and applications. <i>Current Opinion in Genetics and Development</i> , 2022, 75, 101944.	3.3	10
4	Evaluation of the Permeability of Cell Barriers Constituted of Kidney Organoid-Derived Glomerulus. , 2021, , .		1
5	Proximal Tubule On A Chip For Evaluating P-Glycoprotein Transport Property. , 2021, , .		0
6	Effect of Perfusion Culture on Localization, Intensity, and Functionality of Transporter Proteins in a Bilayer Proximal Tubule-on-a Chip. , 2021, , .		0
7	Reprogramming epiblast stem cells into pre-implantation blastocyst cell-like cells. <i>Stem Cell Reports</i> , 2021, 16, 1197-1209.	4.8	6
8	Perspective: Extending the Utility of Three-Dimensional Organoids by Tissue Clearing Technologies. <i>Frontiers in Cell and Developmental Biology</i> , 2021, 9, 679226.	3.7	12
9	27 α -Hydroxycholesterol regulates human <i>SLC22A12</i> gene expression through estrogen receptor action. <i>FASEB Journal</i> , 2021, 35, e21262.	0.5	10
10	Multivariate patterning of human pluripotent cells under perfusion reveals critical roles of induced paracrine factors in kidney organoid development. <i>Science Advances</i> , 2020, 6, eaaw2746.	10.3	21
11	An In Vitro Differentiation Protocol for Human Embryonic Bipotential Gonad and Testis Cell Development. <i>Stem Cell Reports</i> , 2020, 15, 1377-1391.	4.8	22
12	Generation of two human induced pluripotent stem cell lines derived from two juvenile nephronophthisis patients with NPHP1 deletion. <i>Stem Cell Research</i> , 2020, 45, 101815.	0.7	5
13	Development of an exon skipping therapy for X-linked Alport syndrome with truncating variants in COL4A5. <i>Nature Communications</i> , 2020, 11, 2777.	12.8	46
14	Control and design of biosystems. <i>Development Growth and Differentiation</i> , 2020, 62, 149-149.	1.5	0
15	Challenges to future regenerative applications using kidney organoids. <i>Current Opinion in Biomedical Engineering</i> , 2020, 13, 144-151.	3.4	9
16	Genetic background, recent advances in molecular biology, and development of novel therapy in Alport syndrome. <i>Kidney Research and Clinical Practice</i> , 2020, 39, 402-413.	2.2	13
17	Advice for the Next Generation: Minoru Takasato. <i>Cell Stem Cell</i> , 2019, 24, 688-689.	11.1	0
18	Evaluation of variability in human kidney organoids. <i>Nature Methods</i> , 2019, 16, 79-87.	19.0	176

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19	Renal Subcapsular Transplantation of PSC-Derived Kidney Organoids Induces Neo-vasculogenesis and Significant Glomerular and Tubular Maturation In Vivo. <i>Stem Cell Reports</i> , 2018, 10, 751-765.	4.8	304
20	Making a Kidney Organoid Using the Directed Differentiation of Human Pluripotent Stem Cells. <i>Methods in Molecular Biology</i> , 2017, 1597, 195-206.	0.9	34
21	Recapitulating Development to Generate Kidney Organoid Cultures. , 2017, , 193-222.		0
22	Challenges to regenerate the kidney. <i>Japanese Journal of Nephrology</i> , 2017, 59, 11-15.	0.0	0
23	A strategy for generating kidney organoids: Recapitulating the development in human pluripotent stem cells. <i>Developmental Biology</i> , 2016, 420, 210-220.	2.0	42
24	Understanding kidney morphogenesis to guide renal tissue regeneration. <i>Nature Reviews Nephrology</i> , 2016, 12, 624-635.	9.6	38
25	Generation of kidney organoids from human pluripotent stem cells. <i>Nature Protocols</i> , 2016, 11, 1681-1692.	12.0	243
26	A protocol for the identification and validation of novel genetic causes of kidney disease. <i>BMC Nephrology</i> , 2015, 16, 152.	1.8	8
27	The origin of the mammalian kidney: implications for recreating the kidney <i>in vitro</i> . <i>Development (Cambridge)</i> , 2015, 142, 1937-1947.	2.5	98
28	Generating a self-organizing kidney from pluripotent cells. <i>Current Opinion in Organ Transplantation</i> , 2015, 20, 178-186.	1.6	16
29	Kidney organoids from human iPS cells contain multiple lineages and model human nephrogenesis. <i>Nature</i> , 2015, 526, 564-568.	27.8	1,210
30	Recreating kidney progenitors from pluripotent cells. <i>Pediatric Nephrology</i> , 2014, 29, 543-552.	1.7	22
31	Directing human embryonic stem cell differentiation towards a renal lineage generates a self-organizing kidney. <i>Nature Cell Biology</i> , 2014, 16, 118-126.	10.3	640
32	Reprogramming Somatic Cells to a Kidney Fate. <i>Seminars in Nephrology</i> , 2014, 34, 462-480.	1.6	7
33	Direct Transcriptional Reprogramming of Adult Cells to Embryonic Nephron Progenitors. <i>Journal of the American Society of Nephrology: JASN</i> , 2013, 24, 1424-1434.	6.1	119
34	Induction of intermediate mesoderm by retinoic acid receptor signaling from differentiating mouse embryonic stem cells. <i>International Journal of Developmental Biology</i> , 2013, 57, 383-389.	0.6	28
35	Trb2, a mouse homolog of tribbles, is dispensable for kidney and mouse development. <i>Biochemical and Biophysical Research Communications</i> , 2008, 373, 648-652.	2.1	11
36	Identification of multipotent progenitors in the embryonic mouse kidney by a novel colony-forming assay. <i>Development (Cambridge)</i> , 2006, 133, 151-161.	2.5	172

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37	The murine homolog of <i>SALL4</i> , a causative gene in Okihiro syndrome, is essential for embryonic stem cell proliferation, and cooperates with <i>Sall1</i> in anorectal, heart, brain and kidney development. <i>Development (Cambridge)</i> , 2006, 133, 3005-3013.	2.5	241
38	Essential roles of <i>Sall1</i> in kidney development. <i>Kidney International</i> , 2005, 68, 1948-1950.	5.2	32
39	Identification of kidney mesenchymal genes by a combination of microarray analysis and <i>Sall1</i> -GFP knockin mice. <i>Mechanisms of Development</i> , 2004, 121, 547-557.	1.7	64
40	Generation of kidney organoids from human pluripotent stem cells. <i>Protocol Exchange</i> , 0, , .	0.3	3