

Sanjay Jain

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/8523724/publications.pdf>

Version: 2024-02-01

80
papers

6,210
citations

71102

41
h-index

74163

75
g-index

89
all docs

89
docs citations

89
times ranked

10155
citing authors

#	ARTICLE	IF	CITATIONS
1	Pathogenic Germline Variants in 10,389 Adult Cancers. <i>Cell</i> , 2018, 173, 355-370.e14.	28.9	620
2	Glial Cell-Line Derived Neurotrophic Factor-Mediated RET Signaling Regulates Spermatogonial Stem Cell Fate. <i>Biology of Reproduction</i> , 2006, 74, 314-321.	2.7	347
3	Isolation and Characterization of PBP, a Protein That Interacts with Peroxisome Proliferator-activated Receptor. <i>Journal of Biological Chemistry</i> , 1997, 272, 25500-25506.	3.4	313
4	Expression of ARNT, ARNT2, HIF1 β , HIF2 β and Ah receptor mRNAs in the developing mouse. <i>Mechanisms of Development</i> , 1998, 73, 117-123.	1.7	311
5	Overexpression of ABCA1 reduces amyloid deposition in the PDAPP mouse model of Alzheimer disease. <i>Journal of Clinical Investigation</i> , 2008, 118, 671-82.	8.2	301
6	Adventitial MSC-like Cells Are Progenitors of Vascular Smooth Muscle Cells and Drive Vascular Calcification in Chronic Kidney Disease. <i>Cell Stem Cell</i> , 2016, 19, 628-642.	11.1	254
7	A single-nucleus RNA-sequencing pipeline to decipher the molecular anatomy and pathophysiology of human kidneys. <i>Nature Communications</i> , 2019, 10, 2832.	12.8	206
8	Etv4 and Etv5 are required downstream of GDNF and Ret for kidney branching morphogenesis. <i>Nature Genetics</i> , 2009, 41, 1295-1302.	21.4	199
9	Characterization of the Ah Receptor-associated Protein, ARA9. <i>Journal of Biological Chemistry</i> , 1998, 273, 33580-33587.	3.4	180
10	Computational Segmentation and Classification of Diabetic Glomerulosclerosis. <i>Journal of the American Society of Nephrology: JASN</i> , 2019, 30, 1953-1967.	6.1	142
11	Critical and distinct roles for key RET tyrosine docking sites in renal development. <i>Genes and Development</i> , 2006, 20, 321-333.	5.9	137
12	Mice expressing a dominant-negative Ret mutation phenocopy human Hirschsprung disease and delineate a direct role of Ret in spermatogenesis. <i>Development (Cambridge)</i> , 2004, 131, 5503-5513.	2.5	112
13	Conditional ablation of GFR β 1 in postmigratory enteric neurons triggers unconventional neuronal death in the colon and causes a Hirschsprung's disease phenotype. <i>Development (Cambridge)</i> , 2007, 134, 2171-2181.	2.5	112
14	Single cell transcriptomics identifies focal segmental glomerulosclerosis remission endothelial biomarker. <i>JCI Insight</i> , 2020, 5, .	5.0	108
15	An integrated functional genomics and metabolomics approach for defining poor prognosis in human neuroendocrine cancers. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2005, 102, 9901-9906.	7.1	102
16	Differential Expression of the Peroxisome Proliferator-Activated Receptor β (PPAR β) and Its Coactivators Steroid Receptor Coactivator-1 and PPAR-Binding Protein PBP in the Brown Fat, Urinary Bladder, Colon, and Breast of the Mouse. <i>American Journal of Pathology</i> , 1998, 153, 349-354.	3.8	100
17	An integrated iterative annotation technique for easing neural network training in medical image analysis. <i>Nature Machine Intelligence</i> , 2019, 1, 112-119.	16.0	96
18	Mice lacking sister chromatid cohesion protein PDS5B exhibit developmental abnormalities reminiscent of Cornelia de Lange syndrome. <i>Development (Cambridge)</i> , 2007, 134, 3191-3201.	2.5	94

#	ARTICLE	IF	CITATIONS
19	Rationale and design of the Kidney Precision Medicine Project. <i>Kidney International</i> , 2021, 99, 498-510.	5.2	94
20	Angiotensin II Triggers Peripheral Macrophage-to-Sensory Neuron Redox Crosstalk to Elicit Pain. <i>Journal of Neuroscience</i> , 2018, 38, 7032-7057.	3.6	92
21	RET Is Dispensable for Maintenance of Midbrain Dopaminergic Neurons in Adult Mice. <i>Journal of Neuroscience</i> , 2006, 26, 11230-11238.	3.6	88
22	Traditional and targeted exome sequencing reveals common, rare and novel functional deleterious variants in RET-signaling complex in a cohort of living US patients with urinary tract malformations. <i>Human Genetics</i> , 2012, 131, 1725-1738.	3.8	84
23	RET Signaling Is Required for Survival and Normal Function of Nonpeptidergic Nociceptors. <i>Journal of Neuroscience</i> , 2010, 30, 3983-3994.	3.6	80
24	Multi-radial LBP Features as a Tool for Rapid Glomerular Detection and Assessment in Whole Slide Histopathology Images. <i>Scientific Reports</i> , 2018, 8, 2032.	3.3	74
25	Dosage Effects of Cohesin Regulatory Factor PDS5 on Mammalian Development: Implications for Cohesinopathies. <i>PLoS ONE</i> , 2009, 4, e5232.	2.5	74
26	Plasticity of distal nephron epithelia from human kidney organoids enables the induction of ureteric tip and stalk. <i>Cell Stem Cell</i> , 2021, 28, 671-684.e6.	11.1	72
27	To bud or not to bud: the RET perspective in CAKUT. <i>Pediatric Nephrology</i> , 2014, 29, 597-608.	1.7	68
28	Anatomical structures, cell types and biomarkers of the Human Reference Atlas. <i>Nature Cell Biology</i> , 2021, 23, 1117-1128.	10.3	68
29	A reference tissue atlas for the human kidney. <i>Science Advances</i> , 2022, 8, .	10.3	67
30	Expression Profiles Provide Insights into Early Malignant Potential and Skeletal Abnormalities in Multiple Endocrine Neoplasia Type 2B Syndrome Tumors. <i>Cancer Research</i> , 2004, 64, 3907-3913.	0.9	66
31	The many faces of RET dysfunction in kidney. <i>Organogenesis</i> , 2009, 5, 177-190.	1.2	66
32	Injury-induced actin cytoskeleton reorganization in podocytes revealed by super-resolution microscopy. <i>JCI Insight</i> , 2017, 2, .	5.0	65
33	Novel mechanisms of early upper and lower urinary tract patterning regulated by RetY1015 docking tyrosine in mice. <i>Development (Cambridge)</i> , 2012, 139, 2405-2415.	2.5	64
34	Developmental pathology of congenital kidney and urinary tract anomalies. <i>CKJ: Clinical Kidney Journal</i> , 2019, 12, 382-399.	2.9	63
35	A multimodal and integrated approach to interrogate human kidney biopsies with rigor and reproducibility: guidelines from the Kidney Precision Medicine Project. <i>Physiological Genomics</i> , 2021, 53, 1-11.	2.3	59
36	Targeted Exome Sequencing Integrated with Clinicopathological Information Reveals Novel and Rare Mutations in Atypical, Suspected and Unknown Cases of Alport Syndrome or Proteinuria. <i>PLoS ONE</i> , 2013, 8, e76360.	2.5	59

#	ARTICLE	IF	CITATIONS
37	(Re)Building a Kidney. <i>Journal of the American Society of Nephrology: JASN</i> , 2017, 28, 1370-1378.	6.1	58
38	Automated Computational Detection of Interstitial Fibrosis, Tubular Atrophy, and Glomerulosclerosis. <i>Journal of the American Society of Nephrology: JASN</i> , 2021, 32, 837-850.	6.1	52
39	Organotypic specificity of key RET adaptor-docking sites in the pathogenesis of neurocristopathies and renal malformations in mice. <i>Journal of Clinical Investigation</i> , 2010, 120, 778-790.	8.2	50
40	B cell-derived IL-4 acts on podocytes to induce proteinuria and foot process effacement. <i>JCI Insight</i> , 2017, 2, .	5.0	48
41	Loss of Sprouty1 Rescues Renal Agenesis Caused by Ret Mutation. <i>Journal of the American Society of Nephrology: JASN</i> , 2009, 20, 255-259.	6.1	45
42	A Toolbox to Characterize Human Induced Pluripotent Stem Cell-Derived Kidney Cell Types and Organoids. <i>Journal of the American Society of Nephrology: JASN</i> , 2019, 30, 1811-1823.	6.1	45
43	Modelling kidney disease using ontology: insights from the Kidney Precision Medicine Project. <i>Nature Reviews Nephrology</i> , 2020, 16, 686-696.	9.6	45
44	Interaction between DMRT1 function and genetic background modulates signaling and pluripotency to control tumor susceptibility in the fetal germ line. <i>Developmental Biology</i> , 2013, 377, 67-78.	2.0	44
45	Yap and Taz are required for Ret-dependent urinary tract morphogenesis. <i>Development (Cambridge)</i> , 2015, 142, 2696-2703.	2.5	44
46	Genome-wide significance testing of variation from single case exomes. <i>Nature Genetics</i> , 2016, 48, 1455-1461.	21.4	43
47	Dopamine-Dependent Compensation Maintains Motor Behavior in Mice with Developmental Ablation of Dopaminergic Neurons. <i>Journal of Neuroscience</i> , 2013, 33, 17095-17107.	3.6	41
48	A role for genetic susceptibility in sporadic focal segmental glomerulosclerosis. <i>Journal of Clinical Investigation</i> , 2016, 126, 1067-1078.	8.2	41
49	The Overlap and Distinction of Self-Reported Symptoms between Interstitial Cystitis/Bladder Pain Syndrome and Overactive Bladder: A Questionnaire Based Analysis. <i>Journal of Urology</i> , 2014, 192, 1679-1686.	0.4	35
50	Expression regulation and function of heparan sulfate 6-O-endosulfatases in the spermatogonial stem cell niche. <i>Glycobiology</i> , 2011, 21, 152-161.	2.5	34
51	Expression profiles of congenital renal dysplasia reveal new insights into renal development and disease. <i>Pediatric Nephrology</i> , 2007, 22, 962-974.	1.7	32
52	Midline signaling regulates kidney positioning but not nephrogenesis through Shh. <i>Developmental Biology</i> , 2010, 340, 518-527.	2.0	30
53	Peroxisome Proliferator-Activated Receptor α -Responsive Genes Induced in the Newborn but Not Prenatal Liver of Peroxisomal Fatty Acyl-CoA Oxidase Null Mice. <i>Experimental Cell Research</i> , 2001, 268, 70-76.	2.6	28
54	Neurturin-Mediated Ret Activation Is Required for Retinal Function. <i>Journal of Neuroscience</i> , 2008, 28, 4123-4135.	3.6	28

#	ARTICLE	IF	CITATIONS
55	Neural Crest Cell Origin and Signals for Intrinsic Neurogenesis in the Mammalian Respiratory Tract. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2011, 44, 293-301.	2.9	28
56	FAT4 Fine-Tunes Kidney Development by Regulating RET Signaling. <i>Developmental Cell</i> , 2019, 48, 780-792.e4.	7.0	27
57	Stage specific requirement of Gfr1±1 in the ureteric epithelium during kidney development. <i>Mechanisms of Development</i> , 2013, 130, 506-518.	1.7	26
58	Centrosome amplification disrupts renal development and causes cystogenesis. <i>Journal of Cell Biology</i> , 2018, 217, 2485-2501.	5.2	24
59	Development of an Immunoassay for the Kidney-Specific Protein myo-Inositol Oxygenase, a Potential Biomarker of Acute Kidney Injury. <i>Clinical Chemistry</i> , 2014, 60, 747-757.	3.2	23
60	Claudin 1 and nephrin label cellular crescents in diabetic glomerulosclerosis. <i>Human Pathology</i> , 2014, 45, 628-635.	2.0	23
61	A multiplexed analysis approach identifies new association of inflammatory proteins in patients with overactive bladder. <i>American Journal of Physiology - Renal Physiology</i> , 2016, 311, F28-F34.	2.7	21
62	Reduced endothelin converting enzyme-1 and endothelin-3 mRNA in the developing bowel of male mice may increase expressivity and penetrance of Hirschsprung disease-like distal intestinal aganglionosis. <i>Developmental Dynamics</i> , 2007, 236, 106-117.	1.8	20
63	The Med1 Subunit of the Mediator Complex Induces Liver Cell Proliferation and Is Phosphorylated by AMP Kinase. <i>Journal of Biological Chemistry</i> , 2013, 288, 27898-27911.	3.4	19
64	Expression profiles of podocytes exposed to high glucose reveal new insights into early diabetic glomerulopathy. <i>Laboratory Investigation</i> , 2011, 91, 488-498.	3.7	18
65	Reciprocal Spatiotemporally Controlled Apoptosis Regulates Wolffian Duct Cloaca Fusion. <i>Journal of the American Society of Nephrology: JASN</i> , 2018, 29, 775-783.	6.1	18
66	Systemic Nonurological Symptoms in Patients with Overactive Bladder. <i>Journal of Urology</i> , 2016, 196, 467-472.	0.4	17
67	Clinical phenotype of APOL1 nephropathy in young relatives of patients with end-stage renal disease. <i>Pediatric Nephrology</i> , 2015, 30, 983-989.	1.7	15
68	Quantification of vascular damage in acute kidney injury with fluorine magnetic resonance imaging and spectroscopy. <i>Magnetic Resonance in Medicine</i> , 2018, 79, 3144-3153.	3.0	14
69	Glial cell line-derived neurotrophic factor protects against high-fat diet-induced hepatic steatosis by suppressing hepatic PPAR- β expression. <i>American Journal of Physiology - Renal Physiology</i> , 2016, 310, G103-G116.	3.4	11
70	Integrated single-cell sequencing and histopathological analyses reveal diverse injury and repair responses in a participant with acute kidney injury: a clinical-molecular-pathologic correlation. <i>Kidney International</i> , 2022, 101, 1116-1125.	5.2	11
71	7-dehydrocholesterol efficiently supports Ret signaling in a mouse model of Smith-Opitz-Lemli syndrome. <i>Scientific Reports</i> , 2016, 6, 28534.	3.3	9
72	WNT pathway signaling is associated with microvascular injury and predicts kidney transplant failure. <i>American Journal of Transplantation</i> , 2019, 19, 2833-2845.	4.7	7

#	ARTICLE	IF	CITATIONS
73	The Interface of Genetics with Pathology in Alport Nephritis. Journal of the American Society of Nephrology: JASN, 2013, 24, 1925-1927.	6.1	6
74	Routine use of clinical exome-based next-generation sequencing for evaluation of patients with thrombotic microangiopathies. Modern Pathology, 2017, 30, 1739-1747.	5.5	6
75	Patient perspectives and involvement in precision medicine research. Kidney International, 2021, 99, 511-514.	5.2	5
76	A Pilot Study of Urine Proteomics in COVID-19-associated Acute Kidney Injury. Kidney International Reports, 2021, 6, 3064-3069.	0.8	5
77	Imaging centrosomes and cilia in the mouse kidney. Methods in Cell Biology, 2015, 127, 1-17.	1.1	4
78	Validating single-cell genomics for the study of renal development. Kidney International, 2014, 86, 1049-1055.	5.2	3
79	Peroxidasin Is a Novel Target of Autoantibodies in Lupus Nephritis. Kidney International Reports, 2019, 4, 1004-1006.	0.8	3
80	Characterization of Human iPSC RET Reporter Cell Line Differentiation to Kidney and Neural Crest Lineages. FASEB Journal, 2018, 32, 649.3.	0.5	0