

John J Bissler

List of Publications by Year in descending order

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114
papers

8,529
citations

87888

38
h-index

45317

90
g-index

115
all docs

115
docs citations

115
times ranked

6986
citing authors

#	ARTICLE	IF	CITATIONS
1	Tsc2 mutation induces renal tubular cell nonautonomous disease. <i>Genes and Diseases</i> , 2022, 9, 187-200.	3.4	9
2	Acute kidney injury in pediatric hematopoietic cell transplantation: critical appraisal and consensus. <i>Pediatric Nephrology</i> , 2022, 37, 1179-1203.	1.7	10
3	Urinary pCO ₂ Monitoring System with a Planar Severinghaus Type Sensor. <i>Electroanalysis</i> , 2022, 34, 1587-1597.	2.9	1
4	Single Gene Mutations in Pkd1 or Tsc2 Alter Extracellular Vesicle Production and Trafficking. <i>Biology</i> , 2022, 11, 709.	2.8	3
5	Cutaneous Angiomyolipoma of the Eyelid in a 2-Year-Old With Tuberous Sclerosis Complex. <i>Journal of Neuro-Ophthalmology</i> , 2021, 41, e69-e70.	0.8	2
6	Continuous Renal Replacement Therapy: A Review of Use and Application in Pediatric Hematopoietic Stem Cell Transplant Recipients. <i>Frontiers in Oncology</i> , 2021, 11, 632263.	2.8	5
7	Kidney intercalated cells and the transcription factor FOXi1 drive cystogenesis in tuberous sclerosis complex. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2021, 118, .	7.1	14
8	Tsc Gene Locus Disruption and Differences in Renal Epithelial Extracellular Vesicles. <i>Frontiers in Physiology</i> , 2021, 12, 630933.	2.8	5
9	Racial Health Disparity and COVID-19. <i>Journal of NeuroImmune Pharmacology</i> , 2021, 16, 729-742.	4.1	7
10	Renal cystic disease in tuberous sclerosis complex. <i>Experimental Biology and Medicine</i> , 2021, 246, 2111-2117.	2.4	4
11	Updated International Tuberous Sclerosis Complex Diagnostic Criteria and Surveillance and Management Recommendations. <i>Pediatric Neurology</i> , 2021, 123, 50-66.	2.1	230
12	Leptin receptor defect with diabetes causes skeletal muscle atrophy in female obese Zucker rats where peculiar depots networked with mitochondrial damages. <i>Ultrastructural Pathology</i> , 2021, 45, 346-375.	0.9	1
13	Renal Involvement in Tuberous Sclerosis Complex. , 2021, , 1-12.		0
14	Congenital Lymphatic Malformation and Aortic Aneurysm in a Patient with TSC2 Mutation. <i>Neuropediatrics</i> , 2020, 51, 057-061.	0.6	12
15	Tuberous Sclerosis Complex Axis Controls Renal Extracellular Vesicle Production and Protein Content. <i>International Journal of Molecular Sciences</i> , 2020, 21, 1729.	4.1	16
16	Tuberous sclerosis complex and the kidney. , 2020, , 251-258.e3.		0
17	Effect of everolimus on renal function in patients with tuberous sclerosis complex: evidence from EXIST-1 and EXIST-2. <i>Nephrology Dialysis Transplantation</i> , 2019, 34, 1000-1008.	0.7	31
18	A step-wise approach for establishing a multidisciplinary team for the management of tuberous sclerosis complex: a Delphi consensus report. <i>Orphanet Journal of Rare Diseases</i> , 2019, 14, 91.	2.7	36

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19	Tuberous sclerosis complex exhibits a new renal cystogenic mechanism. <i>Physiological Reports</i> , 2019, 7, e13983.	1.7	23
20	Rapalog resistance is associated with mesenchymal-type changes in Tsc2-null cells. <i>Scientific Reports</i> , 2019, 9, 3015.	3.3	15
21	Everolimus compliance and persistence among tuberous sclerosis complex patients with renal angiomyolipoma or subependymal giant cell astrocytoma. <i>Current Medical Research and Opinion</i> , 2019, 35, 1103-1110.	1.9	4
22	Problems of nephrooncology. Proceedings from the 1st Scientific and Training Conference Nephrooncology 5-6 October 2018, Gdańsk, Poland. <i>Polish Archives of Internal Medicine</i> , 2019, 129, 1-74.	0.4	2
23	The effect of everolimus on renal angiomyolipoma in pediatric patients with tuberous sclerosis being treated for subependymal giant cell astrocytoma. <i>Pediatric Nephrology</i> , 2018, 33, 101-109.	1.7	37
24	Acute Kidney Injury in Pediatric Patients Receiving Allogeneic Hematopoietic Cell Transplantation: Incidence, Risk Factors, and Outcomes. <i>Biology of Blood and Marrow Transplantation</i> , 2018, 24, 758-764.	2.0	39
25	Renal manifestation of tuberous sclerosis complex. <i>American Journal of Medical Genetics, Part C: Seminars in Medical Genetics</i> , 2018, 178, 338-347.	1.6	48
26	Angiomyolipoma rebound tumor growth after discontinuation of everolimus in patients with tuberous sclerosis complex or sporadic lymphangiomyomatosis. <i>PLoS ONE</i> , 2018, 13, e0201005.	2.5	27
27	Generation, clearance, toxicity, and monitoring possibilities of unaccounted uremic toxins for improved dialysis prescriptions. <i>American Journal of Physiology - Renal Physiology</i> , 2018, 315, F890-F902.	2.7	5
28	Cystic Kidney Diseases Associated with Increased Cancer Risk: Tuberous Sclerosis Complex, Von Hippel-Lindau, and Birt-Hogg-Dubé. , 2018, , 51-66.		3
29	Outcomes of angioembolization and nephrectomy for renal angiomyolipoma associated with tuberous sclerosis complex: a real-world US national study. <i>Current Medical Research and Opinion</i> , 2017, 33, 821-827.	1.9	13
30	Healthcare utilization and costs in patients with tuberous sclerosis complex-related renal angiomyolipoma. <i>Journal of Medical Economics</i> , 2017, 20, 388-394.	2.1	7
31	Primary cilia regulate the osmotic stress response of renal epithelial cells through TRPM3. <i>American Journal of Physiology - Renal Physiology</i> , 2017, 312, F791-F805.	2.7	23
32	Natural history of patients with tuberous sclerosis complex related renal angiomyolipoma. <i>Current Medical Research and Opinion</i> , 2017, 33, 1277-1282.	1.9	8
33	Everolimus long-term use in patients with tuberous sclerosis complex: Four-year update of the EXIST-2 study. <i>PLoS ONE</i> , 2017, 12, e0180939.	2.5	128
34	Pooled analysis of menstrual irregularities from three major clinical studies evaluating everolimus for the treatment of tuberous sclerosis complex. <i>PLoS ONE</i> , 2017, 12, e0186235.	2.5	10
35	Measurement of glomerular filtration rate by dynamic contrast-enhanced magnetic resonance imaging using a subject-specific two-compartment model. <i>Physiological Reports</i> , 2016, 4, e12755.	1.7	9
36	Pharmacokinetics and pharmacodynamics of everolimus in patients with renal angiomyolipoma and tuberous sclerosis complex or lymphangiomyomatosis. <i>British Journal of Clinical Pharmacology</i> , 2016, 81, 958-970.	2.4	23

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37	Advances and Future Directions for Tuberous Sclerosis Complex Research: Recommendations From the 2015 Strategic Planning Conference. <i>Pediatric Neurology</i> , 2016, 60, 1-12.	2.1	43
38	Long-term Clinical Morbidity in Patients With Renal Angiomyolipoma Associated With Tuberous Sclerosis Complex. <i>Urology</i> , 2016, 95, 80-87.	1.0	28
39	Review of the Tuberous Sclerosis Renal Guidelines from the 2012 Consensus Conference: Current Data and Future Study. <i>Nephron</i> , 2016, 134, 51-58.	1.8	58
40	Official American Thoracic Society/Japanese Respiratory Society Clinical Practice Guidelines: Lymphangioleiomyomatosis Diagnosis and Management. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2016, 194, 748-761.	5.6	236
41	Optimal treatment of tuberous sclerosis complex associated renal angiomyolipomata: a systematic review. <i>Therapeutic Advances in Urology</i> , 2016, 8, 279-290.	2.0	39
42	Everolimus for renal angiomyolipoma in patients with tuberous sclerosis complex or sporadic lymphangioleiomyomatosis: extension of a randomized controlled trial. <i>Nephrology Dialysis Transplantation</i> , 2016, 31, 111-119.	0.7	120
43	Renal function after treatment for childhood cancer.. <i>Journal of Clinical Oncology</i> , 2016, 34, 10571-10571.	1.6	0
44	Response to everolimus is seen in TSC-associated SEGAs and angiomyolipomas independent of mutation type and site in TSC1 and TSC2. <i>European Journal of Human Genetics</i> , 2015, 23, 1665-1672.	2.8	29
45	Therapies for polycystic kidney disease. <i>Current Opinion in Pediatrics</i> , 2015, 27, 227-232.	2.0	1
46	Intracranial arterial aneurysms in children and young adults. <i>Journal of Pediatric Neuroradiology</i> , 2015, 02, 203-235.	0.1	0
47	Rates of interventional procedures in patients with tuberous sclerosis complex-related renal angiomyolipoma. <i>Current Medical Research and Opinion</i> , 2015, 31, 1501-1507.	1.9	15
48	Insight into response to mTOR inhibition when PKD1 and TSC2 are mutated. <i>BMC Medical Genetics</i> , 2015, 16, 39.	2.1	15
49	Hyperglycemia in the absence of cilia accelerates cystogenesis and induces renal damage. <i>American Journal of Physiology - Renal Physiology</i> , 2015, 309, F79-F87.	2.7	16
50	Evidence for pericyte origin of TSC-associated renal angiomyolipomas and implications for angiotensin receptor inhibition therapy. <i>American Journal of Physiology - Renal Physiology</i> , 2014, 307, F560-F570.	2.7	44
51	Tuberous sclerosis complex, mTOR, and the kidney: report of an NIDDK-sponsored workshop. <i>American Journal of Physiology - Renal Physiology</i> , 2014, 306, F279-F283.	2.7	17
52	Characterization of Renal Toxicity in Mice Administered the Marine Biotoxin Domoic Acid. <i>Journal of the American Society of Nephrology: JASN</i> , 2014, 25, 1187-1197.	6.1	24
53	The effect of everolimus on renal angiomyolipoma in patients with tuberous sclerosis complex being treated for subependymal giant cell astrocytoma: subgroup results from the randomized, placebo-controlled, Phase 3 trial EXIST-1. <i>Nephrology Dialysis Transplantation</i> , 2014, 29, 1203-1210.	0.7	79
54	Consensus Expert Recommendations for the Diagnosis and Management of Autosomal Recessive Polycystic Kidney Disease: Report of an International Conference. <i>Journal of Pediatrics</i> , 2014, 165, 611-617.	1.8	138

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55	Performance of cystatin C-based equations in a pediatric cohort at high risk of kidney injury. <i>Pediatric Nephrology</i> , 2013, 28, 453-461.	1.7	36
56	Tuberous Sclerosis Complex Diagnostic Criteria Update: Recommendations of the 2012 International Tuberous Sclerosis Complex Consensus Conference. <i>Pediatric Neurology</i> , 2013, 49, 243-254.	2.1	1,185
57	Tuberous Sclerosis Complex Surveillance and Management: Recommendations of the 2012 International Tuberous Sclerosis Complex Consensus Conference. <i>Pediatric Neurology</i> , 2013, 49, 255-265.	2.1	693
58	Everolimus for angiomyolipoma associated with tuberous sclerosis complex or sporadic lymphangioleiomyomatosis (EXIST-2): a multicentre, randomised, double-blind, placebo-controlled trial. <i>Lancet, The</i> , 2013, 381, 817-824.	13.7	712
59	Lymphangioleiomyomatosis Screening in Women With Tuberous Sclerosis. <i>Chest</i> , 2013, 144, 578-585.	0.8	129
60	Human TSC-associated renal angiomyolipoma cells are hypersensitive to ER stress. <i>American Journal of Physiology - Renal Physiology</i> , 2012, 303, F831-F844.	2.7	33
61	Impact of Ultrasound-Guided Kidney Biopsy Simulation on Trainee Confidence and Biopsy Outcomes. <i>American Journal of Nephrology</i> , 2012, 36, 570-574.	3.1	23
62	Replication Fork Stalling and Checkpoint Activation by a PKD1 Locus Mirror Repeat Polypurine-Polypyrimidine (Pu-Py) Tract. <i>Journal of Biological Chemistry</i> , 2012, 287, 33412-33423.	3.4	38
63	Real-world treatment profile for patients with tuberous sclerosis complex related angiomyolipoma: A U.S. Healthcare Claims Database study.. <i>Journal of Clinical Oncology</i> , 2012, 30, e15096-e15096.	1.6	0
64	Prevalence of angiomyolipoma among patients with tuberous sclerosis complex: A U.S. Healthcare Claims Database study.. <i>Journal of Clinical Oncology</i> , 2012, 30, e12042-e12042.	1.6	0
65	Updated safety results from EXIST-2: Everolimus therapy for angiomyolipoma (AML) associated with tuberous sclerosis complex (TSC) or sporadic lymphangioleiomyomatosis (sLAM).. <i>Journal of Clinical Oncology</i> , 2012, 30, 4632-4632.	1.6	1
66	Effect of everolimus on angiogenic biomarkers in patients with tuberous sclerosis complex (TSC): Results from EXIST-1 and EXIST-2.. <i>Journal of Clinical Oncology</i> , 2012, 30, 10619-10619.	1.6	2
67	Cell Cycle Control and DNA Damage Response of Conditionally Immortalized Urothelial Cells. <i>PLoS ONE</i> , 2011, 6, e16595.	2.5	9
68	Clinical and Molecular Insights into Tuberous Sclerosis Complex Renal Disease. <i>Pediatric Nephrology</i> , 2011, 26, 839-852.	1.7	51
69	Loss of Primary Cilia Upregulates Renal Hypertrophic Signaling and Promotes Cystogenesis. <i>Journal of the American Society of Nephrology: JASN</i> , 2011, 22, 839-848.	6.1	79
70	The human DEK oncogene regulates DNA damage response signaling and repair. <i>Nucleic Acids Research</i> , 2011, 39, 7465-7476.	14.5	82
71	Polycystic Kidney Disease. , 2011, , 703-712.		0
72	Serum Vascular Endothelial Growth Factor-D Prospectively Distinguishes Lymphangioleiomyomatosis From Other Diseases. <i>Chest</i> , 2010, 138, 674-681.	0.8	188

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73	Glomerulocystic kidney disease. <i>Pediatric Nephrology</i> , 2010, 25, 2049-2059.	1.7	48
74	Replication-dependent instability at (CTG) _n (CAG) repeat hairpins in human cells. <i>Nature Chemical Biology</i> , 2010, 6, 652-659.	8.0	135
75	Simulation of real-time ultrasound-guided renal biopsy. <i>Kidney International</i> , 2010, 78, 705-707.	5.2	22
76	Tuberous Sclerosis Complex Renal Disease. <i>Nephron Experimental Nephrology</i> , 2010, 118, e15-e20.	2.2	134
77	Automated Algorithm for Quantifying the Extent of Cystic Change on Volumetric Chest CT: Initial Results in Lymphangiomyomatosis. <i>American Journal of Roentgenology</i> , 2009, 192, 1037-1044.	2.2	31
78	Renal involvement in tuberous sclerosis complex and von Hippel-Lindau disease: shared disease mechanisms?. <i>Nature Reviews Nephrology</i> , 2009, 5, 143-156.	9.6	13
79	Increased cancer risk of augmentation cystoplasty: Possible role for hyperosmolal microenvironment on DNA damage recognition. <i>Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis</i> , 2009, 670, 88-95.	1.0	19
80	Nephritic factor and recurrence in the renal transplant of membranoproliferative glomerulonephritis type II. <i>Pediatric Nephrology</i> , 2008, 23, 1867-1876.	1.7	8
81	RecQ and RecG helicases have distinct roles in maintaining the stability of polypurine-polypyrimidine sequences. <i>Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis</i> , 2008, 643, 20-28.	1.0	15
82	Sirolimus for Angiomyolipoma in Tuberous Sclerosis Complex or Lymphangiomyomatosis. <i>New England Journal of Medicine</i> , 2008, 358, 140-151.	27.0	1,138
83	Analysis of PKD1 for genomic deletion by multiplex ligation-dependent probe assay: Absence of hot spots. <i>Genomics</i> , 2008, 91, 203-208.	2.9	19
84	The Tuberous Sclerosis Complex Regulates Trafficking of Glucose Transporters and Glucose Uptake. <i>American Journal of Pathology</i> , 2008, 172, 1748-1756.	3.8	46
85	Characterization of Fabry Disease in 352 Pediatric Patients in the Fabry Registry. <i>Pediatric Research</i> , 2008, 64, 550-555.	2.3	235
86	Replication Protein A is Required for Etoposide-Induced Assembly of MRE11/RAD50/NBS1 Complex Repair Foci. <i>Cell Cycle</i> , 2007, 6, 2408-2416.	2.6	12
87	Cell Cycle- and Proteasome-Dependent Formation of Etoposide-Induced Replication Protein A (RPA) or Mre11/Rad50/Nbs1 (MRN) Complex Repair Foci. <i>Cell Cycle</i> , 2007, 6, 2399-2407.	2.6	10
88	Unstable Spinocerebellar Ataxia Type 10 (ATTCT) _n (AGAAT) Repeats Are Associated with Aberrant Replication at the ATX10 Locus and Replication Origin-Dependent Expansion at an Ectopic Site in Human Cells. <i>Molecular and Cellular Biology</i> , 2007, 27, 7828-7838.	2.3	55
89	Identification of 54 large deletions/duplications in TSC1 and TSC2 using MLPA, and genotype-phenotype correlations. <i>Human Genetics</i> , 2007, 121, 389-400.	3.8	162
90	Triplex DNA and human disease. <i>Frontiers in Bioscience - Landmark</i> , 2007, 12, 4536.	3.0	53

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91	Embolization of Renal Angiomyolipomata in Patients With Tuberous Sclerosis Complex. American Journal of Kidney Diseases, 2006, 47, 95-102.	1.9	95
92	DNA Structures and Genetic Instabilities Associated with Spinocerebellar Ataxia Type 10 (ATTCT) n A-(AGAAT) n Repeats Suggest a DNA Amplification Model for Repeat Expansion. , 2006, , 447-460.		2
93	A mechanistic approach to inherited polycystic kidney disease. Pediatric Nephrology, 2005, 20, 558-566.	1.7	23
94	DNA Lesion-specific Co-localization of the Mre11/Rad50/Nbs1 (MRN) Complex and Replication Protein A (RPA) to Repair Foci. Journal of Biological Chemistry, 2005, 280, 12927-12934.	3.4	37
95	Frequency and Imaging Appearance of Hepatic Angiomyolipomas in Pediatric and Adult Patients with Tuberous Sclerosis. American Journal of Roentgenology, 2004, 182, 1027-1030.	2.2	68
96	PKD1 intron 21: triplex DNA formation and effect on replication. Nucleic Acids Research, 2004, 32, 1460-1468.	14.5	41
97	Renal angiomyolipomata. Kidney International, 2004, 66, 924-934.	5.2	266
98	Renal, hepatic, and marrow dysfunction in a patient with chronic renal insufficiency. Pediatric Nephrology, 2003, 18, 293-296.	1.7	3
99	Unpaired Structures in SCA10 (ATTCT)nA-(AGAAT)n Repeats. Journal of Molecular Biology, 2003, 326, 1095-1111.	4.2	90
100	Comparative evaluation of Î±-galactosidase A infusions for treatment of Fabry disease. Genetics in Medicine, 2003, 5, 144-153.	2.4	25
101	Apolipoprotein J/Clusterin Prevents a Progressive Glomerulopathy of Aging. Molecular and Cellular Biology, 2002, 22, 1893-1902.	2.3	99
102	Tuberous Sclerosis Complex: Renal Imaging Findings. Radiology, 2002, 225, 451-456.	7.3	161
103	Reduction of postembolization syndrome after ablation of renal angiomyolipoma. American Journal of Kidney Diseases, 2002, 39, 966-971.	1.9	112
104	Association between a High-Expressing Interferon-Î³ Allele and a Lower Frequency of Kidney Angiomyolipomas in TSC2 Patients. American Journal of Human Genetics, 2002, 71, 750-758.	6.2	48
105	Optimizing DpnI Digestion Conditions to Detect Replicated DNA. BioTechniques, 2002, 33, 316-318.	1.8	22
106	Infantile Dilated X-Linked Cardiomyopathy, G4.5 Mutations, Altered Lipids, and Ultrastructural Malformations of Mitochondria in Heart, Liver, and Skeletal Muscle. Laboratory Investigation, 2002, 82, 335-344.	3.7	57
107	DNA inverted repeats and human disease. Frontiers in Bioscience - Landmark, 1998, 3, d408-418.	3.0	55
108	C1 Inhibitor Gene Sequence Facilitates Frameshift Mutations. Molecular Medicine, 1998, 4, 795-806.	4.4	9

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109	Antibody to C1 β -inhibitor in a patient receiving C1 β -inhibitor infusions for treatment of hereditary angioneurotic edema with systemic lupus erythematosus reacts with a normal allotype of residue 458 of C1 β -inhibitor. <i>Translational Research</i> , 1996, 128, 438-443.	2.3	17
110	Contiguous deletion and duplication mutations resulting in type 1 hereditary angioneurotic edema. <i>Human Genetics</i> , 1994, 93, 265-269.	3.8	31
111	[6] C1 Inhibitor. <i>Methods in Enzymology</i> , 1993, 223, 97-120.	1.0	13
112	C1 inhibitor hinge region mutations produce dysfunction by different mechanisms. <i>Nature Genetics</i> , 1992, 1, 354-358.	21.4	68
113	Paradoxical hypertension in hypovolemic children. <i>Pediatric Emergency Care</i> , 1991, 7, 350-352.	0.9	5
114	Alimentary Tract Duplications in Children. <i>Clinical Pediatrics</i> , 1988, 27, 152-157.	0.8	45