Bradley K Yoder

List of Publications by Year
in descending order

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		44042	51562
94	10,754	48	86
papers	citations	h-index	g-index

95	95	95	9380
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	The Polycystic Kidney Disease Proteins, Polycystin-1, Polycystin-2, Polaris, and Cystin, Are Co-Localized in Renal Cilia. Journal of the American Society of Nephrology: JASN, 2002, 13, 2508-2516.	3.0	835
2	Gli2 and Gli3 Localize to Cilia and Require the Intraflagellar Transport Protein Polaris for Processing and Function. PLoS Genetics, 2005, 1, e53.	1.5	815
3	Comparative Genomics Identifies a Flagellar and Basal Body Proteome that Includes the BBS5 Human Disease Gene. Cell, 2004, 117, 541-552.	13.5	721
4	Cilia-driven fluid flow in the zebrafish pronephros, brain and Kupffer's vesicle is required for normal organogenesis. Development (Cambridge), 2005, 132, 1907-1921.	1.2	600
5	Disruption of Intraflagellar Transport in Adult Mice Leads to Obesity and Slow-Onset Cystic Kidney Disease. Current Biology, 2007, 17, 1586-1594.	1.8	425
6	Intraflagellar transport is essential for endochondral bone formation. Development (Cambridge), 2007, 134, 307-316.	1.2	343
7	Dysfunctional cilia lead to altered ependyma and choroid plexus function,and result in the formation of hydrocephalus. Development (Cambridge), 2005, 132, 5329-5339.	1.2	319
8	THM1 negatively modulates mouse sonic hedgehog signal transduction and affects retrograde intraflagellar transport in cilia. Nature Genetics, 2008, 40, 403-410.	9.4	313
9	Ciliary proteins link basal body polarization to planar cell polarity regulation. Nature Genetics, 2008, 40, 69-77.	9.4	306
10	Polaris, a Protein Involved in Left-Right Axis Patterning, Localizes to Basal Bodies and Cilia. Molecular Biology of the Cell, 2001, 12, 589-599.	0.9	296
11	An incredible decade for the primary cilium: a look at a once-forgotten organelle. American Journal of Physiology - Renal Physiology, 2005, 289, F1159-F1169.	1.3	289
12	Role of Primary Cilia in the Pathogenesis of Polycystic Kidney Disease: Figure 1.. Journal of the American Society of Nephrology: JASN, 2007, 18, 1381-1388.	3.0	257
13	Primary Cilia and Signaling Pathways in Mammalian Development, Health and Disease. Nephron Physiology, 2009, 111, p39-p53.	1.5	241
14	Directional Cell Migration and Chemotaxis in Wound Healing Response to PDGF-AA are Coordinated by the Primary Cilium in Fibroblasts. Cellular Physiology and Biochemistry, 2010, 25, 279-292.	1.1	226
15	Polaris, a protein disrupted inorpkmutant mice, is required for assembly of renal cilium. American Journal of Physiology - Renal Physiology, 2002, 282, F541-F552.	1.3	218
16	Chapter 13 Ciliary Dysfunction in Developmental Abnormalities and Diseases. Current Topics in Developmental Biology, 2008, 85, 371-427.	1.0	213
17	Cilia Proteins Control Cerebellar Morphogenesis by Promoting Expansion of the Granule Progenitor Pool. Journal of Neuroscience, 2007, 27, 9780-9789.	1.7	186
18	The Primary Cilium in Cell Signaling and Cancer: Figure 1.. Cancer Research, 2006, 66, 6463-6467.	0.4	181

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19	Cystin, a novel cilia-associated protein, is disrupted in the cpk mouse model of polycystic kidney disease. Journal of Clinical Investigation, 2002, 109, 533-540.	3.9	176
20	Lack of Primary Cilia Primes Shear-Induced Endothelial-to-Mesenchymal Transition. Circulation Research, 2011, 108, 1093-1101.	2.0	173
21	Soluble levels of cytosolic tubulin regulate ciliary length control. Molecular Biology of the Cell, 2011, 22, 806-816.	0.9	150
22	Development of the post-natal growth plate requires intraflagellar transport proteins. Developmental Biology, 2007, 305, 202-216.	0.9	145
23	Mechanoregulation of intracellular Ca2+ concentration is attenuated in collecting duct of monocilium-impaired orpk mice. American Journal of Physiology - Renal Physiology, 2005, 289, F978-F988.	1.3	144
24	The Tumor-Associated Glycosyltransferase ST6Gal-I Regulates Stem Cell Transcription Factors and Confers a Cancer Stem Cell Phenotype. Cancer Research, 2016, 76, 3978-3988.	0.4	134
25	Cystin, a novel cilia-associated protein, is disrupted in the cpk mouse model of polycystic kidney disease. Journal of Clinical Investigation, 2002, 109, 533-540.	3.9	131
26	Single-Cell RNA Sequencing Identifies Candidate Renal Resident Macrophage Gene Expression Signatures across Species. Journal of the American Society of Nephrology: JASN, 2019, 30, 767-781.	3.0	126
27	Loss of theTg737 protein results in skeletal patterning defects. Developmental Dynamics, 2003, 227, 78-90.	0.8	121
28	XBX-1 Encodes a Dynein Light Intermediate Chain Required for Retrograde Intraflagellar Transport and Cilia Assembly inCaenorhabditis elegans. Molecular Biology of the Cell, 2003, 14, 2057-2070.	0.9	120
29	Primary cilia regulate Shh activity in the control of molar tooth number. Development (Cambridge), 2009, 136, 897-903.	1.2	113
30	The Oak Ridge Polycystic Kidney mouse: Modeling ciliopathies of mice and men. Developmental Dynamics, 2008, 237, 1960-1971.	0.8	112
31	High-Throughput Genome Editing and Phenotyping Facilitated by High Resolution Melting Curve Analysis. PLoS ONE, 2014, 9, e114632.	1.1	112
32	The C. elegans homologs of nephrocystin-1 and nephrocystin-4 are cilia transition zone proteins involved in chemosensory perception. Journal of Cell Science, 2005, 118, 5575-5587.	1.2	103
33	Gene therapy rescues cilia defects and restores olfactory function in a mammalian ciliopathy model. Nature Medicine, 2012, 18, 1423-1428.	15.2	103
34	The primary cilium coordinates early cardiogenesis and hedgehog signaling in cardiomyocyte differentiation. Journal of Cell Science, 2009, 122, 3070-3082.	1.2	91
35	Loss of primary cilia results in deregulated and unabated apical calcium entry in ARPKD collecting duct cells. American Journal of Physiology - Renal Physiology, 2006, 290, F1320-F1328.	1.3	86
36	Leptin resistance is a secondary consequence of the obesity in ciliopathy mutant mice. Proceedings of the National Academy of Sciences of the United States of America, 2013, 110, 7796-7801.	3.3	82

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37	Primary cilia enhance kisspeptin receptor signaling on gonadotropin-releasing hormone neurons. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 10335-10340.	3.3	81
38	Identification of CHE-13, a novel intraflagellar transport protein required for cilia formation. Experimental Cell Research, 2003, 284, 249-261.	1.2	80
39	Disruption of IFT results in both exocrine and endocrine abnormalities in the pancreas of Tg737orpk mutant mice. Laboratory Investigation, 2005, 85, 45-64.	1.7	80
40	The zebrafish foxj1a transcription factor regulates cilia function in response to injury and epithelial stretch. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 18499-18504.	3.3	80
41	Cystic Kidney Diseases: All Roads Lead to the Cilium. Physiology, 2004, 19, 225-230.	1.6	77
42	GMAP210 and IFT88 are present in the spermatid golgi apparatus and participate in the development of the acrosomeâ€“acroplaxome complex, headâ€“tail coupling apparatus and tail. Developmental Dynamics, 2011, 240, 723-736.	0.8	77
43	Resident macrophages reprogram toward a developmental state after acute kidney injury. JCI Insight, 2019, 4, .	2.3	75
44	Characterization of primary cilia and Hedgehog signaling during development of the human pancreas and in human pancreatic duct cancer cell lines. Developmental Dynamics, 2008, 237, 2039-2052.	0.8	69
45	An inducible CiliaGFP mouse model for in vivo visualization and analysis of cilia in live tissue. Cilia, 2013, 2, 8.	1.8	68
46	Inflammation and Fibrosis in Polycystic Kidney Disease. Results and Problems in Cell Differentiation, 2017, 60, 323-344.	0.2	68
47	Mutations in Traf3ip1 reveal defects in ciliogenesis, embryonic development, and altered cell size regulation. Developmental Biology, 2011, 360, 66-76.	0.9	59
48	Role of epidermal primary cilia in the homeostasis of skin and hair follicles. Development (Cambridge), 2011, 138, 1675-1685.	1.2	58
49	Hippocampal and Cortical Primary Cilia Are Required for Aversive Memory in Mice. PLoS ONE, 2014, 9, e106576.	1.1	58
50	Coiled-coil domain containing 42 (Ccdc 42) is necessary for proper sperm development and male fertility in the mouse. Developmental Biology, 2016, 412, 208-218.	0.9	54
51	Heterotrimeric Kinesin-2 (KIF3) Mediates Transition Zone and Axoneme Formation of Mouse Photoreceptors. Journal of Biological Chemistry, 2015, 290, 12765-12778.	1.6	53
52	Altered pHi regulation and Na+/HCO3â^' transporter activity in choroid plexus of cilia-defective Tg737orpk mutant mouse. American Journal of Physiology - Cell Physiology, 2007, 292, C1409-C1416.	2.1	52
53	Tissue-Resident Macrophages Promote Renal Cystic Disease. Journal of the American Society of Nephrology: JASN, 2019, 30, 1841-1856.	3.0	51
54	Molecular pathogenesis of autosomal dominant polycystic kidney disease. Expert Reviews in Molecular Medicine, 2006, 8, 1-22.	1.6	46

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55	Proximal Tubule Proliferation Is Insufficient to Induce Rapid Cyst Formation after Cilia Disruption. Journal of the American Society of Nephrology: JASN, 2013, 24, 456-464.	3.0	44
56	Heightened epithelial Na+channel-mediated Na+absorption in a murine polycystic kidney disease model epithelium lacking apical monocilia. American Journal of Physiology - Cell Physiology, 2006, 290, C952-C963.	2.1	43
57	Microtubule modifications and stability are altered by cilia perturbation and in cystic kidney disease. Cytoskeleton, 2013, 70, 24-31.	1.0	42
58	Functional correction of renal defects in a mouse model for ARPKD through expression of the cloned wild-type Tg737 cDNA. Kidney International, 1996, 50, 1240-1248.	2.6	34
59	Mutation of Growth Arrest Specific 8 Reveals a Role in Motile Cilia Function and Human Disease. PLoS Genetics, 2016, 12, e1006220.	1.5	33
60	SnapShot: Sensing and Signaling by Cilia. Cell, 2015, 161, 692-692.e1.	13.5	27
61	Mammalian Clusterin associated protein 1 is an evolutionarily conserved protein required for ciliogenesis. Cilia, 2012, 1, 20.	1.8	26
62	Urinary T cells correlate with rate of renal function loss in autosomal dominant polycystic kidney disease. Physiological Reports, 2019, 7, e13951.	0.7	25
63	Primary cilia disruption differentially affects the infiltrating and resident macrophage compartment in the liver. American Journal of Physiology - Renal Physiology, 2018, 314, G677-G689.	1.6	23
64	Mks6 mutations reveal tissueâ€•and cell typeâ€specific roles for the cilia transition zone. FASEB Journal, 2019, 33, 1440-1455.	0.2	19
65	BBSome Component BBS5 Is Required for Cone Photoreceptor Protein Trafficking and Outer Segment Maintenance. , 2020, 61, 17.		19
66	Interferon Regulatory Factorâ€5 in Resident Macrophage Promotes Polycystic Kidney Disease. Kidney360, 2020, 1, 179-190.	0.9	19
67	Role for primary cilia in the regulation of mouse ovarian function. Developmental Dynamics, 2008, 237, 2053-2060.	0.8	18
68	Quantitative Peptidomics of Purkinje Cell Degeneration Mice. PLoS ONE, 2013, 8, e60981.	1.1	18
69	Heterozygous Pkhd1^{C642*} mice develop cystic liver disease and proximal tubule ectasia that mimics radiographic signs of medullary sponge kidney. American Journal of Physiology - Renal Physiology, 2019, 316, F463-F472.	1.3	17
70	A Screen for Modifiers of Cilia Phenotypes Reveals Novel MKS Alleles and Uncovers a Specific Genetic Interaction between osm-3 and nphp-4. PLoS Genetics, 2016, 12, e1005841.	1.5	17
71	Truncating PKHD1 and PKD2 mutations alter energy metabolism. American Journal of Physiology - Renal Physiology, 2019, 316, F414-F425.	1.3	16
72	Resident Macrophages in Cystic Kidney Disease. Kidney360, 2021, 2, 167-175.	0.9	16

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73	Genetic and Informatic Analyses Implicate Kif12 as a Candidate Gene within the Mpkd2 Locus That Modulates Renal Cystic Disease Severity in the Cys1cpk Mouse. PLoS ONE, 2015, 10, e0135678.	1.1	13
74	Non-essential role for cilia in coordinating precise alignment of lens fibres. Mechanisms of Development, 2016, 139, 10-17.	1.7	13
75	Increased Na⁺/H⁺exchanger activity on the apical surface of a cilium-deficient cortical collecting duct principal cell model of polycystic kidney disease. American Journal of Physiology - Cell Physiology, 2012, 302, C1436-C1451.	2.1	12
76	Intravital visualization of the primary cilium, tubule flow, and innate immune cells in the kidney utilizing an abdominal window imaging approach. Methods in Cell Biology, 2019, 154, 67-83.	0.5	10
77	A mouse model of BBS identifies developmental and homeostatic effects of BBS5 mutation and identifies novel pituitary abnormalities. Human Molecular Genetics, 2021, 30, 234-246.	1.4	10
78	Ectopic Phosphorylated Creb Marks Dedifferentiated Proximal Tubules in Cystic Kidney Disease. American Journal of Pathology, 2018, 188, 84-94.	1.9	9
79	Human transcription factors responsive to initial reprogramming predominantly undergo legitimate reprogramming during fibroblast conversion to iPSCs. Scientific Reports, 2020, 10, 19710.	1.6	9
80	Kidney resident macrophages in the rat have minimal turnover and replacement by blood monocytes. American Journal of Physiology - Renal Physiology, 2021, 321, F162-F169.	1.3	7
81	Evolutionarily conserved genetic interactions between nphp-4 and bbs-5 mutations exacerbate ciliopathy phenotypes. Genetics, 2022, 220, .	1.2	7
82	Preface. Current Topics in Developmental Biology, 2008, 85, xv-xix.	1.0	4
83	Ly6chi Infiltrating Macrophages Promote Cyst Progression in Injured Conditional Ift88 Mice. Kidney360, 2021, 2, 989-995.	0.9	4
84	Monitoring Endosomal Trafficking of the G Protein-Coupled Receptor Somatostatin Receptor 3. Methods in Enzymology, 2014, 534, 261-280.	0.4	3
85	A transgenic <scp>Alx4â€CreER</scp> mouse to analyze anterior limb and nephric duct development. Developmental Dynamics, 2022, 251, 1524-1534.	0.8	2
86	Autocrine Purinergic Signaling Is Required for Monociliumâ€driven Signaling. FASEB Journal, 2007, 21, A503.	0.2	2
87	NHE Dysregulation in Cilium Deficient Mouse Renal Principal Cells from orpk Mice. FASEB Journal, 2007, 21, A504.	0.2	1
88	Renal Cilia Structure, Function, and Physiology. , 2013, , 319-346.		0
89	Early infiltrating macrophage subtype correlates with late-stage phenotypic outcome in a mouse model of hepatorenal fibrocystic disease. Laboratory Investigation, 2021, 101, 1382-1393.	1.7	0
90	Altered cell volume regulation in a mouse cellâ€model of autosomal recessive polycystic kidney disease. FASEB Journal, 2006, 20, .	0.2	0

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91	Neuronal Cilia and Obesity. , 2013, , 165-191.		0
92	Cilia and Polycystic Kidney Disease. , 2018, , 87-110.		0
93	Renal hypertrophic signaling triggers activation of kidney immune response and accelerates cystogenesis in polycystic kidney disease. FASEB Journal, 2019, 33, 747.1.	0.2	0
94	ATXN10 Is Required for Embryonic Heart Development and Maintenance of Epithelial Cell Phenotypes in the Adult Kidney and Pancreas. Frontiers in Cell and Developmental Biology, 2021, 9, 705182.	1.8	0