

Steven J Pittler

List of Publications by Year in descending order

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51
papers

3,238
citations

236925

25
h-index

223800

46
g-index

51
all docs

51
docs citations

51
times ranked

2365
citing authors

#	ARTICLE	IF	CITATIONS
1	Retinal Degeneration Caused by Rod-Specific Dhdds Ablation Occurs without Concomitant Inhibition of Protein N-Glycosylation. <i>IScience</i> , 2020, 23, 101198.	4.1	14
2	Lack of Overt Retinal Degeneration in a K42E Dhdds Knock-In Mouse Model of RP59. <i>Cells</i> , 2020, 9, 896.	4.1	9
3	Selective Ablation of Dehydrololichyl Diphosphate Synthase in Murine Retinal Pigment Epithelium (RPE) Causes RPE Atrophy and Retinal Degeneration. <i>Cells</i> , 2020, 9, 771.	4.1	10
4	GARP2 accelerates retinal degeneration in rod cGMP-gated cation channel $\hat{1}^2$ -subunit knockout mice. <i>Scientific Reports</i> , 2017, 7, 42545.	3.3	23
5	Role of RDS and Rhodopsin in Cngb1-Related Retinal Degeneration. , 2016, 57, 787.		10
6	The B3 Subunit of the Cone Cyclic Nucleotide-gated Channel Regulates the Light Responses of Cones and Contributes to the Channel Structural Flexibility. <i>Journal of Biological Chemistry</i> , 2016, 291, 8721-8734.	3.4	2
7	Varying the GARP2-to-RDS Ratio Leads to Defects in Rim Formation and Rod and Cone Function. , 2015, 56, 8187.		9
8	Overexpression of rod photoreceptor glutamic acid rich protein 2 (GARP2) increases gain and slows recovery in mouse retina. <i>Cell Communication and Signaling</i> , 2014, 12, 67.	6.5	13
9	Dynamic near-infrared imaging reveals transient phototropic change in retinal rod photoreceptors. <i>Journal of Biomedical Optics</i> , 2013, 18, 1.	2.6	25
10	Comparative intrinsic optical signal imaging of wild-type and mutant mouse retinas. <i>Optics Express</i> , 2012, 20, 7646.	3.4	18
11	Investigation of the hyper-reflective inner/outer segment band in optical coherence tomography of living frog retina. <i>Journal of Biomedical Optics</i> , 2012, 17, 060504.	2.6	39
12	Three-Dimensional Architecture of the Rod Sensory Cilium and Its Disruption in Retinal Neurodegeneration. <i>Cell</i> , 2012, 151, 1029-1041.	28.9	142
13	Age-related changes in Cngb1-X1 knockout mice: prolonged cone survival. <i>Documenta Ophthalmologica</i> , 2012, 124, 163-175.	2.2	6
14	Activation of Retinal Guanylyl Cyclase RetGC1 by GCAP1: Stoichiometry of Binding and Effect of New LCA-Related Mutations. <i>Biochemistry</i> , 2010, 49, 709-717.	2.5	28
15	Knockout of GARPs and the $\hat{1}^2$ -subunit of the rod cGMP-gated channel disrupts disk morphogenesis and rod outer segment structural integrity. <i>Journal of Cell Science</i> , 2009, 122, 1927-1927.	2.0	1
16	Characterization of a Canine Model of Autosomal Recessive Retinitis Pigmentosa due to a <i>PDE6A</i> Mutation. , 2009, 50, 801.		48
17	Knockout of GARPs and the $\hat{1}^2$ -subunit of the rod cGMP-gated channel disrupts disk morphogenesis and rod outer segment structural integrity. <i>Journal of Cell Science</i> , 2009, 122, 1192-1200.	2.0	84
18	Focus on Molecules: Rod cGMP Phosphodiesterase Type 6. <i>Experimental Eye Research</i> , 2007, 84, 1-2.	2.6	23

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19	Focus on Molecules: Rod photoreceptor cGMP-gated cation channel. <i>Experimental Eye Research</i> , 2007, 85, 173-174.	2.6	3
20	Identifying photoreceptors in blind eyes caused by <i>RPE65</i> mutations: Prerequisite for human gene therapy success. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2005, 102, 6177-6182.	7.1	249
21	Functional Analysis of the Rod Photoreceptor cGMP Phosphodiesterase β -Subunit Gene Promoter. <i>Journal of Biological Chemistry</i> , 2004, 279, 19800-19807.	3.4	54
22	Characterization of 3',5' cyclic nucleotide phosphodiesterase activity in Y79 retinoblastoma cells: absence of functional PDE6. <i>Molecular Vision</i> , 2004, 10, 738-49.	1.1	16
23	Electroretinographic Abnormalities in Parents of Patients With Leber Congenital Amaurosis Who Have Heterozygous GUCY2D Mutations. <i>JAMA Ophthalmology</i> , 2002, 120, 1325.	2.4	35
24	A PDE6A Promoter Fragment Directs Transcription Predominantly in the Photoreceptor. <i>Biochemical and Biophysical Research Communications</i> , 2001, 282, 543-547.	2.1	17
25	Reproducible high efficiency gene transfer into Y79 retinoblastoma cells using adenofection. <i>Journal of Neuroscience Methods</i> , 2001, 106, 1-7.	2.5	12
26	Four novel mutations in the RPE65 gene in patients with Leber congenital amaurosis. <i>Human Mutation</i> , 2001, 18, 164-164.	2.5	52
27	Rapid restoration of visual pigment and function with oral retinoid in a mouse model of childhood blindness. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2000, 97, 8623-8628.	7.1	292
28	Genomic organization of the human rod photoreceptor cGMP-gated cation channel β -subunit gene. <i>Gene</i> , 2000, 245, 311-318.	2.2	47
29	Structure and upstream region characterization of the human gene encoding rod photoreceptor cGMP phosphodiesterase β -subunit. <i>Journal of Molecular Neuroscience</i> , 1998, 10, 235-250.	2.3	10
30	Cloning and Expression of the Glucocorticoid Receptor from the Squirrel Monkey (<i>Saimiri boliviensis</i>) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 1997, 82, 465-472.	3.6	27
31	The β subunit of human rod photoreceptor cGMP-gated cation channel is generated from a complex transcription unit. <i>FEBS Letters</i> , 1996, 389, 213-218.	2.8	30
32	Gene Structure and Chromosome Localization to 7q21.3 of the Human Rod Photoreceptor Transducin β -Subunit Gene (GNGT1). <i>Genomics</i> , 1996, 35, 241-243.	2.9	25
33	Retinal-specific guanylate cyclase gene mutations in Leber's congenital amaurosis. <i>Nature Genetics</i> , 1996, 14, 461-464.	21.4	433
34	Autosomal recessive retinitis pigmentosa caused by mutations in the β subunit of rod cGMP phosphodiesterase. <i>Nature Genetics</i> , 1995, 11, 468-471.	21.4	233
35	cDNA, Gene Structure, and Chromosomal Localization of Human GAR1 (CNCG3L), a Homolog of the Third Subunit of Bovine Photoreceptor cGMP-Gated Channel. <i>Genomics</i> , 1995, 28, 32-38.	2.9	22
36	Molecular Analysis of the Human GAR1 Gene. , 1995, , 331-338.		0

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37	A YAC Contig of Approximately 3 Mb from Human Chromosome 5q31 â€ˆ q33. <i>Genomics</i> , 1994, 19, 470-477.	2.9	28
38	Human Retinal Guanylate Cyclase (GUC2D) Maps to Chromosome 17p13.1. <i>Genomics</i> , 1994, 22, 478-481.	2.9	26
39	In vivo biosynthesis of cholesterol in the rat retina. <i>FEBS Letters</i> , 1993, 335, 234-238.	2.8	57
40	PCR analysis of DNA from 70-year-old sections of rodless retina demonstrates identity with the mouse rd defect.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1993, 90, 9616-9619.	7.1	196
41	Syntenic assignments of visual transduction genes in cattle. <i>Genomics</i> , 1992, 14, 699-706.	2.9	17
42	Chromosome mapping of the rod photoreceptor cGMP phosphodiesterase Î²-subunit gene in mouse and human: Tight linkage to the Huntington disease region (4p16.3). <i>Genomics</i> , 1992, 12, 750-754.	2.9	26
43	Novel morphological changes in rat retina induced by intravitreal injection of lovastatin. <i>Experimental Eye Research</i> , 1992, 54, 149-152.	2.6	10
44	Primary structure of frog rhodopsin. <i>FEBS Letters</i> , 1992, 313, 103-108.	2.8	31
45	Complete cDNA sequences of mouse rod photoreceptor cGMP phosphodiesterase Î±- and Î²-subunits, and identification of Î²â€², a putative Î²-subunit isozyme produced by alternative splicing of the Î²-subunit gene. <i>FEBS Letters</i> , 1991, 278, 107-114.	2.8	50
46	Identification of a nonsense mutation in the rod photoreceptor cGMP phosphodiesterase beta-subunit gene of the rd mouse.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1991, 88, 8322-8326.	7.1	581
47	HSV antigens and HSV DNA in avascular and vascularized lesions of human herpes simplex keratitis. <i>Current Eye Research</i> , 1991, 10, 63-68.	1.5	30
48	Molecular characterization of human and bovine rod photoreceptor cGMP phosphodiesterase Î±-subunit and chromosomal localization of the human gene. <i>Genomics</i> , 1990, 6, 272-283.	2.9	105
49	An interchromosomal gene conversion of the <i>Drosophila dunce</i> locus identified with restriction site polymorphisms: A potential involvement of transposable elements in gene conversion. <i>Molecular Genetics and Genomics</i> , 1987, 208, 315-324.	2.4	5
50	A new family of the poly-deoxyadenylated class of <i>Drosophila</i> transposable elements identified by a representative member at the <i>dunce</i> locus. <i>Molecular Genetics and Genomics</i> , 1987, 208, 325-328.	2.4	9
51	In vitro synthesis of rat brain hexokinase. <i>Biochimica Et Biophysica Acta - General Subjects</i> , 1985, 843, 186-192.	2.4	6