

Neal S Peachey

List of Publications by Year in descending order

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180
papers

10,575
citations

36303

51
h-index

40979

93
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187
all docs

187
docs citations

187
times ranked

10962
citing authors

#	ARTICLE	IF	CITATIONS
1	A large genome-wide association study of age-related macular degeneration highlights contributions of rare and common variants. <i>Nature Genetics</i> , 2016, 48, 134-143.	21.4	1,167
2	Seven new loci associated with age-related macular degeneration. <i>Nature Genetics</i> , 2013, 45, 433-439.	21.4	687
3	Genetic variants near <i>TIMP3</i> and high-density lipoprotein-associated loci influence susceptibility to age-related macular degeneration. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2010, 107, 7401-7406.	7.1	475
4	Norrin, Frizzled-4, and Lrp5 Signaling in Endothelial Cells Controls a Genetic Program for Retinal Vascularization. <i>Cell</i> , 2009, 139, 285-298.	28.9	377
5	Mechanisms of Rhodopsin Inactivation in Vivo as Revealed by a COOH-Terminal Truncation Mutant. <i>Science</i> , 1995, 267, 374-377.	12.6	280
6	The <i>nob2</i> mouse, a null mutation in <i>Cacna1f</i> : Anatomical and functional abnormalities in the outer retina and their consequences on ganglion cell visual responses. <i>Visual Neuroscience</i> , 2006, 23, 11-24.	1.0	194
7	A Transient Receptor Potential-Like Channel Mediates Synaptic Transmission in Rod Bipolar Cells. <i>Journal of Neuroscience</i> , 2009, 29, 6088-6093.	3.6	192
8	The Light Peak of the Electroretinogram Is Dependent on Voltage-gated Calcium Channels and Antagonized by Bestrophin (Best-1). <i>Journal of General Physiology</i> , 2006, 127, 577-589.	1.9	186
9	Implantation of silicon chip microphotodiode arrays into the cat subretinal space. <i>IEEE Transactions on Neural Systems and Rehabilitation Engineering</i> , 2001, 9, 86-95.	4.9	183
10	Expression of bestrophin-1, the product of the VMD2 gene, modulates voltage-dependent Ca ²⁺ channels in retinal pigment epithelial cells. <i>FASEB Journal</i> , 2006, 20, 178-180.	0.5	179
11	Neuronal Pentraxins Mediate Synaptic Refinement in the Developing Visual System. <i>Journal of Neuroscience</i> , 2006, 26, 6269-6281.	3.6	156
12	Role of the beta(2) subunit of voltage-dependent calcium channels in the retinal outer plexiform layer. <i>Investigative Ophthalmology and Visual Science</i> , 2002, 43, 1595-603.	3.3	146
13	Proteomics Reveal Cochlin Deposits Associated with Glaucomatous Trabecular Meshwork. <i>Journal of Biological Chemistry</i> , 2005, 280, 6080-6084.	3.4	140
14	Class 5 Transmembrane Semaphorins Control Selective Mammalian Retinal Lamination and Function. <i>Neuron</i> , 2011, 71, 460-473.	8.1	137
15	GPR179 Is Required for Depolarizing Bipolar Cell Function and Is Mutated in Autosomal-Recessive Complete Congenital Stationary Night Blindness. <i>American Journal of Human Genetics</i> , 2012, 90, 331-339.	6.2	131
16	Functional roles of bestrophins in ocular epithelia. <i>Progress in Retinal and Eye Research</i> , 2009, 28, 206-226.	15.5	125
17	Mutations in the RPCR gene cause X-linked cone dystrophy. <i>Human Molecular Genetics</i> , 2002, 11, 605-611.	2.9	115
18	Formation and progression of sub-retinal pigment epithelium deposits in <i>Efemp1</i> mutation knock-in mice: a model for the early pathogenic course of macular degeneration. <i>Human Molecular Genetics</i> , 2007, 16, 2423-2432.	2.9	113

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19	The Effect of Peripherin/rds Haploinsufficiency on Rod and Cone Photoreceptors. Journal of Neuroscience, 1997, 17, 8118-8128.	3.6	111
20	Elimination of the $\alpha 1$ Subunit Abolishes GABA _C Receptor Expression and Alters Visual Processing in the Mouse Retina. Journal of Neuroscience, 2002, 22, 4163-4174.	3.6	109
21	Properties of the mouse cone-mediated electroretinogram during light adaptation. Neuroscience Letters, 1993, 162, 9-11.	2.1	105
22	Early Onset Photoreceptor Abnormalities Induced by Targeted Disruption of the Interphotoreceptor Retinoid-Binding Protein Gene. Journal of Neuroscience, 1998, 18, 4511-4520.	3.6	105
23	Identification of the Gene and the Mutation Responsible for the MousenobPhenotype. , 2003, 44, 378.		105
24	Electrophysiological analysis of visual function in mutant mice. Documenta Ophthalmologica, 2003, 107, 13-35.	2.2	104
25	Subretinal Semiconductor Microphotodiode Array. Ophthalmic Surgery Lasers and Imaging Retina, 1998, 29, 234-241.	0.7	99
26	Suppression of Ca ²⁺ signaling in a mouse model of Best disease. Human Molecular Genetics, 2010, 19, 1108-1118.	2.9	92
27	Age-Related Changes in the Mouse Outer Retina. Optometry and Vision Science, 2001, 78, 425-430.	1.2	90
28	Nyctalopin Expression in Retinal Bipolar Cells Restores Visual Function in a Mouse Model of Complete X-Linked Congenital Stationary Night Blindness. Journal of Neurophysiology, 2007, 98, 3023-3033.	1.8	90
29	The rhodopsin cycle is preserved in IRBP α knockout mice despite abnormalities in retinal structure and function. Visual Neuroscience, 2000, 17, 97-105.	1.0	88
30	Assessing Susceptibility to Age-related Macular Degeneration with Proteomic and Genomic Biomarkers. Molecular and Cellular Proteomics, 2009, 8, 1338-1349.	3.8	88
31	Mutations in Lama1 Disrupt Retinal Vascular Development and Inner Limiting Membrane Formation. Journal of Biological Chemistry, 2010, 285, 7697-7711.	3.4	85
32	The luminance-response function of the dark-adapted human electroretinogram. Vision Research, 1989, 29, 263-270.	1.4	83
33	Expression and Localization of Bestrophin during Normal Mouse Development. , 2003, 44, 3622.		82
34	Immunohistochemical Studies of the Retina Following Long-term Implantation with Subretinal Microphotodiode Arrays. Experimental Eye Research, 2001, 73, 333-343.	2.6	75
35	Ocular abnormalities in Largemyd and Largevls mice, spontaneous models for muscle, eye, and brain diseases. Molecular and Cellular Neurosciences, 2005, 30, 160-172.	2.2	74
36	Autoantibodies in Melanoma-Associated Retinopathy Target TRPM1 Cation Channels of Retinal ON Bipolar Cells. Journal of Neuroscience, 2011, 31, 3962-3967.	3.6	73

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37	Modulating GLUT1 expression in retinal pigment epithelium decreases glucose levels in the retina: impact on photoreceptors and Müller glial cells. <i>American Journal of Physiology - Cell Physiology</i> , 2019, 316, C121-C133.	4.6	73
38	Early retinal pigment epithelium dysfunction is concomitant with hyperglycemia in mouse models of type 1 and type 2 diabetes. <i>Journal of Neurophysiology</i> , 2015, 113, 1085-1099.	1.8	72
39	Ciliary Neurotrophic Factor for Macular Telangiectasia Type 2: Results From a Phase 1 Safety Trial. <i>American Journal of Ophthalmology</i> , 2015, 159, 659-666.e1.	3.3	72
40	Insulin Receptor Substrate 2 Is Essential for Maturation and Survival of Photoreceptor Cells. <i>Journal of Neuroscience</i> , 2005, 25, 1240-1248.	3.6	69
41	Temporal Properties of the Mouse Cone Electroretinogram. <i>Journal of Neurophysiology</i> , 2002, 87, 42-48.	1.8	67
42	Morphological and Physiological Consequences of the Selective Elimination of Rod Photoreceptors in Transgenic Mice. <i>Experimental Eye Research</i> , 1996, 63, 35-49.	2.6	66
43	Light-Evoked Responses of the Mouse Retinal Pigment Epithelium. <i>Journal of Neurophysiology</i> , 2004, 91, 1134-1142.	1.8	64
44	Guidance-Cue Control of Horizontal Cell Morphology, Lamination, and Synapse Formation in the Mammalian Outer Retina. <i>Journal of Neuroscience</i> , 2012, 32, 6859-6868.	3.6	62
45	A Novel Role of Complement in Retinal Degeneration. , 2012, 53, 7684.		61
46	Intravitreal delivery of a novel AAV vector targets ON bipolar cells and restores visual function in a mouse model of complete congenital stationary night blindness. <i>Human Molecular Genetics</i> , 2015, 24, 6229-6239.	2.9	60
47	The Subretinal Microphotodiode Array Retinal Prosthesis. <i>Ophthalmic Research</i> , 1998, 30, 195-196.	1.9	59
48	Retinal Degeneration in a Rodent Model of Smith-Lemli-Opitz Syndrome. <i>JAMA Ophthalmology</i> , 2004, 122, 1190.	2.4	59
49	Voltage-Dependent Calcium Channel CaV1.3 Subunits Regulate the Light Peak of the Electroretinogram. <i>Journal of Neurophysiology</i> , 2007, 97, 3731-3735.	1.8	59
50	Rod and Cone Dysfunction in Carriers of X-linked Retinitis Pigmentosa. <i>Ophthalmology</i> , 1988, 95, 677-685.	5.2	58
51	Properties of the human cone system electroretinogram during light adaptation. <i>Applied Optics</i> , 1989, 28, 1145.	2.1	58
52	Rod phototransduction in transgenic mice expressing a mutant opsin gene. <i>Journal of the Optical Society of America A: Optics and Image Science, and Vision</i> , 1996, 13, 577.	1.5	58
53	GPR179 Is Required for High Sensitivity of the mGluR6 Signaling Cascade in Depolarizing Bipolar Cells. <i>Journal of Neuroscience</i> , 2014, 34, 6334-6343.	3.6	58
54	Pharmacological studies of the mouse cone electroretinogram. <i>Visual Neuroscience</i> , 2005, 22, 631-636.	1.0	56

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55	A Model of Best Vitelliform Macular Dystrophy in Rats. , 2004, 45, 3733.		55
56	Genome-wide association identifies SKIV2L and MYRIP as protective factors for age-related macular degeneration. Genes and Immunity, 2010, 11, 609-621.	4.1	55
57	Abnormal vascularization in mouse retina with dysregulated retinal cholesterol homeostasis. Journal of Clinical Investigation, 2012, 122, 3012-3023.	8.2	54
58	Interaction between the Photoreceptor-Specific Tubby-like Protein 1 and the Neuronal-Specific GTPase Dynamin-1. , 2007, 48, 2837.		52
59	Rare and common variants in extracellular matrix gene Fibrillin 2 (FBN2) are associated with macular degeneration. Human Molecular Genetics, 2014, 23, 5827-5837.	2.9	52
60	Rod and cone system contributions to oscillatory potentials: An explanation for the conditioning flash effect. Vision Research, 1987, 27, 859-866.	1.4	51
61	Immunohistochemical analysis of the outer plexiform layer in the nob mouse shows no abnormalities. Visual Neuroscience, 2003, 20, 267-272.	1.0	51
62	Mutations in CTNNA1 cause butterfly-shaped pigment dystrophy and perturbed retinal pigment epithelium integrity. Nature Genetics, 2016, 48, 144-151.	21.4	50
63	Early Synaptic Defects in <i>tulp1</i> Mice. , 2009, 50, 3074.		48
64	Age-related changes in visual function in cystathionine-beta-synthase mutant mice, a model of hyperhomocysteinemia. Experimental Eye Research, 2012, 96, 124-131.	2.6	47
65	Degeneration of Cone Photoreceptors Induced by Expression of the Mas1 Protooncogene. Experimental Neurology, 2000, 163, 207-219.	4.1	46
66	A 32 kb Critical Region Excluding Y402H in CFH Mediates Risk for Age-Related Macular Degeneration. PLoS ONE, 2011, 6, e25598.	2.5	46
67	Light adaptation, rods, and the human cone flicker ERG. Visual Neuroscience, 1992, 8, 145-150.	1.0	44
68	Polygenic Disease and Retinitis Pigmentosa: Albinism Exacerbates Photoreceptor Degeneration Induced by the Expression of a Mutant Opsin in Transgenic Mice. Journal of Neuroscience, 1996, 16, 7853-7858.	3.6	43
69	Mouse b-wave mutants. Documenta Ophthalmologica, 2014, 128, 77-89.	2.2	43
70	Pharmacological analysis of the rat cone electroretinogram. Visual Neuroscience, 2003, 20, 297-306.	1.0	42
71	Membrane frizzled-related protein is necessary for the normal development and maintenance of photoreceptor outer segments. Visual Neuroscience, 2008, 25, 563-574.	1.0	42
72	Depolarizing bipolar cell dysfunction due to a Trpm1 point mutation. Journal of Neurophysiology, 2012, 108, 2442-2451.	1.8	42

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73	Disruption of murine <i>Adamtsl4</i> results in zonular fiber detachment from the lens and in retinal pigment epithelium dedifferentiation. <i>Human Molecular Genetics</i> , 2015, 24, ddv399.	2.9	41
74	Retinal and Nonocular Abnormalities in <i>Cyp27a1Cyp46a1</i> Mice with Dysfunctional Metabolism of Cholesterol. <i>American Journal of Pathology</i> , 2014, 184, 2403-2419.	3.8	40
75	Comparison of three methods of estimating the parameters of the Naka-Rushton equation. <i>Documenta Ophthalmologica</i> , 1993, 84, 19-30.	2.2	39
76	Genetic loss of function of <i>Ptbp1</i> does not induce glia-to-neuron conversion in retina. <i>Cell Reports</i> , 2022, 39, 110849.	6.4	39
77	Functional consequences of oncogene-induced horizontal cell degeneration in the retinas of transgenic mice. <i>Visual Neuroscience</i> , 1997, 14, 627-632.	1.0	38
78	Cochlin deposits in the trabecular meshwork of the glaucomatous DBA/2J mouse. <i>Experimental Eye Research</i> , 2005, 80, 741-744.	2.6	38
79	Light-induced exacerbation of retinal degeneration in a rat model of Smith-Lemli-Opitz syndrome. <i>Experimental Eye Research</i> , 2006, 82, 496-504.	2.6	38
80	Ankyrin-B is required for coordinated expression of beta-2-spectrin, the Na/K-ATPase and the Na/Ca exchanger in the inner segment of rod photoreceptors. <i>Experimental Eye Research</i> , 2009, 88, 57-64.	2.6	37
81	Mouse models of human ocular disease for translational research. <i>PLoS ONE</i> , 2017, 12, e0183837.	2.5	37
82	Mutations of the Opsin Gene (Y102H and I307N) Lead to Light-induced Degeneration of Photoreceptors and Constitutive Activation of Phototransduction in Mice. <i>Journal of Biological Chemistry</i> , 2010, 285, 14521-14533.	3.4	36
83	Light-Evoked Responses of the Retinal Pigment Epithelium: Changes Accompanying Photoreceptor Loss in the Mouse. <i>Journal of Neurophysiology</i> , 2010, 104, 391-402.	1.8	36
84	The Subretinal Microphotodiode Array Retinal Prosthesis II. <i>Ophthalmic Research</i> , 1999, 31, 246-246.	1.9	35
85	Subretinal implantation of semiconductor-based photodiodes: durability of novel implant designs. <i>Journal of Rehabilitation Research and Development</i> , 2002, 39, 313-21.	1.6	35
86	Microtubule-Associated Protein 1 Light Chain 3B, (LC3B) Is Necessary to Maintain Lipid-Mediated Homeostasis in the Retinal Pigment Epithelium. <i>Frontiers in Cellular Neuroscience</i> , 2018, 12, 351.	3.7	34
87	Contribution of Kir4.1 to the mouse electroretinogram. <i>Molecular Vision</i> , 2004, 10, 650-4.	1.1	34
88	Retinal abnormalities associated with the G90D mutation in opsin. <i>Journal of Comparative Neurology</i> , 2004, 478, 149-163.	1.6	33
89	Noninvasive Electroretinographic Procedures for the Study of the Mouse Retina. <i>Current Protocols in Mouse Biology</i> , 2018, 8, 1-16.	1.2	33
90	Effects of light adaptation on the response characteristics of human oscillatory potentials. <i>Electroencephalography and Clinical Neurophysiology</i> , 1991, 78, 27-34.	0.3	32

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91	Distinct and Atypical Intrinsic and Extrinsic Cell Death Pathways between Photoreceptor Cell Types upon Specific Ablation of Ranbp2 in Cone Photoreceptors. <i>PLoS Genetics</i> , 2013, 9, e1003555.	3.5	32
92	Exclusion of aldose reductase as a mediator of ERG deficits in a mouse model of diabetic eye disease. <i>Visual Neuroscience</i> , 2012, 29, 267-274.	1.0	31
93	Proteomic and Genomic Biomarkers for Age-Related Macular Degeneration. <i>Advances in Experimental Medicine and Biology</i> , 2010, 664, 411-417.	1.6	30
94	Electroretinographic Findings in Human Oculocutaneous Albinism. <i>Ophthalmology</i> , 1989, 96, 1778-1785.	5.2	29
95	Short-term changes in the response characteristics of the human visual evoked potential. <i>Vision Research</i> , 1994, 34, 2823-2831.	1.4	29
96	Partial Rescue of Retinal Function and Sterol Steady-State in a Rat Model of Smith-Lemli-Opitz Syndrome. <i>Pediatric Research</i> , 2007, 61, 273-278.	2.3	29
97	Photoreceptor Degeneration, Azoospermia, Leukoencephalopathy, and Abnormal RPE Cell Function in Mice Expressing an Early Stop Mutation in <i>CLCN2</i> . , 2010, 51, 3264.		29
98	Complement anaphylatoxin receptors C3aR and C5aR are required in the pathogenesis of experimental autoimmune uveitis. <i>Journal of Leukocyte Biology</i> , 2016, 99, 447-454.	3.3	29
99	Association of Smoking, Alcohol Consumption, Blood Pressure, Body Mass Index, and Glycemic Risk Factors With Age-Related Macular Degeneration. <i>JAMA Ophthalmology</i> , 2021, 139, 1299.	2.5	29
100	Retinoid isomerase inhibitors impair but do not block mammalian cone photoreceptor function. <i>Journal of General Physiology</i> , 2018, 150, 571-590.	1.9	28
101	An Intramembrane Glutamic Acid Governs Peripherin/rds Function for Photoreceptor Disk Morphogenesis. , 2007, 48, 2975.		27
102	Glycosylation of rhodopsin is necessary for its stability and incorporation into photoreceptor outer segment discs. <i>Human Molecular Genetics</i> , 2015, 24, 2709-2723.	2.9	27
103	Functional consequences of oncogene-induced photoreceptor degeneration in transgenic mice. <i>Visual Neuroscience</i> , 1995, 12, 513-522.	1.0	26
104	Retinal Degeneration in the nervous Mutant Mouse. III. Electrophysiological Studies of the Visual Pathway. <i>Experimental Eye Research</i> , 2000, 70, 467-473.	2.6	25
105	Role of monocarboxylate transporters in regulating metabolic homeostasis in the outer retina: Insight gained from cell-specific <i>Bsg</i> deletion. <i>FASEB Journal</i> , 2020, 34, 5401-5419.	0.5	25
106	Status of the feline retina 5 years after subretinal implantation. <i>Journal of Rehabilitation Research and Development</i> , 2006, 43, 723.	1.6	25
107	Component-specific effects of physostigmine on the cat visual evoked potential. <i>Experimental Brain Research</i> , 1993, 95, 271-6.	1.5	24
108	Selective Impairment of a Subset of Ran-GTP-binding Domains of Ran-binding Protein 2 (Ranbp2) Suffices to Recapitulate the Degeneration of the Retinal Pigment Epithelium (RPE) Triggered by Ranbp2 Ablation. <i>Journal of Biological Chemistry</i> , 2014, 289, 29767-29789.	3.4	24

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109	Protective Effect of Paraoxonase 1 Gene Variant Gln192Arg in Age-Related Macular Degeneration. American Journal of Ophthalmology, 2010, 149, 513-522.	3.3	23
110	Prevention of Retinal Degeneration in a Rat Model of Smith-Lemli-Opitz Syndrome. Scientific Reports, 2018, 8, 1286.	3.3	23
111	Functional abnormalities in the retinal pigment epithelium of CFTR mutant mice. Experimental Eye Research, 2006, 83, 424-428.	2.6	22
112	Deficiency of CC chemokine ligand 2 and decay-accelerating factor causes retinal degeneration in mice. Experimental Eye Research, 2015, 138, 126-133.	2.6	22
113	Deletion of GLUT1 in mouse lens epithelium leads to cataract formation. Experimental Eye Research, 2018, 172, 45-53.	2.6	22
114	Conditional deletion of <i>Des1</i> in the mouse retina does not impair the visual cycle in cones. FASEB Journal, 2019, 33, 5782-5792.	0.5	22
115	Light adaptation and the luminance-response function of the cone electroretinogram. Documenta Ophthalmologica, 1992, 79, 363-369.	2.2	21
116	Cochlin and glaucoma: A mini-review. Visual Neuroscience, 2005, 22, 605-613.	1.0	21
117	The circadian clock gene <i>Bmal1</i> is required to control the timing of retinal neurogenesis and lamination of Müller glia in the mouse retina. FASEB Journal, 2019, 33, 8745-8758.	0.5	21
118	Rapid and slow changes in the human cone electroretinogram during light and dark adaptation. Vision Research, 1992, 32, 2049-2053.	1.4	19
119	Noninvasive recording and response characteristics of the rat dc-electroretinogram. Visual Neuroscience, 2002, 19, 693-701.	1.0	19
120	Rational Tuning of Visual Cycle Modulator Pharmacodynamics. Journal of Pharmacology and Experimental Therapeutics, 2017, 362, 131-145.	2.5	19
121	A 221-bp fragment of the mouse opsin promoter directs expression specifically to the rod photoreceptors of transgenic mice. Visual Neuroscience, 1997, 14, 617-625.	1.0	18
122	Microglial activation in an amyotrophic lateral sclerosis-like model caused by Ranbp2 loss and nucleocytoplasmic transport impairment in retinal ganglion neurons. Cellular and Molecular Life Sciences, 2019, 76, 3407-3432.	5.4	18
123	Retinal structure and function in an animal model that replicates the biochemical hallmarks of desmosterolosis. Neurochemical Research, 2000, 25, 685-694.	3.3	17
124	Transgenic Bcl-2 Expressed in Photoreceptor Cells Confers Both Death-sparing and Death-inducing Effects. Experimental Eye Research, 2001, 73, 711-721.	2.6	17
125	Electrophysiological responses of the mouse retina to 12C ions. Neuroscience Letters, 2007, 416, 231-235.	2.1	17
126	Light-Regulated Thyroid Hormone Signaling Is Required for Rod Photoreceptor Development in the Mouse Retina. , 2015, 56, 8248.		17

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127	Mouse models of X-linked juvenile retinoschisis have an early onset phenotype, the severity of which varies with genotype. <i>Human Molecular Genetics</i> , 2019, 28, 3072-3090.	2.9	17
128	Visual evoked potentials to infrared stimulation in normal cats and rats. <i>Documenta Ophthalmologica</i> , 2001, 103, 155-162.	2.2	16
129	Visual abnormalities associated with enhanced optic nerve myelination. <i>Brain Research</i> , 2011, 1374, 36-42.	2.2	16
130	Spatial properties of rod-cone interactions in flicker and hue detection. <i>Vision Research</i> , 1990, 30, 1205-1210.	1.4	15
131	Mapping a new genetic locus for X linked retinitis pigmentosa to Xq28. <i>Journal of Medical Genetics</i> , 2006, 43, e27-e27.	3.2	15
132	Differential Loss of Prolyl Isomerase or Chaperone Activity of Ran-binding Protein 2 (Ranbp2) Unveils Distinct Physiological Roles of Its Cyclophilin Domain in Proteostasis. <i>Journal of Biological Chemistry</i> , 2014, 289, 4600-4625.	3.4	15
133	Inducing a Visceral Organ to Protect a Peripheral Capillary Bed. <i>American Journal of Pathology</i> , 2014, 184, 1890-1899.	3.8	14
134	Identification of a new mutant allele, <i>Grm6^{nob7}</i> , for complete congenital stationary night blindness. <i>Visual Neuroscience</i> , 2015, 32, E004.	1.0	14
135	Visual cortical plasticity following unilateral sensorimotor cortical lesions in the neonatal rat. <i>Experimental Neurology</i> , 2006, 199, 122-129.	4.1	13
136	Attenuation of oscillatory potentials in nob2 mice. <i>Documenta Ophthalmologica</i> , 2007, 115, 173-186.	2.2	13
137	A missense mutation in <i>Grm6</i> reduces but does not eliminate mGluR6 expression or rod depolarizing bipolar cell function. <i>Journal of Neurophysiology</i> , 2017, 118, 845-854.	1.8	13
138	Correlation of Electroretinographic Findings and Peripheral Retinal Nonperfusion in Patients With Sickle Cell Retinopathy. <i>JAMA Ophthalmology</i> , 1990, 108, 1106.	2.4	12
139	Retinal Degeneration in the nervous Mutant Mouse. IV. Inner Retinal Changes. <i>Experimental Eye Research</i> , 2001, 72, 243-252.	2.6	12
140	Myosin 6 Is Required for Iris Development and Normal Function of the Outer Retina. , 2013, 54, 7223.		12
141	Interdependence Among Members of the mGluR6 G-protein Mediated Signalplex of Retinal Depolarizing Bipolar Cells. , 2014, , 67-79.		12
142	A Chemical Mutagenesis Screen Identifies Mouse Models with ERG Defects. <i>Advances in Experimental Medicine and Biology</i> , 2016, 854, 177-183.	1.6	12
143	Interocular transfer and dark adaptation to long-wave test lights. <i>Vision Research</i> , 1984, 24, 1043-1048.	1.4	11
144	Effects of heavy ions on visual function and electrophysiology of rodents: the ALTEA-MICE project. <i>Advances in Space Research</i> , 2004, 33, 1347-1351.	2.6	11

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145	Presence of the Gpr179(nob5) allele in a C3H-derived transgenic mouse. <i>Molecular Vision</i> , 2013, 19, 2615-25.	1.1	11
146	Probing inner retinal circuits in the rod pathway: A comparison of c-fos activation in mutant mice. <i>Visual Neuroscience</i> , 2004, 21, 873-881.	1.0	10
147	An inducible Cre mouse for studying roles of the RPE in retinal physiology and disease. <i>JCI Insight</i> , 2021, 6, .	5.0	10
148	Membrane frizzled-related protein is necessary for the normal development and maintenance of photoreceptor outer segments. <i>Visual Neuroscience</i> , 2008, 25, 713-716.	1.0	9
149	Evidence of retinal degeneration in Wolfram syndrome. <i>Ophthalmic Genetics</i> , 2019, 40, 34-38.	1.2	9
150	Transient Receptor Potential Melastatin 1: A Hair Cell Transduction Channel Candidate. <i>PLoS ONE</i> , 2013, 8, e77213.	2.5	9
151	Ultrastructural Localization of GPR179 and the Impact of Mutant Forms on Retinal Function in CSNB1 Patients and a Mouse Model. , 2013, 54, 6973.		8
152	nob: A Mouse Model of CSNB1. , 2001, , 319-328.		8
153	Rod influence on thresholds using different detection criteria during dark adaptation. <i>Acta Psychologica</i> , 1987, 64, 261-270.	1.5	7
154	Loss of bipolar cells resulting from the expression of bcl-2 directed by the IRBP promoter. <i>Experimental Eye Research</i> , 2003, 77, 477-483.	2.6	7
155	Mutation Screen of the Cone-Specific Gene, CLUL1, in 376 Patients with Age-Related Macular Degeneration. <i>Ophthalmic Genetics</i> , 2006, 27, 151-155.	1.2	7
156	A mutagenesis-derived mouse mutant with abnormal retinal vasculature and low bone mineral density. <i>Molecular Vision</i> , 2017, 23, 140-148.	1.1	7
157	Interocular sensitization to a rod-detected test. <i>Vision Research</i> , 1986, 26, 1119-1127.	1.4	6
158	Response properties of slow PIII in the Large vls mutant. <i>Documenta Ophthalmologica</i> , 2012, 125, 203-209.	2.2	6
159	Glucose uptake by <scp>GLUT1</scp> in photoreceptors is essential for outer segment renewal and rod photoreceptor survival. <i>FASEB Journal</i> , 2022, 36, .	0.5	6
160	In vivo gene transfer as a means to study the physiology and morphogenesis of the retinal pigment epithelium in the rat. <i>Methods</i> , 2003, 30, 277-285.	3.8	5
161	Impact of MCT1 Haploinsufficiency on the Mouse Retina. <i>Advances in Experimental Medicine and Biology</i> , 2018, 1074, 375-380.	1.6	5
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