

Jan Wijnholds

List of Publications by Year in descending order

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98
papers

9,213
citations

57719

44
h-index

48277

88
g-index

100
all docs

100
docs citations

100
times ranked

8282
citing authors

#	ARTICLE	IF	CITATIONS
1	CRB1-Associated Retinal Dystrophies: A Prospective Natural History Study in Anticipation of Future Clinical Trials. <i>American Journal of Ophthalmology</i> , 2022, 234, 37-48.	1.7	17
2	Sleep Deprivation Does not Change the Flash Electroretinogram in Wild-type and <i>Opn4^{+/+}Gnat1^{+/+}</i> Mice. <i>Journal of Biological Rhythms</i> , 2022, 37, 216-221.	1.4	2
3	Defining inclusion criteria and endpoints for clinical trials: a prospective cross-sectional study in <i>CRB1</i> -associated retinal dystrophies. <i>Acta Ophthalmologica</i> , 2021, 99, e402-e414.	0.6	10
4	Crumbs2 Is an Essential Slit Diaphragm Protein of the Renal Filtration Barrier. <i>Journal of the American Society of Nephrology: JASN</i> , 2021, 32, 1053-1070.	3.0	17
5	AAV-CRB2 protects against vision loss in an inducible CRB1 retinitis pigmentosa mouse model. <i>Molecular Therapy - Methods and Clinical Development</i> , 2021, 20, 423-441.	1.8	14
6	Defining Phenotype, Tropism, and Retinal Gene Therapy Using Adeno-Associated Viral Vectors (AAVs) in New-Born Brown Norway Rats with a Spontaneous Mutation in <i>Crb1</i> . <i>International Journal of Molecular Sciences</i> , 2021, 22, 3563.	1.8	9
7	CLINICAL CHARACTERISTICS AND NATURAL HISTORY OF RHO-ASSOCIATED RETINITIS PIGMENTOSA. <i>Retina</i> , 2021, 41, 213-223.	1.0	18
8	Research Models and Gene Augmentation Therapy for CRB1 Retinal Dystrophies. <i>Frontiers in Neuroscience</i> , 2020, 14, 860.	1.4	16
9	Novel Therapeutic Approaches for the Treatment of Retinal Degenerative Diseases: Focus on CRISPR/Cas-Based Gene Editing. <i>Frontiers in Neuroscience</i> , 2020, 14, 838.	1.4	12
10	Recombinant Adeno-Associated Viral Vectors (rAAV)-Vector Elements in Ocular Gene Therapy Clinical Trials and Transgene Expression and Bioactivity Assays. <i>International Journal of Molecular Sciences</i> , 2020, 21, 4197.	1.8	54
11	Crumbs2 mediates ventricular layer remodelling to form the spinal cord central canal. <i>PLoS Biology</i> , 2020, 18, e3000470.	2.6	12
12	RPGR-Associated Dystrophies: Clinical, Genetic, and Histopathological Features. <i>International Journal of Molecular Sciences</i> , 2020, 21, 835.	1.8	23
13	Crumbs2 mediates ventricular layer remodelling to form the spinal cord central canal. , 2020, 18, e3000470.		0
14	Crumbs2 mediates ventricular layer remodelling to form the spinal cord central canal. , 2020, 18, e3000470.		0
15	Crumbs2 mediates ventricular layer remodelling to form the spinal cord central canal. , 2020, 18, e3000470.		0
16	Crumbs2 mediates ventricular layer remodelling to form the spinal cord central canal. , 2020, 18, e3000470.		0
17	Crumbs2 mediates ventricular layer remodelling to form the spinal cord central canal. , 2020, 18, e3000470.		0
18	Crumbs2 mediates ventricular layer remodelling to form the spinal cord central canal. , 2020, 18, e3000470.		0

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19	CRB2 Loss in Rod Photoreceptors Is Associated with Progressive Loss of Retinal Contrast Sensitivity. <i>International Journal of Molecular Sciences</i> , 2019, 20, 4069.	1.8	16
20	Human iPSC-Derived Retinas Recapitulate the Fetal CRB1 CRB2 Complex Formation and Demonstrate that Photoreceptors and Müller Glia Are Targets of AAV5. <i>Stem Cell Reports</i> , 2019, 12, 906-919.	2.3	75
21	Retinogenesis of the Human Fetal Retina: An Apical Polarity Perspective. <i>Genes</i> , 2019, 10, 987.	1.0	24
22	Basal Cell Migration in Regeneration of the Corneal Wound-Bed. <i>Stem Cell Reports</i> , 2019, 12, 3-5.	2.3	11
23	Loss of CRB2 in Müller glial cells modifies a CRB1-associated retinitis pigmentosa phenotype into a Leber congenital amaurosis phenotype. <i>Human Molecular Genetics</i> , 2019, 28, 105-123.	1.4	29
24	CLINICAL AND GENETIC CHARACTERISTICS OF MALE PATIENTS WITH RPGR-ASSOCIATED RETINAL DYSTROPHIES. <i>Retina</i> , 2019, 39, 1186-1199.	1.0	56
25	Microglial Cell Dysfunction in CRB1-Associated Retinopathies. <i>Advances in Experimental Medicine and Biology</i> , 2019, 1185, 159-163.	0.8	6
26	AAV Gene Augmentation Therapy for CRB1-Associated Retinitis Pigmentosa. <i>Methods in Molecular Biology</i> , 2018, 1715, 135-151.	0.4	15
27	Production of iPSC-Derived Human Retinal Organoids for Use in Transgene Expression Assays. <i>Methods in Molecular Biology</i> , 2018, 1715, 261-273.	0.4	17
28	AAV Serotype Testing on Cultured Human Donor Retinal Explants. <i>Methods in Molecular Biology</i> , 2018, 1715, 275-288.	0.4	9
29	The Spectrum of Structural and Functional Abnormalities in Female Carriers of Pathogenic Variants in the <i>RPGR</i> Gene. , 2018, 59, 4123.		41
30	CRB2 in immature photoreceptors determines the superior-inferior symmetry of the developing retina to maintain retinal structure and function. <i>Human Molecular Genetics</i> , 2018, 27, 3137-3153.	1.4	26
31	NTPDase2 as a Surface Marker to Isolate Flow Cytometrically a Müller Glial Cell Enriched Population from Dissociated Neural Retinae. <i>Journal of Neuroscience and Neurosurgery</i> , 2018, 1, .	0.1	2
32	Transplantation of NTPDase2-positive Sorted Müller Glial Cells into the Mouse Retina. <i>Journal of Neuroscience and Neurosurgery</i> , 2018, 1, .	0.1	1
33	Genotypic and Phenotypic Characteristics of CRB1 -Associated Retinal Dystrophies. <i>Ophthalmology</i> , 2017, 124, 884-895.	2.5	75
34	Genetic and Molecular Approaches to Study Neuronal Migration in the Developing Cerebral Cortex. <i>Brain Sciences</i> , 2017, 7, 53.	1.1	0
35	The CRB1 Complex: Following the Trail of Crumbs to a Feasible Gene Therapy Strategy. <i>Frontiers in Neuroscience</i> , 2017, 11, 175.	1.4	43
36	Crumbs2 promotes cell ingression during the epithelial-to-mesenchymal transition at gastrulation. <i>Nature Cell Biology</i> , 2016, 18, 1281-1291.	4.6	73

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37	Crumbs 2 prevents cortical abnormalities in mouse dorsal telencephalon. <i>Neuroscience Research</i> , 2016, 108, 12-23.	1.0	25
38	Protein O-Glucosyltransferase 1 (POGLUT1) Promotes Mouse Gastrulation through Modification of the Apical Polarity Protein CRUMBS2. <i>PLoS Genetics</i> , 2015, 11, e1005551.	1.5	34
39	Gene therapy into photoreceptors and Müller glial cells restores retinal structure and function in CRB1 retinitis pigmentosa mouse models. <i>Human Molecular Genetics</i> , 2015, 24, 3104-3118.	1.4	65
40	A New CRB1 Rat Mutation Links Müller Glial Cells to Retinal Telangiectasia. <i>Journal of Neuroscience</i> , 2015, 35, 6093-6106.	1.7	54
41	Targeted ablation of Crb2 in photoreceptor cells induces retinitis pigmentosa. <i>Human Molecular Genetics</i> , 2014, 23, 3384-3401.	1.4	41
42	CRB2 acts as a modifying factor of CRB1-related retinal dystrophies in mice. <i>Human Molecular Genetics</i> , 2014, 23, 3759-3771.	1.4	44
43	The CRB1 and adherens junction complex proteins in retinal development and maintenance. <i>Progress in Retinal and Eye Research</i> , 2014, 40, 35-52.	7.3	75
44	Specific tools for targeting and expression in Müller glial cells. <i>Molecular Therapy - Methods and Clinical Development</i> , 2014, 1, 14009.	1.8	46
45	The multi-PDZ domain protein-1 (MUPP-1) expression regulates cellular levels of the PALS-1/PAT polarity complex. <i>Experimental Cell Research</i> , 2013, 319, 2514-2525.	1.2	30
46	MPP3 Is Required for Maintenance of the Apical Junctional Complex, Neuronal Migration, and Stratification in the Developing Cortex. <i>Journal of Neuroscience</i> , 2013, 33, 8518-8527.	1.7	10
47	Targeted Ablation of Crb1 and Crb2 in Retinal Progenitor Cells Mimics Leber Congenital Amaurosis. <i>PLoS Genetics</i> , 2013, 9, e1003976.	1.5	64
48	Loss of CRB2 in the mouse retina mimics human retinitis pigmentosa due to mutations in the CRB1 gene. <i>Human Molecular Genetics</i> , 2013, 22, 35-50.	1.4	74
49	MPP3 regulates levels of PALS1 and adhesion between photoreceptors and Müller cells. <i>Glia</i> , 2013, 61, 1629-1644.	2.5	12
50	Microarray and Morphological Analysis of Early Postnatal CRB2 Mutant Retinas on a Pure C57BL/6J Genetic Background. <i>PLoS ONE</i> , 2013, 8, e82532.	1.1	35
51	PALS1 Is Essential for Retinal Pigment Epithelium Structure and Neural Retina Stratification. <i>Journal of Neuroscience</i> , 2011, 31, 17230-17241.	1.7	48
52	Bone spicule pigment formation in retinitis pigmentosa: insights from a mouse model. <i>Graefes's Archive for Clinical and Experimental Ophthalmology</i> , 2010, 248, 1063-1070.	1.0	44
53	The Apical Complex Couples Cell Fate and Cell Survival to Cerebral Cortical Development. <i>Neuron</i> , 2010, 66, 69-84.	3.8	97
54	GFAP-Driven GFP Expression in Activated Mouse Müller Glial Cells Aligning Retinal Blood Vessels Following Intravitreal Injection of AAV2/6 Vectors. <i>PLoS ONE</i> , 2010, 5, e12387.	1.1	39

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55	Noninvasive, In Vivo Assessment of Mouse Retinal Structure Using Optical Coherence Tomography. PLoS ONE, 2009, 4, e7507.	1.1	183
56	TMEM16B, A Novel Protein with Calcium-Dependent Chloride Channel Activity, Associates with a Presynaptic Protein Complex in Photoreceptor Terminals. Journal of Neuroscience, 2009, 29, 6809-6818.	1.7	184
57	PSD95 ^{Δ2} regulates plasma membrane Ca(2+) pump localization at the photoreceptor synapse. Molecular and Cellular Neurosciences, 2009, 41, 156-165.	1.0	20
58	Contribution of the drug transporter ABCG2 (breast cancer resistance protein) to resistance against anticancer nucleosides. Molecular Cancer Therapeutics, 2008, 7, 3092-3102.	1.9	68
59	A Single Amino Acid Substitution (Cys249Trp) in Crb1 Causes Retinal Degeneration and Deregulates Expression of Pituitary Tumor Transforming Gene Pttg1. Journal of Neuroscience, 2007, 27, 564-573.	1.7	77
60	Crb1 is a determinant of retinal apical Müller glia cell features. Glia, 2007, 55, 1486-1497.	2.5	62
61	cGMP transport by vesicles from human and mouse erythrocytes. FEBS Journal, 2007, 274, 439-450.	2.2	61
62	Mice lacking Mrp3 (Abcc3) have normal bile salt transport, but altered hepatic transport of endogenous glucuronides. Journal of Hepatology, 2006, 44, 768-775.	1.8	158
63	The multidrug resistance protein ρ 1 (Mrp1), but not Mrp5, mediates export of glutathione and glutathione disulfide from brain astrocytes. Journal of Neurochemistry, 2006, 97, 373-384.	2.1	165
64	MPP3 is recruited to the MPP5 protein scaffold at the retinal outer limiting membrane. FEBS Journal, 2006, 273, 1152-1165.	2.2	31
65	Towards understanding CRUMBS function in retinal dystrophies. Human Molecular Genetics, 2006, 15, R235-R243.	1.4	112
66	Pals1/Mpp5 is required for correct localization of Crb1 at the subapical region in polarized Müller glia cells. Human Molecular Genetics, 2006, 15, 2659-2672.	1.4	98
67	Opposite Effects of PSD-95 and MPP3 PDZ Proteins on Serotonin 5-Hydroxytryptamine _{2C} Receptor Desensitization and Membrane Stability. Molecular Biology of the Cell, 2006, 17, 4619-4631.	0.9	70
68	Mpp4 recruits Psd95 and Veli3 towards the photoreceptor synapse. Human Molecular Genetics, 2006, 15, 1291-1302.	1.4	46
69	MPP5 Recruits MPP4 to the CRB1 Complex in Photoreceptors. , 2005, 46, 2192.		62
70	PATJ connects and stabilizes apical and lateral components of tight junctions in human intestinal cells. Journal of Cell Science, 2005, 118, 4049-4057.	1.2	127
71	The Human Multidrug Resistance Protein MRP5 Transports Folates and Can Mediate Cellular Resistance against Antifolates. Cancer Research, 2005, 65, 4425-4430.	0.4	114
72	In vivo confocal imaging of the retina in animal models using scanning laser ophthalmoscopy. Vision Research, 2005, 45, 3512-3519.	0.7	172

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73	Crumbs homologue 1 is required for maintenance of photoreceptor cell polarization and adhesion during light exposure. <i>Journal of Cell Science</i> , 2004, 117, 4169-4177.	1.2	220
74	ABCC6/MRP6 mutations: further insight into the molecular pathology of pseudoxanthoma elasticum. <i>European Journal of Human Genetics</i> , 2003, 11, 215-224.	1.4	57
75	Pseudoxanthoma elasticum: a clinical, histopathological, and molecular update. <i>Survey of Ophthalmology</i> , 2003, 48, 424-438.	1.7	149
76	The human multidrug resistance protein MRP4 functions as a prostaglandin efflux transporter and is inhibited by nonsteroidal antiinflammatory drugs. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2003, 100, 9244-9249.	3.3	478
77	Characterization of the MRP4- and MRP5-mediated Transport of Cyclic Nucleotides from Intact Cells. <i>Journal of Biological Chemistry</i> , 2003, 278, 17664-17671.	1.6	233
78	Characterization of the Transport of Nucleoside Analog Drugs by the Human Multidrug Resistance Proteins MRP4 and MRP5. <i>Molecular Pharmacology</i> , 2003, 63, 1094-1103.	1.0	346
79	Isolation of Crb1, a mouse homologue of <i>Drosophila</i> crumbs, and analysis of its expression pattern in eye and brain. <i>Mechanisms of Development</i> , 2002, 110, 203-207.	1.7	98
80	MRP6 (ABCC6) Detection in Normal Human Tissues and Tumors. <i>Laboratory Investigation</i> , 2002, 82, 515-518.	1.7	458
81	Differential susceptibility of multidrug resistance protein-1 deficient mice to DSS and TNBS-induced colitis. <i>Digestive Diseases and Sciences</i> , 2002, 47, 2056-2063.	1.1	37
82	Mice Lacking the Multidrug Resistance Protein 1 Are Resistant to <i>Streptococcus pneumoniae</i> -Induced Pneumonia. <i>Journal of Immunology</i> , 2001, 166, 4059-4064.	0.4	64
83	A Family of Drug Transporters: the Multidrug Resistance-Associated Proteins. <i>Journal of the National Cancer Institute</i> , 2000, 92, 1295-1302.	3.0	1,579
84	Multidrug resistance protein 1 protects the choroid plexus epithelium and contributes to the blood-cerebrospinal fluid barrier. <i>Journal of Clinical Investigation</i> , 2000, 105, 279-285.	3.9	334
85	The multidrug resistance protein family. <i>Biochimica Et Biophysica Acta - Biomembranes</i> , 1999, 1461, 347-357.	1.4	550
86	Multidrug Resistance Protein 1 Protects the Oropharyngeal Mucosal Layer and the Testicular Tubules against Drug-induced Damage. <i>Journal of Experimental Medicine</i> , 1998, 188, 797-808.	4.2	197
87	Transport of glutathione prostaglandin A conjugates by the multidrug resistance protein 1. <i>FEBS Letters</i> , 1997, 419, 112-116.	1.3	130
88	Increased sensitivity to anticancer drugs and decreased inflammatory response in mice lacking the multidrug resistance-associated protein. <i>Nature Medicine</i> , 1997, 3, 1275-1279.	15.2	409
89	Segment-Specific Expression of the neuronatin Gene during Early Hindbrain Development. <i>Developmental Biology</i> , 1995, 171, 73-84.	0.9	83
90	Pax-3-DNA interaction: flexibility in the DNA binding and induction of DNA conformational changes by paired domains. <i>Nucleic Acids Research</i> , 1994, 22, 3131-3137.	6.5	44

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91	Characterization of Pax-6 and Hoxa-1 Binding to the Promoter Region of the Neural Cell Adhesion Molecule L1. <i>DNA and Cell Biology</i> , 1994, 13, 891-900.	0.9	74
92	Binding of a bZip protein to the estrogen-inducible apoVLDL II promoter. <i>Biochimica Et Biophysica Acta Gene Regulatory Mechanisms</i> , 1994, 1219, 115-120.	2.4	7
93	Pax: Gene regulators in the developing nervous system. <i>Journal of Neurobiology</i> , 1993, 24, 1367-1384.	3.7	190
94	Estrogen-inducible and liver-specific expression of the chicken Very Low Density Apolipoprotein II gene locus in transgenic mice. <i>Nucleic Acids Research</i> , 1993, 21, 1629-1635.	6.5	5
95	cC/EPB, a chicken transcription factor of the leucinezipper c/EBP family. <i>Nucleic Acids Research</i> , 1992, 20, 4093-4093.	6.5	21
96	Oestrogen facilitates the binding of ubiquitous and liver-enriched nuclear proteins to the apoVLDL II promoter in vivo. <i>Nucleic Acids Research</i> , 1991, 19, 33-41.	6.5	43
97	Regulatory elements and DNA-binding proteins mediating transcription from the chicken very-low-density apolipoprotein II gene. <i>Nucleic Acids Research</i> , 1991, 19, 5371-5377.	6.5	36
98	AAV-Mediated Gene Therapy for CRB1-Hereditary Retinopathies. , 0, , .		1