

# Peter M J Quinn

## List of Publications by Year in descending order

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Version: 2024-02-01

23  
papers

745  
citations

687335

13  
h-index

642715

23  
g-index

24  
all docs

24  
docs citations

24  
times ranked

719  
citing authors

#	ARTICLE	IF	CITATIONS
1	PINK1/PARKIN signalling in neurodegeneration and neuroinflammation. <i>Acta Neuropathologica Communications</i> , 2020, 8, 189.	5.2	204
2	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.	4.8	75
3	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.	2.9	65
4	Targeted Ablation of Crb1 and Crb2 in Retinal Progenitor Cells Mimics Leber Congenital Amaurosis. <i>PLoS Genetics</i> , 2013, 9, e1003976.	3.5	64
5	Organoids and organ chips in ophthalmology. <i>Ocular Surface</i> , 2021, 19, 1-15.	4.4	45
6	The CRB1 Complex: Following the Trail of Crumbs to a Feasible Gene Therapy Strategy. <i>Frontiers in Neuroscience</i> , 2017, 11, 175.	2.8	43
7	Role of Oxidative Stress in Ocular Diseases Associated with Retinal Ganglion Cells Degeneration. <i>Antioxidants</i> , 2021, 10, 1948.	5.1	34
8	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.	2.9	29
9	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.	2.9	26
10	Retinogenesis of the Human Fetal Retina: An Apical Polarity Perspective. <i>Genes</i> , 2019, 10, 987.	2.4	24
11	The Role of Small Molecules and Their Effect on the Molecular Mechanisms of Early Retinal Organoid Development. <i>International Journal of Molecular Sciences</i> , 2021, 22, 7081.	4.1	23
12	Phase transition specified by a binary code patterns the vertebrate eye cup. <i>Science Advances</i> , 2021, 7, eabj9846.	10.3	19
13	Production of iPSC-Derived Human Retinal Organoids for Use in Transgene Expression Assays. <i>Methods in Molecular Biology</i> , 2018, 1715, 261-273.	0.9	17
14	Prime Editing for Inherited Retinal Diseases. <i>Frontiers in Genome Editing</i> , 2021, 3, 775330.	5.2	17
15	CRISPR genome surgery in a novel humanized model for autosomal dominant retinitis pigmentosa. <i>Molecular Therapy</i> , 2022, 30, 1407-1420.	8.2	16
16	Precision metabolome reprogramming for imprecision therapeutics in retinitis pigmentosa. <i>Journal of Clinical Investigation</i> , 2020, 130, 3971-3973.	8.2	11
17	Impaired cholesterol efflux in retinal pigment epithelium of individuals with juvenile macular degeneration. <i>American Journal of Human Genetics</i> , 2021, 108, 903-918.	6.2	10
18	Defining Phenotype, Tropism, and Retinal Gene Therapy Using Adeno-Associated Viral Vectors (AAVs) in New-Born Brown Norway Rats with a Spontaneous Mutation in Crb1. <i>International Journal of Molecular Sciences</i> , 2021, 22, 3563.	4.1	9

#	ARTICLE	IF	CITATIONS
19	CRISPR/Cas therapeutic strategies for autosomal dominant disorders. Journal of Clinical Investigation, 2022, 132, .	8.2	8
20	Mouse Models of Achromatopsia in Addressing Temporal "Point of No Return" in Gene-Therapy. International Journal of Molecular Sciences, 2021, 22, 8069.	4.1	2
21	NTPDase2 as a Surface Marker to Isolate Flow Cytometrically a Müller Glial Cell Enriched Population from Dissociated Neural Retinae. Journal of Neuroscience and Neurosurgery, 2018, 1, .	0.1	2
22	Overcoming translational barriers in modeling macular degenerations. Cell Stem Cell, 2021, 28, 781-783.	11.1	1
23	Transplantation of NTPDase2-positive Sorted Müller Glial Cells into the Mouse Retina. Journal of Neuroscience and Neurosurgery, 2018, 1, .	0.1	1