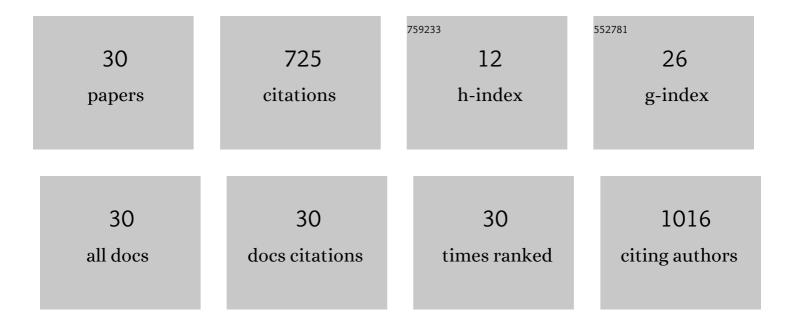
## Naihong Yan

List of Publications by Year in descending order

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Νλιμονς Υλν

#	Article	IF	CITATIONS
1	A microarray for microRNA profiling in mouse testis tissues. Reproduction, 2007, 134, 73-79.	2.6	167
2	Microarray profiling of microRNAs expressed in testis tissues of developing primates. Journal of Assisted Reproduction and Genetics, 2009, 26, 179-186.	2.5	93
3	Tetrahedral framework nucleic acids-based delivery of microRNA-155 inhibits choroidal neovascularization by regulating the polarization of macrophages. Bioactive Materials, 2022, 14, 134-144.	15.6	77
4	Hypoxia-Inducible Factor-1α Target Genes Contribute to Retinal Neuroprotection. Frontiers in Cellular Neuroscience, 2017, 11, 20.	3.7	61
5	Exome Sequencing Identifies Compound Heterozygous Mutations in CYP4V2 in a Pedigree with Retinitis Pigmentosa. PLoS ONE, 2012, 7, e33673.	2.5	40
6	DZNep inhibits H3K27me3 deposition and delays retinal degeneration in the rd1 mice. Cell Death and Disease, 2018, 9, 310.	6.3	37
7	The Neuroprotective Effect of MicroRNAâ€22â€3p Modified Tetrahedral Framework Nucleic Acids on Damaged Retinal Neurons Via TrkB/BDNF Signaling Pathway. Advanced Functional Materials, 2021, 31, 2104141.	14.9	36
8	Neuroprotective Effects of C-Type Natriuretic Peptide on Rat Retinal Ganglion Cells. , 2010, 51, 3544.		26
9	Postnatal onset of retinal degeneration by loss of embryonic Ezh2 repression of Six1. Scientific Reports, 2016, 6, 33887.	3.3	26
10	E2f1 mediates high glucose-induced neuronal death in cultured mouse retinal explants. Cell Cycle, 2017, 16, 1824-1834.	2.6	23
11	Myopia: A collagen disease?. Medical Hypotheses, 2009, 73, 485-487.	1.5	20
12	Effects of nerve growth factor on nerve regeneration after corneal nerve damage. International Journal of Clinical and Experimental Medicine, 2014, 7, 4584-9.	1.3	16
13	A Novel Nonsense Mutation of the GPR143 Gene Identified in a Chinese Pedigree with Ocular Albinism. PLoS ONE, 2012, 7, e43177.	2.5	13
14	Ezh2 does not mediate retinal ganglion cell homeostasis or their susceptibility to injury. PLoS ONE, 2018, 13, e0191853.	2.5	10
15	Investigation of indoleamine 2,3-dioxygenase 1 expression in uveal melanoma. Experimental Eye Research, 2019, 181, 112-119.	2.6	10
16	Mutations of GNAQ, GNA11, SF3B1, EIF1AX, PLCB4 and CYSLTR in Uveal Melanoma in Chinese Patients. Ophthalmic Research, 2020, 63, 358-368.	1.9	10
17	A novel mitochondrial tRNA(Val) T1658C mutation identified in a CPEO family. Molecular Vision, 2010, 16, 1736-42.	1.1	9
18	Antiviral Activity of a Cloned Peptide RC28 Isolated from the Higher Basidiomycetes Mushroom Rozites caperata in a Mouse Model of HSV-1 Keratitis. International Journal of Medicinal Mushrooms, 2015, 17, 819-828.	1.5	7

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#	Article	IF	CITATIONS
19	Molecular genetics of familial nystagmus complicated with cataract and iris anomalies. Molecular Vision, 2011, 17, 2612-7.	1.1	7
20	DZNep protects against retinal ganglion cell death in an NMDA-induced mouse model of retinal degeneration. Experimental Eye Research, 2021, 212, 108785.	2.6	6
21	Purification, partial characterization and molecular cloning of the novel antiviral protein RC28. Peptides, 2009, 30, 654-659.	2.4	5
22	X-linked inheritances recessive of congenital nystagmus and autosomal dominant inheritances of congenital cataracts coexist in a Chinese family: a case report and literature review. BMC Medical Genetics, 2019, 20, 41.	2.1	5
23	Profiling MicroRNAs Differentially Expressed in Rabbit Retina. Advances in Experimental Medicine and Biology, 2010, 664, 203-209.	1.6	5
24	Possible mechanism for the gastro-intestinal adverse effects upon topical application of Prostaglandin F2α analogs. Medical Hypotheses, 2013, 80, 32-35.	1.5	4
25	Retinal Degeneration: Molecular Mechanisms and Therapeutic Strategies. Current Medicinal Chemistry, 2022, 29, 6125-6140.	2.4	4
26	Dihydroartemisinin Inhibits Laser-Induced Choroidal Neovascularization in a Mouse Model of Neovascular AMD. Frontiers in Pharmacology, 2022, 13, 838263.	3.5	3
27	Heterogeneity of GNAQ/11 mutation inversely correlates with the metastatic rate in uveal melanoma. British Journal of Ophthalmology, 2021, 105, 587-592.	3.9	2
28	A Novel Mutation in the Membrane Frizzled-Related Protein Gene for Posterior Microphthalmia, Non-pigmented Retinitis Pigmentosa, Optic Nerve Drusen, and Retinoschisis in a Consanguineous Family. Frontiers in Medicine, 2022, 9, 835621.	2.6	2
29	Case Report: A Novel Mutation in the CRYGD Gene Causing Congenital Cataract Associated with Nystagmus in a Chinese Family. Frontiers in Genetics, 2022, 13, 824550.	2.3	1
30	22q12.3-q13.1 microdeletion including <i>SOX10</i> causes atypical Waardenburg syndrome. European Journal of Ophthalmology, 2021, 31, 2127-2134.	1.3	0