## Ping Liu

## List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/8224034/publications.pdf Version: 2024-02-01



DINCLU

#	Article	IF	CITATIONS
1	The novel mutation P36R in LRP5L contributes to congenital membranous cataract via inhibition of laminin γ1 and c-MAF. Graefe's Archive for Clinical and Experimental Ophthalmology, 2020, 258, 2737-2751.	1.9	2
2	Impairment of the autophagy-lysosomal pathway and activation of pyroptosis in macular corneal dystrophy. Cell Death Discovery, 2020, 6, 85.	4.7	23
3	Laminin α4 overexpression in the anterior lens capsule may contribute to the senescence of human lens epithelial cells in age-related cataract. Aging, 2019, 11, 2699-2723.	3.1	18
4	Laminins in an inÃ <sup>-</sup> Â;½vitro anterior lens capsule model established using HLE B-3 cells. Molecular Medicine Reports, 2018, 17, 5726-5733.	2.4	6
5	<i>CHST6</i> mutation screening and endoplasmatic reticulum stress in macular corneal dystrophy. Oncotarget, 2017, 8, 96301-96312.	1.8	14
6	A C-terminal fragment BIGH3 protein with an RGDRGD motif inhibits corneal neovascularization inÂvitro and inÂvivo. Experimental Eye Research, 2013, 112, 10-20.	2.6	11
7	A new locus in chromosome 2q37-qter is associated with posterior polar cataract. Graefe's Archive for Clinical and Experimental Ophthalmology, 2012, 250, 907-913.	1.9	4
8	Comparison of the antiangiogenic activity of modified RGDRGD-endostatin to endostatin delivered by gene transfer in vivo rabbit neovascularization model. Molecular Vision, 2011, 17, 1918-28.	1.1	11
9	ModifiedBIGH3with anRGDRGDMotif Promotes Human Corneal Epithelial Cell Adhesion and MigrationIn Vitro. Current Eye Research, 2008, 33, 215-223.	1.5	4
10	The E233del mutation in BFSP2 causes a progressive autosomal dominant congenital cataract in a Chinese family. Molecular Vision, 2007, 13, 2023-9.	1.1	18
11	A novel locus of coralliform cataract mapped to chromosome 2p24-pter. Journal of Human Genetics, 2005, 50, 305-310.	2.3	11