

# Shengping Hou

## List of Publications by Year in descending order

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14  
papers

343  
citations

840776

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1058476

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15  
docs citations

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times ranked

298  
citing authors

#	ARTICLE	IF	CITATIONS
1	Galectin-3 regulates microglial activation and promotes inflammation through TLR4/MyD88/NF- $\kappa$ B in experimental autoimmune uveitis. <i>Clinical Immunology</i> , 2022, 236, 108939.	3.2	45
2	Aryl Hydrocarbon Receptor Regulates Apoptosis and Inflammation in a Murine Model of Experimental Autoimmune Uveitis. <i>Frontiers in Immunology</i> , 2018, 9, 1713.	4.8	43
3	Genetic Variations of IL17F and IL23A Show Associations with Behçet's Disease and Vogt-Koyanagi-Harada Syndrome. <i>Ophthalmology</i> , 2015, 122, 518-523.	5.2	40
4	Small molecules targeting ROR $\gamma$ t inhibit autoimmune disease by suppressing Th17 cell differentiation. <i>Cell Death and Disease</i> , 2020, 11, 697.	6.3	32
5	Association between C4, C4A, and C4B copy number variations and susceptibility to autoimmune diseases: a meta-analysis. <i>Scientific Reports</i> , 2017, 7, 42628.	3.3	31
6	Icariin alleviates uveitis by targeting peroxiredoxin 3 to modulate retinal microglia M1/M2 phenotypic polarization. <i>Redox Biology</i> , 2022, 52, 102297.	9.0	25
7	Retinal microglia: Functions and diseases. <i>Immunology</i> , 2022, 166, 268-286.	4.4	24
8	Monocyte chemoattractant protein-1 $\sim$ 2518 A/G single nucleotide polymorphism in Chinese Han patients with ocular Behçet's disease. <i>Human Immunology</i> , 2010, 71, 79-82.	2.4	22
9	Small ubiquitin-like modifier 4 (SUMO4) polymorphisms and Vogt-Koyanagi-Harada (VKH) syndrome in the Chinese Han population. <i>Molecular Vision</i> , 2008, 14, 2597-603.	1.1	20
10	Low Expression of YTH Domain-Containing 1 Promotes Microglial M1 Polarization by Reducing the Stability of Sirtuin 1 mRNA. <i>Frontiers in Cellular Neuroscience</i> , 2021, 15, 774305.	3.7	20
11	A novel mutation of <i>PANK4</i> causes autosomal dominant congenital posterior cataract. <i>Human Mutation</i> , 2019, 40, 380-391.	2.5	12
12	Methyltransferase-like (METTL)14-mediated N6-methyladenosine modification modulates retinal pigment epithelial (RPE) activity by regulating the methylation of microtubule-associated protein (MAP)2. <i>Bioengineered</i> , 2022, 13, 4773-4785.	3.2	12
13	The role of the inflammasomes in the pathogenesis of uveitis. <i>Experimental Eye Research</i> , 2021, 208, 108618.	2.6	10
14	Monocyte chemoattractant protein (MCP)-1 -2518 A/G SNP in Chinese Han patients with VKH syndrome. <i>Molecular Vision</i> , 2009, 15, 1537-41.	1.1	6