Shengping Hou

List of Publications by Year
in descending order

14	343	840776 11	1058476 14
papers	citations	h-index	g-index

15	15	15	298
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Galectin-3 regulates microglial activation and promotes inflammation through TLR4/MyD88/NF-kB in experimental autoimmune uveitis. Clinical Immunology, 2022, 236, 108939.	3.2	45
2	Methyltransferase-like (METTL)14-mediated N6-methyladenosine modification modulates retinal pigment epithelial (RPE) activity by regulating the methylation of microtubule-associated protein (MAP)2. Bioengineered, 2022, 13, 4773-4785.	3.2	12
3	Icariin alleviates uveitis by targeting peroxiredoxin 3 to modulate retinal microglia M1/M2 phenotypic polarization. Redox Biology, 2022, 52, 102297.	9.0	25
4	Retinal microglia: Functions and diseases. Immunology, 2022, 166, 268-286.	4.4	24
5	The role of the inflammasomes in the pathogenesis of uveitis. Experimental Eye Research, 2021, 208, 108618.	2.6	10
6	Low Expression of YTH Domain-Containing 1 Promotes Microglial M1 Polarization by Reducing the Stability of Sirtuin 1 mRNA. Frontiers in Cellular Neuroscience, 2021, 15, 774305.	3.7	20
7	Small molecules targeting RORÎ³t inhibit autoimmune disease by suppressing Th17 cell differentiation. Cell Death and Disease, 2020, 11, 697.	6.3	32
8	A novel mutation ofPANK4causes autosomal dominant congenital posterior cataract. Human Mutation, 2019, 40, 380-391.	2.5	12
9	Aryl Hydrocarbon Receptor Regulates Apoptosis and Inflammation in a Murine Model of Experimental Autoimmune Uveitis. Frontiers in Immunology, 2018, 9, 1713.	4.8	43
10	Association between C4, C4A, and C4B copy number variations and susceptibility to autoimmune diseases: a meta-analysis. Scientific Reports, 2017, 7, 42628.	3.3	31
11	Genetic Variations of IL17F and IL23A ShowÂAssociations with BehÃ§etâ€™s Disease andÂVogt-Koyanagi-Harada Syndrome. Ophthalmology, 2015, 122, 518-523.	5.2	40
12	Monocyte chemoattractant proteinâ€“1 â^’2518 A/G single nucleotide polymorphism in Chinese Han patients with ocular BehÃ§et's disease. Human Immunology, 2010, 71, 79-82.	2.4	22
13	Monocyte chemoattractant protein (MCP)-1 -2518 A/G SNP in Chinese Han patients with VKH syndrome. Molecular Vision, 2009, 15, 1537-41.	1.1	6
14	Small ubiquitin-like modifier 4 (SUMO4) polymorphisms and Vogt-Koyanagi-Harada (VKH) syndrome in the Chinese Han population. Molecular Vision, 2008, 14, 2597-603.	1.1	20