Peizeng Yang

List of Publications by Year
in descending order

259 papers	7,281 citations	108046 37 h-index	129628 63 g-index
280 all docs	280 docs citations	280 times ranked	5576 citing authors

#	Article	IF	Citations
1	Outcome Measures for Disease Monitoring in Intraocular Inflammatory and Infectious Diseases (OCTOMERIA): Understanding the Choroid in Uveitis with Optical Coherence Tomography (OCT). Ocular Immunology and Inflammation, 2023, 31, 374-392.	1.0	4
2	Succinic acid exacerbates experimental autoimmune uveitis by stimulating neutrophil extracellular traps formation via SUCNR1 receptor. British Journal of Ophthalmology, 2023, 107, 1744-1749.	2.1	5
3	Association between temperature changes and uveitis onset in mainland China. British Journal of Ophthalmology, 2022, 106, 91-96.	2.1	10
4	Development of revised diagnostic criteria for Fuchsâ€™ uveitis syndrome in a Chinese population. British Journal of Ophthalmology, 2022, 106, 1678-1683.	2.1	7
5	Identification of differently expressed mRNAs by peripheral blood mononuclear cells in Vogt-Koyanagi-Harada disease. Genes and Diseases, 2022, 9, 1378-1388.	1.5	4
6	Association between Fine Particulate Air Pollution and the Onset of Uveitis in Mainland China. Ocular Immunology and Inflammation, 2022, 30, 1810-1815.	1.0	4
7	Identification of Novel Risk Loci for BehÃ§et's Diseaseâ€“Related Uveitis in a <scp>Genomeâ€Wide</scp> Association Study. Arthritis and Rheumatology, 2022, 74, 671-681.	2.9	14
8	Surveillance of Liver Function in Uveitis with or without Chronic HBV Infection. Ophthalmic Research, 2022, 65, 94-103.	1.0	0
9	Evaluation of sensitivity and specificity of diagnostic criteria for BehÃ§etâ€™s disease in the absence of a gold standard. Rheumatology, 2022, 61, 3667-3676.	0.9	7
10	Progranulin Suppressed Autoimmune Uveitis and Autoimmune Neuroinflammation by Inhibiting Th1/Th17 Cells and Promoting Treg Cells and M2 Macrophages. Neurology: Neuroimmunology and NeuroInflammation, 2022, 9, .	3.1	20
11	Visualizing lipid behind the retina in aging and age-related macular degeneration, via indocyanine green angiography (ASHS-LIA). Eye, 2022, 36, 1735-1746.	1.1	12
12	Genetic and Clinical Features of Blau Syndrome among Chinese Patients with Uveitis. Ophthalmology, 2022, 129, 821-828.	2.5	5
13	Genetically predicted fasting blood glucose level plays a causal role in intraocular pressure: A Mendelian randomisation study. Clinical and Experimental Ophthalmology, 2022, , .	1.3	0
14	Average corticosteroid dose and risk for HBV reactivation and hepatitis flare in patients with resolved hepatitis B infection. Annals of the Rheumatic Diseases, 2022, 81, 584-591.	0.5	11
15	SNP rs7130280 in lncRNA NONHSAT159216.1 confers susceptibility to BehÃ§etâ€™s disease uveitis in a Chinese Han population. Rheumatology, 2022, 62, 384-396.	0.9	3
16	PD-1 Targeted Nanoparticles Inhibit Activated T Cells and Alleviate Autoimmunity via Suppression of Cellular Energy Metabolism Mediated by PKM2. International Journal of Nanomedicine, 2022, Volume 17, 1711-1724.	3.3	5
17	Prevalence, risk factors and management of ocular hypertension or glaucoma in patients with Vogt-Koyanagi-Harada disease. British Journal of Ophthalmology, 2021, 105, 1678-1682.	2.1	7
18	Prevalence and clinical features of systemic diseases in Chinese patients with uveitis. British Journal of Ophthalmology, 2021, 105, 75-82.	2.1	37

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19	Activation of the interleukin-23/interleukin-17 signalling pathway in autoinflammatory and autoimmune uveitis. Progress in Retinal and Eye Research, 2021, 80, 100866.	7.3	104
20	Higher 25-hydroxyvitamin D level is associated with increased risk for BehÃ§et's disease. Clinical Nutrition, 2021, 40, 518-524.	2.3	12
21	Linoleic acid inhibits in vitro function of human and murine dendritic cells, CD4+T cells and retinal pigment epithelial cells. Graefe's Archive for Clinical and Experimental Ophthalmology, 2021, 259, 987-998.	1.0	8
22	Association of a CARD9 Gene Haplotype with Behcetâ€™s Disease in a Chinese Han Population. Ocular Immunology and Inflammation, 2021, 29, 219-227.	1.0	5
23	Metabolomic Analysis of Aqueous Humor Identifies Aberrant Amino Acid and Fatty Acid Metabolism in Vogt-Koyanagi-Harada and Behcetâ€™s Disease. Frontiers in Immunology, 2021, 12, 587393.	2.2	11
24	Identification of Urine Metabolic Biomarkers for Vogt-Koyanagi-Harada Disease. Frontiers in Cell and Developmental Biology, 2021, 9, 637489.	1.8	10
25	High Ambient Temperature Aggravates Experimental Autoimmune Uveitis Symptoms. Frontiers in Cell and Developmental Biology, 2021, 9, 629306.	1.8	8
26	Tuberculosis Exposure With Risk of BehÃ§et Disease Among Patients With Uveitis. JAMA Ophthalmology, 2021, 139, 415.	1.4	12
27	Ocular involvement in extranodal natural-killer T-cell lymphoma. Lancet Haematology,the, 2021, 8, e382.	2.2	2
28	Specific sweat metabolite profile in ocular Behcetâ€™s disease. International Immunopharmacology, 2021, 97, 107812.	1.7	12
29	Changes in the Gut Microbiome Contribute to the Development of Behcetâ€™s Disease via Adjuvant Effects. Frontiers in Cell and Developmental Biology, 2021, 9, 716760.	1.8	9
30	Auxiliary Ocular Examinations. , 2021, , 81-138.		1
31	Meta-Analysis of miRNA Variants Associated with Susceptibility to Autoimmune Disease. Disease Markers, 2021, 2021, 1-21.	0.6	4
32	SNP-mediated binding of TBX1 to the enhancer element of IL-10 reduces the risk of BehÃ§etâ€™s disease. Epigenomics, 2021, 13, 1523-1537.	1.0	1
33	Complications and Their Management. , 2021, , 165-168.		0
34	Acute Anterior Uveitis. , 2021, , 171-185.		0
35	Intermediate Uveitis. , 2021, , 187-193.		0
36	Uveitis-associated with Juvenile Idiopathic Arthritis. , 2021, , 307-325.		0

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37	Vogtâ€“Koyanagiâ€“Harada Disease. , 2021, , 411-537.		2
38	Retinal Vasculitis. , 2021, , 563-587.		0
39	Ocular Sarcoidosis. , 2021, , 611-625.		0
40	Ocular Examinations. , 2021, , 19-79.		0
41	Scleritis. , 2021, , 825-857.		0
42	Steroid-sparing Immunosuppressive Agents. , 2021, , 155-161.		0
43	Optical Coherence Tomographic Features and Prognostic Values of Macular Edema in Vogt-Koyanagi-Harada Disease. Frontiers in Medicine, 2021, 8, 772439.	1.2	1
44	A Single-Cell Transcriptome Atlas of the Human Retinal Pigment Epithelium. Frontiers in Cell and Developmental Biology, 2021, 9, 802457.	1.8	15
45	Analyses of circRNA and mRNA Profiles in Vogtâ€“Koyanagiâ€“Harada Disease. Frontiers in Immunology, 2021, 12, 738760.	2.2	7
46	Decreased Expression of TGR5 in Vogt-Koyanagi-Harada (VKH) Disease. Ocular Immunology and Inflammation, 2020, 28, 200-208.	1.0	5
47	Genetic aspects of idiopathic paediatric uveitis and juvenile idiopathic arthritis associated uveitis in Chinese Han. British Journal of Ophthalmology, 2020, 104, 443-447.	2.1	4
48	Association of toll-like receptor 10 polymorphisms with paediatric idiopathic uveitis in Han Chinese. British Journal of Ophthalmology, 2020, 104, 1467-1471.	2.1	5
49	Uveitis genetics. Experimental Eye Research, 2020, 190, 107853.	1.2	36
50	Association of apoptosis genes in PDCD1 but not PDCD1LG2, FAS, and FASLG with pediatric idiopathic uveitis in Han Chinese. Pediatric Research, 2020, 87, 634-638.	1.1	6
51	Altered gut microbiome composition in patients with Vogt-Koyanagi-Harada disease. Gut Microbes, 2020, 11, 539-555.	4.3	52
52	Aqueous cytokine levels in four common uveitis entities. International Immunopharmacology, 2020, 78, 106021.	1.7	18
53	The haplotypes of various TNF related genes associated with scleritis in Chinese Han. Human Genomics, 2020, 14, 46.	1.4	3
54	Sharing of Genetic Association Signals by Age-Related Macular Degeneration and Alzheimerâ€™s Disease at Multiple Levels. Molecular Neurobiology, 2020, 57, 4488-4499.	1.9	7

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55	Small molecules targeting RORÎ³t inhibit autoimmune disease by suppressing Th17 cell differentiation. Cell Death and Disease, 2020, 11, 697.	2.7	32
56	The Rs12569232 SNP Association with Vogt-Koyanagi-Harada Disease and Behcetâ€™s Disease is Probably Mediated by Regulation of Linc00467 Expression. Ocular Immunology and Inflammation, 2020, , 1-7.	1.0	3
57	Increased Expression of Indoleamine 2,3-Dioxygenase (IDO) in Vogt-Koyanagi-Harada (VKH) Disease May Lead to a Shift of T Cell Responses Toward a Treg Population. Inflammation, 2020, 43, 1780-1788.	1.7	3
58	Effect of berberine on spleen transcriptome and gut microbiota composition in experimental autoimmune uveitis. International Immunopharmacology, 2020, 81, 106270.	1.7	14
59	Integrated omics analysis of sweat reveals an aberrant amino acid metabolism pathway in Vogtâ€“Koyanagiâ€“Harada disease. Clinical and Experimental Immunology, 2020, 200, 250-259.	1.1	12
60	Identification of Ribosomal Protein S4, Y-Linked 1 as a cyclosporin A plus corticosteroid resistance gene. Journal of Autoimmunity, 2020, 112, 102465.	3.0	10
61	Plasma metabolomics study of Vogt-Koyanagi-Harada disease identifies potential diagnostic biomarkers. Experimental Eye Research, 2020, 196, 108070.	1.2	13
62	Analysis of the role of palmitoleic acid in acute anterior uveitis. International Immunopharmacology, 2020, 84, 106552.	1.7	8
63	Vogtâ€“Koyanagiâ€“Harada Disease. Retina Atlas, 2020, , 67-75.	0.0	1
64	Integrated Analysis of Key Pathways and Drug Targets Associated With Vogt-Koyanagi-Harada Disease. Frontiers in Immunology, 2020, 11, 587443.	2.2	11
65	Weak association of a TNFRSF1A polymorphism with Behcet's disease in Chinese Han. Experimental Eye Research, 2020, 196, 108045.	1.2	1
66	Macular Abnormalities in Vogt-Koyanagi-Harada Disease. Ocular Immunology and Inflammation, 2019, 27, 1195-1202.	1.0	15
67	Association of TLR2 Gene Polymorphisms with Presumed Viral-Induced Anterior Uveitis in male Han Chinese. Experimental Eye Research, 2019, 187, 107777.	1.2	1
68	Diagnosis and treatment of human sparganosis. Lancet Infectious Diseases, The, 2019, 19, 577-578.	4.6	3
69	Different Methylation of CpG-SNPs in Behcetâ€™s Disease. BioMed Research International, 2019, 2019, 1-7.	0.9	3
70	Comparison of Clinical Features and Visual Outcome between Sympathetic Ophthalmia and Vogtâ€“Koyanagiâ€“Harada Disease in Chinese Patients. Ophthalmology, 2019, 126, 1297-1305.	2.5	30
71	Epigenome-wide association study identifies BehÃ§etâ€™s disease-associated methylation loci in Han Chinese. Rheumatology, 2019, 58, 1574-1584.	0.9	21
72	Identification of susceptibility SNPs in CTLA-4 and PTPN22 for scleritis in Han Chinese. Clinical and Experimental Immunology, 2019, 197, 230-236.	1.1	6

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73	A 50-year-old woman with a recurrent eyelid swelling. Lancet Infectious Diseases, The, 2019, 19, 338.	4.6	4
74	Replication of Genome-Wide Association Analysis Identifies New Susceptibility Loci at Long Noncoding RNA Regions for Vogt-Koyanagi-Harada Disease. , 2019, 60, 4820.		7
75	UVEOGENE: An SNP database for investigations on genetic factors associated with uveitis and their relationship with other systemic autoimmune diseases. Human Mutation, 2019, 40, 258-266.	1.1	12
76	Clinical features of Chinese patients with relapsing polychondritis. British Journal of Ophthalmology, 2019, 103, 1129-1132.	2.1	8
77	Causes of Visual Impairment and Blindness in the 2006 and 2014 Nine-Province Surveys in Rural China. American Journal of Ophthalmology, 2019, 197, 80-87.	1.7	32
78	Long-Term Efficacy and Safety of Interferon Alpha-2a in the Treatment of Chinese Patients with BehÃ§etâ€™s Uveitis Not Responding to Conventional Therapy. Ocular Immunology and Inflammation, 2019, 27, 7-14.	1.0	20
79	Decreased interleukin(IL)-35 Expression is Associated with Active Intraocular Inflammation in Vogt-Koyanagi-Harada (VKH) Disease. Ocular Immunology and Inflammation, 2019, 27, 595-601.	1.0	6
80	ERAP1/ERAP2 and RUNX3 polymorphisms are not associated with ankylosing spondylitis susceptibility in Chinese Han. Clinical and Experimental Immunology, 2018, 193, 95-102.	1.1	13
81	Association of genetic variations in PTPN2 and CD122 with ocular Behcetâ€™s disease. British Journal of Ophthalmology, 2018, 102, 996-1002.	2.1	7
82	Longitudinal Study of Visual Function in Vogt-Koyanagi-Harada Disease Using Full-Field Electroretinography. American Journal of Ophthalmology, 2018, 191, 92-99.	1.7	6
83	Decreased expression of A20 is associated with ocular Behcetâ€™s disease (BD) but not with Vogt-Koyanagi-HaradaÂ(VKH)Âdisease. British Journal of Ophthalmology, 2018, 102, 1167-1172.	2.1	11
84	Analysis of the association between Fc receptor family gene polymorphisms and ocular BehÃ§etâ€™s disease in Han Chinese. Scientific Reports, 2018, 8, 4850.	1.6	5
85	Clinical Features and Complications of Scleritis in Chinese Patients. Ocular Immunology and Inflammation, 2018, 26, 387-396.	1.0	38
86	Clinical features of HLA-B27-positive acute anterior uveitis with or without ankylosing spondylitis in a Chinese cohort. British Journal of Ophthalmology, 2018, 102, 215-219.	2.1	50
87	Prevalence of Vision Impairment in Older Adults in Rural China in 2014 and Comparisons With the 2006 China Nine-Province Survey. American Journal of Ophthalmology, 2018, 185, 81-93.	1.7	48
88	MicroRNA-20a-5p suppresses IL-17 production by targeting OSM and CCL1 in patients with Vogt-Koyanagi-Harada disease. British Journal of Ophthalmology, 2018, 102, 282-290.	2.1	31
89	The Choroidal Vascularity Index Decreases and Choroidal Thickness Increases in Vogtâ€“Koyanagiâ€“Harada Disease Patients During a Recurrent Anterior Uveitis Attack. Ocular Immunology and Inflammation, 2018, 26, 1237-1243.	1.0	33
90	Multispectral image analysis in Vogtâ€“Koyanagiâ€“Harada disease. Acta Ophthalmologica, 2018, 96, 411-419.	0.6	13

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91	Novel treatment regimen of Vogtâ€“Koyanagiâ€“Harada disease with a reduced dose of corticosteroids combined with immunosuppressive agents. Current Eye Research, 2018, 43, 254-261.	0.7	39
92	How To Deal With Uveitis Patients?. Current Molecular Medicine, 2018, 17, 468-470.	0.6	30
93	Case-Control Study and Meta-Analysis Show a Weak Association between ANTXR2 Polymorphisms and Ankylosing Spondylitis in Chinese Han. BioMed Research International, 2018, 2018, 1-7.	0.9	1
94	Disabled-2 (DAB2) Overexpression Inhibits Monocyte-Derived Dendritic Cells' Function in Vogt-Koyanagi-Harada Disease. , 2018, 59, 4662.		10
95	Dynamic DNA Methylation Changes ofTbx21andRorcduring Experimental Autoimmune Uveitis in Mice. Mediators of Inflammation, 2018, 2018, 1-13.	1.4	16
96	Outcome and Prognostic Factors of Phacoemulsification Cataract Surgery in Vogt-Koyanagi-Harada Uveitis. American Journal of Ophthalmology, 2018, 196, 121-128.	1.7	17
97	Development and Evaluation of Diagnostic Criteria for Vogt-Koyanagi-Harada Disease. JAMA Ophthalmology, 2018, 136, 1025.	1.4	83
98	Association of LACC1, CEBPB-PTPN1, RIPK2 and ADO-EGR2 with ocular Behcetâ€™s disease in a Chinese Han population. British Journal of Ophthalmology, 2018, 102, 1308-1314.	2.1	16
99	Association of Long Noncoding RNAs Polymorphisms With Ankylosing Spondylitis, Vogt-Koyanagi-Harada Disease, and Behcet's Disease. , 2018, 59, 1158.		12
100	A metagenomic study of the gut microbiome in Behcetâ€™s disease. Microbiome, 2018, 6, 135.	4.9	173
101	Gut Microbiota Composition and Fecal Metabolic Phenotype in Patients With Acute Anterior Uveitis. , 2018, 59, 1523.		77
102	Aryl Hydrocarbon Receptor Regulates Apoptosis and Inflammation in a Murine Model of Experimental Autoimmune Uveitis. Frontiers in Immunology, 2018, 9, 1713.	2.2	43
103	Increased Expression of IL-23 Receptor (IL-23R) in Vogtâ€“Koyanagiâ€“Harada (VKH) Disease. Current Eye Research, 2018, 43, 1369-1373.	0.7	7
104	Hypermethylation of Interferon Regulatory Factor 8 (IRF8) Confers Risk to Vogt-Koyanagi-Harada Disease. Scientific Reports, 2017, 7, 1007.	1.6	23
105	Genetic Background of Uveitis in Chinese Population. Essentials in Ophthalmology, 2017, , 425-436.	0.0	0
106	Increased Complement 3a Receptor is Associated with Behcetâ€™s disease and Vogt-Koyanagi-Harada disease. Scientific Reports, 2017, 7, 15579.	1.6	10
107	Genetic polymorphisms of C-type lectin receptors in Behcetâ€™s disease in a Chinese Han population. Scientific Reports, 2017, 7, 5348.	1.6	9
108	Uveitis in Chinese Patients with Psoriasis. Ocular Immunology and Inflammation, 2017, 25, 855-865.	1.0	14

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109	Identification of susceptibility SNPs in IL10 and IL23R-IL12RB2 for BehÃ§et's disease in Han Chinese. Journal of Allergy and Clinical Immunology, 2017, 139, 621-627.	1.5	36
110	miRNA Copy Number Variants Confer Susceptibility to Acute Anterior Uveitis With or Without Ankylosing Spondylitis. , 2017, 58, 1991.		11
111	Association of a PDCD1 Polymorphism With Sympathetic Ophthalmia in Han Chinese. , 2017, 58, 4218.		12
112	The Association of Chemokine Gene Polymorphisms with VKH and Behcetâ€™s Disease in a Chinese Han Population. BioMed Research International, 2017, 2017, 1-8.	0.9	16
113	Propofol inhibits lung cancer cell viability and induces cell apoptosis by upregulating microRNA-486 expression. Brazilian Journal of Medical and Biological Research, 2017, 50, e5794.	0.7	55
114	Promoter Hypermethylation of GATA3, IL-4, and TGF-Î² Confers Susceptibility to Vogt-Koyanagi-Harada Disease in Han Chinese. , 2017, 58, 1529.		18
115	Ocular Behcetâ€™s disease is associated with aberrant methylation of interferon regulatory factor 8 (IRF8) in monocyte-derived dendritic cells. Oncotarget, 2017, 8, 51277-51287.	0.8	9
116	Aberrant DNA methylation of GATA binding protein 3 (GATA3), interleukin-4 (IL-4), and transforming growth factor-Î² (TGF-Î²) promoters in Behcet's disease. Oncotarget, 2017, 8, 64263-64272.	0.8	14
117	The Role of Mitochondria-Associated Reactive Oxygen Species in the Amyloid Î² Induced Production of Angiogenic Factors b y ARPE-19 Cells. Current Molecular Medicine, 2017, 17, 140-148.	0.6	18
118	Association of T-Bet, GATA-3, RORC, and FOXP3 Copy Number Variations With Acute Anterior Uveitis With or Without Ankylosing Spondylitis in Chinese Han. , 2016, 57, 1847.		9
119	Decreased B and T lymphocyte attenuator in Behcetâ€™s disease may trigger abnormal Th17 and Th1 immune responses. Scientific Reports, 2016, 6, 20401.	1.6	26
120	Genetic polymorphisms of cell adhesion molecules in Behcetâ€™s disease in a Chinese Han population. Scientific Reports, 2016, 6, 24974.	1.6	21
121	Analysis of receptor tyrosine kinase genetics identifies two novel risk loci in GAS6 and PROS1 in BehÃ§etâ€™s disease. Scientific Reports, 2016, 6, 26662.	1.6	10
122	Association of TNFSF4 Polymorphisms with Vogt-Koyanagi-Harada and Behcetâ€™s Disease in Han Chinese. Scientific Reports, 2016, 6, 37257.	1.6	16
123	miR-23a, miR-146a and miR-301a confer predisposition to Vogt-Koyanagi-Harada syndrome but not to Behcetâ€™s disease. Scientific Reports, 2016, 6, 20057.	1.6	22
124	Genetic analysis of innate immunity in Behcetâ€™s disease identifies an association with IL-37 and IL-18RAP. Scientific Reports, 2016, 6, 35802.	1.6	36
125	Two Genetic Variations in the IRF8 region are associated with BehÃ§etâ€™s disease in Han Chinese. Scientific Reports, 2016, 6, 19651.	1.6	22
126	Genetic Variations of NLR family genes in Behcetâ€™s Disease. Scientific Reports, 2016, 6, 20098.	1.6	21

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127	Vogt-Koyanagi-Harada disease: Novel insights into pathophysiology, diagnosis and treatment. Progress in Retinal and Eye Research, 2016, 52, 84-111.	7.3	168
128	Investigation of the association of Vogtâ€“Koyanagiâ€“Harada syndrome with IL23R-C1orf141 in Han Chinese Singaporean and ADO-ZNF365-EGR2 in Thai. British Journal of Ophthalmology, 2016, 100, 436-442.	2.1	12
129	Higher Expression of NOD1 and NOD2 is Associated with Vogt-Koyanagi-Harada (VKH) Syndrome But Not Behcet's Disease (BD). Current Molecular Medicine, 2016, 16, 424-435.	0.6	13
130	Association of a NOS3 gene polymorphism with BehÃ§et's disease but not with Vogt-Koyanagi-Harada syndrome in Han Chinese. Molecular Vision, 2016, 22, 311-8.	1.1	5
131	Copy Number Variants and Genetic Polymorphisms in TBX21, GATA3, Rorc, Foxp3 and Susceptibility to Behcet's Disease and Vogt-Koyanagi-Harada Syndrome. Scientific Reports, 2015, 5, 9511.	1.6	14
132	Copy number variations and gene polymorphisms of Complement components in ocular Behcetâ€™s disease and Vogt-Koyanagi-Harada syndrome. Scientific Reports, 2015, 5, 12989.	1.6	9
133	FASGene Copy Numbers are Associated with Susceptibility to BehÃ§et Disease and VKH Syndrome in Han Chinese. Human Mutation, 2015, 36, 1064-1069.	1.1	15
134	Whole exome sequencing confirms the clinical diagnosis of Marfan syndrome combined with X-linked hypophosphatemia. Journal of Translational Medicine, 2015, 13, 179.	1.8	5
135	Association ofERAP1Gene Polymorphisms With BehÃ§et's Disease in Han Chinese. , 2015, 56, 6029.		22
136	Shock wave treatment enhances endothelial proliferation via autocrine vascular endothelial growth factor. Genetics and Molecular Research, 2015, 14, 19203-19210.	0.3	18
137	Association ofATG5Gene Polymorphisms With BehÃ§et's Disease andATG10Gene Polymorphisms With VKH Syndrome in a Chinese Han Population. , 2015, 56, 8280.		24
138	Editorial (Thematic Issue: Uveitis: Pathology, Molecular Mechanisms and Therapy). Current Molecular Medicine, 2015, 15, 510-510.	0.6	4
139	Genetic Variations of IL17F and IL23A ShowÂAssociations with BehÃ§etâ€™s Disease andÂVogt-Koyanagi-Harada Syndrome. Ophthalmology, 2015, 122, 518-523.	2.5	40
140	Association Between Copy Number Variations of TLR7 and Ocular Behcet's Disease in a Chinese Han Population. Investigative Ophthalmology and Visual Science, 2015, 56, 1517-1523.	3.3	18
141	A variant of CLEC16A gene confers protection for Vogtâ€“Koyanagiâ€“Harada syndrome but not for Behcet's disease in a Chinese Han population. Experimental Eye Research, 2015, 132, 225-230.	1.2	6
142	No association between Bach2 gene polymorphisms with Vogtâ€“Koyanagiâ€“Harada syndrome (VKH) and Behcet's disease (BD) in a Chinese Han population. British Journal of Ophthalmology, 2015, 99, 1150-1154.	2.1	1
143	Molecular Genetic Advances in Uveitis. Progress in Molecular Biology and Translational Science, 2015, 134, 283-298.	0.9	28
144	Decreased Interleukin-37 Expression in Vogt-Koyanagi-Harada Disease and Upregulation Following Immunosuppressive Treatment. Journal of Interferon and Cytokine Research, 2015, 35, 265-272.	0.5	19

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145	TLR3 and TLR4 But not TLR2 are Involved in Vogt-Koyanagi- Harada Disease by Triggering Proinflammatory Cytokines Production Through Promoting the Production of Mitochondrial Reactive Oxygen Species. Current Molecular Medicine, 2015, 15, 529-542.	0.6	15
146	Interleukin-10 gene polymorphisms are associated with Behcet's disease but not with Vogt-Koyanagi-Harada syndrome in the Chinese Han population. Molecular Vision, 2015, 21, 589-603.	1.1	25
147	TRAF5 and TRAF3IP2 Gene Polymorphisms Are Associated with BehÃ§et's Disease and Vogt-Koyanagi-Harada Syndrome: A Case-Control Study. PLoS ONE, 2014, 9, e84214.	1.1	22
148	The Role of Interleukin-1 Receptor-Associated Kinases in Vogt-Koyanagi-Harada Disease. PLoS ONE, 2014, 9, e93214.	1.1	11
149	Association of a TNIP1 Polymorphism with Vogt-Koyanagi-Harada Syndrome but Not with Ocular Behcetâ€™s Disease in Han Chinese. PLoS ONE, 2014, 9, e95573.	1.1	7
150	A Functional Variant of PTPN22 Confers Risk for Vogt-Koyanagi-Harada Syndrome but Not for Ankylosing Spondylitis. PLoS ONE, 2014, 9, e96943.	1.1	19
151	Genetic Variations of IL-12B, IL-12RÎ²1, IL-12RÎ²2 in Behcet's Disease and VKH Syndrome. PLoS ONE, 2014, 9, e98373.	1.1	21
152	Activation of Liver X Receptor Alleviates Ocular Inflammation in Experimental Autoimmune Uveitis. , 2014, 55, 2795.		40
153	Inhibition of Proinflammatory Cytokine by IL-25 in Vogt-Koyanagi-Harada Syndrome. Ocular Immunology and Inflammation, 2014, 22, 294-299.	1.0	8
154	FoxO1 Gene Confers Genetic Predisposition to Acute Anterior Uveitis With Ankylosing Spondylitis. Investigative Ophthalmology and Visual Science, 2014, 55, 7970-7974.	3.3	29
155	High C4 gene copy numbers protects against Vogt-Koyanagi-Harada syndrome in Chinese Han. British Journal of Ophthalmology, 2014, 98, 1733-1737.	2.1	21
156	Activation of the aryl hydrocarbon receptor affects activation and function of human monocyte-derived dendritic cells. Clinical and Experimental Immunology, 2014, 177, 521-530.	1.1	66
157	MicroRNA-146aandEts-1gene polymorphisms in ocular BehÃ§et's disease and Vogtâ€“Koyanagiâ€“Harada syndrome. Annals of the Rheumatic Diseases, 2014, 73, 170-176.	0.5	77
158	Increased Notch pathway activation in BehÃ§etâ€™s disease. Rheumatology, 2014, 53, 810-820.	0.9	33
159	Polymorphisms in Genetics of Vitamin D Metabolism Confer Susceptibility to Ocular BehÃ§et Disease in a Chinese Han Population. American Journal of Ophthalmology, 2014, 157, 488-494.e6.	1.7	18
160	Genome-wide association analysis of Vogt-Koyanagi-Harada syndrome identifies two new susceptibility loci at 1p31.2 and 10q21.3. Nature Genetics, 2014, 46, 1007-1011.	9.4	88
161	Decreased interleukin 27 expression is associated with active uveitis in BehÃ§etâ€™s disease. Arthritis Research and Therapy, 2014, 16, R117.	1.6	36
162	Predisposition to BehÃ§etâ€™s disease and VKH syndrome by genetic variants of miR-182. Journal of Molecular Medicine, 2014, 92, 961-967.	1.7	56

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164	A Possible Role for Interleukin 37 in the Pathogenesis of Behcet's Disease. Current Molecular Medicine, 2014, 14, 535-542.	0.6	53
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