Gangxiang Yuan

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/5007555/publications.pdf

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14	197	1306789 7	1372195 10
papers	citations	h-index	g-index

14	14	14	345
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	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
2	Hypermethylation of Interferon Regulatory Factor 8 (IRF8) Confers Risk to Vogt-Koyanagi-Harada Disease. Scientific Reports, 2017, 7, 1007.	1.6	23
3	Genetic polymorphisms of cell adhesion molecules in Behcetâ€™s disease in a Chinese Han population. Scientific Reports, 2016, 6, 24974.	1.6	21
4	Epigenome-wide association study identifies BehÃ§etâ€™s disease-associated methylation loci in Han Chinese. Rheumatology, 2019, 58, 1574-1584.	0.9	21
5	Association of Genetic Variations inTNFSF15With Acute Anterior Uveitis in Chinese Han. , 2015, 56, 4605.		18
6	Promoter Hypermethylation of GATA3, IL-4, and TGF-Î² Confers Susceptibility to Vogt-Koyanagi-Harada Disease in Han Chinese. , 2017, 58, 1529.		18
7	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
8	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
9	Disabled-2 (DAB2) Overexpression Inhibits Monocyte-Derived Dendritic Cells' Function in Vogt-Koyanagi-Harada Disease. , 2018, 59, 4662.		10
10	Genetic polymorphisms of C-type lectin receptors in Behcetâ€™s disease in a Chinese Han population. Scientific Reports, 2017, 7, 5348.	1.6	9
11	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
12	Interleukin-17 induces angiogenesis inÃ¯Â¿Â½vitro via CXCL8 and CCL2 in retinal pigment epithelium. Molecular Medicine Reports, 2018, 17, 4627-4632.	1.1	6
13	Different Methylation of CpG-SNPs in Behcetâ€™s Disease. BioMed Research International, 2019, 2019, 1-7.	0.9	3
14	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