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List of Publications by Year
in descending order

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29	804	759233 12	552781 26
papers	citations	h-index	g-index

30	30	30	1404
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Genome-wide association study identifies five new susceptibility loci for primary angle closure glaucoma. Nature Genetics, 2016, 48, 556-562.	21.4	147
2	Genetic association study of exfoliation syndrome identifies a protective rare variant at LOXL1 and five new susceptibility loci. Nature Genetics, 2017, 49, 993-1004.	21.4	114
3	Three susceptible loci associated with primary open-angle glaucoma identified by genome-wide association study in a Japanese population. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 12838-12842.	7.1	106
4	A common variant mapping to CACNA1A is associated with susceptibility to exfoliation syndrome. Nature Genetics, 2015, 47, 387-392.	21.4	97
5	Common Variants in CDKN2B-AS1 Associated with Optic-Nerve Vulnerability of Glaucoma Identified by Genome-Wide Association Studies in Japanese. PLoS ONE, 2012, 7, e33389.	2.5	88
6	Aberrant astrocyte Ca²⁺ signals â€œAxCa signalsâ€•exacerbate pathological alterations in an Alexander disease model. Glia, 2018, 66, 1053-1067.	4.9	24
7	Differential expression of individual transcript variants of PD-1 and PD-L2 genes on Th-1/Th-2 status is guaranteed for prognosis prediction in PCNSL. Scientific Reports, 2019, 9, 10004.	3.3	24
8	Novel common variants and susceptible haplotype for exfoliation glaucoma specific to Asian population. Scientific Reports, 2015, 4, 5340.	3.3	23
9	Stageâ€specific reference genes significant for quantitative <scp>PCR</scp> during mouse retinal development. Genes To Cells, 2015, 20, 625-635.	1.2	19
10	C/EBPÎ² is a critical mediator of IFN-Î±â€“induced exhaustion of chronic myeloid leukemia stem cells. Blood Advances, 2019, 3, 476-488.	5.2	17
11	Mitochondria as a Platform for Dictating the Cell Fate of Cultured Human Corneal Endothelial Cells. , 2020, 61, 10.		16
12	Association of Rare CYP39A1 Variants With Exfoliation Syndrome Involving the Anterior Chamber of the Eye. JAMA - Journal of the American Medical Association, 2021, 325, 753.	7.4	16
13	Effect of Trinucleotide Repeat Expansion on the Expression ofTCF4mRNA in Fuchs' Endothelial Corneal Dystrophy. , 2019, 60, 779.		14
14	Transcriptome dataset of human corneal endothelium based on ribosomal RNA-depleted RNA-Seq data. Scientific Data, 2020, 7, 407.	5.3	14
15	High-risk follicular lymphomas harbour more somatic mutations including those in the AID-motif. Scientific Reports, 2017, 7, 14039.	3.3	13
16	Efficient and reliable establishment of lymphoblastoid cell lines by Epstein-Barr virus transformation from a limited amount of peripheral blood. Scientific Reports, 2017, 7, 43833.	3.3	12
17	Association of rs613872 and Trinucleotide Repeat Expansion in the TCF4 Gene of German Patients With Fuchs Endothelial Corneal Dystrophy. Cornea, 2019, 38, 799-805.	1.7	12
18	Detection of novel and recurrent conjoined genes in non-Hodgkin B-cell lymphoma. Journal of Clinical and Experimental Hematopathology: JCEH, 2021, 61, 71-77.	0.8	11

#	Article	IF	Citations
19	Towards genomic database of Alexander disease to identify variations modifying disease phenotype. Scientific Reports, 2019, 9, 14763.	3.3	10
20	Trinucleotide repeat expansion in the transcription factor 4 (TCF4) gene in Thai patients with Fuchs endothelial corneal dystrophy. Eye, 2020, 34, 880-885.	2.1	7
21	Adult-onset leukoencephalopathy with homozygous LAMB1 missense mutation. Neurology: Genetics, 2020, 6, e442.	1.9	4
22	Seasonal Variation and Trend of Intraocular Pressure Decrease Over a 20-Year Period in Normal-Tension Glaucoma Patients. American Journal of Ophthalmology, 2022, 234, 235-240.	3.3	4
23	Association of the CYP39A1 G204E Genetic Variant with Increased Risk of Glaucoma and Blindness in Patients with Exfoliation Syndrome. Ophthalmology, 2022, 129, 406-413.	5.2	4
24	Longitudinal seasonal variations of intraocular pressure in primary openâ€angle glaucoma patients as revealed by realâ€world data. Acta Ophthalmologica, 2020, 98, e657.	1.1	2
25	Expression of activated B-cell gene signature is predictive of the outcome of follicular lymphoma. Blood Advances, 2022, 6, 1932-1936.	5.2	2
26	Association studies getting broader: A commentary on A polymorphism of the POLG2 gene is genetically associated with the invasiveness of urinary bladder cancer in Japanese males. Journal of Human Genetics, 2011, 56, 550-551.	2.3	1
27	Downregulation of EGFR in a metastatic brain lesion of EGFR-mutated non-small cell lung cancer using a tyrosine kinase inhibitor: A case report. Oncology Letters, 2017, 13, 2085-2088.	1.8	1
28	C/EBPÎ² Is a Critical Regulator of CML Stem Cell Differentiation and Exhaustion Induced By Interferon-Î±. Blood, 2016, 128, 1120-1120.	1.4	1
29	More Somatic Mutations Including Those in the Aid-Motif with High-Risk, Histologically Non-Transformed Follicular Lymphomas. Blood, 2016, 128, 4116-4116.	1.4	0