

Masakazu Nakano

List of Publications by Year in descending order

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Version: 2024-02-01

29
papers

804
citations

759233

12
h-index

552781

26
g-index

30
all docs

30
docs citations

30
times ranked

1404
citing authors

#	ARTICLE	IF	CITATIONS
1	Seasonal Variation and Trend of Intraocular Pressure Decrease Over a 20-Year Period in Normal-Tension Glaucoma Patients. <i>American Journal of Ophthalmology</i> , 2022, 234, 235-240.	3.3	4
2	Association of the CYP39A1 G204E Genetic Variant with Increased Risk of Glaucoma and Blindness in Patients with Exfoliation Syndrome. <i>Ophthalmology</i> , 2022, 129, 406-413.	5.2	4
3	Expression of activated B-cell gene signature is predictive of the outcome of follicular lymphoma. <i>Blood Advances</i> , 2022, 6, 1932-1936.	5.2	2
4	Association of Rare <i>CYP39A1</i> Variants With Exfoliation Syndrome Involving the Anterior Chamber of the Eye. <i>JAMA - Journal of the American Medical Association</i> , 2021, 325, 753.	7.4	16
5	Detection of novel and recurrent conjoined genes in non-Hodgkin B-cell lymphoma. <i>Journal of Clinical and Experimental Hematopathology: JCEH</i> , 2021, 61, 71-77.	0.8	11
6	Trinucleotide repeat expansion in the transcription factor 4 (TCF4) gene in Thai patients with Fuchs endothelial corneal dystrophy. <i>Eye</i> , 2020, 34, 880-885.	2.1	7
7	Transcriptome dataset of human corneal endothelium based on ribosomal RNA-depleted RNA-Seq data. <i>Scientific Data</i> , 2020, 7, 407.	5.3	14
8	Mitochondria as a Platform for Dictating the Cell Fate of Cultured Human Corneal Endothelial Cells. , 2020, 61, 10.		16
9	Adult-onset leukoencephalopathy with homozygous LAMB1 missense mutation. <i>Neurology: Genetics</i> , 2020, 6, e442.	1.9	4
10	Longitudinal seasonal variations of intraocular pressure in primary open-angle glaucoma patients as revealed by real-world data. <i>Acta Ophthalmologica</i> , 2020, 98, e657.	1.1	2
11	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. <i>Scientific Reports</i> , 2019, 9, 10004.	3.3	24
12	Towards genomic database of Alexander disease to identify variations modifying disease phenotype. <i>Scientific Reports</i> , 2019, 9, 14763.	3.3	10
13	Effect of Trinucleotide Repeat Expansion on the Expression of TCF4 mRNA in Fuchs' Endothelial Corneal Dystrophy. , 2019, 60, 779.		14
14	C/EBP β is a critical mediator of IFN- γ -induced exhaustion of chronic myeloid leukemia stem cells. <i>Blood Advances</i> , 2019, 3, 476-488.	5.2	17
15	Association of rs613872 and Trinucleotide Repeat Expansion in the TCF4 Gene of German Patients With Fuchs Endothelial Corneal Dystrophy. <i>Cornea</i> , 2019, 38, 799-805.	1.7	12
16	Aberrant astrocyte Ca ²⁺ signals exacerbate pathological alterations in an Alexander disease model. <i>Glia</i> , 2018, 66, 1053-1067.	4.9	24
17	Downregulation of EGFR in a metastatic brain lesion of EGFR-mutated non-small cell lung cancer using a tyrosine kinase inhibitor: A case report. <i>Oncology Letters</i> , 2017, 13, 2085-2088.	1.8	1
18	Genetic association study of exfoliation syndrome identifies a protective rare variant at LOXL1 and five new susceptibility loci. <i>Nature Genetics</i> , 2017, 49, 993-1004.	21.4	114

#	ARTICLE	IF	CITATIONS
19	High-risk follicular lymphomas harbour more somatic mutations including those in the AID-motif. Scientific Reports, 2017, 7, 14039.	3.3	13
20	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
21	Genome-wide association study identifies five new susceptibility loci for primary angle closure glaucoma. Nature Genetics, 2016, 48, 556-562.	21.4	147
22	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
23	C/EBP β Is a Critical Regulator of CML Stem Cell Differentiation and Exhaustion Induced By Interferon- γ . Blood, 2016, 128, 1120-1120.	1.4	1
24	Novel common variants and susceptible haplotype for exfoliation glaucoma specific to Asian population. Scientific Reports, 2015, 4, 5340.	3.3	23
25	A common variant mapping to CACNA1A is associated with susceptibility to exfoliation syndrome. Nature Genetics, 2015, 47, 387-392.	21.4	97
26	Stage-specific reference genes significant for quantitative PCR during mouse retinal development. Genes To Cells, 2015, 20, 625-635.	1.2	19
27	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
28	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
29	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