

Jonas Carlsson AlmlÃ¶f

List of Publications by Year in descending order

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Version: 2024-02-01

17
papers

3,067
citations

686830

13
h-index

887659

17
g-index

17
all docs

17
docs citations

17
times ranked

8049
citing authors

#	ARTICLE	IF	CITATIONS
1	Transcriptome and genome sequencing uncovers functional variation in humans. <i>Nature</i> , 2013, 501, 506-511.	13.7	1,857
2	Transancestral mapping and genetic load in systemic lupus erythematosus. <i>Nature Communications</i> , 2017, 8, 16021.	5.8	314
3	Reproducibility of high-throughput mRNA and small RNA sequencing across laboratories. <i>Nature Biotechnology</i> , 2013, 31, 1015-1022.	9.4	251
4	Genome-wide DNA methylation analysis in multiple tissues in primary Sjögren's syndrome reveals regulatory effects at interferon-induced genes. <i>Annals of the Rheumatic Diseases</i> , 2016, 75, 2029-2036.	0.5	180
5	DNA methylation mapping identifies gene regulatory effects in patients with systemic lupus erythematosus. <i>Annals of the Rheumatic Diseases</i> , 2018, 77, 736-743.	0.5	135
6	Whole-genome sequencing identifies complex contributions to genetic risk by variants in genes causing monogenic systemic lupus erythematosus. <i>Human Genetics</i> , 2019, 138, 141-150.	1.8	63
7	Novel risk genes for systemic lupus erythematosus predicted by random forest classification. <i>Scientific Reports</i> , 2017, 7, 6236.	1.6	54
8	Shared and Unique Patterns of DNA Methylation in Systemic Lupus Erythematosus and Primary Sjögren's Syndrome. <i>Frontiers in Immunology</i> , 2019, 10, 1686.	2.2	39
9	Powerful Identification of Cis-regulatory SNPs in Human Primary Monocytes Using Allele-Specific Gene Expression. <i>PLoS ONE</i> , 2012, 7, e52260.	1.1	36
10	Deep targeted sequencing in pediatric acute lymphoblastic leukemia unveils distinct mutational patterns between genetic subtypes and novel relapse-associated genes. <i>Oncotarget</i> , 2016, 7, 64071-64088.	0.8	36
11	CopyNumber450kCancer: baseline correction for accurate copy number calling from the 450k methylation array. <i>Bioinformatics</i> , 2016, 32, 1080-1082.	1.8	31
12	Allelic expression mapping across cellular lineages to establish impact of non-coding <sc>SNPs</sc>s. <i>Molecular Systems Biology</i> , 2014, 10, 754.	3.2	21
13	Basic Tilted Helix Bundle – A new protein fold in human FKBP25/FKBP3 and HectD1. <i>Biochemical and Biophysical Research Communications</i> , 2014, 447, 26-31.	1.0	14
14	Function of multiple sclerosis-protective HLA class I alleles revealed by genome-wide protein-quantitative trait loci mapping of interferon signalling. <i>PLoS Genetics</i> , 2020, 16, e1009199.	1.5	12
15	Variants in BANK1 are associated with lupus nephritis of European ancestry. <i>Genes and Immunity</i> , 2021, 22, 194-202.	2.2	9
16	Single Nucleotide Polymorphisms with Cis-Regulatory Effects on Long Non-Coding Transcripts in Human Primary Monocytes. <i>PLoS ONE</i> , 2014, 9, e102612.	1.1	9
17	Contributions of de novo variants to systemic lupus erythematosus. <i>European Journal of Human Genetics</i> , 2021, 29, 184-193.	1.4	6