

Philipp Maass

List of Publications by Year in descending order

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Version: 2024-02-01

19
papers

1,570
citations

516215

16
h-index

752256

20
g-index

23
all docs

23
docs citations

23
times ranked

3071
citing authors

#	ARTICLE	IF	CITATIONS
1	Whole genome sequencing delineates regulatory, copy number, and cryptic splice variants in early onset cardiomyopathy. <i>Npj Genomic Medicine</i> , 2022, 7, 18.	1.7	14
2	Phosphodiesterase 3A and Arterial Hypertension. <i>Circulation</i> , 2020, 142, 133-149.	1.6	35
3	High-throughput functional analysis of lncRNA core promoters elucidates rules governing tissue specificity. <i>Genome Research</i> , 2019, 29, 344-355.	2.4	100
4	Interchromosomal interactions: A genomic love story of kissing chromosomes. <i>Journal of Cell Biology</i> , 2019, 218, 27-38.	2.3	98
5	Inter-chromosomal Contact Properties in Live-Cell Imaging and in Hi-C. <i>Molecular Cell</i> , 2018, 69, 1039-1045.e3.	4.5	60
6	A TAD boundary is preserved upon deletion of the CTCF-rich Firre locus. <i>Nature Communications</i> , 2018, 9, 1444.	5.8	97
7	High-throughput identification of <i>scn</i> nuclear enrichment sequences. <i>EMBO Journal</i> , 2018, 37, .	3.5	99
8	Spatiotemporal allele organization by allele-specific CRISPR live-cell imaging (SNP-CLING). <i>Nature Structural and Molecular Biology</i> , 2018, 25, 176-184.	3.6	75
9	Reorganization of <i>inter</i> chromosomal interactions in the 2q37 deletion syndrome. <i>EMBO Journal</i> , 2018, 37, .	3.5	13
10	Gene silencing and a novel monoallelic expression pattern in distinct CD177 neutrophil subsets. <i>Journal of Experimental Medicine</i> , 2017, 214, 2089-2101.	4.2	53
11	A map of human circular RNAs in clinically relevant tissues. <i>Journal of Molecular Medicine</i> , 2017, 95, 1179-1189.	1.7	286
12	PDE3A mutations cause autosomal dominant hypertension with brachydactyly. <i>Nature Genetics</i> , 2015, 47, 647-653.	9.4	146
13	Clinical Effects of Phosphodiesterase 3A Mutations in Inherited Hypertension With Brachydactyly. <i>Hypertension</i> , 2015, 66, 800-808.	1.3	39
14	Long non-coding RNA in health and disease. <i>Journal of Molecular Medicine</i> , 2014, 92, 337-346.	1.7	221
15	A misplaced lncRNA causes brachydactyly in humans. <i>Journal of Clinical Investigation</i> , 2012, 122, 3990-4002.	3.9	108
16	A cis-regulatory site downregulates PTHLH in translocation t(8;12)(q13;p11.2) and leads to Brachydactyly Type E. <i>Human Molecular Genetics</i> , 2010, 19, 848-860.	1.4	67
17	Childhood Hypertension in Autosomal-Dominant Hypertension With Brachydactyly. <i>Hypertension</i> , 2010, 56, 988-994.	1.3	17
18	Inversion Region for Hypertension and Brachydactyly on Chromosome 12p Features Multiple Splicing and Noncoding RNA. <i>Hypertension</i> , 2008, 51, 426-431.	1.3	25

#	ARTICLE	IF	CITATIONS
19	A gene expression analysis in rat kidney following high and low salt intake. Journal of Hypertension, 2002, 20, 1115-1120.	0.3	11