Philipp Maass

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/3708817/publications.pdf

Version: 2024-02-01

19	1,570	516215	752256
papers	citations	16	20
		h-index	g-index

23	23	23	3071
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	A map of human circular RNAs in clinically relevant tissues. Journal of Molecular Medicine, 2017, 95, 1179-1189.	1.7	286
2	Long non-coding RNA in health and disease. Journal of Molecular Medicine, 2014, 92, 337-346.	1.7	221
3	PDE3A mutations cause autosomal dominant hypertension with brachydactyly. Nature Genetics, 2015, 47, 647-653.	9.4	146
4	A misplaced lncRNA causes brachydactyly in humans. Journal of Clinical Investigation, 2012, 122, 3990-4002.	3.9	108
5	High-throughput functional analysis of lncRNA core promoters elucidates rules governing tissue specificity. Genome Research, 2019, 29, 344-355.	2.4	100
6	Highâ€throughput identification of <scp>RNA</scp> nuclear enrichment sequences. EMBO Journal, 2018, 37, .	3.5	99
7	Interchromosomal interactions: A genomic love story of kissing chromosomes. Journal of Cell Biology, 2019, 218, 27-38.	2.3	98
8	A TAD boundary is preserved upon deletion of the CTCF-rich Firre locus. Nature Communications, 2018, 9, 1444.	5.8	97
9	Spatiotemporal allele organization by allele-specific CRISPR live-cell imaging (SNP-CLING). Nature Structural and Molecular Biology, 2018, 25, 176-184.	3.6	75
10	A cis-regulatory site downregulates PTHLH in translocation t(8;12)(q13;p11.2) and leads to Brachydactyly Type E. Human Molecular Genetics, 2010, 19, 848-860.	1.4	67
11	Inter-chromosomal Contact Properties in Live-Cell Imaging and in Hi-C. Molecular Cell, 2018, 69, 1039-1045.e3.	4.5	60
12	Gene silencing and a novel monoallelic expression pattern in distinct CD177 neutrophil subsets. Journal of Experimental Medicine, 2017, 214, 2089-2101.	4.2	53
13	Clinical Effects of Phosphodiesterase 3A Mutations in Inherited Hypertension With Brachydactyly. Hypertension, 2015, 66, 800-808.	1.3	39
14	Phosphodiesterase 3A and Arterial Hypertension. Circulation, 2020, 142, 133-149.	1.6	35
15	Inversion Region for Hypertension and Brachydactyly on Chromosome 12p Features Multiple Splicing and Noncoding RNA. Hypertension, 2008, 51, 426-431.	1.3	25
16	Childhood Hypertension in Autosomal-Dominant Hypertension With Brachydactyly. Hypertension, 2010, 56, 988-994.	1.3	17
17	Whole genome sequencing delineates regulatory, copy number, and cryptic splice variants in early onset cardiomyopathy. Npj Genomic Medicine, 2022, 7, 18.	1.7	14
18	Reorganization of interâ€ chromosomal interactions in the 2q37â€deletion syndrome. EMBO Journal, 2018, 37, .	3.5	13

#	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