

Iouri Chepelev

List of Publications by Year in descending order

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Version: 2024-02-01

25
papers

8,051
citations

393982

19
h-index

580395

25
g-index

25
all docs

25
docs citations

25
times ranked

13935
citing authors

#	ARTICLE	IF	CITATIONS
1	High-Resolution Profiling of Histone Methylations in the Human Genome. <i>Cell</i> , 2007, 129, 823-837.	13.5	6,036
2	Intragenic DNA methylation modulates alternative splicing by recruiting MeCP2 to promote exon recognition. <i>Cell Research</i> , 2013, 23, 1256-1269.	5.7	489
3	Characterization of genome-wide enhancer-promoter interactions reveals co-expression of interacting genes and modes of higher order chromatin organization. <i>Cell Research</i> , 2012, 22, 490-503.	5.7	238
4	Epigenome Mapping in Normal and Disease States. <i>Circulation Research</i> , 2010, 107, 327-339.	2.0	164
5	Pol II and its associated epigenetic marks are present at Pol III-transcribed noncoding RNA genes. <i>Nature Structural and Molecular Biology</i> , 2010, 17, 629-634.	3.6	161
6	Detection of single nucleotide variations in expressed exons of the human genome using RNA-Seq. <i>Nucleic Acids Research</i> , 2009, 37, e106-e106.	6.5	152
7	Critical role of histone demethylase Jmjd3 in the regulation of CD4+ T-cell differentiation. <i>Nature Communications</i> , 2014, 5, 5780.	5.8	136
8	Dynamic regulation of alternative splicing and chromatin structure in Drosophila gonads revealed by RNA-seq. <i>Cell Research</i> , 2010, 20, 763-783.	5.7	107
9	Cell Fate Determination Factor Dachshund Reprograms Breast Cancer Stem Cell Function. <i>Journal of Biological Chemistry</i> , 2011, 286, 2132-2142.	1.6	74
10	Common inversion polymorphism at 17q21.31 affects expression of multiple genes in tissue-specific manner. <i>BMC Genomics</i> , 2012, 13, 458.	1.2	62
11	Attenuation of Forkhead signaling by the retinal determination factor DACH1. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2010, 107, 6864-6869.	3.3	58
12	CTCF boundary remodels chromatin domain and drives aberrant HOX gene transcription in acute myeloid leukemia. <i>Blood</i> , 2018, 132, 837-848.	0.6	56
13	Mapping of INS promoter interactions reveals its role in long-range regulation of SYT8 transcription. <i>Nature Structural and Molecular Biology</i> , 2011, 18, 372-378.	3.6	55
14	Stage-Dependent and Locus-Specific Role of Histone Demethylase Jumonji D3 (JMJD3) in the Embryonic Stages of Lung Development. <i>PLoS Genetics</i> , 2014, 10, e1004524.	1.5	50
15	Long-distance interactions of D-brane bound states and longitudinal five-brane in M(atr)ix theory. <i>Physical Review D</i> , 1997, 56, 3672-3685.	1.6	32
16	Response: Mapping Nucleosome Positions Using ChIP-Seq Data. <i>Cell</i> , 2007, 131, 832-833.	13.5	32
17	Epstein-Barr virus nuclear antigen 2 extensively rewires the human chromatin landscape at autoimmune risk loci. <i>Genome Research</i> , 2021, 31, 2185-2198.	2.4	24
18	The Effect of Inversion at 8p23 on BLK Association with Lupus in Caucasian Population. <i>PLoS ONE</i> , 2014, 9, e115614.	1.1	23

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19	Alternative splicing switching in stem cell lineages. <i>Frontiers in Biology</i> , 2013, 8, 50-59.	0.7	22
20	Cyclin D1 integrates G9a-mediated histone methylation. <i>Oncogene</i> , 2019, 38, 4232-4249.	2.6	20
21	A Barrier-Only Boundary Element Delimits the Formation of Facultative Heterochromatin in <i>Drosophila melanogaster</i> and Vertebrates. <i>Molecular and Cellular Biology</i> , 2011, 31, 2729-2741.	1.1	19
22	Alteration of CTCF-associated chromatin neighborhood inhibits TAL1-driven oncogenic transcription program and leukemogenesis. <i>Nucleic Acids Research</i> , 2020, 48, 3119-3133.	6.5	19
23	Genome-wide chromatin occupancy of BRDT and gene expression analysis suggest transcriptional partners and specific epigenetic landscapes that regulate gene expression during spermatogenesis. <i>Molecular Reproduction and Development</i> , 2021, 88, 141-157.	1.0	9
24	Tissue-specific expression of IL-15RA alternative splicing transcripts and its regulation by DNA methylation. <i>European Cytokine Network</i> , 2010, 21, 308-18.	1.1	8
25	Validation of low-coverage whole-genome sequencing for mitochondrial DNA variants suggests mitochondrial DNA as a genetic cause of preterm birth. <i>Human Mutation</i> , 2021, 42, 1602-1614.	1.1	5