Alvaro Rada-Iglesias

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Identification and analysis of functional elements in 1% of the human genome by the ENCODE pilot project. Nature, 2007, 447, 799-816.	13.7	4,709
2	A unique chromatin signature uncovers early developmental enhancers in humans. Nature, 2011, 470, 279-283.	13.7	1,949
3	CHD7 cooperates with PBAF to control multipotent neural crest formation. Nature, 2010, 463, 958-962.	13.7	527
4	Epigenomic Annotation of Enhancers Predicts Transcriptional Regulators of Human Neural Crest. Cell Stem Cell, 2012, 11, 633-648.	5.2	283
5	Nucleosomes are well positioned in exons and carry characteristic histone modifications. Genome Research, 2009, 19, 1732-1741.	2.4	274
6	Monte Carlo feature selection for supervised classification. Bioinformatics, 2008, 24, 110-117.	1.8	262
7	PRC2 Facilitates the Regulatory Topology Required for Poised Enhancer Function during Pluripotent Stem Cell Differentiation. Cell Stem Cell, 2017, 20, 689-705.e9.	5.2	198
8	Sequence-specific regulator Prdm14 safeguards mouse ESCs from entering extraembryonic endoderm fates. Nature Structural and Molecular Biology, 2011, 18, 120-127.	3.6	178
9	The homeodomain-leucine zipper (HD-Zip) class I transcription factors ATHB7 and ATHB12 modulate abscisic acid signalling by regulating protein phosphatase 2C and abscisic acid receptor gene activities. Plant Molecular Biology, 2012, 80, 405-418.	2.0	144
10	Butyrate mediates decrease of histone acetylation centered on transcription start sites and down-regulation of associated genes. Genome Research, 2007, 17, 708-719.	2.4	130
11	Modeling the Pathological Long-Range Regulatory Effects of Human Structural Variation with Patient-Specific hiPSCs. Cell Stem Cell, 2019, 24, 736-752.e12.	5.2	90
12	Histone Acetylation and Methylation at Sites Initiating Divergent Polycistronic Transcription in Trypanosoma cruzi. Journal of Biological Chemistry, 2008, 283, 15884-15892.	1.6	89
13	Whole-genome maps of USF1 and USF2 binding and histone H3 acetylation reveal new aspects of promoter structure and candidate genes for common human disorders. Genome Research, 2008, 18, 380-392.	2.4	85
14	Imputation of Orofacial Clefting Data Identifies Novel Risk Loci and Sheds Light on the Genetic Background of Cleft Lip ± Cleft Palate and Cleft Palate Only Human Molecular Genetics, 2017, 26, ddx012.	1.4	84
15	Forces driving the threeâ€dimensional folding of eukaryotic genomes. Molecular Systems Biology, 2018, 14, e8214.	3.2	75
16	Foxd3 Promotes Exit from Naive Pluripotency through Enhancer Decommissioning and Inhibits Germline Specification. Cell Stem Cell, 2016, 18, 118-133.	5.2	73
17	Binding sites for metabolic disease related transcription factors inferred at base pair resolution by chromatin immunoprecipitation and genomic microarrays. Human Molecular Genetics, 2005, 14, 3435-3447.	1.4	71
18	In vitro analysis of DNA-protein interactions by proximity ligation. Proceedings of the National Academy of Sciences of the United States of America, 2007, 104, 3067-3072.	3.3	68

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19	Is H3K4me1 at enhancers correlative or causative?. Nature Genetics, 2018, 50, 4-5.	9.4	67
20	Dosage analysis of the 7q11.23 Williams region identifies <i>BAZ1B</i> as a major human gene patterning the modern human face and underlying self-domestication. Science Advances, 2019, 5, eaaw7908.	4.7	67
21	Orphan CpG islands amplify poised enhancer regulatory activity and determine target gene responsiveness. Nature Genetics, 2021, 53, 1036-1049.	9.4	56
22	Niche stiffening compromises hair follicle stem cell potential during ageing by reducing bivalent promoter accessibility. Nature Cell Biology, 2021, 23, 771-781.	4.6	51
23	The chromatin, topological and regulatory properties of pluripotency-associated poised enhancers are conserved in vivo. Nature Communications, 2021, 12, 4344.	5.8	50
24	Epigenomics of human embryonic stem cells and induced pluripotent stem cells: insights into pluripotency and implications for disease. Genome Medicine, 2011, 3, 36.	3.6	49
25	yylncT Defines a Class of Divergently Transcribed lncRNAs and Safeguards the T-mediated Mesodermal Commitment of Human PSCs. Cell Stem Cell, 2019, 24, 318-327.e8.	5.2	44
26	Epigenomic and Transcriptomic Changes During Human RPE EMT in a Stem Cell Model of Epiretinal Membrane Pathogenesis and Prevention by Nicotinamide. Stem Cell Reports, 2020, 14, 631-647.	2.3	43
27	Identification of candidate regulatory SNPs by combination of transcription-factor-binding site prediction, SNP genotyping and haploChIP. Nucleic Acids Research, 2009, 37, e85-e85.	6.5	34
28	Integrative epigenomic and genomic analysis of malignant pheochromocytoma. Experimental and Molecular Medicine, 2010, 42, 484.	3.2	32
29	Epigenomics-Based Identification of Major Cell Identity Regulators within Heterogeneous Cell Populations. Cell Reports, 2016, 17, 3062-3076.	2.9	29
30	The promoter of inducible nitric oxide synthase implicated in glaucoma based on genetic analysis and nuclear factor binding. Molecular Vision, 2005, 11, 950-7.	1.1	28
31	Histone H3 lysine 27 trimethylation in adult differentiated colon associated to cancer DNA hypermethylation. Epigenetics, 2009, 4, 107-113.	1.3	25
32	Lineage specific transcription factors and epigenetic regulators mediate TGFβ-dependent enhancer activation. Nucleic Acids Research, 2018, 46, 3351-3365.	6.5	24
33	Mechanism suppressing H3K9 trimethylation in pluripotent stem cells and its demise by polyQ-expanded huntingtin mutations. Human Molecular Genetics, 2018, 27, 4117-4134.	1.4	21
34	Enhancer-associated H3K4 methylation safeguards in vitro germline competence. Nature Communications, 2021, 12, 5771.	5.8	20
35	Human genetic variation within neural crest enhancers: molecular and phenotypic implications. Philosophical Transactions of the Royal Society B: Biological Sciences, 2013, 368, 20120360.	1.8	18
36	The ubiquitin-conjugating enzyme UBE2K determines neurogenic potential through histone H3 in human embryonic stem cells. Communications Biology, 2020, 3, 262.	2.0	18

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37	Polycomb proteins as organizers of 3D genome architecture in embryonic stem cells. Briefings in Functional Genomics, 2019, 18, 358-366.	1.3	16
38	Pathological ASXL1 Mutations and Protein Variants Impair Neural Crest Development. Stem Cell Reports, 2019, 12, 861-868.	2.3	16
39	Overarching control of autophagy and DNA damage response by CHD6 revealed by modeling a rare human pathology. Nature Communications, 2021, 12, 3014.	5.8	16
40	The formation of the thumb requires direct modulation of <i>Gli3</i> transcription by Hoxa13. Proceedings of the National Academy of Sciences of the United States of America, 2020, 117, 1090-1096.	3.3	15
41	Enhancer-gene specificity in development and disease. Development (Cambridge), 2022, 149, .	1.2	15
42	Pioneering Barren Land: Mitotic Bookmarking by Transcription Factors. Developmental Cell, 2013, 24, 342-344.	3.1	12
43	Rare or Overlooked? Structural Disruption of Regulatory Domains in Human Neurocristopathies. Frontiers in Genetics, 2020, 11, 688.	1.1	12
44	Transcriptional and epigenetic control of germline competence and specification. Current Opinion in Cell Biology, 2019, 61, 1-8.	2.6	11
45	MAPRE2 mutations result in altered human cranial neural crest migration, underlying craniofacial malformations in CSC-KT syndrome. Scientific Reports, 2021, 11, 4976.	1.6	10
46	Two polypyrimidine tracts in the nitric oxide synthase 2 gene: similar regulatory sequences with different properties. Molecular Biology Reports, 2010, 37, 2021-2030.	1.0	9
47	Integrative approaches generate insights into the architecture of non-syndromic cleft lip ± cleft palate. Human Genetics and Genomics Advances, 2021, 2, 100038.	1.0	8
48	Genetic variation within transcriptional regulatory elements and its implications for human disease. Biological Chemistry, 2014, 395, 1453-1460.	1.2	7
49	JMJD3 intrinsically disordered region links the 3D-genome structure to TGFÎ ² -dependent transcription activation. Nature Communications, 2022, 13, .	5.8	6
50	Chromatin Immunoprecipitation (ChIP) Protocol for Low-abundance Embryonic Samples. Journal of Visualized Experiments, 2017, , .	0.2	4
51	Pioneering of Enhancer Landscapes during Pluripotent State Transitions. Cell Stem Cell, 2018, 23, 149-151.	5.2	2
52	GARLIC: a bioinformatic toolkit for aetiologically connecting diseases and cell type-specific regulatory maps. Human Molecular Genetics, 2016, 26, ddw423.	1.4	1
53	Disruption of the TFAP2A Regulatory Domain Causes Banchio-Oculo-Facial Syndrome (BOFS) and Illuminates Pathomechanisms for Other Human Neurocristopathies. SSRN Electronic Journal, 0, , .	0.4	1
54	Enhancer Remodeling During Early Mammalian Embryogenesis: Lessons for Somatic Reprogramming, Rejuvenation, and Aging. Current Stem Cell Reports, 2016, 2, 263-272.	0.7	0

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55	Ready, Set…Poised!: Polycomb targetÂgenes are bound by poised <scp>RNA</scp> Âpolymerase <scp>II</scp> throughout differentiation. Molecular Systems Biology, 2017, 13, 950.	3.2	0
56	Protocol to study sufficiency of cis-regulatory elements in mouse embryonic stem cells using a CRISPR-mediated knockin approach. STAR Protocols, 2022, 3, 101492.	0.5	0