

Paola Benaglio

List of Publications by Year in descending order

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Version: 2024-02-01

20
papers

1,848
citations

516710

16
h-index

713466

21
g-index

27
all docs

27
docs citations

27
times ranked

5794
citing authors

#	ARTICLE	IF	CITATIONS
1	The Influence of Age and Sex on Genetic Associations with Adult Body Size and Shape: A Large-Scale Genome-Wide Interaction Study. <i>PLoS Genetics</i> , 2015, 11, e1005378.	3.5	331
2	Identification of heart rate-associated loci and their effects on cardiac conduction and rhythm disorders. <i>Nature Genetics</i> , 2013, 45, 621-631.	21.4	282
3	Interpreting type 1 diabetes risk with genetics and single-cell epigenomics. <i>Nature</i> , 2021, 594, 398-402.	27.8	170
4	Large-Scale Profiling Reveals the Influence of Genetic Variation on Gene Expression in Human Induced Pluripotent Stem Cells. <i>Cell Stem Cell</i> , 2017, 20, 533-546.e7.	11.1	157
5	iPSCORE: A Resource of 222 iPSC Lines Enabling Functional Characterization of Genetic Variation across a Variety of Cell Types. <i>Stem Cell Reports</i> , 2017, 8, 1086-1100.	4.8	147
6	Genomewide Association Study Using a High-Density Single Nucleotide Polymorphism Array and Case-Control Design Identifies a Novel Essential Hypertension Susceptibility Locus in the Promoter Region of Endothelial NO Synthase. <i>Hypertension</i> , 2012, 59, 248-255.	2.7	144
7	Subtle changes in chromatin loop contact propensity are associated with differential gene regulation and expression. <i>Nature Communications</i> , 2019, 10, 1054.	12.8	100
8	Systematic analysis of binding of transcription factors to noncoding variants. <i>Nature</i> , 2021, 591, 147-151.	27.8	89
9	Insights into the Mutational Burden of Human Induced Pluripotent Stem Cells from an Integrative Multi-Omics Approach. <i>Cell Reports</i> , 2018, 24, 883-894.	6.4	85
10	Comparative genome analysis of <i>Pseudomonas knackmussii</i> ... <i>B</i> 13, the first bacterium known to degrade chloroaromatic compounds. <i>Environmental Microbiology</i> , 2015, 17, 91-104.	3.8	52
11	Target Sequencing, Cell Experiments, and a Population Study Establish Endothelial Nitric Oxide Synthase (<i>eNOS</i>) Gene as Hypertension Susceptibility Gene. <i>Hypertension</i> , 2013, 62, 844-852.	2.7	48
12	Next generation sequencing of pooled samples reveals new SNRNP200 mutations associated with retinitis pigmentosa. <i>Human Mutation</i> , 2011, 32, E2246-E2258.	2.5	42
13	Pgltools: a genomic arithmetic tool suite for manipulation of Hi-C peak and other chromatin interaction data. <i>BMC Bioinformatics</i> , 2017, 18, 207.	2.6	35
14	Allele-specific NKX2-5 binding underlies multiple genetic associations with human electrocardiographic traits. <i>Nature Genetics</i> , 2019, 51, 1506-1517.	21.4	35
15	Sequencing and characterizing the genome of <i>Estrella lausannensis</i> as an undergraduate project: training students and biological insights. <i>Frontiers in Microbiology</i> , 2015, 6, 101.	3.5	32
16	Human iPSC-Derived Retinal Pigment Epithelium: A Model System for Prioritizing and Functionally Characterizing Causal Variants at AMD Risk Loci. <i>Stem Cell Reports</i> , 2019, 12, 1342-1353.	4.8	32
17	Glucocorticoid signaling in pancreatic islets modulates gene regulatory programs and genetic risk of type 2 diabetes. <i>PLoS Genetics</i> , 2021, 17, e1009531.	3.5	13
18	Ultra High Throughput Sequencing in Human DNA Variation Detection: A Comparative Study on the NDUF3-PRPF31 Region. <i>PLoS ONE</i> , 2010, 5, e13071.	2.5	11

#	ARTICLE	IF	CITATIONS
19	Mutational screening of splicing factor genes in cases with autosomal dominant retinitis pigmentosa. <i>Molecular Vision</i> , 2014, 20, 843-51.	1.1	11
20	FOXO1 mitigates the SMAD3/FOXL2C134W transcriptomic effect in a model of human adult granulosa cell tumor. <i>Journal of Translational Medicine</i> , 2021, 19, 90.	4.4	5