

Ivana Persico

List of Publications by Year in descending order

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Version: 2024-02-01

28
papers

1,650
citations

687363

13
h-index

552781

26
g-index

29
all docs

29
docs citations

29
times ranked

4867
citing authors

#	ARTICLE	IF	CITATIONS
1	Crisponi/coldâ€induced sweating syndrome: Differential diagnosis, pathogenesis and treatment concepts. <i>Clinical Genetics</i> , 2020, 97, 209-221.	2.0	12
2	Infant developmental profile of Crisponi syndrome due to compound heterozygosity for CRLF1 deletion. <i>Clinical Dysmorphology</i> , 2020, 29, 141-143.	0.3	11
3	Crisponi/Cold Induced Sweating Syndrome Type 1 With a Private Cytokine Receptor Like Factor 1 (CRLF1) Mutation in an Indian Family. <i>Indian Pediatrics</i> , 2020, 57, 1075-1077.	0.4	0
4	Landscape of transcriptome variations uncovering known and novel driver events in colorectal carcinoma. <i>Scientific Reports</i> , 2020, 10, 432.	3.3	16
5	Crisponi/Cold Induced Sweating Syndrome Type 1 With a Private Cytokine Receptor Like Factor 1 (CRLF1) Mutation in an Indian Family. <i>Indian Pediatrics</i> , 2020, 57, 1075-1076.	0.4	0
6	Exome sequencing in Crisponi/coldâ€induced sweating syndromeâ€like individuals reveals unpredicted alternative diagnoses. <i>Clinical Genetics</i> , 2019, 95, 607-614.	2.0	7
7	Confirmation of a new phenotype in an individual with a variant in the last part of exon 30 of <i>CREBBP</i> . <i>American Journal of Medical Genetics, Part A</i> , 2019, 179, 634-638.	1.2	13
8	Novel ANKRD11 gene mutation in an individual with a mild phenotype of KBG syndrome associated to a GEFS+ phenotypic spectrum: a case report. <i>BMC Medical Genetics</i> , 2019, 20, 16.	2.1	14
9	Novel <i>NALCN</i> biallelic truncating mutations in siblings with IHPRF1 syndrome. <i>Clinical Genetics</i> , 2018, 93, 1245-1247.	2.0	14
10	A new case series of Crisponi syndrome in a Turkish family and review of the literature. <i>Clinical Dysmorphology</i> , 2017, 26, 66-72.	0.3	5
11	Molecular Characterization of β^2 -Thalassemia Mutations in Central Vietnam. <i>Hemoglobin</i> , 2017, 41, 96-99.	0.8	7
12	Bi-allelic Mutations in KLHL7 Cause a Crisponi/CISS1-like Phenotype Associated with Early-Onset Retinitis Pigmentosa. <i>American Journal of Human Genetics</i> , 2016, 99, 236-245.	6.2	28
13	Genetic associations at 53 loci highlight cell types and biological pathways relevant for kidney function. <i>Nature Communications</i> , 2016, 7, 10023.	12.8	412
14	Genome-wide association analyses identify 18 new loci associated with serum urate concentrations. <i>Nature Genetics</i> , 2013, 45, 145-154.	21.4	675
15	A strategy analysis for genetic association studies with known inbreeding. <i>BMC Genetics</i> , 2011, 12, 63.	2.7	8
16	Application of a New Method for GWAS in a Related Case/Control Sample with Known Pedigree Structure: Identification of New Loci for Nephrolithiasis. <i>PLoS Genetics</i> , 2011, 7, e1001281.	3.5	10
17	High Differentiation among Eight Villages in a Secluded Area of Sardinia Revealed by Genome-Wide High Density SNPs Analysis. <i>PLoS ONE</i> , 2009, 4, e4654.	2.5	30
18	Microsatellites and SNPs linkage analysis in a Sardinian genetic isolate confirms several essential hypertension loci previously identified in different populations. <i>BMC Medical Genetics</i> , 2009, 10, 81.	2.1	8

#	ARTICLE	IF	CITATIONS
19	EDA2R Is Associated with Androgenetic Alopecia. <i>Journal of Investigative Dermatology</i> , 2008, 128, 2268-2270.	0.7	79
20	Patterns of Linkage Disequilibrium between SNPs in a Sardinian Population Isolate and the Selection of Markers for Association Studies. <i>Human Heredity</i> , 2008, 65, 9-22.	0.8	14
21	Methylenetetrahydrofolate reductase gene polymorphisms in Burkina Faso. <i>Clinica Chimica Acta</i> , 2005, 360, 199-200.	1.1	0
22	A Genomewide Search Using an Original Pairwise Sampling Approach for Large Genealogies Identifies a New Locus for Total and Low-Density Lipoprotein Cholesterol in Two Genetically Differentiated Isolates of Sardinia. <i>American Journal of Human Genetics</i> , 2004, 75, 1015-1031.	6.2	48
23	A 24-bp duplication in exon 10 of human chitotriosidase gene from the sub-Saharan to the Mediterranean area: role of parasitic diseases and environmental conditions. <i>Genes and Immunity</i> , 2003, 4, 570-574.	4.1	88
24	Extent of linkage disequilibrium in a Sardinian sub-isolate: sampling and methodological considerations. <i>Human Molecular Genetics</i> , 2003, 13, 25-33.	2.9	42
25	BRCA1 and BRCA2 germline mutations in Sardinian breast cancer families and their implications for genetic counseling. <i>Annals of Oncology</i> , 2002, 13, 1899-1907.	1.2	20
26	Not all isolates are equal: linkage disequilibrium analysis on Xq13.3 reveals different patterns in Sardinian sub-populations. <i>Human Genetics</i> , 2002, 111, 9-15.	3.8	39
27	Identification of a founder BRCA2 mutation in Sardinia. <i>British Journal of Cancer</i> , 2000, 82, 553-559.	6.4	42
28	Molecular basis of open-angle glaucoma in Italy. <i>Acta Ophthalmologica</i> , 1998, 76, 16-17.	0.3	2