

Christopher J Cardinale

List of Publications by Year in descending order

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Version: 2024-02-01

22
papers

1,223
citations

687363

13
h-index

794594

19
g-index

23
all docs

23
docs citations

23
times ranked

2755
citing authors

#	ARTICLE	IF	CITATIONS
1	A novel <i>MBTPS2</i> variant associated with BRESHECK syndrome impairs sterol-regulated transcription and the endoplasmic reticulum stress response. American Journal of Medical Genetics, Part A, 2022, 188, 463-472.	1.2	4
2	Genetic Underpinnings of Asthma and Related Traits. , 2020, , 341-360.		0
3	Rare copy number variants in over 100,000 European ancestry subjects reveal multiple disease associations. Nature Communications, 2020, 11, 255.	12.8	48
4	Regulation of Janus Kinase 2 by an Inflammatory Bowel Disease Causal Non-coding Single Nucleotide Polymorphism. Journal of Crohn's and Colitis, 2020, 14, 646-653.	1.3	5
5	Multiple Epistasis Interactions Within MHC Are Associated With Ulcerative Colitis. Frontiers in Genetics, 2019, 10, 257.	2.3	7
6	P027 DIFFERENTIAL BINDING OF RBPJ AND CUX1 TO IBD CAUSAL SNP RS1887428 MODIFIES EXPRESSION OF JAK2. Gastroenterology, 2018, 154, S14-S15.	1.3	0
7	Genetics of Inflammatory Bowel Diseases. , 2017, , 3-14.		1
8	Pathway-based Genome-wide Association Studies Reveal the Association Between Growth Factor Activity and Inflammatory Bowel Disease. Inflammatory Bowel Diseases, 2016, 22, 1540-1551.	1.9	8
9	Association of a rare NOTCH4 coding variant with systemic sclerosis: a family-based whole exome sequencing study. BMC Musculoskeletal Disorders, 2016, 17, 462.	1.9	12
10	CNV Analysis Associates AKNAD1 with Type-2 Diabetes in Jordan Subpopulations. Scientific Reports, 2015, 5, 13391.	3.3	18
11	Genetic sharing and heritability of paediatric age of onset autoimmune diseases. Nature Communications, 2015, 6, 8442.	12.8	58
12	Meta-analysis of shared genetic architecture across ten pediatric autoimmune diseases. Nature Medicine, 2015, 21, 1018-1027.	30.7	212
13	A Novel <i>BHLHE41</i> Variant is Associated with Short Sleep and Resistance to Sleep Deprivation in Humans. Sleep, 2014, 37, 1327-1336.	1.1	104
14	Transcriptome Profiling of Human Ulcerative Colitis Mucosa Reveals Altered Expression of Pathways Enriched in Genetic Susceptibility Loci. PLoS ONE, 2014, 9, e96153.	2.5	8
15	Large Sample Size, Wide Variant Spectrum, and Advanced Machine-Learning Technique Boost Risk Prediction for Inflammatory Bowel Disease. American Journal of Human Genetics, 2013, 92, 1008-1012.	6.2	162
16	Whole-genome DNA/RNA sequencing identifies truncating mutations in RBCK1 in a novel Mendelian disease with neuromuscular and cardiac involvement. Genome Medicine, 2013, 5, 67.	8.2	87
17	Targeted resequencing identifies defective variants of decoy receptor 3 in pediatric-onset inflammatory bowel disease. Genes and Immunity, 2013, 14, 447-452.	4.1	16
18	Extended haplotype association study in Crohn's disease identifies a novel, Ashkenazi Jewish-specific missense mutation in the NF- κ B pathway gene, HEATR3. Genes and Immunity, 2013, 14, 310-316.	4.1	31

#	ARTICLE	IF	CITATIONS
19	Impact of exome sequencing in inflammatory bowel disease. World Journal of Gastroenterology, 2013, 19, 6721.	3.3	20
20	Termination Factor Rho and Its Cofactors NusA and NusG Silence Foreign DNA in <i>E. coli</i> . Science, 2008, 320, 935-938.	12.6	266
21	An Allosteric Path to Transcription Termination. Molecular Cell, 2007, 28, 991-1001.	9.7	114
22	The B Cell SH2/PH Domain-Containing Adaptor Bam32/DAPP1 Is Required for T Cell-Independent II Antigen Responses. Current Biology, 2003, 13, 1858-1866.	3.9	42