

Carmen Oleaga-Quintas

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/5294683/publications.pdf>

Version: 2024-02-01

16
papers

740
citations

758635

12
h-index

887659

17
g-index

17
all docs

17
docs citations

17
times ranked

1452
citing authors

#	ARTICLE	IF	CITATIONS
1	Mendelian susceptibility to mycobacterial disease: 2014–2018 update. <i>Immunology and Cell Biology</i> , 2019, 97, 360-367.	1.0	163
2	Inherited p40phox deficiency differs from classic chronic granulomatous disease. <i>Journal of Clinical Investigation</i> , 2018, 128, 3957-3975.	3.9	99
3	Inherited human IFN- γ deficiency underlies mycobacterial disease. <i>Journal of Clinical Investigation</i> , 2020, 130, 3158-3171.	3.9	89
4	Human T-bet Governs Innate and Innate-like Adaptive IFN- γ Immunity against Mycobacteria. <i>Cell</i> , 2020, 183, 1826-1847.e31.	13.5	83
5	IFN- γ and CD25 drive distinct pathologic features during hemophagocytic lymphohistiocytosis. <i>Journal of Allergy and Clinical Immunology</i> , 2019, 143, 2215-2226.e7.	1.5	49
6	A Variety of Alu-Mediated Copy Number Variations Can Underlie IL-12R β 1 Deficiency. <i>Journal of Clinical Immunology</i> , 2018, 38, 617-627.	2.0	45
7	Laboratory evaluation of the IFN- γ circuit for the molecular diagnosis of Mendelian susceptibility to mycobacterial disease. <i>Critical Reviews in Clinical Laboratory Sciences</i> , 2018, 55, 184-204.	2.7	43
8	Genetic, Immunological, and Clinical Features of 32 Patients with Autosomal Recessive STAT1 Deficiency. <i>Journal of Immunology</i> , 2021, 207, 133-152.	0.4	33
9	Inherited GATA2 Deficiency Is Dominant by Haploinsufficiency and Displays Incomplete Clinical Penetrance. <i>Journal of Clinical Immunology</i> , 2021, 41, 639-657.	2.0	30
10	Lethal Influenza in Two Related Adults with Inherited GATA2 Deficiency. <i>Journal of Clinical Immunology</i> , 2018, 38, 513-526.	2.0	29
11	Patient iPSC-Derived Macrophages to Study Inborn Errors of the IFN- γ Responsive Pathway. <i>Cells</i> , 2020, 9, 483.	1.8	16
12	Infectious diseases, autoimmunity and midline defect in a patient with a novel bi-allelic mutation in IL12RB1 gene. <i>Turkish Journal of Pediatrics</i> , 2016, 58, 331-336.	0.3	15
13	A purely quantitative form of partial recessive IFN- γ R2 deficiency caused by mutations of the initiation or second codon. <i>Human Molecular Genetics</i> , 2018, 27, 3919-3935.	1.4	14
14	Severe Enteropathy and Hypogammaglobulinemia Complicating Refractory Mycobacterium tuberculosis Complex Disseminated Disease in a Child with IL-12R β 1 Deficiency. <i>Journal of Clinical Immunology</i> , 2017, 37, 732-738.	2.0	10
15	Disseminated BCG Infectious Disease and Hyperferritinemia in a Patient With a Novel NEMO Mutation. <i>Journal of Investigational Allergology and Clinical Immunology</i> , 2016, 26, 268-271.	0.6	10
16	LINE-1-Mediated AluYa5 Insertion Underlying Complete Autosomal Recessive IFN- γ R1 Deficiency. <i>Journal of Clinical Immunology</i> , 2019, 39, 739-742.	2.0	5