Carmen Oleaga-Quintas

List of Publications by Year
in descending order

16 papers	740 citations	759233 12 h-index	888059 17 g-index
17 all docs	17 docs citations	17 times ranked	1452 citing authors

#	Article	IF	Citations
1	Mendelian susceptibility to mycobacterial disease: 2014â€“2018 update. Immunology and Cell Biology, 2019, 97, 360-367.	2.3	163
2	Inherited p40phox deficiency differs from classic chronic granulomatous disease. Journal of Clinical Investigation, 2018, 128, 3957-3975.	8.2	99
3	Inherited human IFN-Î³ deficiency underlies mycobacterial disease. Journal of Clinical Investigation, 2020, 130, 3158-3171.	8.2	89
4	Human T-bet Governs Innate and Innate-like Adaptive IFN-Î³ Immunity against Mycobacteria. Cell, 2020, 183, 1826-1847.e31.	28.9	83
5	IFN-Î³ and CD25 drive distinct pathologic features during hemophagocytic lymphohistiocytosis. Journal of Allergy and Clinical Immunology, 2019, 143, 2215-2226.e7.	2.9	49
6	A Variety of Alu-Mediated Copy Number Variations Can Underlie IL-12RÎ²1 Deficiency. Journal of Clinical Immunology, 2018, 38, 617-627.	3.8	45
7	Laboratory evaluation of the IFN-Î³ circuit for the molecular diagnosis of Mendelian susceptibility to mycobacterial disease. Critical Reviews in Clinical Laboratory Sciences, 2018, 55, 184-204.	6.1	43
8	Genetic, Immunological, and Clinical Features of 32 Patients with Autosomal Recessive STAT1 Deficiency. Journal of Immunology, 2021, 207, 133-152.	0.8	33
9	Inherited GATA2 Deficiency Is Dominant by Haploinsufficiency and Displays Incomplete Clinical Penetrance. Journal of Clinical Immunology, 2021, 41, 639-657.	3.8	30
10	Lethal Influenza in Two Related Adults with Inherited GATA2 Deficiency. Journal of Clinical Immunology, 2018, 38, 513-526.	3.8	29
11	Patient iPSC-Derived Macrophages to Study Inborn Errors of the IFN-Î³ Responsive Pathway. Cells, 2020, 9, 483.	4.1	16
12	Infectious diseases, autoimmunity and midline defect in a patient with a novel bi-allelic mutation in IL12RB1 gene. Turkish Journal of Pediatrics, 2016, 58, 331-336.	0.6	15
13	A purely quantitative form of partial recessive IFN-Î³R2 deficiency caused by mutations of the initiation or second codon. Human Molecular Genetics, 2018, 27, 3919-3935.	2.9	14
14	Severe Enteropathy and Hypogammaglobulinemia Complicating Refractory Mycobacterium tuberculosis Complex Disseminated Disease in a Child with IL-12RÎ²1 Deficiency. Journal of Clinical Immunology, 2017, 37, 732-738.	3.8	10
15	Disseminated BCG Infectious Disease and Hyperferritinemia in a Patient With a Novel NEMO Mutation. Journal of Investigational Allergology and Clinical Immunology, 2016, 26, 268-271.	1.3	10
16	LINE-1-Mediated AluYa5 Insertion Underlying Complete Autosomal Recessive IFN-Î³R1 Deficiency. Journal of Clinical Immunology, 2019, 39, 739-742.	3.8	5