

Marcela Moncada

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/3867254/publications.pdf>

Version: 2024-02-01

13
papers

3,371
citations

840776

11
h-index

1125743

13
g-index

15
all docs

15
docs citations

15
times ranked

8084
citing authors

#	ARTICLE	IF	CITATIONS
1	Inborn errors of type I IFN immunity in patients with life-threatening COVID-19. <i>Science</i> , 2020, 370, .	12.6	1,749
2	Autoantibodies neutralizing type I IFNs are present in ~4% of uninfected individuals over 70 years old and account for ~20% of COVID-19 deaths. <i>Science Immunology</i> , 2021, 6, .	11.9	357
3	Human TYK2 deficiency: Mycobacterial and viral infections without hyper-IgE syndrome. <i>Journal of Experimental Medicine</i> , 2015, 212, 1641-1662.	8.5	293
4	X-linked recessive TLR7 deficiency in ~1% of men under 60 years old with life-threatening COVID-19. <i>Science Immunology</i> , 2021, 6, .	11.9	267
5	Human genetic and immunological determinants of critical COVID-19 pneumonia. <i>Nature</i> , 2022, 603, 587-598.	27.8	216
6	The human gene damage index as a gene-level approach to prioritizing exome variants. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2015, 112, 13615-13620.	7.1	213
7	The risk of COVID-19 death is much greater and age dependent with type I IFN autoantibodies. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2022, 119, e2200413119.	7.1	110
8	Recessive inborn errors of type I IFN immunity in children with COVID-19 pneumonia. <i>Journal of Experimental Medicine</i> , 2022, 219, .	8.5	59
9	Partial IFN- β 2 deficiency is due to protein misfolding and can be rescued by inhibitors of glycosylation. <i>Blood</i> , 2013, 122, 2390-2401.	1.4	34
10	Inherited GATA2 Deficiency Is Dominant by Haploinsufficiency and Displays Incomplete Clinical Penetrance. <i>Journal of Clinical Immunology</i> , 2021, 41, 639-657.	3.8	30
11	Respiratory viral infections in otherwise healthy humans with inherited IRF7 deficiency. <i>Journal of Experimental Medicine</i> , 2022, 219, .	8.5	21
12	Severe Mycobacterial Diseases in a Patient with GOF β 2 Mutation Without EDA. <i>Journal of Clinical Immunology</i> , 2016, 36, 12-15.	3.8	11
13	Autosomal Dominant IFN- β 1 Deficiency Presenting with both Atypical Mycobacteriosis and Tuberculosis in a BCG-Vaccinated South African Patient. <i>Journal of Clinical Immunology</i> , 2018, 38, 460-463.	3.8	8