Jamila El-Baghdadi

List of Publications by Year in descending order

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#	Article	IF	Citations
1	Recessive inborn errors of type I IFN immunity in children with COVID-19 pneumonia. Journal of Experimental Medicine, 2022, 219, .	8.5	59
2	Human <i>STAT3</i> variants underlie autosomal dominant hyper-IgE syndrome by negative dominance. Journal of Experimental Medicine, 2021, 218, .	8.5	30
3	Biochemically deleterious human <i>NFKB1</i> variants underlie an autosomal dominant form of common variable immunodeficiency. Journal of Experimental Medicine, 2021, 218, .	8.5	32
4	Identification of p.Met215lle mutation of the MC4R gene in a Moroccan woman with obesity. Clinical Case Reports (discontinued), 2021, 9, e05059.	0.5	0
5	A prognostic approach on a case of pediatric Philadelphia chromosomeâ€positive acute lymphoblastic leukemia with monosomyâ€7. Clinical Case Reports (discontinued), 2021, 9, e05207.	0.5	O
6	Human T-bet Governs Innate and Innate-like Adaptive IFN- \hat{l}^3 Immunity against Mycobacteria. Cell, 2020, 183, 1826-1847.e31.	28.9	83
7	Prevalence and risk factors for latent tuberculosis infection among healthcare workers in Morocco. PLoS ONE, 2019, 14, e0221081.	2.5	17
8	Tuberculosis and impaired IL-23–dependent IFN-γ immunity in humans homozygous for a common <i>TYK2</i> missense variant. Science Immunology, 2018, 3, .	11.9	148
9	Whole-exome sequencing to analyze population structure, parental inbreeding, and familial linkage. Proceedings of the National Academy of Sciences of the United States of America, 2016, 113, 6713-6718.	7.1	53
10	Inherited and acquired immunodeficiencies underlying tuberculosis in childhood. Immunological Reviews, 2015, 264, 103-120.	6.0	180
11	Human TYK2 deficiency: Mycobacterial and viral infections without hyper-lgE syndrome. Journal of Experimental Medicine, 2015, 212, 1641-1662.	8.5	293
12	Human genetics of tuberculosis: a long and winding road. Philosophical Transactions of the Royal Society B: Biological Sciences, 2014, 369, 20130428.	4.0	144
13	Chronic Mucocutaneous Candidiasis in Humans with Inborn Errors of Interleukin-17 Immunity. Science, 2011, 332, 65-68.	12.6	1,482
14	IL- $12R\hat{l}^21$ Deficiency in Two of Fifty Children with Severe Tuberculosis from Iran, Morocco, and Turkey. PLoS ONE, 2011, 6, e18524.	2.5	111
15	Revisiting Human IL-12RÎ ² 1 Deficiency. Medicine (United States), 2010, 89, 381-402.	1.0	367
16	Association of TP53 gene polymorphisms with the risk of acute lymphoblastic leukemia in Moroccan children. Molecular Biology Reports, 0, , .	2.3	1