Jamila El-Baghdadi

List of Publications by Year
in descending order

16	3,003	687363 13	1125743 13
papers	citations	h-index	g-index

17	17	17	3963
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Chronic Mucocutaneous Candidiasis in Humans with Inborn Errors of Interleukin-17 Immunity. Science, 2011, 332, 65-68.	12.6	1,482
2	Revisiting Human IL-12RÎ²1 Deficiency. Medicine (United States), 2010, 89, 381-402.	1.0	367
3	Human TYK2 deficiency: Mycobacterial and viral infections without hyper-IgE syndrome. Journal of Experimental Medicine, 2015, 212, 1641-1662.	8.5	293
4	Inherited and acquired immunodeficiencies underlying tuberculosis in childhood. Immunological Reviews, 2015, 264, 103-120.	6.0	180
5	Tuberculosis and impaired IL-23â€“dependent IFN-Î³ immunity in humans homozygous for a common TYK2 missense variant. Science Immunology, 2018, 3, .	11.9	148
6	Human genetics of tuberculosis: a long and winding road. Philosophical Transactions of the Royal Society B: Biological Sciences, 2014, 369, 20130428.	4.0	144
7	IL-12RÎ²1 Deficiency in Two of Fifty Children with Severe Tuberculosis from Iran, Morocco, and Turkey. PLoS ONE, 2011, 6, e18524.	2.5	111
8	Human T-bet Governs Innate and Innate-like Adaptive IFN-Î³ Immunity against Mycobacteria. Cell, 2020, 183, 1826-1847.e31.	28.9	83
9	Recessive inborn errors of type I IFN immunity in children with COVID-19 pneumonia. Journal of Experimental Medicine, 2022, 219, .	8.5	59
10	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
11	Biochemically deleterious human NFKB1 variants underlie an autosomal dominant form of common variable immunodeficiency. Journal of Experimental Medicine, 2021, 218, .	8.5	32
12	Human STAT3 variants underlie autosomal dominant hyper-IgE syndrome by negative dominance. Journal of Experimental Medicine, 2021, 218, .	8.5	30
13	Prevalence and risk factors for latent tuberculosis infection among healthcare workers in Morocco. PLoS ONE, 2019, 14, e0221081.	2.5	17
14	Association of TP53 gene polymorphisms with the risk of acute lymphoblastic leukemia in Moroccan children. Molecular Biology Reports, 0, , .	2.3	1
15	Identification of p.Met215Ile mutation of the MC4R gene in a Moroccan woman with obesity. Clinical Case Reports (discontinued), 2021, 9, e05059.	0.5	0
16	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	0