

Romain LÃ©vy

List of Publications by Year in descending order

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Version: 2024-02-01

25
papers

5,490
citations

393982

19
h-index

552369

26
g-index

27
all docs

27
docs citations

27
times ranked

10146
citing authors

#	ARTICLE	IF	CITATIONS
1	Autoantibodies against type I IFNs in patients with life-threatening COVID-19. <i>Science</i> , 2020, 370, .	6.0	1,983
2	Inborn errors of type I IFN immunity in patients with life-threatening COVID-19. <i>Science</i> , 2020, 370, .	6.0	1,749
3	Human genetic and immunological determinants of critical COVID-19 pneumonia. <i>Nature</i> , 2022, 603, 587-598.	13.7	216
4	Inherited CARD9 deficiency in otherwise healthy children and adults with <i>Candida</i> species-induced meningoencephalitis, colitis, or both. <i>Journal of Allergy and Clinical Immunology</i> , 2015, 135, 1558-1568.e2.	1.5	208
5	Preexisting autoantibodies to type I IFNs underlie critical COVID-19 pneumonia in patients with APS-1. <i>Journal of Experimental Medicine</i> , 2021, 218, .	4.2	185
6	Phenotypic and genotypic characteristics of cryopyrin-associated periodic syndrome: a series of 136 patients from the Eurofever Registry. <i>Annals of the Rheumatic Diseases</i> , 2015, 74, 2043-2049.	0.5	180
7	Inherited IL-17RC deficiency in patients with chronic mucocutaneous candidiasis. <i>Journal of Experimental Medicine</i> , 2015, 212, 619-631.	4.2	162
8	Genetic, immunological, and clinical features of patients with bacterial and fungal infections due to inherited IL-17RA deficiency. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2016, 113, E8277-E8285.	3.3	137
9	A recessive form of hyper-IgE syndrome by disruption of ZNF341-dependent STAT3 transcription and activity. <i>Science Immunology</i> , 2018, 3, .	5.6	132
10	Haploidentical Hematopoietic Stem Cell Transplantation with Post-Transplant Cyclophosphamide for Primary Immunodeficiencies and Inherited Disorders in Children. <i>Biology of Blood and Marrow Transplantation</i> , 2019, 25, 1363-1373.	2.0	78
11	Herpes simplex encephalitis in a patient with a distinctive form of inherited IFNAR1 deficiency. <i>Journal of Clinical Investigation</i> , 2021, 131, .	3.9	64
12	Profound symptomatic hypogammaglobulinemia: A rare late complication after rituximab treatment for immune thrombocytopenia. Report of 3 cases and systematic review of the literature.. <i>Autoimmunity Reviews</i> , 2014, 13, 1055-1063.	2.5	61
13	Chronic mucocutaneous candidiasis and connective tissue disorder in humans with impaired JNK1-dependent responses to IL-17A/F and TGF- β 2. <i>Science Immunology</i> , 2019, 4, .	5.6	45
14	Interferon- β Therapy in a Patient with Incontinentia Pigmenti and Autoantibodies against Type I IFNs Infected with SARS-CoV-2. <i>Journal of Clinical Immunology</i> , 2021, 41, 931-933.	2.0	40
15	IFN- β 2a Therapy in Two Patients with Inborn Errors of TLR3 and IRF3 Infected with SARS-CoV-2. <i>Journal of Clinical Immunology</i> , 2021, 41, 26-27.	2.0	40
16	Effectiveness and safety of ruxolitinib for the treatment of refractory systemic idiopathic juvenile arthritis like associated with interstitial lung disease : a case report. <i>Annals of the Rheumatic Diseases</i> , 2022, 81, e20-e20.	0.5	36
17	Efficacy of Dupilumab for Controlling Severe Atopic Dermatitis in a Patient with Hyper-IgE Syndrome. <i>Journal of Clinical Immunology</i> , 2020, 40, 418-420.	2.0	28
18	Monoclonal antibody-mediated neutralization of SARS-CoV-2 in an IRF9-deficient child. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2021, 118, .	3.3	24

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19	Impaired respiratory burst contributes to infections in PKC δ -deficient patients. Journal of Experimental Medicine, 2021, 218, .	4.2	23
20	Efficacy of ruxolitinib in subcutaneous panniculitis-like T-cell lymphoma and hemophagocytic lymphohistiocytosis. Blood Advances, 2020, 4, 1383-1387.	2.5	21
21	Inherited human c-Rel deficiency disrupts myeloid and lymphoid immunity to multiple infectious agents. Journal of Clinical Investigation, 2021, 131, .	3.9	21
22	Somatic reversion of pathogenic DOCK8 variants alters lymphocyte differentiation and function to effectively cure DOCK8 deficiency. Journal of Clinical Investigation, 2021, 131, .	3.9	18
23	Differential Expression of Interferon-Alpha Protein Provides Clues to Tissue Specificity Across Type I Interferonopathies. Journal of Clinical Immunology, 2021, 41, 603-609.	2.0	16
24	Donor-targeted serotherapy as a rescue therapy for steroid-resistant acute GVHD after HLA-mismatched kidney transplantation. American Journal of Transplantation, 2020, 20, 2243-2253.	2.6	11
25	Inherited TNFSF9 deficiency causes broad Epsteinâ€Barr virus infection with EBV+ smooth muscle tumors. Journal of Experimental Medicine, 2022, 219, .	4.2	7