Jacques G RiviÃ¨re

List of Publications by Year
in descending order

26 papers	1,422 citations	759233 12 h-index	580821 25 g-index
29 all docs	29 docs citations	29 times ranked	2409 citing authors

#	Article	IF	Citations
1	Autoantibodies neutralizing type I IFNs are present in ~4% of uninfected individuals over 70 years old and account for ~20% of COVID-19 deaths. Science Immunology, 2021, 6, .	11.9	357
2	X-linked recessive TLR7 deficiency in ~1% of men under 60 years old with life-threatening COVID-19. Science Immunology, 2021, 6, .	11.9	267
3	Long-term outcome of LRBA deficiency in 76 patients after various treatment modalities as evaluated by the immune deficiency and dysregulation activity (IDDA) score. Journal of Allergy and Clinical Immunology, 2020, 145, 1452-1463.	2.9	112
4	The risk of COVID-19 death is much greater and age dependent with type I IFN autoantibodies. Proceedings of the National Academy of Sciences of the United States of America, 2022, 119, e2200413119.	7.1	110
5	SARS-CoV-2â€“related MIS-C: A key to the viral and genetic causes of Kawasaki disease?. Journal of Experimental Medicine, 2021, 218, .	8.5	100
6	Multi-inflammatory Syndrome in Children Related to Severe Acute Respiratory Syndrome Coronavirus 2 (SARS-CoV-2) in Spain. Clinical Infectious Diseases, 2021, 72, e397-e401.	5.8	98
7	Overview of STING-Associated Vasculopathy with Onset in Infancy (SAVI) Among 21 Patients. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 803-818.e11.	3.8	98
8	Recessive inborn errors of type I IFN immunity in children with COVID-19 pneumonia. Journal of Experimental Medicine, 2022, 219, .	8.5	59
9	First Universal Newborn Screening Program for Severe Combined Immunodeficiency in Europe. Two-Years' Experience in Catalonia (Spain). Frontiers in Immunology, 2019, 10, 2406.	4.8	45
10	LRBA Deficiency in a Patient With a Novel Homozygous Mutation Due to Chromosome 4 Segmental Uniparental Isodisomy. Frontiers in Immunology, 2018, 9, 2397.	4.8	37
11	Neuroinflammatory Disease as an Isolated Manifestation of Hemophagocytic Lymphohistiocytosis. Journal of Clinical Immunology, 2020, 40, 901-916.	3.8	33
12	Curation and expansion of Human Phenotype Ontology for defined groups of inborn errors of immunity. Journal of Allergy and Clinical Immunology, 2022, 149, 369-378.	2.9	16
13	Dynamics of Reverse Transcription-Polymerase Chain Reaction and Serologic Test Results in Children with SARS-CoV-2 Infection. Journal of Pediatrics, 2022, 241, 126-132.e3.	1.8	12
14	Inborn errors of TLR3- or MDA5-dependent type I IFN immunity in children with enterovirus rhombencephalitis. Journal of Experimental Medicine, 2021, 218, .	8.5	12
15	Epigenetic profiling linked to multisystem inflammatory syndrome in children (MIS-C): A multicenter, retrospective study. EClinicalMedicine, 2022, 50, 101515.	7.1	11
16	Atypical Inflammatory Syndrome Triggered by SARS-CoV-2 in Infants with Down Syndrome. Journal of Clinical Immunology, 2021, 41, 1457-1462.	3.8	9
17	Executive Summary of the Consensus Document on the Diagnosis and Management of Patients with Primary Immunodeficiencies. Journal of Allergy and Clinical Immunology: in Practice, 2020, 8, 3342-3347.	3.8	7
18	Tocilizumab in a child with acute lymphoblastic leukaemia and COVID-19-related cytokine release syndrome. Anales De PediatrÃa (English Edition), 2020, 93, 132-133.	0.2	6

#	Article	IF	Citations
19	Health-Related Quality of Life and Multidimensional Fatigue Scale in Children with Primary Immunodeficiencies. Journal of Clinical Immunology, 2020, 40, 602-609.	3.8	6
20	Uncovering Low-Level Maternal Gonosomal Mosaicism in X-Linked Agammaglobulinemia: Implications for Genetic Counseling. Frontiers in Immunology, 2020, 11, 46.	4.8	5
21	Activation-induced deaminase is critical for the establishment of DNA methylation patterns prior to the germinal center reaction. Nucleic Acids Research, 2021, 49, 5057-5073.	14.5	5
22	FHLdb: A Comprehensive Database on the Molecular Basis of Familial Hemophagocytic Lymphohistiocytosis. Frontiers in Immunology, 2020, 11, 107.	4.8	4
23	Newborn Screening for SCID: Experience in Spain (Catalonia). International Journal of Neonatal Screening, 2021, 7, 46.	3.2	4
24	Influenza-Associated Disseminated Aspergillosis in a 9-Year-Old Girl Requiring ECMO Support. Journal of Fungi (Basel, Switzerland), 2021, 7, 726.	3.5	4
25	Early Diagnosis and Treatment of Purine Nucleoside Phosphorylase (PNP) Deficiency through TREC-Based Newborn Screening. International Journal of Neonatal Screening, 2021, 7, 62.	3.2	2
26	Executive Summary of the Consensus Document on the Diagnosis and Management of Patients with Primary Immunodeficiencies. Enfermedades Infecciosas Y MicrobiologÃa ClÃnica, 2020, 38, 438-443.	0.5	0