

Jacques G Rivière

List of Publications by Year in descending order

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Version: 2024-02-01

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	Curation and expansion of Human Phenotype Ontology for defined groups of inborn errors of immunity. <i>Journal of Allergy and Clinical Immunology</i> , 2022, 149, 369-378.	2.9	16
2	Dynamics of Reverse Transcription-Polymerase Chain Reaction and Serologic Test Results in Children with SARS-CoV-2 Infection. <i>Journal of Pediatrics</i> , 2022, 241, 126-132.e3.	1.8	12
3	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
4	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
5	Epigenetic profiling linked to multisystem inflammatory syndrome in children (MIS-C): A multicenter, retrospective study. <i>EClinicalMedicine</i> , 2022, 50, 101515.	7.1	11
6	Multi-inflammatory Syndrome in Children Related to Severe Acute Respiratory Syndrome Coronavirus 2 (SARS-CoV-2) in Spain. <i>Clinical Infectious Diseases</i> , 2021, 72, e397-e401.	5.8	98
7	Overview of STING-Associated Vasculopathy with Onset in Infancy (SAVI) Among 21 Patients. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2021, 9, 803-818.e11.	3.8	98
8	SARS-CoV-2-related MIS-C: A key to the viral and genetic causes of Kawasaki disease?. <i>Journal of Experimental Medicine</i> , 2021, 218, .	8.5	100
9	Activation-induced deaminase is critical for the establishment of DNA methylation patterns prior to the germinal center reaction. <i>Nucleic Acids Research</i> , 2021, 49, 5057-5073.	14.5	5
10	Atypical Inflammatory Syndrome Triggered by SARS-CoV-2 in Infants with Down Syndrome. <i>Journal of Clinical Immunology</i> , 2021, 41, 1457-1462.	3.8	9
11	Newborn Screening for SCID: Experience in Spain (Catalonia). <i>International Journal of Neonatal Screening</i> , 2021, 7, 46.	3.2	4
12	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
13	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
14	Influenza-Associated Disseminated Aspergillosis in a 9-Year-Old Girl Requiring ECMO Support. <i>Journal of Fungi (Basel, Switzerland)</i> , 2021, 7, 726.	3.5	4
15	Early Diagnosis and Treatment of Purine Nucleoside Phosphorylase (PNP) Deficiency through TREC-Based Newborn Screening. <i>International Journal of Neonatal Screening</i> , 2021, 7, 62.	3.2	2
16	Inborn errors of TLR3- or MDA5-dependent type I IFN immunity in children with enterovirus rhombencephalitis. <i>Journal of Experimental Medicine</i> , 2021, 218, .	8.5	12
17	Long-term outcome of LRBA deficiency in 76 patients after various treatment modalities as evaluated by the immune deficiency and dysregulation activity (IDDA) score. <i>Journal of Allergy and Clinical Immunology</i> , 2020, 145, 1452-1463.	2.9	112
18	Tocilizumab in a child with acute lymphoblastic leukaemia and COVID-19-related cytokine release syndrome. <i>Anales De Pediatr�a (English Edition)</i> , 2020, 93, 132-133.	0.2	6

#	ARTICLE	IF	CITATIONS
19	Executive Summary of the Consensus Document on the Diagnosis and Management of Patients with Primary Immunodeficiencies. <i>Enfermedades Infecciosas Y Microbiología Clínica</i> , 2020, 38, 438-443.	0.5	0
20	Executive Summary of the Consensus Document on the Diagnosis and Management of Patients with Primary Immunodeficiencies. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2020, 8, 3342-3347.	3.8	7
21	Neuroinflammatory Disease as an Isolated Manifestation of Hemophagocytic Lymphohistiocytosis. <i>Journal of Clinical Immunology</i> , 2020, 40, 901-916.	3.8	33
22	FHLdb: A Comprehensive Database on the Molecular Basis of Familial Hemophagocytic Lymphohistiocytosis. <i>Frontiers in Immunology</i> , 2020, 11, 107.	4.8	4
23	Uncovering Low-Level Maternal Gonosomal Mosaicism in X-Linked Agammaglobulinemia: Implications for Genetic Counseling. <i>Frontiers in Immunology</i> , 2020, 11, 46.	4.8	5
24	Health-Related Quality of Life and Multidimensional Fatigue Scale in Children with Primary Immunodeficiencies. <i>Journal of Clinical Immunology</i> , 2020, 40, 602-609.	3.8	6
25	First Universal Newborn Screening Program for Severe Combined Immunodeficiency in Europe. Two-Years' Experience in Catalonia (Spain). <i>Frontiers in Immunology</i> , 2019, 10, 2406.	4.8	45
26	LRBA Deficiency in a Patient With a Novel Homozygous Mutation Due to Chromosome 4 Segmental Uniparental Isodisomy. <i>Frontiers in Immunology</i> , 2018, 9, 2397.	4.8	37