

David Boutboul

List of Publications by Year in descending order

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Version: 2024-02-01

43
papers

1,820
citations

331670

21
h-index

276875

41
g-index

45
all docs

45
docs citations

45
times ranked

3132
citing authors

#	ARTICLE	IF	CITATIONS
1	Convalescent plasma therapy for B-cell-depleted patients with protracted COVID-19. <i>Blood</i> , 2020, 136, 2290-2295.	1.4	251
2	Clinical and immunologic phenotype associated with activated phosphoinositide 3-kinase γ syndrome 2: A cohort study. <i>Journal of Allergy and Clinical Immunology</i> , 2016, 138, 210-218.e9.	2.9	215
3	Late-Onset Combined Immune Deficiency: A Subset of Common Variable Immunodeficiency with Severe T Cell Defect. <i>Clinical Infectious Diseases</i> , 2009, 49, 1329-1338.	5.8	192
4	The full spectrum of Castleman disease: 273 patients studied over 20 years. <i>British Journal of Haematology</i> , 2018, 180, 206-216.	2.5	137
5	Dominant-negative IKZF1 mutations cause a T, B, and myeloid cell combined immunodeficiency. <i>Journal of Clinical Investigation</i> , 2018, 128, 3071-3087.	8.2	133
6	Synergistic convergence of microbiota-specific systemic IgG and secretory IgA. <i>Journal of Allergy and Clinical Immunology</i> , 2019, 143, 1575-1585.e4.	2.9	86
7	Good Syndrome: An Adult-Onset Immunodeficiency Remarkable for Its High Incidence of Invasive Infections and Autoimmune Complications. <i>Clinical Infectious Diseases</i> , 2015, 61, e13-e19.	5.8	81
8	Acute Kidney Injury in Adults With Hemophagocytic Lymphohistiocytosis. <i>American Journal of Kidney Diseases</i> , 2015, 65, 851-859.	1.9	77
9	Dominant-negative mutations in human <i>IL6ST</i> underlie hyper-IgE syndrome. <i>Journal of Experimental Medicine</i> , 2020, 217, .	8.5	64
10	Exclusion of Patients with a Severe T-Cell Defect Improves the Definition of Common Variable Immunodeficiency. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2016, 4, 1147-1157.	3.8	45
11	PROMIS1: A T-cell receptor γ signature associated with immunodeficiencies caused by V(D)J recombination defects. <i>Journal of Allergy and Clinical Immunology</i> , 2019, 143, 325-334.e2.	2.9	43
12	Hematopoietic Cell Transplantation Cures Adenosine Deaminase 2 Deficiency: Report on 30 Patients. <i>Journal of Clinical Immunology</i> , 2021, 41, 1633-1647.	3.8	43
13	Kaposi sarcoma-associated herpesvirus/human herpesvirus 8-associated lymphoproliferative disorders. <i>Blood</i> , 2019, 133, 1186-1190.	1.4	38
14	Rapid identification and characterization of infected cells in blood during chronic active Epstein-Barr virus infection. <i>Journal of Experimental Medicine</i> , 2020, 217, .	8.5	37
15	Treatment and outcome of Unicentric Castleman Disease: a retrospective analysis of 71 cases. <i>British Journal of Haematology</i> , 2019, 186, 269-273.	2.5	36
16	Severe COVID-19 in Patients with B Cell Lymphocytosis and Response to Convalescent Plasma Therapy. <i>Journal of Clinical Immunology</i> , 2021, 41, 356-361.	3.8	35
17	Vaccine breakthrough hypoxemic COVID-19 pneumonia in patients with auto-Abs neutralizing type I IFNs. <i>Science Immunology</i> , 2023, 8, .	11.9	35
18	Allogeneic hematopoietic stem cell transplant outcomes for patients with dominant negative IKZF1/IKAROS mutations. <i>Journal of Allergy and Clinical Immunology</i> , 2019, 144, 339-342.	2.9	28

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19	Autoimmune cytopenias associated with inflammatory bowel diseases: Insights from a multicenter retrospective cohort. <i>Digestive and Liver Disease</i> , 2017, 49, 397-404.	0.9	27
20	Gain-of-function <i>IKZF1</i> variants in humans cause immune dysregulation associated with abnormal T/B cell late differentiation. <i>Science Immunology</i> , 2022, 7, eabi7160.	11.9	27
21	A comprehensive analysis of Lymphoma-associated haemophagocytic syndrome in a large French multicentre cohort detects some clues to improve prognosis. <i>British Journal of Haematology</i> , 2018, 183, 68-75.	2.5	23
22	Accessible LAMP-Enabled Rapid Test (ALERT) for Detecting SARS-CoV-2. <i>Viruses</i> , 2021, 13, 742.	3.3	23
23	Hemophagocytic Lymphohistiocytosis Associated With <i>Bartonella henselae</i> Infection in an HIV-Infected Patient. <i>Clinical Infectious Diseases</i> , 2016, 62, 804-806.	5.8	17
24	Epidemiology, Risk Factors, and Outcomes of Opportunistic Infections after Kidney Allograft Transplantation in the Era of Modern Immunosuppression: A Monocentric Cohort Study. <i>Journal of Clinical Medicine</i> , 2019, 8, 594.	2.4	17
25	Improving the diagnostic efficiency of primary immunodeficiencies with targeted next-generation sequencing. <i>Journal of Allergy and Clinical Immunology</i> , 2021, 147, 734-737.	2.9	17
26	Late-Onset EBV Susceptibility and Refractory Pure Red Cell Aplasia Revealing DADA2. <i>Journal of Clinical Immunology</i> , 2020, 40, 948-953.	3.8	14
27	DADA2 diagnosed in adulthood versus childhood: A comparative study on 306 patients including a systematic literature review and 12 French cases. <i>Seminars in Arthritis and Rheumatism</i> , 2021, 51, 1170-1179.	3.4	14
28	Autoimmune thrombotic thrombocytopenic purpura associated with HHV-8-related Multicentric Castleman disease. <i>British Journal of Haematology</i> , 2017, 178, 486-488.	2.5	13
29	Epidemiology of Castleman disease associated with AA amyloidosis: description of 2 new cases and literature review. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2019, 26, 197-202.	3.0	7
30	Identification and characterization of two novel Gammapapillomavirus genomes in skin of an immunosuppressed Epidermodysplasia Verruciformis patient. <i>Virus Research</i> , 2018, 249, 66-68.	2.2	6
31	Monoclonal Gammopathy, Arthralgias, and Recurrent Fever Syndrome: A New Autoinflammatory Syndrome?. <i>Journal of Rheumatology</i> , 2019, 46, 1535-1539.	2.0	6
32	Talc pleurodesis allows long-term remission in HIV-unrelated Human Herpesvirus 8-associated primary effusion lymphoma. <i>Leukemia and Lymphoma</i> , 2017, 58, 1993-1998.	1.3	5
33	Characteristics of thrombocytopenia, anasarca, fever, reticulin fibrosis and organomegaly syndrome: a retrospective study from a large Western cohort. <i>British Journal of Haematology</i> , 2022, 196, 599-605.	2.5	5
34	Pseudo-Sarcoidosis Revealing MonoMAC Syndrome. <i>Journal of Clinical Immunology</i> , 2018, 38, 739-741.	3.8	4
35	Uterine intravascular lymphoma as a cause of fever of unknown origin. <i>Annals of Hematology</i> , 2017, 96, 1891-1896.	1.8	3
36	Understanding therapeutic emergencies in acute hemolysis. <i>Intensive Care Medicine</i> , 2018, 44, 482-485.	8.2	3

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37	Extreme gonococcal susceptibility associated with acquired complement deficiency secondary to hypocomplementemic urticarial vasculitis and systemic lupus erythematosus. <i>Journal of Infection and Chemotherapy</i> , 2021, , .	1.7	3
38	Cutaneous Granulomatosis Revealing Whipple's Disease: Value of <i>Tropheryma whipplei</i> Polymerase Chain Reaction Assay for the Diagnosis. <i>Pathogens</i> , 2021, 10, 1438.	2.8	3
39	Epstein-Barr Virus Genome Deletions in Epstein-Barr Virus-Positive T/NK Cell Lymphoproliferative Diseases. <i>Journal of Virology</i> , 2022, 96, .	3.4	3
40	Autoimmune hypoglycemia expands the biological spectrum of HHV8+ multicentric Castleman disease. <i>Blood Advances</i> , 2021, 5, 1848-1852.	5.2	2
41	Chronic Disseminated Aspergillosis, a Novel Fungal Immune Reconstitution Inflammatory Syndrome. <i>Open Forum Infectious Diseases</i> , 2020, 7, ofaa175.	0.9	2
42	Characteristics and mid-term follow-up of COVID-19 patients with hematological diseases: a retrospective study from a French tertiary care hospital. <i>Blood Cancer Journal</i> , 2021, 11, 129.	6.2	0
43	Leg-type form of idiopathic multicentric Castleman disease associated with severe lower extremity chronic venous/lymphatic disease. <i>EJHaem</i> , 2022, 3, 175-179.	1.0	0