David Boutboul

List of Publications by Year in descending order

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	Vaccine breakthrough hypoxemic COVID-19 pneumonia in patients with auto-Abs neutralizing type I IFNs. Science Immunology, 2023, 8, .	11.9	35
2	Characteristics of thrombocytopenia, anasarca, fever, reticulin fibrosis and organomegaly syndrome: a retrospective study from a large Western cohort. British Journal of Haematology, 2022, 196, 599-605.	2.5	5
3	Gain-of-function IKZF1 variants in humans cause immune dysregulation associated with abnormal T/B cell late differentiation. Science Immunology, 2022, 7, eabi7160.	11.9	27
4	Legâ€type form of idiopathic multicentric Castleman disease associated with severe lower extremity chronic venous/lymphatic disease. EJHaem, 2022, 3, 175-179.	1.0	0
5	Epstein-Barr Virus Genome Deletions in Epstein-Barr Virus-Positive T/NK Cell Lymphoproliferative Diseases. Journal of Virology, 2022, 96, .	3.4	3
6	Improving the diagnostic efficiency of primary immunodeficiencies with targeted next-generation sequencing. Journal of Allergy and Clinical Immunology, 2021, 147, 734-737.	2.9	17
7	Severe COVID-19 in Patients with B Cell Alymphocytosis and Response to Convalescent Plasma Therapy. Journal of Clinical Immunology, 2021, 41, 356-361.	3.8	35
8	Autoimmune hypoglycemia expands the biological spectrum of HHV8+ multicentric Castleman disease. Blood Advances, 2021, 5, 1848-1852.	5.2	2
9	Accessible LAMP-Enabled Rapid Test (ALERT) for Detecting SARS-CoV-2. Viruses, 2021, 13, 742.	3.3	23
10	Characteristics and mid-term follow-up of COVID-19 patients with hematological diseases: a retrospective study from a French tertiary care hospital. Blood Cancer Journal, 2021, 11, 129.	6.2	0
11	Hematopoietic Cell Transplantation Cures Adenosine Deaminase 2 Deficiency: Report on 30 Patients. Journal of Clinical Immunology, 2021, 41, 1633-1647.	3.8	43
12	DADA2 diagnosed in adulthood versus childhood: A comparative study on 306 patients including a systematic literature review and 12 French cases. Seminars in Arthritis and Rheumatism, 2021, 51, 1170-1179.	3.4	14
13	Extreme gonococcal susceptibility associated with acquired complement deficiency secondary to hypocomplementemic urticarial vasculitis and systemic lupus erythematosus. Journal of Infection and Chemotherapy, 2021, , .	1.7	3
14	Cutaneous Granulomatosis Revealing Whippleâ€™s Disease: Value of Tropheryma whipplei Polymerase Chain Reaction Assay for the Diagnosis. Pathogens, 2021, 10, 1438.	2.8	3
15	Late-Onset EBV Susceptibility and Refractory Pure Red Cell Aplasia Revealing DADA2. Journal of Clinical Immunology, 2020, 40, 948-953.	3.8	14
16	Rapid identification and characterization of infected cells in blood during chronic active Epstein-Barr virus infection. Journal of Experimental Medicine, 2020, 217, .	8.5	37
17	Dominant-negative mutations in human IL6ST underlie hyper-IgE syndrome. Journal of Experimental Medicine, 2020, 217, .	8.5	64
18	Convalescent plasma therapy for B-cellâ€“depleted patients with protracted COVID-19. Blood, 2020, 136, 2290-2295.	1.4	251

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19	â€œChronic Disseminated Aspergillosis,â€•a Novel Fungal Immune Reconstitution Inflammatory Syndrome. Open Forum Infectious Diseases, 2020, 7, ofaa175.	0.9	2
20	PROMIDISÎ±: AÂT-cell receptor Î± signature associated with immunodeficiencies caused by V(D)J recombination defects. Journal of Allergy and Clinical Immunology, 2019, 143, 325-334.e2.	2.9	43
21	Epidemiology of Castleman disease associated with AA amyloidosis: description of 2 new cases and literature review. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2019, 26, 197-202.	3.0	7
22	Epidemiology, Risk Factors, and Outcomes of Opportunistic Infections after Kidney Allograft Transplantation in the Era of Modern Immunosuppression: A Monocentric Cohort Study. Journal of Clinical Medicine, 2019, 8, 594.	2.4	17
23	Allogeneic hematopoietic stem cell transplant outcomes for patients with dominant negative IKZF1/IKAROS mutations. Journal of Allergy and Clinical Immunology, 2019, 144, 339-342.	2.9	28
24	Treatment and outcome of Unicentric Castleman Disease: a retrospective analysis of 71 cases. British Journal of Haematology, 2019, 186, 269-273.	2.5	36
25	Monoclonal Gammopathy, Arthralgias, and Recurrent Fever Syndrome: A New Autoinflammatory Syndrome?. Journal of Rheumatology, 2019, 46, 1535-1539.	2.0	6
26	Synergistic convergence of microbiota-specific systemic IgG and secretory IgA. Journal of Allergy and Clinical Immunology, 2019, 143, 1575-1585.e4.	2.9	86
27	Kaposi sarcomaâ€“associated herpesvirus/human herpesvirus 8â€“associated lymphoproliferative disorders. Blood, 2019, 133, 1186-1190.	1.4	38
28	Identification and characterization of two novel Gammapapillomavirus genomes in skin of an immunosuppressed Epidermodysplasia Verruciformis patient. Virus Research, 2018, 249, 66-68.	2.2	6
29	The full spectrum of Castleman disease: 273 patients studied over 20Âyears. British Journal of Haematology, 2018, 180, 206-216.	2.5	137
30	Pseudo-Sarcoidosis Revealing MonoMAC Syndrome. Journal of Clinical Immunology, 2018, 38, 739-741.	3.8	4
31	A comprehensive analysis of Lymphomaâ€associated haemophagocytic syndrome in a large French multicentre cohort detects some clues to improve prognosis. British Journal of Haematology, 2018, 183, 68-75.	2.5	23
32	Understanding therapeutic emergencies in acute hemolysis. Intensive Care Medicine, 2018, 44, 482-485.	8.2	3
33	Dominant-negative IKZF1 mutations cause a T, B, and myeloid cell combined immunodeficiency. Journal of Clinical Investigation, 2018, 128, 3071-3087.	8.2	133
34	Autoimmune cytopenias associated with inflammatory bowel diseases: Insights from a multicenter retrospective cohort. Digestive and Liver Disease, 2017, 49, 397-404.	0.9	27
35	Talc pleurodesis allows long-term remission in HIV-unrelated Human Herpesvirus 8-associated primary effusion lymphoma. Leukemia and Lymphoma, 2017, 58, 1993-1998.	1.3	5
36	Uterine intravascular lymphoma as a cause of fever of unknown origin. Annals of Hematology, 2017, 96, 1891-1896.	1.8	3

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37	Autoimmune thrombotic thrombocytopenic purpura associated with <scp>HHV</scp>8â€related Multicentric Castleman disease. British Journal of Haematology, 2017, 178, 486-488.	2.5	13
38	Exclusion of Patients with a Severe T-Cell Defect Improves the Definition of Common Variable Immunodeficiency. Journal of Allergy and Clinical Immunology: in Practice, 2016, 4, 1147-1157.	3.8	45
39	Clinical and immunologic phenotype associated with activated phosphoinositide 3-kinase Î´ syndrome 2: AÂcohort study. Journal of Allergy and Clinical Immunology, 2016, 138, 210-218.e9.	2.9	215
40	Hemophagocytic Lymphohistiocytosis Associated WithBartonella henselaeInfection in an HIV-Infected Patient. Clinical Infectious Diseases, 2016, 62, 804-806.	5.8	17
41	Good Syndrome: An Adult-Onset Immunodeficiency Remarkable for Its High Incidence of Invasive Infections and Autoimmune Complications. Clinical Infectious Diseases, 2015, 61, e13-e19.	5.8	81
42	Acute Kidney Injury in Adults With Hemophagocytic Lymphohistiocytosis. American Journal of Kidney Diseases, 2015, 65, 851-859.	1.9	77
43	Lateâ€Onset Combined Immune Deficiency: A Subset of Common Variable Immunodeficiency with Severe T Cell Defect. Clinical Infectious Diseases, 2009, 49, 1329-1338.	5.8	192