Eric Oksenhendler

List of Publications by Year
in descending order

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70 papers	8,584 citations	109321 35 h-index	102487 66 g-index
71 all docs	71 docs citations	71 times ranked	8909 citing authors

#	Article	IF	Citations
1	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
2	First Extensive Analysis of ¹⁸F-Labeled Fluorodeoxyglucose Positron Emission Tomographyâ€“Computed Tomography in a Large Cohort of Patients With HIV-Associated Hodgkin Lymphoma: Baseline Total Metabolic Tumor Volume Affects Prognosis. Journal of Clinical Oncology, 2022, 40, 1346-1355.	1.6	11
3	Hepatosplenic Candidiasis in Patients With Hematological Malignancies: A 13-Year Retrospective Cohort Study. Open Forum Infectious Diseases, 2022, 9, ofac088.	0.9	4
4	Inflammatory WaldenstrÃ¶m's macroglobulinaemia: A French monocentric retrospective study of 67 patients. British Journal of Haematology, 2022, 197, 728-735.	2.5	3
5	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
6	Hepatitis E infection in adults with primary immunodeficiency with or without immunoglobulin replacement therapy.. Blood Transfusion, 2022, , .	0.4	0
7	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
8	The Ever-Increasing Array of Novel Inborn Errors of Immunity: an Interim Update by the IUIS Committee. Journal of Clinical Immunology, 2021, 41, 666-679.	3.8	165
9	Autoimmune hypoglycemia expands the biological spectrum of HHV8+ multicentric Castleman disease. Blood Advances, 2021, 5, 1848-1852.	5.2	2
10	Validated international definition of the thrombocytopenia, anasarca, fever, reticulin fibrosis, renal insufficiency, and organomegaly clinical subtype (TAFRO) of idiopathic multicentric <scp>Castleman</scp> disease. American Journal of Hematology, 2021, 96, 1241-1252.	4.1	47
11	Phenocopies of Inborn Errors of Immunity. , 2021, , .		0
12	A Novel Predictive Model for Idiopathic Multicentric Castleman Disease: The International Castleman Disease Consortium Study. Oncologist, 2020, 25, 963-973.	3.7	11
13	Insufficient evidence exists to use histopathologic subtype to guide treatment of idiopathic multicentric Castleman disease. American Journal of Hematology, 2020, 95, 1553-1561.	4.1	18
14	International evidence-based consensus diagnostic and treatment guidelines for unicentric Castleman disease. Blood Advances, 2020, 4, 6039-6050.	5.2	94
15	From Dysgammaglobulinemia to Autosomal-Dominant Activation-Induced Cytidine Deaminase Deficiency: Unraveling an Inherited ImmunodeficiencyÂafter 50ÂYears. Journal of Pediatrics, 2020, 223, 207-211.e1.	1.8	4
16	Autoimmune Lymphoproliferative Syndrome Presenting with Invasive Streptococcus pneumoniae Infection. Journal of Clinical Immunology, 2020, 40, 543-546.	3.8	3
17	Human Inborn Errors of Immunity: 2019 Update on the Classification from the International Union of Immunological Societies Expert Committee. Journal of Clinical Immunology, 2020, 40, 24-64.	3.8	881
18	Characterization of the clinical and immunologic phenotype and management of 157 individuals with 56 distinct heterozygous NFKB1 mutations. Journal of Allergy and Clinical Immunology, 2020, 146, 901-911.	2.9	78

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19	The French paediatric cohort of Castleman disease: a retrospective report of 23 patients. Orphanet Journal of Rare Diseases, 2020, 15, 95.	2.7	10
20	Human Inborn Errors of Immunity: 2019 Update of the IUIS Phenotypical Classification. Journal of Clinical Immunology, 2020, 40, 66-81.	3.8	525
21	ACCELERATE: A Patient-Powered Natural History Study Design Enabling Clinical and Therapeutic Discoveries in a Rare Disorder. Cell Reports Medicine, 2020, 1, 100158.	6.5	18
22	Campylobacter infection in adult patients with primary antibody deficiency. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 1038-1041.e4.	3.8	7
23	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
24	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
25	HIVâ€1 Tat protein induces aberrant activation of AICDA in human Bâ€lymphocytes from peripheral blood. Journal of Cellular Physiology, 2019, 234, 15678-15685.	4.1	16
26	Bortezomib and dexamethasone, an original approach for treating multiâ€refractory warm autoimmune haemolytic anaemia. British Journal of Haematology, 2019, 187, 124-128.	2.5	29
27	Treatment and outcome of Unicentric Castleman Disease: a retrospective analysis of 71 cases. British Journal of Haematology, 2019, 186, 269-273.	2.5	36
28	CD89 Is a Potent Innate Receptor for Bacteria and Mediates Host Protection from Sepsis. Cell Reports, 2019, 27, 762-775.e5.	6.4	19
29	Synergistic convergence of microbiota-specific systemic IgG and secretory IgA. Journal of Allergy and Clinical Immunology, 2019, 143, 1575-1585.e4.	2.9	86
30	Kaposi sarcomaâ€“associated herpesvirus/human herpesvirus 8â€“associated lymphoproliferative disorders. Blood, 2019, 133, 1186-1190.	1.4	38
31	Loss of ARHGEF1 causes a human primary antibody deficiency. Journal of Clinical Investigation, 2019, 129, 1047-1060.	8.2	32
32	Microbial ecology perturbation in human IgA deficiency. Science Translational Medicine, 2018, 10, .	12.4	206
33	Loss-of-function nuclear factor ÎºB subunit 1 (NFKB1) variants are the most common monogenic cause of common variable immunodeficiency in Europeans. Journal of Allergy and Clinical Immunology, 2018, 142, 1285-1296.	2.9	185
34	International Union of Immunological Societies: 2017 Primary Immunodeficiency Diseases Committee Report on Inborn Errors of Immunity. Journal of Clinical Immunology, 2018, 38, 96-128.	3.8	732
35	The 2017 IUIS Phenotypic Classification for Primary Immunodeficiencies. Journal of Clinical Immunology, 2018, 38, 129-143.	3.8	488
36	HIV-1 Tat protein induces DNA damage in human peripheral blood B-lymphocytes via mitochondrial ROS production. Redox Biology, 2018, 15, 97-108.	9.0	62

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37	The full spectrum of Castleman disease: 273 patients studied over 20Âyears. British Journal of Haematology, 2018, 180, 206-216.	2.5	137
38	International, evidence-based consensus treatment guidelines for idiopathic multicentric Castleman disease. Blood, 2018, 132, 2115-2124.	1.4	232
39	Mutations in the SRP54 gene cause severe congenital neutropenia as well as Shwachman-Diamondâ€“like syndrome. Blood, 2018, 132, 1318-1331.	1.4	85
40	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
41	International, evidence-based consensus diagnostic criteria for HHV-8â€“negative/idiopathic multicentric Castleman disease. Blood, 2017, 129, 1646-1657.	1.4	381
42	Autoimmune cytopenias associated with inflammatory bowel diseases: Insights from a multicenter retrospective cohort. Digestive and Liver Disease, 2017, 49, 397-404.	0.9	27
43	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
44	Neutropenia in Patients with Common Variable Immunodeficiency: a Rare Event Associated with Severe Outcome. Journal of Clinical Immunology, 2017, 37, 715-726.	3.8	11
45	Uterine intravascular lymphoma as a cause of fever of unknown origin. Annals of Hematology, 2017, 96, 1891-1896.	1.8	3
46	Persistent risk of adult T-cell leukemia/lymphoma after neonatal HTLV-1 infection through exchange transfusion. International Journal of Hematology, 2017, 105, 859-862.	1.6	0
47	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
48	Classic and extracavitary primary effusion lymphoma in 51 <scp>HIV</scp>â€infected patients from a single institution. American Journal of Hematology, 2016, 91, 233-237.	4.1	89
49	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
50	Hemophagocytic Lymphohistiocytosis Associated WithBartonella henselaeInfection in an HIV-Infected Patient. Clinical Infectious Diseases, 2016, 62, 804-806.	5.8	17
51	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
52	Primary Immunodeficiency Diseases: an Update on the Classification from the International Union of Immunological Societies Expert Committee for Primary Immunodeficiency 2015. Journal of Clinical Immunology, 2015, 35, 696-726.	3.8	621
53	The 2015 IUIS Phenotypic Classification for Primary Immunodeficiencies. Journal of Clinical Immunology, 2015, 35, 727-738.	3.8	199
54	The ratio of mean daily IgG increment/mean daily dose in immunoglobulin replacement therapy in primary antibody deficiencies. Journal of Allergy and Clinical Immunology: in Practice, 2015, 3, 998-1000.e2.	3.8	7

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55	Clinical picture and treatment of 2212 patients with common variable immunodeficiency. Journal of Allergy and Clinical Immunology, 2014, 134, 116-126.e11.	2.9	512
56	Human herpesvirus 8+ polyclonal <scp>I</scp>g<scp>M</scp>Î» <scp>B</scp>â€cell lymphocytosis mimicking plasmablastic leukemia/lymphoma in <scp>HIV</scp>â€infected patients. European Journal of Haematology, 2013, 91, 497-503.	2.2	18
57	Rituximab decreases the risk of lymphoma in patients with HIV-associated multicentric Castleman disease. Blood, 2012, 119, 2228-2233.	1.4	98
58	Confirmation and improvement of criteria for clinical phenotyping in common variable immunodeficiency disorders in replicate cohorts. Journal of Allergy and Clinical Immunology, 2012, 130, 1197-1198.e9.	2.9	129
59	HHV-8 Related Castleman Disease in the Absence of HIV Infection. Blood, 2012, 120, 1578-1578.	1.4	0
60	A survey of 90 patients with autoimmune lymphoproliferative syndrome related to TNFRSF6 mutation. Blood, 2011, 118, 4798-4807.	1.4	153
61	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
62	HIV-associated multicentric Castleman disease. Current Opinion in HIV and AIDS, 2009, 4, 16-21.	3.8	60
63	Infections in 252 Patients with Common Variable Immunodeficiency. Clinical Infectious Diseases, 2008, 46, 1547-1554.	5.8	394
64	Prospective Study of Rituximab in Chemotherapy-Dependent Human Immunodeficiency Virusâ€“Associated Multicentric Castleman's Disease: ANRS 117 CastlemaB Trial. Journal of Clinical Oncology, 2007, 25, 3350-3356.	1.6	220
65	HIV-Associated Non-Hodgkin Lymphoma in 114 Patients with Undetectable HIV Viral Load at NHL Diagnosis.. Blood, 2007, 110, 1578-1578.	1.4	23
66	Improved survival in HIV-related Hodgkin's lymphoma since the introduction of highly active antiretroviral therapy. Aids, 2003, 17, 81-87.	2.2	88
67	Castleman's disease and lymphoma: Report of eight cases in HIV-negative patients and literature review. American Journal of Hematology, 2002, 69, 119-126.	4.1	118
68	Diffuse large B-cell non-Hodgkin's lymphoma in a patient with autoimmune lymphoproliferative syndrome. British Journal of Haematology, 2001, 113, 432-434.	2.5	20
69	High levels of human herpesvirus 8 viral load, human interleukin-6, interleukin-10, and C reactive protein correlate with exacerbation of multicentric Castleman disease in HIV-infected patients. Blood, 2000, 96, 2069-2073.	1.4	325
70	Multicentric CastlemanÊ¼s disease in HIV infection. Aids, 1996, 10, 61-68.	2.2	317