

Eric Oksenhendler

List of Publications by Year in descending order

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Version: 2024-02-01

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	Human Inborn Errors of Immunity: 2019 Update on the Classification from the International Union of Immunological Societies Expert Committee. <i>Journal of Clinical Immunology</i> , 2020, 40, 24-64.	3.8	881
2	International Union of Immunological Societies: 2017 Primary Immunodeficiency Diseases Committee Report on Inborn Errors of Immunity. <i>Journal of Clinical Immunology</i> , 2018, 38, 96-128.	3.8	732
3	Primary Immunodeficiency Diseases: an Update on the Classification from the International Union of Immunological Societies Expert Committee for Primary Immunodeficiency 2015. <i>Journal of Clinical Immunology</i> , 2015, 35, 696-726.	3.8	621
4	Human Inborn Errors of Immunity: 2019 Update of the IUIS Phenotypic Classification. <i>Journal of Clinical Immunology</i> , 2020, 40, 66-81.	3.8	525
5	Clinical picture and treatment of 2212 patients with common variable immunodeficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2014, 134, 116-126.e11.	2.9	512
6	The 2017 IUIS Phenotypic Classification for Primary Immunodeficiencies. <i>Journal of Clinical Immunology</i> , 2018, 38, 129-143.	3.8	488
7	Infections in 252 Patients with Common Variable Immunodeficiency. <i>Clinical Infectious Diseases</i> , 2008, 46, 1547-1554.	5.8	394
8	International, evidence-based consensus diagnostic criteria for HHV-8-negative/idiopathic multicentric Castleman disease. <i>Blood</i> , 2017, 129, 1646-1657.	1.4	381
9	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. <i>Blood</i> , 2000, 96, 2069-2073.	1.4	325
10	Multicentric Castleman's disease in HIV infection. <i>Aids</i> , 1996, 10, 61-68.	2.2	317
11	International, evidence-based consensus treatment guidelines for idiopathic multicentric Castleman disease. <i>Blood</i> , 2018, 132, 2115-2124.	1.4	232
12	Prospective Study of Rituximab in Chemotherapy-Dependent Human Immunodeficiency Virus-associated Multicentric Castleman's Disease: ANRS 117 CastlemaB Trial. <i>Journal of Clinical Oncology</i> , 2007, 25, 3350-3356.	1.6	220
13	Microbial ecology perturbation in human IgA deficiency. <i>Science Translational Medicine</i> , 2018, 10, .	12.4	206
14	The 2015 IUIS Phenotypic Classification for Primary Immunodeficiencies. <i>Journal of Clinical Immunology</i> , 2015, 35, 727-738.	3.8	199
15	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
16	Loss-of-function nuclear factor κ B subunit 1 (NFKB1) variants are the most common monogenic cause of common variable immunodeficiency in Europeans. <i>Journal of Allergy and Clinical Immunology</i> , 2018, 142, 1285-1296.	2.9	185
17	The Ever-Increasing Array of Novel Inborn Errors of Immunity: an Interim Update by the IUIS Committee. <i>Journal of Clinical Immunology</i> , 2021, 41, 666-679.	3.8	165
18	A survey of 90 patients with autoimmune lymphoproliferative syndrome related to TNFRSF6 mutation. <i>Blood</i> , 2011, 118, 4798-4807.	1.4	153

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19	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
20	Confirmation and improvement of criteria for clinical phenotyping in common variable immunodeficiency disorders in replicate cohorts. <i>Journal of Allergy and Clinical Immunology</i> , 2012, 130, 1197-1198.e9.	2.9	129
21	Castleman's disease and lymphoma: Report of eight cases in HIV-negative patients and literature review. <i>American Journal of Hematology</i> , 2002, 69, 119-126.	4.1	118
22	Rituximab decreases the risk of lymphoma in patients with HIV-associated multicentric Castleman disease. <i>Blood</i> , 2012, 119, 2228-2233.	1.4	98
23	International evidence-based consensus diagnostic and treatment guidelines for unicentric Castleman disease. <i>Blood Advances</i> , 2020, 4, 6039-6050.	5.2	94
24	Classic and extracavitary primary effusion lymphoma in 51 HIV-infected patients from a single institution. <i>American Journal of Hematology</i> , 2016, 91, 233-237.	4.1	89
25	Improved survival in HIV-related Hodgkin's lymphoma since the introduction of highly active antiretroviral therapy. <i>Aids</i> , 2003, 17, 81-87.	2.2	88
26	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
27	Mutations in the SRP54 gene cause severe congenital neutropenia as well as Shwachman-Diamond-like syndrome. <i>Blood</i> , 2018, 132, 1318-1331.	1.4	85
28	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
29	Characterization of the clinical and immunologic phenotype and management of 157 individuals with 56 distinct heterozygous NFKB1 mutations. <i>Journal of Allergy and Clinical Immunology</i> , 2020, 146, 901-911.	2.9	78
30	HIV-1 Tat protein induces DNA damage in human peripheral blood B-lymphocytes via mitochondrial ROS production. <i>Redox Biology</i> , 2018, 15, 97-108.	9.0	62
31	HIV-associated multicentric Castleman disease. <i>Current Opinion in HIV and AIDS</i> , 2009, 4, 16-21.	3.8	60
32	Validated international definition of the thrombocytopenia, anasarca, fever, reticulin fibrosis, renal insufficiency, and organomegaly clinical subtype (TAFRO) of idiopathic multicentric Castleman disease. <i>American Journal of Hematology</i> , 2021, 96, 1241-1252.	4.1	47
33	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
34	PROMIS1: A T-cell receptor 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
35	Kaposi sarcoma-associated herpesvirus/human herpesvirus 8-associated lymphoproliferative disorders. <i>Blood</i> , 2019, 133, 1186-1190.	1.4	38
36	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

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37	Loss of ARHGEF1 causes a human primary antibody deficiency. <i>Journal of Clinical Investigation</i> , 2019, 129, 1047-1060.	8.2	32
38	Bortezomib and dexamethasone, an original approach for treating multi-refractory warm autoimmune haemolytic anaemia. <i>British Journal of Haematology</i> , 2019, 187, 124-128.	2.5	29
39	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
40	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
41	HIV-Associated Non-Hodgkin Lymphoma in 114 Patients with Undetectable HIV Viral Load at NHL Diagnosis. <i>Blood</i> , 2007, 110, 1578-1578.	1.4	23
42	Diffuse large B-cell non-Hodgkin's lymphoma in a patient with autoimmune lymphoproliferative syndrome. <i>British Journal of Haematology</i> , 2001, 113, 432-434.	2.5	20
43	CD89 Is a Potent Innate Receptor for Bacteria and Mediates Host Protection from Sepsis. <i>Cell Reports</i> , 2019, 27, 762-775.e5.	6.4	19
44	Human herpesvirus 8+ polyclonal <i>IgM</i> <i>B</i> -cell lymphocytosis mimicking plasmablastic leukemia/lymphoma in HIV-infected patients. <i>European Journal of Haematology</i> , 2013, 91, 497-503.	2.2	18
45	Insufficient evidence exists to use histopathologic subtype to guide treatment of idiopathic multicentric Castleman disease. <i>American Journal of Hematology</i> , 2020, 95, 1553-1561.	4.1	18
46	ACCELERATE: A Patient-Powered Natural History Study Design Enabling Clinical and Therapeutic Discoveries in a Rare Disorder. <i>Cell Reports Medicine</i> , 2020, 1, 100158.	6.5	18
47	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
48	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
49	HIV-1 Tat protein induces aberrant activation of AICDA in human <i>B</i> -lymphocytes from peripheral blood. <i>Journal of Cellular Physiology</i> , 2019, 234, 15678-15685.	4.1	16
50	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
51	Neutropenia in Patients with Common Variable Immunodeficiency: a Rare Event Associated with Severe Outcome. <i>Journal of Clinical Immunology</i> , 2017, 37, 715-726.	3.8	11
52	A Novel Predictive Model for Idiopathic Multicentric Castleman Disease: The International Castleman Disease Consortium Study. <i>Oncologist</i> , 2020, 25, 963-973.	3.7	11
53	First Extensive Analysis of ¹⁸ F-Labeled Fluorodeoxyglucose Positron Emission Tomography-Computed Tomography in a Large Cohort of Patients With HIV-Associated Hodgkin Lymphoma: Baseline Total Metabolic Tumor Volume Affects Prognosis. <i>Journal of Clinical Oncology</i> , 2022, 40, 1346-1355.	1.6	11
54	The French paediatric cohort of Castleman disease: a retrospective report of 23 patients. <i>Orphanet Journal of Rare Diseases</i> , 2020, 15, 95.	2.7	10

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55	The ratio of mean daily IgG increment/mean daily dose in immunoglobulin replacement therapy in primary antibody deficiencies. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2015, 3, 998-1000.e2.	3.8	7
56	Campylobacter infection in adult patients with primary antibody deficiency. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2019, 7, 1038-1041.e4.	3.8	7
57	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
58	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
59	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
60	From Dysgammaglobulinemia to Autosomal-Dominant Activation-Induced Cytidine Deaminase Deficiency: Unraveling an Inherited Immunodeficiency After 50 Years. <i>Journal of Pediatrics</i> , 2020, 223, 207-211.e1.	1.8	4
61	Hepatosplenic Candidiasis in Patients With Hematological Malignancies: A 13-Year Retrospective Cohort Study. <i>Open Forum Infectious Diseases</i> , 2022, 9, ofac088.	0.9	4
62	Uterine intravascular lymphoma as a cause of fever of unknown origin. <i>Annals of Hematology</i> , 2017, 96, 1891-1896.	1.8	3
63	Autoimmune Lymphoproliferative Syndrome Presenting with Invasive Streptococcus pneumoniae Infection. <i>Journal of Clinical Immunology</i> , 2020, 40, 543-546.	3.8	3
64	Inflammatory Waldenström's macroglobulinaemia: A French monocentric retrospective study of 67 patients. <i>British Journal of Haematology</i> , 2022, 197, 728-735.	2.5	3
65	Autoimmune hypoglycemia expands the biological spectrum of HHV8+ multicentric Castleman disease. <i>Blood Advances</i> , 2021, 5, 1848-1852.	5.2	2
66	Persistent risk of adult T-cell leukemia/lymphoma after neonatal HTLV-1 infection through exchange transfusion. <i>International Journal of Hematology</i> , 2017, 105, 859-862.	1.6	0
67	HHV-8 Related Castleman Disease in the Absence of HIV Infection. <i>Blood</i> , 2012, 120, 1578-1578.	1.4	0
68	Phenocopies of Inborn Errors of Immunity. , 2021, , .		0
69	Legãtype 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
70	Hepatitis E infection in adults with primary immunodeficiency with or without immunoglobulin replacement therapy. <i>Blood Transfusion</i> , 2022, , .	0.4	0