

# Eric Jeziorski

## List of Publications by Year in descending order

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Version: 2024-02-01

86  
papers

2,776  
citations

236612

25  
h-index

189595

50  
g-index

120  
all docs

120  
docs citations

120  
times ranked

3951  
citing authors

#	ARTICLE	IF	CITATIONS
1	<i>BRAF</i> Mutation Correlates With High-Risk Langerhans Cell Histiocytosis and Increased Resistance to First-Line Therapy. <i>Journal of Clinical Oncology</i> , 2016, 34, 3023-3030.	0.8	233
2	Autoimmune and inflammatory manifestations occur frequently in patients with primary immunodeficiencies. <i>Journal of Allergy and Clinical Immunology</i> , 2017, 140, 1388-1393.e8.	1.5	222
3	Efficacy of the Janus kinase 1/2 inhibitor ruxolitinib in the treatment of vasculopathy associated with TMEM173 -activating mutations in 3 children. <i>Journal of Allergy and Clinical Immunology</i> , 2016, 138, 1752-1755.	1.5	192
4	A new autoinflammatory and autoimmune syndrome associated with NLRP1 mutations: NAIAD ( <i>NLRP1</i> -associated autoinflammation with arthritis and dyskeratosis). <i>Annals of the Rheumatic Diseases</i> , 2017, 76, 1191-1198.	0.5	181
5	A survey of 90 patients with autoimmune lymphoproliferative syndrome related to TNFRSF6 mutation. <i>Blood</i> , 2011, 118, 4798-4807.	0.6	153
6	Expansion of Regulatory T Cells in Patients with Langerhans Cell Histiocytosis. <i>PLoS Medicine</i> , 2007, 4, e253.	3.9	128
7	Cladribine and cytarabine in refractory multisystem Langerhans cell histiocytosis: results of an international phase 2 study. <i>Blood</i> , 2015, 126, 1415-1423.	0.6	117
8	Pediatric Evans syndrome is associated with a high frequency of potentially damaging variants in immune genes. <i>Blood</i> , 2019, 134, 9-21.	0.6	102
9	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.	2.0	98
10	Medical management of langerhans cell histiocytosis from diagnosis to treatment. <i>Expert Opinion on Pharmacotherapy</i> , 2012, 13, 1309-1322.	0.9	86
11	Langerhans cell histiocytosis: therapeutic strategy and outcome in a 30-year nationwide cohort of 1478 patients under 18 years of age. <i>British Journal of Haematology</i> , 2016, 174, 887-898.	1.2	83
12	Invasive Pneumococcal Disease in Children Can Reveal a Primary Immunodeficiency. <i>Clinical Infectious Diseases</i> , 2014, 59, 244-251.	2.9	75
13	Diagnostic Yield of Next-generation Sequencing in Very Early-onset Inflammatory Bowel Diseases: A Multicentre Study. <i>Journal of Crohn's and Colitis</i> , 2018, 12, 1104-1112.	0.6	68
14	Risk of autoimmune diseases and human papilloma virus (HPV) vaccines: Six years of case-referent surveillance. <i>Journal of Autoimmunity</i> , 2017, 79, 84-90.	3.0	67
15	Circulating cell-free <i>BRAF</i> <sup>V600E</sup> as a biomarker in children with Langerhans cell histiocytosis. <i>British Journal of Haematology</i> , 2017, 178, 457-467.	1.2	57
16	Incidence and risk factors for clinical neurodegenerative Langerhans cell histiocytosis: a longitudinal cohort study. <i>British Journal of Haematology</i> , 2018, 183, 608-617.	1.2	54
17	Pediatric-onset Evans syndrome: Heterogeneous presentation and high frequency of monogenic disorders including LRBA and CTLA4 mutations. <i>Clinical Immunology</i> , 2018, 188, 52-57.	1.4	53
18	Evans Syndrome in Children: Long-Term Outcome in a Prospective French National Observational Cohort. <i>Frontiers in Pediatrics</i> , 2015, 3, 79.	0.9	49

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19	Clinical spectrum and long-term follow-up of 14 cases with G6PC3 mutations from the French severe congenital neutropenia registry. <i>Orphanet Journal of Rare Diseases</i> , 2014, 9, 183.	1.2	48
20	Herpes-Virus Infection in Patients with Langerhans Cell Histiocytosis: A Case-Controlled Sero-Epidemiological Study, and In Situ Analysis. <i>PLoS ONE</i> , 2008, 3, e3262.	1.1	48
21	Thymus and mediastinal node involvement in childhood langerhans cell histiocytosis: Long-term follow-up from the French national cohort. <i>Pediatric Blood and Cancer</i> , 2013, 60, 1759-1765.	0.8	35
22	Inherited GATA2 Deficiency Is Dominant by Haploinsufficiency and Displays Incomplete Clinical Penetrance. <i>Journal of Clinical Immunology</i> , 2021, 41, 639-657.	2.0	30
23	KI and WU Polyomaviruses in Children, France. <i>Emerging Infectious Diseases</i> , 2008, 14, 523-525.	2.0	29
24	Bier anaemic spots, cyanosis with urticaria-like eruption (<scp>BASCULE</scp>) syndrome: a new entity?. <i>British Journal of Dermatology</i> , 2016, 175, 218-220.	1.4	27
25	Physical health conditions and quality of life in adults with primary immunodeficiency diagnosed during childhood: A French Reference Center for PIDs (CEREDIH) study. <i>Journal of Allergy and Clinical Immunology</i> , 2017, 139, 1275-1281.e7.	1.5	26
26	B-cell polyclonal activation and Epstein-Barr viral abortive lytic cycle are two key features in acute infectious mononucleosis. <i>Journal of Clinical Virology</i> , 2011, 52, 33-37.	1.6	25
27	Benefits of rituximab as a second-line treatment for autoimmune haemolytic anaemia in children: a prospective French cohort study. <i>British Journal of Haematology</i> , 2017, 177, 751-758.	1.2	24
28	Current insights in invasive group A streptococcal infections in pediatrics. <i>European Journal of Pediatrics</i> , 2012, 171, 1589-1598.	1.3	22
29	Genetic diagnosis of primary immunodeficiencies: A survey of the French national registry. <i>Journal of Allergy and Clinical Immunology</i> , 2019, 143, 1646-1649.e10.	1.5	20
30	No evidence for XMRV association in pediatric idiopathic diseases in France. <i>Retrovirology</i> , 2010, 7, 63.	0.9	18
31	Synovial-Fluid miRNA Signature for Diagnosis of Juvenile Idiopathic Arthritis. <i>Cells</i> , 2019, 8, 1521.	1.8	18
32	Alemtuzumab as First Line Treatment in Children with Familial Lymphohistiocytosis. <i>Blood</i> , 2019, 134, 80-80.	0.6	18
33	Differential Expression of Interferon-Alpha Protein Provides Clues to Tissue Specificity Across Type I Interferonopathies. <i>Journal of Clinical Immunology</i> , 2021, 41, 603-609.	2.0	16
34	BASCULE syndrome, orthostatic cyanosis and postural orthostatic tachycardia syndrome: time for decanting old wine?. <i>British Journal of Dermatology</i> , 2016, 175, 1110-1111.	1.4	15
35	Burden of Poor Health Conditions and Quality of Life in 656 Children with Primary Immunodeficiency. <i>Journal of Pediatrics</i> , 2018, 194, 211-217.e5.	0.9	15
36	Intravenous Immunoglobulins for Neonatal Alloimmune Neutropenia Refractory to Recombinant Human Granulocyte Colony-Stimulating Factor. <i>American Journal of Perinatology</i> , 2011, 28, 461-466.	0.6	14

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37	Long-term follow-up of children with risk organ-negative Langerhans cell histiocytosis after 2-chlorodeoxyadenosine treatment. <i>British Journal of Haematology</i> , 2020, 191, 825-834.	1.2	14
38	Clinical, functional and genetic characterization of 16 patients suffering from chronic granulomatous disease variants—Identification of 11 novel mutations in CYBB. <i>Clinical and Experimental Immunology</i> , 2021, 203, 247-266.	1.1	14
39	Relevance of Human Parechovirus Detection in Cerebrospinal Fluid Samples From Young Infants With Sepsis-Like Illness. <i>Journal of Clinical Laboratory Analysis</i> , 2015, 29, 112-115.	0.9	13
40	Short-course antibiotic treatment of bone and joint infections in children: a retrospective study at Montpellier University Hospital from 2009 to 2013. <i>Journal of Antimicrobial Chemotherapy</i> , 2019, 74, 3579-3587.	1.3	13
41	Differential Accumulation and Activation of Monocyte and Dendritic Cell Subsets in Inflamed Synovial Fluid Discriminates Between Juvenile Idiopathic Arthritis and Septic Arthritis. <i>Frontiers in Immunology</i> , 2020, 11, 1716.	2.2	13
42	Intrafamilial Cluster of Pulmonary Tuberculosis Due to <i>Mycobacterium bovis</i> of the African 1 Clonal Complex. <i>Journal of Clinical Microbiology</i> , 2010, 48, 4680-4683.	1.8	12
43	Second-line treatment trends and long-term outcomes of 392 children with chronic immune thrombocytopenic purpura: the French experience over the past 25 years. <i>British Journal of Haematology</i> , 2020, 189, 931-942.	1.2	12
44	Gene editing rescue of a novel MPL mutant associated with congenital amegakaryocytic thrombocytopenia. <i>Blood Advances</i> , 2017, 1, 1815-1826.	2.5	11
45	Mucosal relapse of visceral leishmaniasis in a child treated with anti-TNF. <i>International Journal of Infectious Diseases</i> , 2015, 33, 135-136.	1.5	10
46	The French paediatric cohort of Castleman disease: a retrospective report of 23 patients. <i>Orphanet Journal of Rare Diseases</i> , 2020, 15, 95.	1.2	10
47	Long term follow-up of pediatric-onset Evans syndrome: broad immunopathological manifestations and high treatment burden. <i>Haematologica</i> , 2022, 107, 457-466.	1.7	9
48	Risk factors of clinical dysimmune manifestations in a cohort of 86 children with 22q11.2 deletion syndrome: A retrospective study in France. <i>American Journal of Medical Genetics, Part A</i> , 2019, 179, 2207-2213.	0.7	8
49	Two neurologic facets of CTLA4-related haploinsufficiency. <i>Neurology: Neuroimmunology and Neuroinflammation</i> , 2020, 7, .	3.1	8
50	An appraisal of the frequency and severity of noninfectious manifestations in primary immunodeficiencies: A study of a national retrospective cohort of 1375 patients over 10 years. <i>Journal of Allergy and Clinical Immunology</i> , 2022, 149, 2116-2125.	1.5	7
51	Atypical Pneumonia Linked to Community-Acquired <i>Staphylococcus aureus</i> ; Cross-Transmission in the Nursery. <i>Neonatology</i> , 2013, 104, 156-160.	0.9	6
52	What to expect from molecular tools for non-documented pediatric infectious diseases. <i>Expert Review of Molecular Diagnostics</i> , 2015, 15, 1645-1656.	1.5	6
53	Implementation of an organizational infrastructure paediatric plan adapted to bronchiolitis epidemics. <i>Journal of Infection and Public Health</i> , 2020, 13, 167-172.	1.9	6
54	Determinants of long-term outcomes of splenectomy in pediatric autoimmune cytopenias. <i>Blood</i> , 2022, 140, 253-261.	0.6	6

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55	Safety of the Northern Hemisphere 2014/2015 formulation of the inactivated split-virion intramuscular trivalent influenza vaccine. <i>Vaccine Reports</i> , 2016, 6, 1-7.	1.2	5
56	Survey of <i>Staphylococcus aureus</i> in a general pediatric population and focus on isolates with three clinically relevant toxin-encoding genes. <i>World Journal of Pediatrics</i> , 2018, 14, 35-43.	0.8	5
57	Clinical, virological and immunological features of HIV-positive children internationally adopted in France from 2005-2015. <i>PLoS ONE</i> , 2018, 13, e0203438.	1.1	5
58	Pediatric deep neck infections: Clinical description and analysis of therapeutic management. <i>Archives De Pediatrie</i> , 2022, 29, 128-132.	0.4	5
59	Treatment with cyclosporin in auto-immune cytopenias in children: The experience from the French cohort OBS'CEREVANCE. <i>American Journal of Hematology</i> , 2018, 93, E196.	2.0	4
60	Epidemiological survey in a day care center following toddler sudden death due to human metapneumovirus infection. <i>Archives De Pediatrie</i> , 2019, 26, 479-482.	0.4	4
61	Idiopathic purpura fulminans with anti-PS antibodies in children: a multicenter case series and systematic review. <i>Blood Advances</i> , 2021, , .	2.5	4
62	Complications in the Subacute Phase of Invasive <i>Streptococcus pyogenes</i> Infections in Pediatrics. <i>Clinical Pediatrics</i> , 2014, 53, 191-193.	0.4	3
63	Torticollis in Children. <i>Clinical Pediatrics</i> , 2016, 55, 459-462.	0.4	3
64	Pediatric angiostrongyliasis. <i>Medecine Et Sante Tropicales</i> , 2018, 28, 76-81.	0.3	3
65	A 1-Year Prospective French Nationwide Study of Emergency Hospital Admissions in Children and Adults with Primary Immunodeficiency. <i>Journal of Clinical Immunology</i> , 2019, 39, 702-712.	2.0	3
66	When Familial Hearing Loss Means Genetic Heterogeneity: A Model Case Report. <i>Diagnostics</i> , 2021, 11, 1636.	1.3	3
67	Lymphoma-Like Syndrome. <i>Medicine (United States)</i> , 2015, 94, e855.	0.4	2
68	Thrombopoietin receptor agonists as an emergency treatment for severe newly diagnosed immune thrombocytopenia in children. <i>Blood</i> , 2021, 137, 138-141.	0.6	2
69	Outpatient central venous access device insertion in very young children with severe haemophilia. <i>Blood Coagulation and Fibrinolysis</i> , 2020, 31, 490-492.	0.5	2
70	Searching for Common Mammalian Retroviruses in Pediatric Idiopathic Diseases. <i>Viruses</i> , 2016, 8, 86.	1.5	1
71	Case Report: Persistency Pneumococcal Polysaccharide in Cerebrospinal Fluid During a Post Pneumococcal Chronic Aseptic Meningitis: Coincidental or (Auto-)Inflammatory Embers. <i>Frontiers in Pediatrics</i> , 2022, 10, 762457.	0.9	1
72	SFP CO-03 - Syndrome de Di George et manifestations auto immunes, À propos de 15 cas. <i>Archives De Pediatrie</i> , 2014, 21, 656.	0.4	0

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73	FRI0509â€¦Chromosome 22Q11.2 Deletion Syndrome (Digeorge Syndrome) and Autoimmunity: A French Retrospective Pediatric Study of 15 Cases. <i>Annals of the Rheumatic Diseases</i> , 2015, 74, 613.2-613.	0.5	0
74	CO-30 â€œ Couverture vaccinale de l'entourage des enfants immunodÃ©primÃ©s. <i>Archives De Pediatrie</i> , 2015, 22, 217-218.	0.4	0
75	Identification of rare genetic variants in Juvenile Idiopathic Arthritis using whole exome sequencing. <i>Pediatric Rheumatology</i> , 2015, 13, .	0.9	0
76	CO-63 â€œ Ã‰volution de 'Ã©pidÃ©miologie de la bronchiolite aiguÃ© du nourrisson au CHU de Montpellier de 2007 Ã 2013. <i>Archives De Pediatrie</i> , 2015, 22, 226.	0.4	0
77	Severe hypotonia and respiratory failure in a one month old boy with complete recovery. <i>European Journal of Paediatric Neurology</i> , 2017, 21, e225-e226.	0.7	0
78	P165â€¦Molecular and cellular biomarkers that discriminate juvenile idiopathic arthritis from septic arthritis. , 2019, , .		0
79	SAT0488â€¦THE FRENCH PAEDIATRIC COHORT OF CASTLEMAN DISEASE. , 2019, , .		0
80	FiÃ©vre chez l'enfant sous chimiothÃ©rapie. <i>Perfectionnement En PÃ©diatrie</i> , 2021, 4, S31-S33.	0.0	0
81	Langerhans cell histiocytosis in children: Correlation of <i>BRAF</i> status with clinical characteristic.. <i>Journal of Clinical Oncology</i> , 2015, 33, 10003-10003.	0.8	0
82	In Vitro Functional Rescue of a Double MPL K39N/W272R Mutant Associated with Congenital Amegakaryocytic Thrombocytopenia (CAMT) Using Crispr/Cas9. <i>Blood</i> , 2015, 126, 1207-1207.	0.6	0
83	Treatment-Resistant Actinomycosis Reveals Concomitant Systemic Vasculitis. <i>Scholars Journal of Medical Case Reports</i> , 2020, 8, 623-626.	0.0	0
84	Evaluation of Post-Infectious Inflammatory Reactions in a Retrospective Study for Three Common Invasive Bacterial Infections in Pediatrics. <i>SSRN Electronic Journal</i> , 0, , .	0.4	0
85	Impact and Dynamics of <i>TP53</i> Mutated Clones in Shwachman Diamond Syndrome in a Series of 80 Patients. <i>Blood</i> , 2020, 136, 22-23.	0.6	0
86	Pediatric-Onset Evans Syndrome Is Associated with Broad Immunopathological Manifestations, High Treatment Burden and Mortality in Long-Term Follow-up. <i>Blood</i> , 2020, 136, 20-22.	0.6	0