Eric Jeziorski

List of Publications by Year
in descending order

86 papers	2,776 citations	236925 25 h-index	189892 50 g-index
120 all docs	120 docs citations	120 times ranked	3951 citing authors

#	Article	IF	Citations
1	Long term follow-up of pediatric-onset Evans syndrome: broad immunopathological manifestations and high treatment burden. Haematologica, 2022, 107, 457-466.	3.5	9
2	An appraisal of the frequency and severity of noninfectious manifestations in primary immunodeficiencies: AÂstudy of a national retrospective cohort of 1375 patients over 10 years. Journal of Allergy and Clinical Immunology, 2022, 149, 2116-2125.	2.9	7
3	Pediatric deep neck infections: Clinical description and analysis of therapeutic management. Archives De Pediatrie, 2022, 29, 128-132.	1.0	5
4	Case Report: Persistency Pneumococcal Polysaccharide in Cerebrospinal Fluid During a Post Pneumococcal Chronic Aseptic Meningitis: Coincidental or (Auto-)Inflammatory Embers. Frontiers in Pediatrics, 2022, 10, 762457.	1.9	1
5	Determinants of long-term outcomes of splenectomy in pediatric autoimmune cytopenias. Blood, 2022, 140, 253-261.	1.4	6
6	Thrombopoietin receptor agonists as an emergency treatment for severe newly diagnosed immune thrombocytopenia in children. Blood, 2021, 137, 138-141.	1.4	2
7	Clinical, functional and genetic characterization of 16 patients suffering from chronic granulomatous disease variantsÂâ€“Âidentification of 11 novel mutations in CYBB. Clinical and Experimental Immunology, 2021, 203, 247-266.	2.6	14
8	Overview of STING-Associated Vasculopathy with Onset in Infancy (SAVI) Among 21 Patients. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 803-818.e11.	3.8	98
9	Inherited GATA2 Deficiency Is Dominant by Haploinsufficiency and Displays Incomplete Clinical Penetrance. Journal of Clinical Immunology, 2021, 41, 639-657.	3.8	30
10	Differential Expression of Interferon-Alpha Protein Provides Clues to Tissue Specificity Across Type I Interferonopathies. Journal of Clinical Immunology, 2021, 41, 603-609.	3.8	16
11	FiÃ¨vre chez l'enfant sous chimiothÃ©rapie. Perfectionnement En PÃ©diatrie, 2021, 4, S31-S33.	0.0	0
12	When Familial Hearing Loss Means Genetic Heterogeneity: A Model Case Report. Diagnostics, 2021, 11, 1636.	2.6	3
13	Idiopathic purpura fulminans with anti-PS antibodies in children: a multicenter case series and systematic review. Blood Advances, 2021, , .	5.2	4
14	Implementation of an organizational infrastructure paediatric plan adapted to bronchiolitis epidemics. Journal of Infection and Public Health, 2020, 13, 167-172.	4.1	6
15	Longâ€term followâ€up of children with risk organâ€negative Langerhans cell histiocytosis after 2â€chlorodeoxyadenosine treatment. British Journal of Haematology, 2020, 191, 825-834.	2.5	14
16	Differential Accumulation and Activation of Monocyte and Dendritic Cell Subsets in Inflamed Synovial Fluid Discriminates Between Juvenile Idiopathic Arthritis and Septic Arthritis. Frontiers in Immunology, 2020, 11, 1716.	4.8	13
17	Two neurologic facets of CTLA4-related haploinsufficiency. Neurology: Neuroimmunology and NeuroInflammation, 2020, 7, .	6.0	8
18	Secondâ€line treatment trends and longâ€term outcomes of 392 children with chronic immune thrombocytopenic purpura: the French experience over the past 25Âyears. British Journal of Haematology, 2020, 189, 931-942.	2.5	12

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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	Treatment-Resistant Actinomycosis Reveals Concomitant Systemic Vasculitis. Scholars Journal of Medical Case Reports, 2020, 8, 623-626.	0.0	0
21	Outpatient central venous access device insertion in very young children with severe haemophilia. Blood Coagulation and Fibrinolysis, 2020, 31, 490-492.	1.0	2
22	Impact and Dynamics of TP53 Mutated Clones in Shwachman Diamond Syndrome in a Series of 80 Patients. Blood, 2020, 136, 22-23.	1.4	0
23	Pediatric-Onset Evans Syndrome Is Associated with Broad Immunopathological Manifestations, High Treatment Burden and Mortality in Long-Term Follow-up. Blood, 2020, 136, 20-22.	1.4	0
24	A 1-Year Prospective French Nationwide Study of Emergency Hospital Admissions in Children and Adults with Primary Immunodeficiency. Journal of Clinical Immunology, 2019, 39, 702-712.	3.8	3
25	Risk factors of clinical dysimmune manifestations in a cohort of 86 children with 22q11.2 deletion syndrome: A retrospective study in France. American Journal of Medical Genetics, Part A, 2019, 179, 2207-2213.	1.2	8
26	Short-course antibiotic treatment of bone and joint infections in children: a retrospective study at Montpellier University Hospital from 2009 to 2013. Journal of Antimicrobial Chemotherapy, 2019, 74, 3579-3587.	3.0	13
27	Pediatric Evans syndrome is associated with a high frequency of potentially damaging variants in immune genes. Blood, 2019, 134, 9-21.	1.4	102
28	P165â€...Molecular and cellular biomarkers that discriminate juvenile idiopathic arthritis from septic arthritis. , 2019, , .		0
29	SAT0488â€...THE FRENCH PAEDIATRIC COHORT OF CASTLEMAN DISEASE. , 2019, , .		0
30	Epidemiological survey in a day care center following toddler sudden death due to human metapneumovirus infection. Archives De Pediatrie, 2019, 26, 479-482.	1.0	4
31	Synovial-Fluid miRNA Signature for Diagnosis of Juvenile Idiopathic Arthritis. Cells, 2019, 8, 1521.	4.1	18
32	Genetic diagnosis of primary immunodeficiencies: AÂsurvey of the French national registry. Journal of Allergy and Clinical Immunology, 2019, 143, 1646-1649.e10.	2.9	20
33	Alemtuzumab as First Line Treatment in Children with Familial Lymphohistiocytosis. Blood, 2019, 134, 80-80.	1.4	18
34	Survey of Staphylococcus aureus in a general pediatric population and focus on isolates with three clinically relevant toxin-encoding genes. World Journal of Pediatrics, 2018, 14, 35-43.	1.8	5
35	Pediatric-onset Evans syndrome: Heterogeneous presentation and high frequency of monogenic disorders including LRBA and CTLA4 mutations. Clinical Immunology, 2018, 188, 52-57.	3.2	53
36	Incidence and risk factors for clinical neurodegenerative Langerhans cell histiocytosis: a longitudinal cohort study. British Journal of Haematology, 2018, 183, 608-617.	2.5	54

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37	Clinical, virological and immunological features of HIV-positive children internationally adopted in France from 2005-2015. PLoS ONE, 2018, 13, e0203438.	2.5	5
38	Diagnostic Yield of Next-generation Sequencing in Very Early-onset Inflammatory Bowel Diseases: A Multicentre Study. Journal of Crohn's and Colitis, 2018, 12, 1104-1112.	1.3	68
39	Treatment with cyclosporin in autoâ€immune cytopenias in children: The experience from the French cohort OBS'CEREVANCE. American Journal of Hematology, 2018, 93, E196.	4.1	4
40	Pediatric angiostrongyliasis. Medecine Et Sante Tropicales, 2018, 28, 76-81.	0.3	3
41	Burden of Poor Health Conditions and Quality of Life in 656 Children with Primary Immunodeficiency. Journal of Pediatrics, 2018, 194, 211-217.e5.	1.8	15
42	Autoimmune and inflammatory manifestations occur frequently in patients with primary immunodeficiencies. Journal of Allergy and Clinical Immunology, 2017, 140, 1388-1393.e8.	2.9	222
43	Risk of autoimmune diseases and human papilloma virus (HPV) vaccines: Six years of case-referent surveillance. Journal of Autoimmunity, 2017, 79, 84-90.	6.5	67
44	Circulating cellâ€free BRAF^{V600E} as a biomarker in children with Langerhans cell histiocytosis. British Journal of Haematology, 2017, 178, 457-467.	2.5	57
45	Benefits of rituximab as a secondâ€line treatment for autoimmune haemolytic anaemia in children: a prospective French cohort study. British Journal of Haematology, 2017, 177, 751-758.	2.5	24
46	A new autoinflammatory and autoimmune syndrome associated with NLRP1 mutations: NAIAD (NLRP1-associated autoinflammation with arthritis and dyskeratosis). Annals of the Rheumatic Diseases, 2017, 76, 1191-1198.	0.9	181
47	Severe hypotonia and respiratory failure in a one month old boy with complete recovery. European Journal of Paediatric Neurology, 2017, 21, e225-e226.	1.6	0
48	Physical health conditions and quality of life in adults with primary immunodeficiency diagnosed during childhood: AÂFrench Reference Center for PIDs (CEREDIH) study. Journal of Allergy and Clinical Immunology, 2017, 139, 1275-1281.e7.	2.9	26
49	Gene editing rescue of a novel MPL mutant associated with congenital amegakaryocytic thrombocytopenia. Blood Advances, 2017, 1, 1815-1826.	5.2	11
50	Searching for Common Mammalian Retroviruses in Pediatric Idiopathic Diseases. Viruses, 2016, 8, 86.	3.3	1
51	BRAF Mutation Correlates With High-Risk Langerhans Cell Histiocytosis and Increased Resistance to First-Line Therapy. Journal of Clinical Oncology, 2016, 34, 3023-3030.	1.6	233
52	Langerhans cell histiocytosis: therapeutic strategy and outcome in a 30â€year nationwide cohort of 1478 patients under 18Âyears of age. British Journal of Haematology, 2016, 174, 887-898.	2.5	83
53	Safety of the Northern Hemisphere 2014/2015 formulation of the inactivated split-virion intramuscular trivalent influenza vaccine. Vaccine Reports, 2016, 6, 1-7.	1.2	5
54	BASCULE syndrome, orthostatic cyanosis and postural orthostatic tachycardia syndrome: time for decanting old wine?. British Journal of Dermatology, 2016, 175, 1110-1111.	1.5	15

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55	Efficacy of the Janus kinase 1/2 inhibitor ruxolitinib in the treatment of vasculopathy associated with TMEM173 -activating mutations in 3 children. Journal of Allergy and Clinical Immunology, 2016, 138, 1752-1755.	2.9	192
56	Bier anaemic spots, cyanosis with urticariaâ€like eruption (<scp>BASCULE</scp>) syndrome: a new entity?. British Journal of Dermatology, 2016, 175, 218-220.	1.5	27
57	Torticollis in Children. Clinical Pediatrics, 2016, 55, 459-462.	0.8	3
58	Cladribine and cytarabine in refractory multisystem Langerhans cell histiocytosis: results of an international phase 2 study. Blood, 2015, 126, 1415-1423.	1.4	117
59	FRI0509â€…Chromosome 22Q11.2 Deletion Syndrome (Digeorge Syndrome) and Autoimmunity: A French Retrospective Pediatric Study of 15 Cases. Annals of the Rheumatic Diseases, 2015, 74, 613.2-613.	0.9	0
60	CO-30 â€“ Couverture vaccinale de l'entourage des enfants immunodÃ©primÃ©s. Archives De Pediatrie, 2015, 22, 217-218.	1.0	0
61	Lymphoma-Like Syndrome. Medicine (United States), 2015, 94, e855.	1.0	2
62	Evans Syndrome in Children: Long-Term Outcome in a Prospective French National Observational Cohort. Frontiers in Pediatrics, 2015, 3, 79.	1.9	49
63	Identification of rare genetic variants in Juvenile Idiopathic Arthritis using whole exome sequencing. Pediatric Rheumatology, 2015, 13, .	2.1	0
64	Relevance of Human Parechovirus Detection in Cerebrospinal Fluid Samples From Young Infants With Sepsis-Like Illness. Journal of Clinical Laboratory Analysis, 2015, 29, 112-115.	2.1	13
65	What to expect from molecular tools for non-documented pediatric infectious diseases. Expert Review of Molecular Diagnostics, 2015, 15, 1645-1656.	3.1	6
66	Mucosal relapse of visceral leishmaniasis in a child treated with anti-TNFÎ±. International Journal of Infectious Diseases, 2015, 33, 135-136.	3.3	10
67	CO-63 â€“ Ã‰volution de 'Ã©pidÃ©miologie de la bronchiolite aiguÃ« du nourrisson au CHU de Montpellier de 2007 Ã 2013. Archives De Pediatrie, 2015, 22, 226.	1.0	0
68	Langerhans cell histiocytosis in children: Correlation of BRAF status with clinical characteristic.. Journal of Clinical Oncology, 2015, 33, 10003-10003.	1.6	0
69	In Vitro Functional Rescue of a Double MPL K39N/W272R Mutant Associated with Congenital Amegakaryocytic Thrombocytopenia (CAMT) Using Crispr/Cas9. Blood, 2015, 126, 1207-1207.	1.4	0
70	Clinical spectrum and long-term follow-up of 14 cases with G6PC3 mutations from the French severe congenital neutropenia registry. Orphanet Journal of Rare Diseases, 2014, 9, 183.	2.7	48
71	Complications in the Subacute Phase of Invasive Streptococcus pyogenes Infections in Pediatrics. Clinical Pediatrics, 2014, 53, 191-193.	0.8	3
72	Invasive Pneumococcal Disease in Children Can Reveal a Primary Immunodeficiency. Clinical Infectious Diseases, 2014, 59, 244-251.	5.8	75

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73	SFP CO-03 - Syndrome de Di George et manifestations auto immunes, Ã propos de 15 cas. Archives De Pediatrie, 2014, 21, 656.	1.0	0
74	Atypical Pneumonia Linked to Community-Acquired <i>Staphylococcus aureus</i> Cross-Transmission in the Nursery. Neonatology, 2013, 104, 156-160.	2.0	6
75	Thymus and mediastinal node involvement in childhood langerhans cell histiocytosis: Longâ€term followâ€up from the French national cohort. Pediatric Blood and Cancer, 2013, 60, 1759-1765.	1.5	35
76	Current insights in invasive group A streptococcal infections in pediatrics. European Journal of Pediatrics, 2012, 171, 1589-1598.	2.7	22
77	Medical management of langerhans cell histiocytosis from diagnosis to treatment. Expert Opinion on Pharmacotherapy, 2012, 13, 1309-1322.	1.8	86
78	B-cell polyclonal activation and Epstein-Barr viral abortive lytic cycle are two key features in acute infectious mononucleosis. Journal of Clinical Virology, 2011, 52, 33-37.	3.1	25
79	A survey of 90 patients with autoimmune lymphoproliferative syndrome related to TNFRSF6 mutation. Blood, 2011, 118, 4798-4807.	1.4	153
80	Intravenous Immunoglobulins for Neonatal Alloimmune Neutropenia Refractory to Recombinant Human Granulocyte Colony-Stimulating Factor. American Journal of Perinatology, 2011, 28, 461-466.	1.4	14
81	No evidence for XMRV association in pediatric idiopathic diseases in France. Retrovirology, 2010, 7, 63.	2.0	18
82	Intrafamilial Cluster of Pulmonary Tuberculosis Due to Mycobacterium bovis of the African 1 Clonal Complex. Journal of Clinical Microbiology, 2010, 48, 4680-4683.	3.9	12
83	KI and WU Polyomaviruses in Children, France. Emerging Infectious Diseases, 2008, 14, 523-525.	4.3	29
84	Herpes-Virus Infection in Patients with Langerhans Cell Histiocytosis: A Case-Controlled Sero-Epidemiological Study, and In Situ Analysis. PLoS ONE, 2008, 3, e3262.	2.5	48
85	Expansion of Regulatory T Cells in Patients with Langerhans Cell Histiocytosis. PLoS Medicine, 2007, 4, e253.	8.4	128
86	Evaluation of Post-Infectious Inflammatory Reactions in a Retrospective Study for Three Common Invasive Bacterial Infections in Pediatrics. SSRN Electronic Journal, 0, , .	0.4	0