

Claire Fieschi

List of Publications by Year in descending order

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Version: 2024-02-01

24
papers

1,175
citations

516710

16
h-index

677142

22
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docs citations

24
times ranked

2314
citing authors

#	ARTICLE	IF	CITATIONS
1	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
2	Legã€type 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
3	Hepatitis E infection in adults with primary immunodeficiency with or without immunoglobulin replacement therapy.. <i>Blood Transfusion</i> , 2022, , .	0.4	0
4	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
5	Coronavirus disease 2019 in patients with inborn errors of immunity: An international study. <i>Journal of Allergy and Clinical Immunology</i> , 2021, 147, 520-531.	2.9	278
6	Inherited GATA2 Deficiency Is Dominant by Haploinsufficiency and Displays Incomplete Clinical Penetrance. <i>Journal of Clinical Immunology</i> , 2021, 41, 639-657.	3.8	30
7	Autoimmune hypoglycemia expands the biological spectrum of HHV8+ multicentric Castleman disease. <i>Blood Advances</i> , 2021, 5, 1848-1852.	5.2	2
8	Recurrent bacterial infections, but not fungal infections, characterise patients with <i>ELANE</i>-related neutropenia: a French Severe Chronic Neutropenia Registry study. <i>British Journal of Haematology</i> , 2021, 194, 908-920.	2.5	11
9	Biochemically deleterious human <i>NFKB1</i> variants underlie an autosomal dominant form of common variable immunodeficiency. <i>Journal of Experimental Medicine</i> , 2021, 218, .	8.5	32
10	The expansion of human T-bet ^{high} CD21 ^{low} B cells is T cell dependent. <i>Science Immunology</i> , 2021, 6, eabh0891.	11.9	82
11	Human IgA binds a diverse array of commensal bacteria. <i>Journal of Experimental Medicine</i> , 2020, 217, .	8.5	65
12	Dominant-negative mutations in human <i>IL6ST</i> underlie hyper-IgE syndrome. <i>Journal of Experimental Medicine</i> , 2020, 217, .	8.5	64
13	PROMIDISÎ±: AÎ±T-cell receptor Î± 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
14	Clinical and Genetic Spectrum of a Large Cohort With Total and Sub-total Complement Deficiencies. <i>Frontiers in Immunology</i> , 2019, 10, 1936.	4.8	34
15	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
16	Chronic mucocutaneous candidiasis and connective tissue disorder in humans with impaired JNK1-dependent responses to IL-17A/F and TGF-Î². <i>Science Immunology</i> , 2019, 4, .	11.9	45
17	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
18	The full spectrum of Castleman disease: 273 patients studied over 20Î±years. <i>British Journal of Haematology</i> , 2018, 180, 206-216.	2.5	137

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19	Mutations in the SRP54 gene cause severe congenital neutropenia as well as Shwachman-Diamondâ€™like syndrome. <i>Blood</i> , 2018, 132, 1318-1331.	1.4	85
20	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
21	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
22	Strains Responsible for Invasive Meningococcal Disease in Patients With Terminal Complement Pathway Deficiencies. <i>Journal of Infectious Diseases</i> , 2017, 215, 1331-1338.	4.0	35
23	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
24	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