Claire Fieschi

List of Publications by Year
in descending order

24 papers	1,175 citations	516710 16 h-index	677142 22 g-index
24 all docs	24 docs citations	24 times ranked	2314 citing authors

#	Article	IF	Citations
1	Coronavirus disease 2019 in patients with inborn errors of immunity: An international study. Journal of Allergy and Clinical Immunology, 2021, 147, 520-531.	2.9	278
2	The full spectrum of Castleman disease: 273 patients studied over 20Âyears. British Journal of Haematology, 2018, 180, 206-216.	2.5	137
3	Synergistic convergence of microbiota-specific systemic IgG and secretory IgA. Journal of Allergy and Clinical Immunology, 2019, 143, 1575-1585.e4.	2.9	86
4	Mutations in the SRP54 gene cause severe congenital neutropenia as well as Shwachman-Diamondâ€“like syndrome. Blood, 2018, 132, 1318-1331.	1.4	85
5	The expansion of human T-bet ^{high} CD21 ^{low} B cells is T cell dependent. Science Immunology, 2021, 6, eabh0891.	11.9	82
6	Human IgA binds a diverse array of commensal bacteria. Journal of Experimental Medicine, 2020, 217, .	8.5	65
7	Dominant-negative mutations in human IL6ST underlie hyper-IgE syndrome. Journal of Experimental Medicine, 2020, 217, .	8.5	64
8	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
9	Chronic mucocutaneous candidiasis and connective tissue disorder in humans with impaired JNK1-dependent responses to IL-17A/F and TGF-Î². Science Immunology, 2019, 4, .	11.9	45
10	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
11	Treatment and outcome of Unicentric Castleman Disease: a retrospective analysis of 71 cases. British Journal of Haematology, 2019, 186, 269-273.	2.5	36
12	Strains Responsible for Invasive Meningococcal Disease in Patients With Terminal Complement Pathway Deficiencies. Journal of Infectious Diseases, 2017, 215, 1331-1338.	4.0	35
13	Clinical and Genetic Spectrum of a Large Cohort With Total and Sub-total Complement Deficiencies. Frontiers in Immunology, 2019, 10, 1936.	4.8	34
14	Biochemically deleterious human NFKB1 variants underlie an autosomal dominant form of common variable immunodeficiency. Journal of Experimental Medicine, 2021, 218, .	8.5	32
15	Inherited GATA2 Deficiency Is Dominant by Haploinsufficiency and Displays Incomplete Clinical Penetrance. Journal of Clinical Immunology, 2021, 41, 639-657.	3.8	30
16	Autoimmune cytopenias associated with inflammatory bowel diseases: Insights from a multicenter retrospective cohort. Digestive and Liver Disease, 2017, 49, 397-404.	0.9	27
17	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
18	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

#	Article	IF	Citations
19	Recurrent bacterial infections, but not fungal infections, characterise patients with ELANEâ€related neutropenia: a French Severe Chronic Neutropenia Registry study. British Journal of Haematology, 2021, 194, 908-920.	2.5	11
20	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
21	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
22	Autoimmune hypoglycemia expands the biological spectrum of HHV8+ multicentric Castleman disease. Blood Advances, 2021, 5, 1848-1852.	5.2	2
23	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
24	Hepatitis E infection in adults with primary immunodeficiency with or without immunoglobulin replacement therapy.. Blood Transfusion, 2022, , .	0.4	0