FrÃ©dÃ©ric Rieux-Laucat

List of Publications by Year
in descending order

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160 papers	18,063 citations	22099 59 h-index	14702 127 g-index
169 all docs	169 docs citations	169 times ranked	23872 citing authors

#	Article	IF	Citations
1	Impaired type I interferon activity and inflammatory responses in severe COVID-19 patients. Science, 2020, 369, 718-724.	6.0	2,374
2	Autoantibodies against type I IFNs in patients with life-threatening COVID-19. Science, 2020, 370, .	6.0	1,983
3	Mutations in Fas associated with human lymphoproliferative syndrome and autoimmunity. Science, 1995, 268, 1347-1349.	6.0	1,221
4	XIAP deficiency in humans causes an X-linked lymphoproliferative syndrome. Nature, 2006, 444, 110-114.	13.7	649
5	Efficacy of Gene Therapy for X-Linked Severe Combined Immunodeficiency. New England Journal of Medicine, 2010, 363, 355-364.	13.9	561
6	STIM1Mutation Associated with a Syndrome of Immunodeficiency and Autoimmunity. New England Journal of Medicine, 2009, 360, 1971-1980.	13.9	459
7	Inherited STING-activating mutation underlies a familial inflammatory syndrome with lupus-like manifestations. Journal of Clinical Investigation, 2014, 124, 5516-5520.	3.9	435
8	Revised diagnostic criteria and classification for the autoimmune lymphoproliferative syndrome (ALPS): report from the 2009 NIH International Workshop. Blood, 2010, 116, e35-e40.	0.6	405
9	The European Society for Immunodeficiencies (ESID) Registry Working Definitions for the ClinicalÂDiagnosis of Inborn Errors of Immunity. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 1763-1770.	2.0	381
10	Clinical spectrum and features of activated phosphoinositide 3-kinase Î´ syndrome: AÂlarge patient cohort study. Journal of Allergy and Clinical Immunology, 2017, 139, 597-606.e4.	1.5	377
11	Detection of interferon alpha protein reveals differential levels and cellular sources in disease. Journal of Experimental Medicine, 2017, 214, 1547-1555.	4.2	288
12	Autoimmune Lymphoproliferative Syndrome with SomaticFasMutations. New England Journal of Medicine, 2004, 351, 1409-1418.	13.9	276
13	MST1 mutations in autosomal recessive primary immunodeficiency characterized by defective naive T-cell survival. Blood, 2012, 119, 3458-3468.	0.6	244
14	Severe Food Allergy as a Variant of IPEX Syndrome Caused by a Deletion in a Noncoding Region of the FOXP3 Gene. Gastroenterology, 2007, 132, 1705-1717.	0.6	236
15	Long-term follow-up of IPEX syndrome patients after different therapeutic strategies: An international multicenter retrospective study. Journal of Allergy and Clinical Immunology, 2018, 141, 1036-1049.e5.	1.5	233
16	Defective IL10 Signaling Defining a Subgroup of Patients With Inflammatory Bowel Disease. American Journal of Gastroenterology, 2011, 106, 1544-1555.	0.2	232
17	Autoimmune and inflammatory manifestations occur frequently in patients with primary immunodeficiencies. Journal of Allergy and Clinical Immunology, 2017, 140, 1388-1393.e8.	1.5	222
18	Autoimmune lymphoproliferative syndromes: genetic defects of apoptosis pathways. Cell Death and Differentiation, 2003, 10, 124-133.	5.0	215

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19	Tartrate-resistant acid phosphatase deficiency causes a bone dysplasia with autoimmunity and a type I interferon expression signature. Nature Genetics, 2011, 43, 127-131.	9.4	214
20	NF-Î¹B: At the Borders of Autoimmunity and Inflammation. Frontiers in Immunology, 2021, 12, 716469.	2.2	214
21	Intrinsic antiproliferative activity of the innate sensor STING in T lymphocytes. Journal of Experimental Medicine, 2017, 214, 1769-1785.	4.2	202
22	Human adenylate kinase 2 deficiency causes a profound hematopoietic defect associated with sensorineural deafness. Nature Genetics, 2009, 41, 106-111.	9.4	198
23	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.	1.5	192
24	Clinical, immunological, and pathological consequences of Fas-deficient conditions. Lancet, The, 1996, 348, 719-723.	6.3	191
25	Whole-Exome-Sequencing-Based Discovery of Human FADD Deficiency. American Journal of Human Genetics, 2010, 87, 873-881.	2.6	171
26	Type I interferon-mediated autoinflammation due to DNase II deficiency. Nature Communications, 2017, 8, 2176.	5.8	164
27	Disease-associated mutations identify a novel region in human STING necessary for the control of type I interferon signaling. Journal of Allergy and Clinical Immunology, 2017, 140, 543-552.e5.	1.5	159
28	Digestive histopathological presentation of IPEX syndrome. Modern Pathology, 2009, 22, 95-102.	2.9	158
29	NATURALLY OCCURRING PRIMARY DEFICIENCIES OF THE IMMUNE SYSTEM. Annual Review of Immunology, 1997, 15, 93-124.	9.5	157
30	A survey of 90 patients with autoimmune lymphoproliferative syndrome related to TNFRSF6 mutation. Blood, 2011, 118, 4798-4807.	0.6	153
31	FAS-L, IL-10, and double-negative CD4â^'CD8â^' TCR Î±/Î²+ T cells are reliable markers of autoimmune lymphoproliferative syndrome (ALPS) associated with FAS loss of function. Blood, 2009, 113, 3027-3030.	0.6	134
32	Lymphoproliferative syndrome with autoimmunity: A possible genetic basis for dominant expression of the clinical manifestations. Blood, 1999, 94, 2575-82.	0.6	122
33	Activation-induced cytidine deaminase (AID) is required for B-cell tolerance in humans. Proceedings of the National Academy of Sciences of the United States of America, 2011, 108, 11554-11559.	3.3	118
34	Cell-death signaling and human disease. Current Opinion in Immunology, 2003, 15, 325-331.	2.4	117
35	A Mendelian predisposition to B-cell lymphoma caused by IL-10R deficiency. Blood, 2013, 122, 3713-3722.	0.6	116
36	Severe Pulmonary Fibrosis as the First Manifestation of Interferonopathy (TMEM173 Mutation). Chest, 2016, 150, e65-e71.	0.4	112

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37	Inherited and SomaticCD3Î¶ Mutations in a Patient with T-Cell Deficiency. New England Journal of Medicine, 2006, 354, 1913-1921.	13.9	111
38	Cutaneous manifestations of immune dysregulation, polyendocrinopathy, enteropathy, X-linked (IPEX) syndrome. British Journal of Dermatology, 2009, 160, 645-651.	1.4	110
39	Onset of autoimmune lymphoproliferative syndrome (ALPS) in humans as a consequence of genetic defect accumulation. Journal of Clinical Investigation, 2011, 121, 106-112.	3.9	110
40	Distinct systemic and mucosal immune responses during acute SARS-CoV-2 infection. Nature Immunology, 2021, 22, 1428-1439.	7.0	110
41	Stimulator of Interferon Genesâ€“Associated Vasculopathy With Onset in Infancy. JAMA Dermatology, 2015, 151, 872.	2.0	108
42	Pediatric Evans syndrome is associated with a high frequency of potentially damaging variants in immune genes. Blood, 2019, 134, 9-21.	0.6	102
43	Defective human interleukin 2 receptor gamma chain in an atypical X chromosome-linked severe combined immunodeficiency with peripheral T cells.. Proceedings of the National Academy of Sciences of the United States of America, 1994, 91, 9466-9470.	3.3	99
44	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.	2.0	98
45	Primary T-cell immunodeficiency with immunodysregulation caused by autosomal recessive LCK deficiency. Journal of Allergy and Clinical Immunology, 2012, 130, 1144-1152.e11.	1.5	96
46	Induction of T lymphocyte apoptosis by sulphasalazine in patients with Crohn's disease. Gut, 2004, 53, 1632-1638.	6.1	89
47	Hypomorphic mutation of ZAP70 in human results in a late onset immunodeficiency and no autoimmunity. European Journal of Immunology, 2009, 39, 1966-1976.	1.6	88
48	Reduced Expression of FOXP3 and Regulatory T-Cell Function in Severe Forms of Early-onset Autoimmune Enteropathy. Gastroenterology, 2010, 139, 770-778.	0.6	88
49	Phenotypic Characterization of Very Early-onset IBD Due to Mutations in the IL10, IL10 Receptor Alpha or Beta Gene. Inflammatory Bowel Diseases, 2013, 19, 2820-2828.	0.9	80
50	Highly restricted human T cell repertoire in peripheral blood and tissue-infiltrating lymphocytes in Omenn's syndrome.. Journal of Clinical Investigation, 1998, 102, 312-321.	3.9	79
51	Severe combined immunodeficiency in stimulator of interferon genes (STING) V154M/wild-type mice. Journal of Allergy and Clinical Immunology, 2019, 143, 712-725.e5.	1.5	74
52	Monogenic lupus: Dissecting heterogeneity. Autoimmunity Reviews, 2019, 18, 102361.	2.5	74
53	Early-onset autoimmunity associated with SOCS1 haploinsufficiency. Nature Communications, 2020, 11, 5341.	5.8	74
54	Human TCR Î±/Î²+ CD4â^'CD8â^' Double-Negative T Cells in Patients with Autoimmune Lymphoproliferative Syndrome Express Restricted Vβ TCR Diversity and Are Clonally Related to CD8+ T Cells. Journal of Immunology, 2008, 181, 440-448.	0.4	70

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55	Deficiency in Mucosaâ€associated Lymphoid Tissue Lymphoma Translocation 1. Journal of Pediatric Gastroenterology and Nutrition, 2017, 64, 378-384.	0.9	69
56	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.	0.6	68
57	Immune deficiencyâ€“related enteropathy-lymphocytopenia-alopecia syndrome results from tetratricopeptide repeat domain 7A deficiency. Journal of Allergy and Clinical Immunology, 2014, 134, 1354-1364.e6.	1.5	66
58	Failure of HY-Specific Thymocytes to Escape Negative Selection by Receptor Editing. Immunity, 2002, 16, 707-718.	6.6	64
59	Long-term immune reconstitution in RAG-1-deficient mice treated by retroviral gene therapy: a balance between efficiency and toxicity. Blood, 2006, 107, 63-72.	0.6	64
60	Autoimmunity by haploinsufficiency. Science, 2014, 345, 1560-1561.	6.0	64
61	ORAI1 mutations abolishing store-operated Ca2+ entry cause anhidrotic ectodermal dysplasia with immunodeficiency. Journal of Allergy and Clinical Immunology, 2018, 142, 1297-1310.e11.	1.5	62
62	The Autoimmune Lymphoproliferative Syndrome with Defective FAS or FAS-Ligand Functions. Journal of Clinical Immunology, 2018, 38, 558-568.	2.0	61
63	Platelet activation in critically ill COVID-19 patients. Annals of Intensive Care, 2021, 11, 113.	2.2	61
64	X-linked primary immunodeficiency associated with hemizygous mutations in the moesin (MSN) gene. Journal of Allergy and Clinical Immunology, 2016, 138, 1681-1689.e8.	1.5	60
65	Germline TET2 loss of function causes childhood immunodeficiency and lymphoma. Blood, 2020, 136, 1055-1066.	0.6	58
66	LRBA deficiency with autoimmunity and early onset chronic erosive polyarthritis. Clinical Immunology, 2016, 168, 88-93.	1.4	57
67	VPS33B regulates protein sorting into and maturation of Î±-granule progenitor organelles in mouse megakaryocytes. Blood, 2015, 126, 133-143.	0.6	56
68	Pediatric-onset Evans syndrome: Heterogeneous presentation and high frequency of monogenic disorders including LRBA and CTLA4 mutations. Clinical Immunology, 2018, 188, 52-57.	1.4	53
69	Correction of Fas (CD95) deficiency by haploidentical bone marrow transplantation. European Journal of Immunology, 1997, 27, 2043-2047.	1.6	51
70	Monitoring Disease Activity in Systemic Lupus Erythematosus With Singleâ€Molecule Array Digital Enzymeâ€Linked Immunosorbent Assay Quantification of Serum Interferonâ€Î±. Arthritis and Rheumatology, 2019, 71, 756-765.	2.9	51
71	Autoimmune lymphoproliferative syndrome caused by a homozygous null FAS ligand (FASLG) mutation. Journal of Allergy and Clinical Immunology, 2013, 131, 486-490.	1.5	50
72	Evans Syndrome in Children: Long-Term Outcome in a Prospective French National Observational Cohort. Frontiers in Pediatrics, 2015, 3, 79.	0.9	49

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73	Evolution of disease activity and biomarkers on and off rapamycin in 28 patients with autoimmune lymphoproliferative syndrome. Haematologica, 2017, 102, e52-e56.	1.7	49
74	<scp>FAS</scp> and <scp>RAS</scp> related Apoptosis defects: From autoimmunity to leukemia. Immunological Reviews, 2019, 287, 50-61.	2.8	49
75	Defective anti-polysaccharide response and splenic marginal zone disorganization in ALPS patients. Blood, 2014, 124, 1597-1609.	0.6	48
76	Are RASopathies new monogenic predisposing conditions to the development of systemic lupus erythematosus? Case report and systematic review of the literature. Seminars in Arthritis and Rheumatism, 2013, 43, 217-219.	1.6	47
77	Clinical and molecular aspects of autoimmune enteropathy and immune dysregulation, polyendocrinopathy autoimmune enteropathy X-linked syndrome. Current Opinion in Gastroenterology, 2008, 24, 742-748.	1.0	42
78	RAS-associated lymphoproliferative disease evolves into severe juvenile myelo-monocytic leukemia. Blood, 2014, 123, 1960-1963.	0.6	41
79	Inactivation of the Fas gene by Alu insertion: retrotransposition in an intron causing splicing variation and autoimmune lymphoproliferative syndrome. Genes and Immunity, 2002, 3, S66-S70.	2.2	38
80	A monocyte/dendritic cell molecular signature of SARS-CoV-2-related multisystem inflammatory syndrome in children with severe myocarditis. Med, 2021, 2, 1072-1092.e7.	2.2	38
81	Normal T cell receptor VÎ² usage in a primary immunodeficiency associated with HLA class II deficiency. European Journal of Immunology, 1993, 23, 928-934.	1.6	37
82	Autoimmune lymphoproliferative syndrome: a multifactorial disorder. Haematologica, 2010, 95, 1805-1807.	1.7	35
83	Clinical Heterogeneity of Immune Dysregulation, Polyendocrinopathy, Enteropathy, X-Linked Syndrome: A French Multicenter Retrospective Study. Clinical and Translational Gastroenterology, 2018, 9, e201.	1.3	35
84	Single-cell analysis of FOXP3 deficiencies in humans and mice unmasks intrinsic and extrinsic CD4+ T cell perturbations. Nature Immunology, 2021, 22, 607-619.	7.0	35
85	Regulation of the acetylcholine/Î±7nAChR anti-inflammatory pathway in COVID-19 patients. Scientific Reports, 2021, 11, 11886.	1.6	35
86	Perforin-dependent apoptosis functionally compensates Fas deficiency in activation-induced cell death of human T lymphocytes. Blood, 2007, 110, 4285-4292.	0.6	34
87	What's up in the ALPS. Current Opinion in Immunology, 2017, 49, 79-86.	2.4	34
88	Type I interferon response and vascular alteration in chilblainâ€like lesions during the COVIDâ€19 outbreak*. British Journal of Dermatology, 2021, 185, 1176-1185.	1.4	33
89	Dyserythropoiesis associated with a Fas-deficient condition in childhood. British Journal of Haematology, 2000, 108, 300-304.	1.2	30
90	RASopathies: From germline mutations to somatic and multigenic diseases. Biomedical Journal, 2021, 44, 422-432.	1.4	28

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91	CD34-positive early human thymocytes: T cell receptor and cytokine receptor gene expression. European Journal of Immunology, 1995, 25, 2471-2478.	1.6	27
92	Immune checkpoint inhibitors increase T cell immunity during SARS-CoV-2 infection. Science Advances, 2021, 7, .	4.7	27
93	Gain-of-function IKZF1 variants in humans cause immune dysregulation associated with abnormal T/B cell late differentiation. Science Immunology, 2022, 7, eabi7160.	5.6	27
94	Lack of selective VÎ² deletion in peripheral CD4+ T cells of human immunodeficiency virus-infected infants. European Journal of Immunology, 1993, 23, 2041-2044.	1.6	26
95	Diagnosis of autoimmune lymphoproliferative syndrome caused by FAS deficiency in adults. Haematologica, 2013, 98, 389-392.	1.7	25
96	Severe COVID-19 is associated with hyperactivation of the alternative complement pathway. Journal of Allergy and Clinical Immunology, 2022, 149, 550-556.e2.	1.5	25
97	FAS/FAS-L dependent killing of activated human monocytes and macrophages by CD4+CD25â^' responder T cells, but not CD4+CD25+ regulatory T cells. Journal of Autoimmunity, 2012, 38, 29-38.	3.0	24
98	Inherited Immunodeficiency: A New Association With Early-Onset Childhood Panniculitis. Pediatrics, 2018, 141, S496-S500.	1.0	24
99	Inherited and Acquired Death Receptor Defects in Human Autoimmune Lymphoproliferative Syndrome. , 2005, 9, 18-36.		22
100	Overexpression of the antiapoptotic gene Bfl-1 in B cells from patients with familial systemic lupus erythematosus. Lupus, 2007, 16, 95-100.	0.8	21
101	Gray platelet syndrome can mimic autoimmune lymphoproliferative syndrome. Blood, 2015, 126, 1967-1969.	0.6	21
102	Diffuse large B-cell non-Hodgkin's lymphoma in a patient with autoimmune lymphoproliferative syndrome. British Journal of Haematology, 2001, 113, 432-434.	1.2	20
103	Childhood linear IgA disease in association with autoimmune lymphoproliferative syndrome. British Journal of Dermatology, 2004, 150, 578-580.	1.4	20
104	Mycophenolate Mofetil as an Alternate Immunosuppressor for Autoimmune Lymphoproliferative Syndrome. Journal of Pediatric Hematology/Oncology, 2006, 28, 824-826.	0.3	20
105	Somatic loss of heterozygosity, but not haploinsufficiency alone, leads to full-blown autoimmune lymphoproliferative syndrome in 1 of 12 family members with FAS start codon mutation. Clinical Immunology, 2013, 147, 61-68.	1.4	20
106	Loss-of-Function Mutation in PTPN2 Causes Aberrant Activation of JAK Signaling Via STAT and Very Early Onset Intestinal Inflammation. Gastroenterology, 2020, 159, 1968-1971.e4.	0.6	20
107	Genetic diagnosis of primary immunodeficiencies: AÂsurvey of the French national registry. Journal of Allergy and Clinical Immunology, 2019, 143, 1646-1649.e10.	1.5	20
108	Expression of the HLA-C2-specific activating killer-cell Ig-like receptor KIR2DS1 on NK and T cells. Clinical Immunology, 2010, 135, 26-32.	1.4	19

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109	Atypical Manifestation of LPS-Responsive Beige-Like Anchor Deficiency Syndrome as an Autoimmune Endocrine Disorder without Enteropathy and Immunodeficiency. Frontiers in Pediatrics, 2016, 4, 98.	0.9	18
110	STING Gain-of-Function Disrupts Lymph Node Organogenesis and Innate Lymphoid Cell Development in Mice. Cell Reports, 2020, 31, 107771.	2.9	18
111	Identification of germline monoallelic mutations in IKZF2 in patients with immune dysregulation. Blood Advances, 2022, 6, 2444-2451.	2.5	18
112	Improving the diagnostic efficiency of primary immunodeficiencies with targeted next-generation sequencing. Journal of Allergy and Clinical Immunology, 2021, 147, 734-737.	1.5	17
113	The genetic landscape of the FAS pathway deficiencies. Biomedical Journal, 2021, 44, 388-399.	1.4	16
114	Characterization of antigen-specific repertoire diversity following in vitro restimulation by a recombinant adenovirus expressing human cytomegalovirus pp65. European Journal of Immunology, 2003, 33, 760-768.	1.6	15
115	Autoimmune Lymphoproliferative Syndrome and Perforin. New England Journal of Medicine, 2005, 352, 306-307.	13.9	15
116	After 95â€¯years, it's time to eRASe JMML. Blood Reviews, 2020, 43, 100652.	2.8	14
117	Immunologic evaluation and genetic defects of apoptosis in patients with autoimmune lymphoproliferative syndrome (ALPS). Critical Reviews in Clinical Laboratory Sciences, 2021, 58, 253-274.	2.7	14
118	Differential sensitivity of Jurkat and primary T cells to caspase-independent cell death triggered upon Fas stimulation. European Journal of Immunology, 2002, 32, 2376.	1.6	13
119	Copy number variations and founder effect underlying complete IL-10RÎ² deficiency in Portuguese kindreds. PLoS ONE, 2018, 13, e0205826.	1.1	13
120	Autoimmune Lymphoproliferative Syndrome-FAS Patients Have an Abnormal Regulatory T Cell (Treg) Phenotype but Display Normal Natural Treg-Suppressive Function on T Cell Proliferation. Frontiers in Immunology, 2018, 9, 718.	2.2	13
121	T cell activation deficiencies. Clinical Immunology and Immunopathology, 1995, 76, S163-S164.	2.1	12
122	Cytomegalovirus infection in infants with autoimmune lymphoproliferative syndrome (ALPS). Clinical and Experimental Immunology, 2000, 121, 353-357.	1.1	12
123	Next Generation Sequencing for Detecting Somatic FAS Mutations in Patients With Autoimmune Lymphoproliferative Syndrome. Frontiers in Immunology, 2021, 12, 656356.	2.2	12
124	Neutropenia in Patients with Common Variable Immunodeficiency: a Rare Event Associated with Severe Outcome. Journal of Clinical Immunology, 2017, 37, 715-726.	2.0	11
125	Chronic granulomatous skin lesions leading to a diagnosis of <scp>TAP</scp>1 deficiency syndrome. Pediatric Dermatology, 2018, 35, e375-e377.	0.5	11
126	Life-Saving, Dose-Adjusted, Targeted Therapy in a Patient with a STAT3 Gain-of-Function Mutation. Journal of Clinical Immunology, 2021, 41, 807-810.	2.0	10

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127	Clonal hematopoiesis is not significantly associated with COVID-19 disease severity. Blood, 2022, 140, 1650-1655.	0.6	10
128	Long term follow-up of pediatric-onset Evans syndrome: broad immunopathological manifestations and high treatment burden. Haematologica, 2022, 107, 457-466.	1.7	9
129	Investigation of common variable immunodeficiency patients and healthy individuals using autoimmune lymphoproliferative syndrome biomarkers. Human Immunology, 2013, 74, 1531-1535.	1.2	8
130	Two neurologic facets of CTLA4-related haploinsufficiency. Neurology: Neuroimmunology and NeuroInflammation, 2020, 7, .	3.1	8
131	Abatacept is useful in autoimmune cytopenia with immunopathologic manifestations caused by CTLA-4 defects. Blood, 2022, 139, 300-304.	0.6	8
132	Integrative genetic and immune cell analysis of plasma proteins in healthy donors identifies novel associations involving primary immune deficiency genes. Genome Medicine, 2022, 14, 28.	3.6	8
133	Familial and syndromic lupus share the same phenotype as other early-onset forms of lupus. Joint Bone Spine, 2017, 84, 589-593.	0.8	7
134	Neurological Involvement in Childhood Evans Syndrome. Journal of Clinical Immunology, 2019, 39, 171-181.	2.0	6
135	Scaling the tips of the ALPS. Biomedical Journal, 2021, 44, 383-387.	1.4	6
136	PD-L1 is expressed on human activated naive effector CD4+ T cells. Regulation by dendritic cells and regulatory CD4+ T cells. PLoS ONE, 2021, 16, e0260206.	1.1	6
137	Clinical effects of mutations to CD95 (Fas): relevance to autoimmunity?. Seminars in Immunopathology, 1998, 19, 301-310.	4.0	5
138	A new peak in the ALPS. Nature Medicine, 1999, 5, 876-877.	15.2	5
139	Expression of Granzyme B in viral hepatitis in patients with ALPS. Hepatology, 2004, 39, 864-865.	3.6	5
140	Early-onset hypogammaglobulinemia: A survey of 44 patients. Journal of Allergy and Clinical Immunology, 2015, 136, 1097-1099.e2.	1.5	5
141	In Vitro Evaluation of the Apoptosis Function in Human Activated T Cells. Methods in Molecular Biology, 2017, 1557, 33-40.	0.4	5
142	Fatal Hypogammaglobulinemia 3 Years after Rituximab in a Patient with Immune Thrombocytopenia: An Underlying Genetic Predisposition?. Case Reports in Immunology, 2019, 2019, 1-6.	0.2	5
143	MHCâ€restricted T cell receptor signaling is required for Î±Î²â€„TCR replacement of the pre T cell receptor. European Journal of Immunology, 2008, 38, 391-399.	1.6	4
144	Live and let die at TEMRA. Blood, 2014, 124, 828-830.	0.6	4

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145	Early IFNÎ² secretion determines variable downstream IL-12p70 responses upon TLR4 activation. Cell Reports, 2022, 39, 110989.	2.9	4
146	Lymphadenopathy driven by TCR-Vî³8Vî´1 T-cell expansion in FAS-related autoimmune lymphoproliferative syndrome. Blood Advances, 2017, 1, 1101-1106.	2.5	3
147	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.	2.0	3
148	Autoimmune Lymphoproliferative Syndrome Presenting with Invasive Streptococcus pneumoniae Infection. Journal of Clinical Immunology, 2020, 40, 543-546.	2.0	3
149	Significance of Interdigitating Reticulum Cells in Omenn's Syndrome. American Journal of Surgical Pathology, 1996, 20, 1032.	2.1	3
150	Generation of an iPSC line (IMAGINi011-A) from a patient carrying a STING mutation. Stem Cell Research, 2021, 50, 102107.	0.3	2
151	Generation of Functional Regulatory T Cells By FOXP3 Gene Transfer into CD4 T Cells from Scurfy Mice and IPEX Patients. Blood, 2016, 128, 2526-2526.	0.6	2
152	Around the V(D)J recombinase machinery. Research in Immunology, 1994, 145, 151-158.	0.9	1
153	Comment on: Monogenic mimics of BehÃ§etâ€™s disease in the young. Rheumatology, 2020, 59, e109-e111.	0.9	1
154	Accessory spleen: Differential diagnosis for lymphoma in autoimmune lymphoproliferative syndrome. Pediatric Blood and Cancer, 2010, 54, 1020-1022.	0.8	0
155	Apoptosis-Related Autoimmune Lymphoproliferative Syndrome. , 2016, , 426-435.		0
156	560. Generation of Functional Regulatory T Cells by FOXP3 Gene Transfer into CD4 T Cell from IPEX Patients. Molecular Therapy, 2016, 24, S224.	3.7	0
157	Life-threatening pulmonary interstitial lung disease complicating pediatric nonhumoral immunodeficiencies. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 2456-2458.e4.	2.0	0
158	Human Adenylate Kinase 2 Deficiency Inhibits Hematopoietic Cell Differentiation towards Neutrophil and T Lymphoid Lineages.. Blood, 2009, 114, 78-78.	0.6	0
159	Il y a un fort parallÃ¨le entre auto-immunitÃ© et cancer. Pourlascience Fr, 2021, NÂ° 531 â€“ janvier, 40-45.	0.0	0
160	Letter to the Editor: Coexistence of Autoimmune Lymphoproliferative Syndrome and Familial Mediterranean Fever. Iranian Journal of Immunology, 2020, 17, 172-174.	0.4	0