Alain Fischer

List of Publications by Year
in descending order

134 papers	25,352 citations	22132 59 h-index	11601 135 g-index
136 all docs	136 docs citations	136 times ranked	20052 citing authors

#	Article	IF	Citations
1	LMO2-Associated Clonal T Cell Proliferation in Two Patients after Gene Therapy for SCID-X1. Science, 2003, 302, 415-419.	6.0	3,264
2	Gene Therapy of Human Severe Combined Immunodeficiency (SCID)-X1 Disease. Science, 2000, 288, 669-672.	6.0	2,451
3	Insertional oncogenesis in 4 patients after retrovirus-mediated gene therapy of SCID-X1. Journal of Clinical Investigation, 2008, 118, 3132-3142.	3.9	1,531
4	Hematopoietic Stem Cell Gene Therapy with a Lentiviral Vector in X-Linked Adrenoleukodystrophy. Science, 2009, 326, 818-823.	6.0	1,368
5	Transfusion independence and HMGA2 activation after gene therapy of human Î²-thalassaemia. Nature, 2010, 467, 318-322.	13.7	1,153
6	Sustained Correction of X-Linked Severe Combined Immunodeficiency by ex Vivo Gene Therapy. New England Journal of Medicine, 2002, 346, 1185-1193.	13.9	1,075
7	Revised classification of histiocytoses and neoplasms of the macrophage-dendritic cell lineages. Blood, 2016, 127, 2672-2681.	0.6	1,040
8	Lymphoid development in mice with a targeted deletion of the interleukin 2 receptor gamma chain.. Proceedings of the National Academy of Sciences of the United States of America, 1995, 92, 377-381.	3.3	834
9	XIAP deficiency in humans causes an X-linked lymphoproliferative syndrome. Nature, 2006, 444, 110-114.	13.7	649
10	Efficacy of Gene Therapy for X-Linked Severe Combined Immunodeficiency. New England Journal of Medicine, 2010, 363, 355-364.	13.9	561
11	Phosphoinositide 3-Kinase Î´ Gene Mutation Predisposes to Respiratory Infection and Airway Damage. Science, 2013, 342, 866-871.	6.0	541
12	Long-term survival and transplantation of haemopoietic stem cells for immunodeficiencies: report of the European experience 1968â€“99. Lancet, The, 2003, 361, 553-560.	6.3	524
13	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
14	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
15	A Modified Î³-Retrovirus Vector for X-Linked Severe Combined Immunodeficiency. New England Journal of Medicine, 2014, 371, 1407-1417.	13.9	358
16	Outcomes Following Gene Therapy in Patients With Severe Wiskott-Aldrich Syndrome. JAMA - Journal of the American Medical Association, 2015, 313, 1550.	3.8	327
17	Reduced-intensity conditioning and HLA-matched haemopoietic stem-cell transplantation in patients with chronic granulomatous disease: a prospective multicentre study. Lancet, The, 2014, 383, 436-448.	6.3	322
18	Long-term outcome and lineage-specific chimerism in 194 patients with Wiskott-Aldrich syndrome treated by hematopoietic cell transplantation in the period 1980-2009: an international collaborative study. Blood, 2011, 118, 1675-1684.	0.6	296

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19	Autoimmunity in Wiskott-Aldrich Syndrome: Risk Factors, Clinical Features, and Outcome in a Single-Center Cohort of 55 Patients. Pediatrics, 2003, 111, e622-e627.	1.0	294
20	Coronavirus disease 2019 in patients with inborn errors of immunity: An international study. Journal of Allergy and Clinical Immunology, 2021, 147, 520-531.	1.5	278
21	Autosomal Dominant STAT3 Deficiency and Hyper-IgE Syndrome. Medicine (United States), 2012, 91, e1-e19.	0.4	274
22	Evidence of innate lymphoid cell redundancy in humans. Nature Immunology, 2016, 17, 1291-1299.	7.0	260
23	Atypical X-Linked Severe Combined Immunodeficiency Due to Possible Spontaneous Reversion of the Genetic Defect in T Cells. New England Journal of Medicine, 1996, 335, 1563-1567.	13.9	259
24	A human immunodeficiency caused by mutations in the PIK3R1 gene. Journal of Clinical Investigation, 2014, 124, 3923-3928.	3.9	239
25	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
26	Long-term outcome after hematopoietic stem cell transplantation of a single-center cohort of 90 patients with severe combined immunodeficiency. Blood, 2009, 113, 4114-4124.	0.6	220
27	Long-term outcome following hematopoietic stem-cell transplantation in Wiskott-Aldrich syndrome: collaborative study of the European Society for Immunodeficiencies and European Group for Blood and Marrow Transplantation. Blood, 2008, 111, 439-445.	0.6	216
28	Clinical and immunologic phenotype associated with activated phosphoinositide 3-kinase Î´ syndrome 2: AÂcohort study. Journal of Allergy and Clinical Immunology, 2016, 138, 210-218.e9.	1.5	215
29	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
30	Incidence, Presentation, and Prognosis of Malignancies in Ataxia-Telangiectasia: A Report From the French National Registry of Primary Immune Deficiencies. Journal of Clinical Oncology, 2015, 33, 202-208.	0.8	176
31	CTP synthase 1 deficiency in humans reveals its central role in lymphocyte proliferation. Nature, 2014, 510, 288-292.	13.7	174
32	Severe combined immunodeficiencies and related disorders. Nature Reviews Disease Primers, 2015, 1, 15061.	18.1	173
33	Therapeutic effect of JAK1/2 blockade on the manifestations of hemophagocytic lymphohistiocytosis in mice. Blood, 2016, 128, 60-71.	0.6	151
34	Germline HAVCR2 mutations altering TIM-3 characterize subcutaneous panniculitis-like T cell lymphomas with hemophagocytic lymphohistiocytic syndrome. Nature Genetics, 2018, 50, 1650-1657.	9.4	151
35	Bone marrow gene transfer in three patients with adenosine deaminase deficiency. Gene Therapy, 1996, 3, 179-83.	2.3	145
36	Inherited defects in lymphocyte cytotoxic activity. Immunological Reviews, 2010, 235, 10-23.	2.8	143

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37	Diversity, functionality, and stability of the T cell repertoire derived in vivo from a single human T cell precursor. Proceedings of the National Academy of Sciences of the United States of America, 2000, 97, 274-278.	3.3	142
38	The CARD11-BCL10-MALT1 (CBM) signalosome complex: Stepping into the limelight of human primary immunodeficiency. Journal of Allergy and Clinical Immunology, 2014, 134, 276-284.	1.5	133
39	Failure of SCID-X1 gene therapy in older patients. Blood, 2005, 105, 4255-4257.	0.6	128
40	Loss-of-function mutations within the IL-2 inducible kinase ITK in patients with EBV-associated lymphoproliferative diseases. Leukemia, 2012, 26, 963-971.	3.3	122
41	Inherited CD70 deficiency in humans reveals a critical role for the CD70â€“CD27 pathway in immunity to Epstein-Barr virus infection. Journal of Experimental Medicine, 2017, 214, 73-89.	4.2	122
42	Dynamics of gene-modified progenitor cells analyzed by tracking retroviral integration sites in a human SCID-X1 gene therapy trial. Blood, 2010, 115, 4356-4366.	0.6	115
43	Role of interleukin-2 (IL-2), IL-7, and IL-15 in natural killer cell differentiation from cord blood hematopoietic progenitor cells and from gamma c transduced severe combined immunodeficiency X1 bone marrow cells. Blood, 1996, 88, 3901-3909.	0.6	113
44	gamma-c gene transfer into SCID X1 patients' B-cell lines restores normal high-affinity interleukin-2 receptor expression and function. Blood, 1996, 87, 3108-3116.	0.6	108
45	Characterization of Crohn disease in X-linked inhibitor of apoptosisâ€“deficient male patients and female symptomatic carriers. Journal of Allergy and Clinical Immunology, 2014, 134, 1131-1141.e9.	1.5	101
46	Occurrence of B-cell lymphomas in patients with activated phosphoinositide 3-kinase Î´ syndrome. Journal of Allergy and Clinical Immunology, 2014, 134, 233-236.e3.	1.5	101
47	Terminal transport of lytic granules to the immune synapse is mediated by the kinesin-1/Slp3/Rab27a complex. Blood, 2012, 119, 3879-3889.	0.6	98
48	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
49	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
50	Signaling pathways involved in the Tâ€cellâ€mediated immunity against Epsteinâ€Barr virus: Lessons from genetic diseases. Immunological Reviews, 2019, 291, 174-189.	2.8	85
51	Characteristics and outcome of early-onset, severe forms of Wiskott-Aldrich syndrome. Blood, 2013, 121, 1510-1516.	0.6	82
52	Prevention of Infections During Primary Immunodeficiency. Clinical Infectious Diseases, 2014, 59, 1462-1470.	2.9	81
53	Survival and Functional Outcomes in Boys with Cerebral Adrenoleukodystrophy with and without Hematopoietic Stem Cell Transplantation. Biology of Blood and Marrow Transplantation, 2019, 25, 538-548.	2.0	81
54	Outcome of hematopoietic cell transplantation for DNA double-strand break repair disorders. Journal of Allergy and Clinical Immunology, 2018, 141, 322-328.e10.	1.5	79

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55	Haploidentical Hematopoietic Stem Cell Transplantation with Post-Transplant Cyclophosphamide for Primary Immunodeficiencies and Inherited Disorders in Children. Biology of Blood and Marrow Transplantation, 2019, 25, 1363-1373.	2.0	78
56	Initial presenting manifestations in 16,486 patients with inborn errors of immunity include infections and noninfectious manifestations. Journal of Allergy and Clinical Immunology, 2021, 148, 1332-1341.e5.	1.5	75
57	Somatic genetic rescue in Mendelian haematopoietic diseases. Nature Reviews Genetics, 2019, 20, 582-598.	7.7	74
58	Pulmonary manifestations in adult patients with chronic granulomatous disease. European Respiratory Journal, 2015, 45, 1613-1623.	3.1	65
59	A novel immunoregulatory role for NK-cell cytotoxicity in protection from HLH-like immunopathology in mice. Blood, 2015, 125, 1427-1434.	0.6	64
60	Faster T-cell development following gene therapy compared with haploidentical HSCT in the treatment of SCID-X1. Blood, 2015, 125, 3563-3569.	0.6	64
61	Long-term safety and efficacy of lentiviral hematopoietic stem/progenitor cell gene therapy for Wiskottâ€“Aldrich syndrome. Nature Medicine, 2022, 28, 71-80.	15.2	64
62	<scp>LYST</scp> Controls the Biogenesis of the Endosomal Compartment Required for Secretory Lysosome Function. Traffic, 2015, 16, 191-203.	1.3	63
63	Gene therapy for severe combined immunodeficiencies and beyond. Journal of Experimental Medicine, 2020, 217, .	4.2	63
64	CD3 deficiencies. Current Opinion in Allergy and Clinical Immunology, 2005, 5, 491-495.	1.1	62
65	AK2 deficiency compromises the mitochondrial energy metabolism required for differentiation of human neutrophil and lymphoid lineages. Cell Death and Disease, 2015, 6, e1856-e1856.	2.7	61
66	Loss of <scp>RASGRP</scp> 1 in humans impairs Tâ€cell expansion leading to Epsteinâ€Barr virus susceptibility. EMBO Molecular Medicine, 2018, 10, 188-199.	3.3	61
67	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
68	Hematopoietic stem cell transplantation for CD40 ligand deficiency: Results from an EBMT/ESID-IEWP-SCETIDE-PIDTC study. Journal of Allergy and Clinical Immunology, 2019, 143, 2238-2253.	1.5	60
69	Concomitant PIK3CD and TNFRSF9 deficiencies cause chronic active Epstein-Barr virus infection of T cells. Journal of Experimental Medicine, 2019, 216, 2800-2818.	4.2	59
70	Stable and Functional Lymphoid Reconstitution in Artemis-deficient Mice Following Lentiviral Artemis Gene Transfer Into Hematopoietic Stem Cells. Molecular Therapy, 2008, 16, 1490-1499.	3.7	58
71	A newly identified isoform of Slp2a associates with Rab27a in cytotoxic T cells and participates to cytotoxic granule secretion. Blood, 2008, 112, 5052-5062.	0.6	57
72	LRBA deficiency with autoimmunity and early onset chronic erosive polyarthritis. Clinical Immunology, 2016, 168, 88-93.	1.4	57

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73	Chronic Granulomatous Disease in Patients Reaching Adulthood: A Nationwide Study in France. Clinical Infectious Diseases, 2017, 64, 767-775.	2.9	57
74	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
75	Î³c Gene Transfer in the Presence of Stem Cell Factor, FLT-3L, Interleukin-7 (IL-7), IL-1îœ€, and IL-15 Cytokines Restores T-Cell Differentiation From Î³c(â¯) X-Linked Severe Combined Immunodeficiency Hematopoietic Progenitor Cells in Murine Fetal Thymic Organ Cultures. Blood, 1998, 92, 4090-4097.	0.6	51
76	The management of chronic granulomatous disease. European Journal of Pediatrics, 1993, 152, 896-899.	1.3	50
77	Clonal tracking in gene therapy patients reveals a diversity of human hematopoietic differentiation programs. Blood, 2020, 135, 1219-1231.	0.6	50
78	Evans Syndrome in Children: Long-Term Outcome in a Prospective French National Observational Cohort. Frontiers in Pediatrics, 2015, 3, 79.	0.9	49
79	Genetic, Cellular and Clinical Features of ICF Syndrome: a French National Survey. Journal of Clinical Immunology, 2016, 36, 149-159.	2.0	48
80	A novel hypomorphic mutation in STIM1 results in a late-onset immunodeficiency. Journal of Allergy and Clinical Immunology, 2015, 136, 816-819.e4.	1.5	47
81	Mild B-cell lymphocytosis in patients with a CARD11 C49Y mutation. Journal of Allergy and Clinical Immunology, 2015, 136, 819-821.e1.	1.5	44
82	Polygenic mutations in the cytotoxicity pathway increase susceptibility to develop HLH immunopathology in mice. Blood, 2016, 127, 2113-2121.	0.6	44
83	RASGRP1 mutation in autoimmune lymphoproliferative syndrome-like disease. Journal of Allergy and Clinical Immunology, 2018, 142, 595-604.e16.	1.5	44
84	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.	1.5	43
85	SYK expression endows human ZAP70-deficient CD8 T cells with residual TCR signaling. Clinical Immunology, 2015, 161, 103-109.	1.4	38
86	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
87	An inherited immunoglobulin class-switch recombination deficiency associated with a defect in the INO80 chromatin remodeling complex. Journal of Allergy and Clinical Immunology, 2015, 135, 998-1007.e6.	1.5	37
88	Rapid identification and characterization of infected cells in blood during chronic active Epstein-Barr virus infection. Journal of Experimental Medicine, 2020, 217, .	4.2	37
89	Kinesin-1 controls mast cell degranulation and anaphylaxis through PI3K-dependent recruitment to the granular Slp3/Rab27b complex. Journal of Cell Biology, 2016, 215, 203-216.	2.3	36
90	Prevalence of primary immunodeficiencies in France is underestimated. Journal of Allergy and Clinical Immunology, 2017, 140, 1731-1733.	1.5	32

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91	Reticular dysgenesis: international survey on clinical presentation, transplantation, and outcome. Blood, 2017, 129, 2928-2938.	0.6	31
92	Mammalian target of rapamycin inhibition counterbalances the inflammatory status of immune cells in patients with chronic granulomatous disease. Journal of Allergy and Clinical Immunology, 2017, 139, 1641-1649.e6.	1.5	30
93	Systematic neonatal screening for severe combined immunodeficiency and severe T-cell lymphopenia: Analysis of cost-effectiveness based on French real field data. Journal of Allergy and Clinical Immunology, 2015, 135, 1589-1593.	1.5	29
94	Cutaneous granulomas with primary immunodeficiency in children: a report of 17 new patients and a review of the literature. Journal of the European Academy of Dermatology and Venereology, 2019, 33, 1412-1420.	1.3	29
95	Impaired lymphocyte function and differentiation in CTPS1-deficient patients result from a hypomorphic homozygous mutation. JCI Insight, 2020, 5, .	2.3	29
96	T cell dynamics and response of the microbiota after gene therapy to treat X-linked severe combined immunodeficiency. Genome Medicine, 2018, 10, 70.	3.6	28
97	Intestinal dysbiosis in inflammatory bowel disease associated with primary immunodeficiency. Journal of Allergy and Clinical Immunology, 2019, 143, 775-778.e6.	1.5	28
98	Role of interleukin-2 (IL-2), IL-7, and IL-15 in natural killer cell differentiation from cord blood hematopoietic progenitor cells and from gamma c transduced severe combined immunodeficiency X1 bone marrow cells. Blood, 1996, 88, 3901-9.	0.6	27
99	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
100	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.	1.5	26
101	Successful haematopoietic stem cell transplantation in a case of pulmonary alveolar proteinosis due to GM-CSF receptor deficiency. Thorax, 2018, 73, 590-592.	2.7	24
102	Cytotoxic granule secretion by lymphocytes and its link to immune homeostasis. F1000Research, 2015, 4, 930.	0.8	23
103	gamma-c gene transfer into SCID X1 patients' B-cell lines restores normal high-affinity interleukin-2 receptor expression and function. Blood, 1996, 87, 3108-16.	0.6	23
104	Outcome of chronic granulomatous disease â€•Conventional treatment vs stem cell transplantation. Pediatric Allergy and Immunology, 2021, 32, 576-585.	1.1	21
105	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
106	Recent advances in understanding the pathophysiology of primary T cell immunodeficiencies. Trends in Molecular Medicine, 2015, 21, 408-416.	3.5	18
107	Alemtuzumab as First Line Treatment in Children with Familial Lymphohistiocytosis. Blood, 2019, 134, 80-80.	0.6	18
108	Kinesin-1 Is a New Actor Involved in Platelet Secretion and Thrombus Stability. Arteriosclerosis, Thrombosis, and Vascular Biology, 2018, 38, 1037-1051.	1.1	17

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109	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
110	Benefit Corporation: a path to affordable gene therapies?. Nature Medicine, 2019, 25, 1813-1814.	15.2	16
111	Primary immunodeficiency-related bronchiectasis in adults: comparison with bronchiectasis of other etiologies in a French reference center. Respiratory Research, 2019, 20, 275.	1.4	16
112	Kinesin-1 regulates antigen cross-presentation through the scission of tubulations from early endosomes in dendritic cells. Nature Communications, 2020, 11, 1817.	5.8	16
113	Burden of Poor Health Conditions and Quality of Life in 656 Children with Primary Immunodeficiency. Journal of Pediatrics, 2018, 194, 211-217.e5.	0.9	15
114	Variable correction of Artemis deficiency by I-Sce1-meganuclease-assisted homologous recombination in murine hematopoietic stem cells. Gene Therapy, 2014, 21, 529-532.	2.3	14
115	Lymphoproliferative disease in patients with Wiskott-Aldrich syndrome: Analysis of the French Registry of Primary Immunodeficiencies. Journal of Allergy and Clinical Immunology, 2019, 143, 2311-2315.e7.	1.5	13
116	Cutaneous barrier leakage and gut inflammation drive skin disease in Omenn syndrome. Journal of Allergy and Clinical Immunology, 2020, 146, 1165-1179.e11.	1.5	13
117	Nocardiosis Associated with Primary Immunodeficiencies (Nocar-DIP): an International Retrospective Study and Literature Review. Journal of Clinical Immunology, 2020, 40, 1144-1155.	2.0	11
118	Platelets are the Achilles' heel of Wiskott-Aldrich syndrome. Journal of Allergy and Clinical Immunology, 2019, 144, 668-670.	1.5	10
119	Chronic Granulomatous Disease with the McLeod Phenotype: a French National Retrospective Case Series. Journal of Clinical Immunology, 2020, 40, 752-762.	2.0	10
120	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
121	gammac gene transfer in the presence of stem cell factor, FLT-3L, interleukin-7 (IL-7), IL-1, and IL-15 cytokines restores T-cell differentiation from gammac(-) X-linked severe combined immunodeficiency hematopoietic progenitor cells in murine fetal thymic organ cultures. Blood, 1998, 92, 4090-7.	0.6	10
122	50th Anniversary of the French Society for Immunology (SFI). European Journal of Immunology, 2016, 46, 1545-1547.	1.6	8
123	Safety and efficacy of brentuximab vedotin as a treatment for lymphoproliferative disorders in primary immunodeficiencies. Haematologica, 2020, 105, e461-464.	1.7	7
124	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.	1.5	7
125	Gene therapy: Myth or reality?. Comptes Rendus - Biologies, 2016, 339, 314-318.	0.1	6
126	Reply to â€˜Comment on: Evidence of innate lymphoid cell redundancy in humansâ€™. Nature Immunology, 2018, 19, 789-790.	7.0	6

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127	Current Spectrum of Infections in Patients with X-Linked Agammaglobulinemia. Journal of Clinical Immunology, 2021, 41, 1266-1271.	2.0	6
128	Early-onset hypogammaglobulinemia: A survey of 44 patients. Journal of Allergy and Clinical Immunology, 2015, 136, 1097-1099.e2.	1.5	5
129	A neuropathological study of cerebrovascular abnormalities in a signal transducer and activator of transcription 3â€“deficient patient. Journal of Allergy and Clinical Immunology, 2015, 136, 1418-1421.e5.	1.5	5
130	Gene therapy for human severe combined immunodeficiencies. Israel Medical Association Journal, 2002, 4, 51-4.	0.1	4
131	Rab44 regulates murine mast cellâ€“driven anaphylaxis through kinesin-1â€“dependent secretory granule translocation. Journal of Allergy and Clinical Immunology, 2022, 150, 676-689.	1.5	4
132	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
133	Antibody-coated microbiota in nasopharynx of healthy individuals and IVIg-treated patients with hypogammaglobulinemia. Journal of Allergy and Clinical Immunology, 2020, 145, 1686-1690.e4.	1.5	3
134	Vaccination of children. Archives De Pediatrie, 2019, 26, 55.	0.4	0