

# Stuart G Tangye

## List of Publications by Year in descending order

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Version: 2024-02-01

248  
papers

31,038  
citations

3525

90  
h-index

4988

167  
g-index

263  
all docs

263  
docs citations

263  
times ranked

30784  
citing authors

#	ARTICLE	IF	CITATIONS
1	Vaccine breakthrough hypoxemic COVID-19 pneumonia in patients with auto-Abs neutralizing type I IFNs. <i>Science Immunology</i> , 2023, 8, .	5.6	35
2	Hyper-IgE Syndrome due to an Elusive Novel Intronic Homozygous Variant in DOCK8. <i>Journal of Clinical Immunology</i> , 2022, 42, 119-129.	2.0	4
3	Human genetic and immunological determinants of critical COVID-19 pneumonia. <i>Nature</i> , 2022, 603, 587-598.	13.7	216
4	Atypical Autosomal Recessive AID Deficiencyâ€”Yet Another Piece of the Hyper-IgM Puzzle. <i>Journal of Clinical Immunology</i> , 2022, , 1.	2.0	0
5	Getting to the (germinal) center of humoral immune responses to SARS-CoV-2. <i>Cell</i> , 2022, 185, 945-948.	13.5	7
6	â€œAre you gonna go my way?â€”Decisions at the Tfh-B cell interface. <i>Immunity</i> , 2022, 55, 377-379.	6.6	1
7	Identification of germline monoallelic mutations in <i>IKZF2</i> in patients with immune dysregulation. <i>Blood Advances</i> , 2022, 6, 2444-2451.	2.5	18
8	Severe COVID-19 represents an undiagnosed primary immunodeficiency in a high proportion of infected individuals. <i>Clinical and Translational Immunology</i> , 2022, 11, e1365.	1.7	7
9	A loss-of-function <i>IFNAR1</i> allele in Polynesia underlies severe viral diseases in homozygotes. <i>Journal of Experimental Medicine</i> , 2022, 219, .	4.2	28
10	STAT5B restrains human B-cell differentiation to maintain humoral immune homeostasis. <i>Journal of Allergy and Clinical Immunology</i> , 2022, 150, 931-946.	1.5	19
11	A Novel Targeted Amplicon Next-Generation Sequencing Gene Panel for the Diagnosis of Common Variable Immunodeficiency Has a High Diagnostic Yield. <i>Journal of Molecular Diagnostics</i> , 2022, 24, 586-599.	1.2	4
12	The risk of COVID-19 death is much greater and age dependent with type I IFN autoantibodies. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2022, 119, e2200413119.	3.3	110
13	Human Inborn Errors of Immunity: 2022 Update on the Classification from the International Union of Immunological Societies Expert Committee. <i>Journal of Clinical Immunology</i> , 2022, 42, 1473-1507.	2.0	389
14	Respiratory viral infections in otherwise healthy humans with inherited IRF7 deficiency. <i>Journal of Experimental Medicine</i> , 2022, 219, .	4.2	21
15	Recessive inborn errors of type I IFN immunity in children with COVID-19 pneumonia. <i>Journal of Experimental Medicine</i> , 2022, 219, .	4.2	59
16	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.	1.5	278
17	Molecular requirements for human lymphopoiesis as defined by inborn errors of immunity. <i>Stem Cells</i> , 2021, 39, 389-402.	1.4	4
18	Somatic reversion of pathogenic DOCK8 variants alters lymphocyte differentiation and function to effectively cure DOCK8 deficiency. <i>Journal of Clinical Investigation</i> , 2021, 131, .	3.9	18

#	ARTICLE	IF	CITATIONS
19	The Ever-Increasing Array of Novel Inborn Errors of Immunity: an Interim Update by the IUIS Committee. <i>Journal of Clinical Immunology</i> , 2021, 41, 666-679.	2.0	165
20	SARS-CoV-2-related MIS-C: A key to the viral and genetic causes of Kawasaki disease?. <i>Journal of Experimental Medicine</i> , 2021, 218, .	4.2	100
21	Genomic Spectrum and Phenotypic Heterogeneity of Human IL-21 Receptor Deficiency. <i>Journal of Clinical Immunology</i> , 2021, 41, 1272-1290.	2.0	25
22	CD8+ T cell landscape in Indigenous and non-Indigenous people restricted by influenza mortality-associated HLA-A*24:02 allomorph. <i>Nature Communications</i> , 2021, 12, 2931.	5.8	20
23	Hematopoietic Stem Cell Transplantation Cures Chronic Aichi Virus Infection in a Patient with X-linked Agammaglobulinemia. <i>Journal of Clinical Immunology</i> , 2021, 41, 1403-1405.	2.0	8
24	Human STAT3 variants underlie autosomal dominant hyper-IgE syndrome by negative dominance. <i>Journal of Experimental Medicine</i> , 2021, 218, .	4.2	30
25	High Th2 cytokine levels and upper airway inflammation in human inherited T-bet deficiency. <i>Journal of Experimental Medicine</i> , 2021, 218, .	4.2	25
26	Phosphatidylinositol 3-kinase signaling and immune regulation: insights into disease pathogenesis and clinical implications. <i>Expert Review of Clinical Immunology</i> , 2021, 17, 905-914.	1.3	4
27	Inherited PD-1 deficiency underlies tuberculosis and autoimmunity in a child. <i>Nature Medicine</i> , 2021, 27, 1646-1654.	15.2	65
28	Humans with inherited T cell CD28 deficiency are susceptible to skin papillomaviruses but are otherwise healthy. <i>Cell</i> , 2021, 184, 3812-3828.e30.	13.5	53
29	Tissue-resident regulatory T cells accumulate at human barrier lymphoid organs. <i>Immunology and Cell Biology</i> , 2021, 99, 894-906.	1.0	6
30	CD4+ T cells that help B cells – a proposal for uniform nomenclature. <i>Trends in Immunology</i> , 2021, 42, 658-669.	2.9	65
31	Autoantibodies neutralizing type I IFNs are present in ~4% of uninfected individuals over 70 years old and account for ~20% of COVID-19 deaths. <i>Science Immunology</i> , 2021, 6, .	5.6	357
32	X-linked recessive TLR7 deficiency in ~1% of men under 60 years old with life-threatening COVID-19. <i>Science Immunology</i> , 2021, 6, .	5.6	267
33	Inherited human c-Rel deficiency disrupts myeloid and lymphoid immunity to multiple infectious agents. <i>Journal of Clinical Investigation</i> , 2021, 131, .	3.9	21
34	Mechanisms underlying host defense and disease pathology in response to severe acute respiratory syndrome (SARS)-CoV2 infection: insights from inborn errors of immunity. <i>Current Opinion in Allergy and Clinical Immunology</i> , 2021, 21, 515-524.	1.1	19
35	Coronavirus disease 2019 in patients with inborn errors of immunity: lessons learned. <i>Current Opinion in Pediatrics</i> , 2021, 33, 648-656.	1.0	42
36	Molecular regulation and dysregulation of T follicular helper cells – learning from inborn errors of immunity. <i>Current Opinion in Immunology</i> , 2021, 72, 249-261.	2.4	6

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37	<i>JEM</i> career launchpad. Journal of Experimental Medicine, 2021, 218, .	4.2	0
38	The expansion of human T-bet <sup>high</sup> CD21 <sup>low</sup> B cells is T cell dependent. Science Immunology, 2021, 6, eabh0891.	5.6	82
39	Intrinsic Defects in B Cell Development and Differentiation, T Cell Exhaustion and Altered Unconventional T Cell Generation Characterize Human Adenosine Deaminase Type 2 Deficiency. Journal of Clinical Immunology, 2021, 41, 1915-1935.	2.0	23
40	The Next Generation of Diagnostic Tests for Primary Immunodeficiency Disorders. Journal of Infectious Diseases, 2020, 221, 1232-1234.	1.9	3
41	Diversity of XMEN Disease: Description of 2 Novel Variants and Analysis of the Lymphocyte Phenotype. Journal of Clinical Immunology, 2020, 40, 299-309.	2.0	25
42	Unresponsiveness to inhaled antigen is governed by conventional dendritic cells and overridden during infection by monocytes. Science Immunology, 2020, 5, .	5.6	12
43	Activated PI3KÎ breaches multiple B cell tolerance checkpoints and causes autoantibody production. Journal of Experimental Medicine, 2020, 217, .	4.2	33
44	Regulation of the germinal center and humoral immunity by interleukin-21. Journal of Experimental Medicine, 2020, 217, .	4.2	74
45	Dominant-negative mutations in human <i>IL6ST</i> underlie hyper-IgE syndrome. Journal of Experimental Medicine, 2020, 217, .	4.2	64
46	Autoantibodies against type I IFNs in patients with life-threatening COVID-19. Science, 2020, 370, .	6.0	1,983
47	Itâ€™s that time of yearâ€™ APRIL promotes humoral immunity in humans. Journal of Allergy and Clinical Immunology, 2020, 146, 1013-1015.	1.5	2
48	Editorial: Human Disorders of PI3K Biology. Frontiers in Immunology, 2020, 11, 617464.	2.2	3
49	Human T-bet Governs Innate and Innate-like Adaptive IFN-Î³ Immunity against Mycobacteria. Cell, 2020, 183, 1826-1847.e31.	13.5	83
50	A Global Effort to Define the Human Genetics of Protective Immunity to SARS-CoV-2 Infection. Cell, 2020, 181, 1194-1199.	13.5	185
51	Three Copies of Four Interferon Receptor Genes Underlie a Mild Type I Interferonopathy in Down Syndrome. Journal of Clinical Immunology, 2020, 40, 807-819.	2.0	44
52	The Clinical Immunogenomics Research Consortium Australasia (CIRCA): a Distributed Network Model for Genomic Healthcare Delivery. Journal of Clinical Immunology, 2020, 40, 763-766.	2.0	5
53	Everolimus-Induced Remission of Classic Kaposiâ€™s Sarcoma Secondary to Cryptic Splicing Mediated CTLA4 Haploinsufficiency. Journal of Clinical Immunology, 2020, 40, 774-779.	2.0	5
54	Genetic susceptibility to EBV infection: insights from inborn errors of immunity. Human Genetics, 2020, 139, 885-901.	1.8	38

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55	Extended clinical and immunological phenotype and transplant outcome in CD27 and CD70 deficiency. <i>Blood</i> , 2020, 136, 2638-2655.	0.6	64
56	Human Inborn Errors of Immunity: 2019 Update on the Classification from the International Union of Immunological Societies Expert Committee. <i>Journal of Clinical Immunology</i> , 2020, 40, 24-64.	2.0	881
57	Systemic Inflammation and Myelofibrosis in a Patient with Takenouchi-Kosaki Syndrome due to CDC42 Tyr64Cys Mutation. <i>Journal of Clinical Immunology</i> , 2020, 40, 567-570.	2.0	29
58	Primary immunodeficiencies reveal the molecular requirements for effective host defense against EBV infection. <i>Blood</i> , 2020, 135, 644-655.	0.6	80
59	Human Inborn Errors of Immunity: 2019 Update of the IUIS Phenotypical Classification. <i>Journal of Clinical Immunology</i> , 2020, 40, 66-81.	2.0	525
60	Human inborn errors of immunity to herpes viruses. <i>Current Opinion in Immunology</i> , 2020, 62, 106-122.	2.4	60
61	Refractory very early-onset inflammatory bowel disease associated with cytosolic isoleucyl-tRNA synthetase deficiency: A case report. <i>World Journal of Gastroenterology</i> , 2020, 26, 1841-1846.	1.4	6
62	Activating PIK3CD mutations impair human cytotoxic lymphocyte differentiation and function and EBV immunity. <i>Journal of Allergy and Clinical Immunology</i> , 2019, 143, 276-291.e6.	1.5	64
63	A deep intronic splice mutation of <i>STAT3</i> underlies hyper IgE syndrome by negative dominance. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2019, 116, 16463-16472.	3.3	17
64	Flow Cytometric-Based Analysis of Defects in Lymphocyte Differentiation and Function Due to Inborn Errors of Immunity. <i>Frontiers in Immunology</i> , 2019, 10, 2108.	2.2	24
65	An essential role for the Zn <sup>2+</sup> transporter ZIP7 in B cell development. <i>Nature Immunology</i> , 2019, 20, 350-361.	7.0	92
66	The FOXP3 <sup>2</sup> isoform supports Treg cell development and protects against severe IPEX syndrome. <i>Journal of Allergy and Clinical Immunology</i> , 2019, 144, 317-320.e8.	1.5	20
67	B cell intrinsic requirement for STK4 in humoral immunity in mice and human subjects. <i>Journal of Allergy and Clinical Immunology</i> , 2019, 143, 2302-2305.	1.5	21
68	Human DOCK2 Deficiency: Report of a Novel Mutation and Evidence for Neutrophil Dysfunction. <i>Journal of Clinical Immunology</i> , 2019, 39, 298-308.	2.0	31
69	Activating mutations in PIK3CD disrupt the differentiation and function of human and murine CD4 <sup>+</sup> T cells. <i>Journal of Allergy and Clinical Immunology</i> , 2019, 144, 236-253.	1.5	44
70	Immune Dysregulation and Disease Pathogenesis due to Activating Mutations in PIK3CD: the Goldilocks™ Effect. <i>Journal of Clinical Immunology</i> , 2019, 39, 148-158.	2.0	26
71	Case Presentation of Hyper-IgE Syndrome With Novel Mutation. <i>Pathology</i> , 2019, 51, S50.	0.3	0
72	Human CD8 <sup>+</sup> T cell cross-reactivity across influenza A, B and C viruses. <i>Nature Immunology</i> , 2019, 20, 613-625.	7.0	180

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73	Human inborn errors of the actin cytoskeleton affecting immunity: way beyond WAS and WIP. <i>Immunology and Cell Biology</i> , 2019, 97, 389-402.	1.0	39
74	Chronic mucocutaneous candidiasis and connective tissue disorder in humans with impaired JNK1-dependent responses to IL-17A/F and TGF- $\beta$ 2. <i>Science Immunology</i> , 2019, 4, .	5.6	45
75	Denisovan, modern human and mouse TNFAIP3 alleles tune A20 phosphorylation and immunity. <i>Nature Immunology</i> , 2019, 20, 1299-1310.	7.0	53
76	What can primary immunodeficiencies teach us about Th9 cell differentiation and function?. <i>Immunology and Cell Biology</i> , 2019, 97, 380-388.	1.0	4
77	Hematopoietic stem cell transplant effectively rescues lymphocyte differentiation and function in DOCK8-deficient patients. <i>JCI Insight</i> , 2019, 4, .	2.3	23
78	STAT3 regulates cytotoxicity of human CD57+ CD4+ T cells in blood and lymphoid follicles. <i>Scientific Reports</i> , 2018, 8, 3529.	1.6	29
79	Combined Immunodeficiency with Ring Chromosome 21. <i>Journal of Clinical Immunology</i> , 2018, 38, 251-256.	2.0	2
80	Circulating T <sub>FH</sub> cells, serological memory, and tissue compartmentalization shape human influenza-specific B cell immunity. <i>Science Translational Medicine</i> , 2018, 10, .	5.8	196
81	Reversible Suppression of Lymphoproliferation and Thrombocytopenia with Rapamycin in a Patient with Common Variable Immunodeficiency. <i>Journal of Clinical Immunology</i> , 2018, 38, 159-162.	2.0	3
82	Human plasma C3 is essential for the development of memory B, but not T, lymphocytes. <i>Journal of Allergy and Clinical Immunology</i> , 2018, 141, 1151-1154.e14.	1.5	26
83	Is it dead or alive? TLR8 can tell. <i>Nature Immunology</i> , 2018, 19, 324-326.	7.0	2
84	International Union of Immunological Societies: 2017 Primary Immunodeficiency Diseases Committee Report on Inborn Errors of Immunity. <i>Journal of Clinical Immunology</i> , 2018, 38, 96-128.	2.0	732
85	The 2017 IUIS Phenotypic Classification for Primary Immunodeficiencies. <i>Journal of Clinical Immunology</i> , 2018, 38, 129-143.	2.0	488
86	Human IFN- $\beta$ immunity to mycobacteria is governed by both IL-12 and IL-23. <i>Science Immunology</i> , 2018, 3, .	5.6	152
87	Tuberculosis and impaired IL-23-dependent IFN- $\beta$ immunity in humans homozygous for a common <i>TYK2</i> missense variant. <i>Science Immunology</i> , 2018, 3, .	5.6	148
88	Chronic Aichi Virus Infection in a Patient with X-Linked Agammaglobulinemia. <i>Journal of Clinical Immunology</i> , 2018, 38, 748-752.	2.0	18
89	IRF4 haploinsufficiency in a family with Whipple's disease. <i>ELife</i> , 2018, 7, .	2.8	43
90	B cells race the clock to get a second wind. <i>Nature Immunology</i> , 2018, 19, 791-793.	7.0	2

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91	Germline-activating mutations in <i>PIK3CD</i> compromise B cell development and function. <i>Journal of Experimental Medicine</i> , 2018, 215, 2073-2095.	4.2	79
92	Mutations affecting the actin regulator WD repeat-containing protein 1 lead to aberrant lymphoid immunity. <i>Journal of Allergy and Clinical Immunology</i> , 2018, 142, 1589-1604.e11.	1.5	64
93	Disruption of an antimycobacterial circuit between dendritic and helper T cells in human SPPL2a deficiency. <i>Nature Immunology</i> , 2018, 19, 973-985.	7.0	96
94	Memory B cells are reactivated in subcapsular proliferative foci of lymph nodes. <i>Nature Communications</i> , 2018, 9, 3372.	5.8	88
95	A recessive form of hyper-IgE syndrome by disruption of ZNF341-dependent STAT3 transcription and activity. <i>Science Immunology</i> , 2018, 3, .	5.6	132
96	Human immunity against EBV lessons from the clinic. <i>Journal of Experimental Medicine</i> , 2017, 214, 269-283.	4.2	132
97	Pathogenic CD4 <sup>+</sup> T cells regulating B cell differentiation in autoimmunity: not exactly Tfh cells. <i>Immunology and Cell Biology</i> , 2017, 95, 419-421.	1.0	3
98	The TORC that Gets the GC Cycling. <i>Immunity</i> , 2017, 46, 974-976.	6.6	1
99	Memory B cells: total recall. <i>Current Opinion in Immunology</i> , 2017, 45, 132-140.	2.4	57
100	Combined immunodeficiency and Epstein-Barr virus-induced B cell malignancy in humans with inherited CD70 deficiency. <i>Journal of Experimental Medicine</i> , 2017, 214, 91-106.	4.2	134
101	Arginine methylation catalyzed by PRMT1 is required for B cell activation and differentiation. <i>Nature Communications</i> , 2017, 8, 891.	5.8	34
102	Cytokine-Mediated Regulation of Human Lymphocyte Development and Function: Insights from Primary Immunodeficiencies. <i>Journal of Immunology</i> , 2017, 199, 1949-1958.	0.4	23
103	DOCK8 Drives Src-Dependent NK Cell Effector Function. <i>Journal of Immunology</i> , 2017, 199, 2118-2127.	0.4	18
104	Defective protein prenylation is a diagnostic biomarker of mevalonate kinase deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2017, 140, 873-875.e6.	1.5	29
105	Low IgE Is Insufficiently Sensitive to Guide Genetic Testing of STAT3 Gain-of-Function Mutations. <i>Clinical Chemistry</i> , 2017, 63, 1539-1540.	1.5	4
106	AD Hyper-IgE Syndrome Due to a Novel Loss-of-Function Mutation in STAT3: a Diagnostic Pursuit Won by Clinical Acuity. <i>Journal of Clinical Immunology</i> , 2017, 37, 12-17.	2.0	5
107	Dedicator of cytokinesis 8-deficient CD4 <sup>+</sup> T cells are biased to a TH2 effector fate at the expense of TH1 and TH17 cells. <i>Journal of Allergy and Clinical Immunology</i> , 2017, 139, 933-949.	1.5	69
108	CCR6 Defines Memory B Cell Precursors in Mouse and Human Germinal Centers, Revealing Light-Zone Location and Predominant Low Antigen Affinity. <i>Immunity</i> , 2017, 47, 1142-1153.e4.	6.6	196

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109	Inherited GINS1 deficiency underlies growth retardation along with neutropenia and NK cell deficiency. <i>Journal of Clinical Investigation</i> , 2017, 127, 1991-2006.	3.9	115
110	Unique and shared signaling pathways cooperate to regulate the differentiation of human CD4+ T cells into distinct effector subsets. <i>Journal of Experimental Medicine</i> , 2016, 213, 1589-1608.	4.2	77
111	Elucidating the effects of disease-causing mutations on STAT3 function in autosomal-dominant hyper-IgE syndrome. <i>Journal of Allergy and Clinical Immunology</i> , 2016, 138, 1210-1213.e5.	1.5	16
112	Naïve and memory B cells exhibit distinct biochemical responses following BCR engagement. <i>Immunology and Cell Biology</i> , 2016, 94, 774-786.	1.0	21
113	Mevalonate kinase deficiency leads to decreased prenylation of Rab GTPases. <i>Immunology and Cell Biology</i> , 2016, 94, 994-999.	1.0	36
114	Cytotoxic T cells that escape exhaustion. <i>Nature</i> , 2016, 537, 312-314.	13.7	6
115	IL-27 Directly Enhances Germinal Center B Cell Activity and Potentiates Lupus in <i>Sanroque</i> Mice. <i>Journal of Immunology</i> , 2016, 197, 3008-3017.	0.4	27
116	Dual T cell and B cell intrinsic deficiency in humans with biallelic <i>RLTPR</i> mutations. <i>Journal of Experimental Medicine</i> , 2016, 213, 2413-2435.	4.2	117
117	B-cell specific STAT3 deficiency: Insight into the molecular basis of autosomal-dominant hyper-IgE syndrome. <i>Journal of Allergy and Clinical Immunology</i> , 2016, 138, 1455-1458.e3.	1.5	28
118	The Integrin LFA-1 Controls T Follicular Helper Cell Generation and Maintenance. <i>Immunity</i> , 2016, 45, 831-846.	6.6	65
119	The Expanding Spectrum of NFκB1 Deficiency. <i>Journal of Clinical Immunology</i> , 2016, 36, 531-532.	2.0	5
120	Genetic cause of immune dysregulation – one gene or two?. <i>Journal of Clinical Investigation</i> , 2016, 126, 4065-4067.	3.9	5
121	Compartmentalization of Total and Virus-Specific Tissue-Resident Memory CD8+ T Cells in Human Lymphoid Organs. <i>PLoS Pathogens</i> , 2016, 12, e1005799.	2.1	74
122	Thucydides and longer-lived plasma cells. <i>Blood</i> , 2015, 125, 1684-1685.	0.6	1
123	Monogenic mutations differentially affect the quantity and quality of T follicular helper cells in patients with human primary immunodeficiencies. <i>Journal of Allergy and Clinical Immunology</i> , 2015, 136, 993-1006.e1.	1.5	181
124	Impairment of immunity to <i>Candida</i> and <i>Mycobacterium</i> in humans with bi-allelic <i>RORC</i> mutations. <i>Science</i> , 2015, 349, 606-613.	6.0	366
125	FAS Inactivation Releases Unconventional Germinal Center B Cells that Escape Antigen Control and Drive IgE and Autoantibody Production. <i>Immunity</i> , 2015, 42, 890-902.	6.6	77
126	STAT3 is a critical cell-intrinsic regulator of human unconventional T cell numbers and function. <i>Journal of Experimental Medicine</i> , 2015, 212, 855-864.	4.2	70



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127	Advances in IL-21 biologyâ€”enhancing our understanding of human disease. <i>Current Opinion in Immunology</i> , 2015, 34, 107-115.	2.4	62
128	T Follicular Helper Cells Have Distinct Modes of Migration and Molecular Signatures in Naive and Memory Immune Responses. <i>Immunity</i> , 2015, 42, 704-718.	6.6	159
129	SnapShot: Interactions between B Cells and T Cells. <i>Cell</i> , 2015, 162, 926-926.e1.	13.5	25
130	Cerebral Vasculitis in X-linked Lymphoproliferative Disease Cured by Matched Unrelated Cord Blood Transplant. <i>Journal of Clinical Immunology</i> , 2015, 35, 604-609.	2.0	17
131	Human TYK2 deficiency: Mycobacterial and viral infections without hyper-IgE syndrome. <i>Journal of Experimental Medicine</i> , 2015, 212, 1641-1662.	4.2	293
132	STAT3 interrupts ATR-Chk1 signaling to allow oncovirus-mediated cell proliferation. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014, 111, 4946-4951.	3.3	72
133	Cytokine-Mediated Regulation of Plasma Cell Generation: IL-21 Takes Center Stage. <i>Frontiers in Immunology</i> , 2014, 5, 65.	2.2	186
134	Human T follicular helper cells in primary immunodeficiencies. <i>Current Opinion in Pediatrics</i> , 2014, 26, 720-726.	1.0	15
135	T cells require DOCK8 for flexibility and function. <i>Journal of Experimental Medicine</i> , 2014, 211, 2482-2483.	4.2	1
136	Dominant-activating germline mutations in the gene encoding the PI(3)K catalytic subunit p110 $\beta$ result in T cell senescence and human immunodeficiency. <i>Nature Immunology</i> , 2014, 15, 88-97.	7.0	575
137	Cell membrane associated free kappa light chains are found on a subset of tonsil and in vitro-derived plasmablasts. <i>Human Immunology</i> , 2014, 75, 986-990.	1.2	7
138	Immune dysregulation in human subjects with heterozygous germline mutations in <i>CTLA4</i> . <i>Science</i> , 2014, 345, 1623-1627.	6.0	745
139	STAT3 is a central regulator of lymphocyte differentiation and function. <i>Current Opinion in Immunology</i> , 2014, 28, 49-57.	2.4	76
140	XLP: Clinical Features and Molecular Etiology due to Mutations in SH2D1A Encoding SAP. <i>Journal of Clinical Immunology</i> , 2014, 34, 772-779.	2.0	105
141	Signaling lymphocytic activation molecule (SLAM)/SLAM-associated protein pathway regulates human B-cell tolerance. <i>Journal of Allergy and Clinical Immunology</i> , 2014, 133, 1149-1161.	1.5	33
142	Editorial overview: Lymphocyte activation and effector functions. <i>Current Opinion in Immunology</i> , 2014, 28, v-vii.	2.4	0
143	The right â€œjobâ€”for STAT3 mutant mice!. <i>Blood</i> , 2014, 123, 2907-2909.	0.6	1
144	Signal transducer and activator of transcription 3 (STAT3) mutations underlying autosomal dominant hyper-IgE syndrome impair human CD8+ T-cell memory formation and function. <i>Journal of Allergy and Clinical Immunology</i> , 2013, 132, 400-411.e9.	1.5	63

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145	IL-21 signalling via STAT3 primes human naïve B cells to respond to IL-2 to enhance their differentiation into plasmablasts. <i>Blood</i> , 2013, 122, 3940-3950.	0.6	121
146	Circulating Precursor CCR7 <sup>lo</sup> PD-1 <sup>hi</sup> CXCR5 <sup>+</sup> CD4 <sup>+</sup> T Cells Indicate Tfh Cell Activity and Promote Antibody Responses upon Antigen Reexposure. <i>Immunity</i> , 2013, 39, 770-781.	6.6	571
147	The good, the bad and the ugly – TFH cells in human health and disease. <i>Nature Reviews Immunology</i> , 2013, 13, 412-426.	10.6	475
148	Transitional B cell subsets in human bone marrow. <i>Clinical and Experimental Immunology</i> , 2013, 174, 53-59.	1.1	29
149	Inherited human OX40 deficiency underlying classic Kaposi sarcoma of childhood. <i>Journal of Experimental Medicine</i> , 2013, 210, 1743-1759.	4.2	119
150	Signal Transducer and Activator of Transcription 3 Limits Epstein-Barr Virus Lytic Activation in B Lymphocytes. <i>Journal of Virology</i> , 2013, 87, 11438-11446.	1.5	42
151	Naive and memory human B cells have distinct requirements for STAT3 activation to differentiate into antibody-secreting plasma cells. <i>Journal of Experimental Medicine</i> , 2013, 210, 2739-2753.	4.2	158
152	To B1 or not to B1: that really is still the question!. <i>Blood</i> , 2013, 121, 5109-5110.	0.6	47
153	DOCK8 is critical for the survival and function of NKT cells. <i>Blood</i> , 2013, 122, 2052-2061.	0.6	68
154	A recurrent dominant negative E47 mutation causes agammaglobulinemia and BCR <sup>hi</sup> , <sup>+</sup> B cells. <i>Journal of Clinical Investigation</i> , 2013, 123, 4781-4785.	3.9	94
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