Stuart G Tangye

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

82 22,419 231 147 h-index g-index citations papers 262 6.85 27,566 12.5 L-index avg, IF ext. citations ext. papers

#	Paper	IF	Citations
231	Human genetic and immunological determinants of critical COVID-19 pneumonia <i>Nature</i> , 2022 ,	50.4	23
230	Getting to the (germinal) center of humoral immune responses to SARS-CoV-2 <i>Cell</i> , 2022 , 185, 945-948	3 56.2	O
229	"Are you gonna go my way?"-Decisions at the Tfh-B cell interface Immunity, 2022, 55, 377-379	32.3	
228	Severe COVID-19 represents an undiagnosed primary immunodeficiency in a high proportion of infected individuals <i>Clinical and Translational Immunology</i> , 2022 , 11, e1365	6.8	O
227	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, e220041311	9 ^{11.5}	3
226	The expansion of human T-betCD21 B cells is T cell dependent. <i>Science Immunology</i> , 2021 , 6, eabh0891	28	11
225	Intrinsic Defects in B Cell Development and Differentiation, T Cell Exhaustion and Altered Unconventional T Cell Generation Characterize Human Adenosine Deaminase Type 2 Deficiency. <i>Journal of Clinical Immunology</i> , 2021 , 41, 1915-1935	5.7	5
224	Hyper-IgE Syndrome due to an Elusive Novel Intronic Homozygous Variant in DOCK8. <i>Journal of Clinical Immunology</i> , 2021 , 1	5.7	O
223	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,	16.6	45
222	Genomic Spectrum and Phenotypic Heterogeneity of Human IL-21 Receptor Deficiency. <i>Journal of Clinical Immunology</i> , 2021 , 41, 1272-1290	5.7	5
221	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	17.4	4
220	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	5.7	1
219	Human STAT3 variants underlie autosomal dominant hyper-IgE syndrome by negative dominance. <i>Journal of Experimental Medicine</i> , 2021 , 218,	16.6	6
218	High Th2 cytokine levels and upper airway inflammation in human inherited T-bet deficiency. <i>Journal of Experimental Medicine</i> , 2021 , 218,	16.6	7
217	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	5.1	2
216	Inherited PD-1 deficiency underlies tuberculosis and autoimmunity in a child. <i>Nature Medicine</i> , 2021 , 27, 1646-1654	50.5	17
215	Humans with inherited Trell CD28 deficiency are susceptible to skin papillomaviruses but are otherwise healthy. <i>Cell</i> , 2021 , 184, 3812-3828.e30	56.2	18

(2020-2021)

214	Tissue-resident regulatory T cells accumulate at human barrier lymphoid organs. <i>Immunology and Cell Biology</i> , 2021 , 99, 894-906	5	О
213	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	11.5	142
212	Molecular requirements for human lymphopoiesis as defined by inborn errors of immunity. <i>Stem Cells</i> , 2021 , 39, 389-402	5.8	1
211	Somatic reversion of pathogenic DOCK8 variants alters lymphocyte differentiation and function to effectively cure DOCK8 deficiency. <i>Journal of Clinical Investigation</i> , 2021 , 131,	15.9	12
210	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	5.7	66
209	CD4 T cells that help B cells - a proposal for uniform nomenclature. <i>Trends in Immunology</i> , 2021 , 42, 658	-669	16
208	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,	28	91
207	X-linked recessive TLR7 deficiency in ~1% of men under 60 years old with life-threatening COVID-19. <i>Science Immunology</i> , 2021 , 6,	28	67
206	Inherited human c-Rel deficiency disrupts myeloid and lymphoid immunity to multiple infectious agents. <i>Journal of Clinical Investigation</i> , 2021 , 131,	15.9	3
205	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	3.3	6
204	Coronavirus disease 2019 in patients with inborn errors of immunity: lessons learned. <i>Current Opinion in Pediatrics</i> , 2021 , 33, 648-656	3.2	10
203	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	7.8	O
202	Identification of Germline Monoallelic Mutations in IKZF2 in Patients with Immune Dysregulation <i>Blood Advances</i> , 2021 ,	7.8	1
201	Human T-bet Governs Innate and Innate-like Adaptive IFN-Immunity against Mycobacteria. <i>Cell</i> , 2020 , 183, 1826-1847.e31	56.2	35
200	A Global Effort to Define the Human Genetics of Protective Immunity to SARS-CoV-2 Infection. <i>Cell</i> , 2020 , 181, 1194-1199	56.2	113
199	Three Copies of Four Interferon Receptor Genes Underlie a Mild Type I Interferonopathy in Down Syndrome. <i>Journal of Clinical Immunology</i> , 2020 , 40, 807-819	5.7	16
198	The Clinical Immunogenomics Research Consortium Australasia (CIRCA): a Distributed Network Model for Genomic Healthcare Delivery. <i>Journal of Clinical Immunology</i> , 2020 , 40, 763-766	5.7	2
197	Everolimus-Induced Remission of Classic KaposiN Sarcoma Secondary to Cryptic Splicing Mediated CTLA4 Haploinsufficiency. <i>Journal of Clinical Immunology</i> , 2020 , 40, 774-779	5.7	2

196	Genetic susceptibility to EBV infection: insights from inborn errors of immunity. <i>Human Genetics</i> , 2020 , 139, 885-901	6.3	22
195	Extended clinical and immunological phenotype and transplant outcome in CD27 and CD70 deficiency. <i>Blood</i> , 2020 , 136, 2638-2655	2.2	32
194	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	5.7	497
193	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	5.7	17
192	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	5.6	2
191	Human Inborn Errors of Immunity: 2019 Update of the IUIS Phenotypical Classification. <i>Journal of Clinical Immunology</i> , 2020 , 40, 66-81	5.7	267
190	Human inborn errors of immunity to herpes viruses. Current Opinion in Immunology, 2020, 62, 106-122	7.8	33
189	Diversity of XMEN Disease: Description of 2 Novel Variants and Analysis of the Lymphocyte Phenotype. <i>Journal of Clinical Immunology</i> , 2020 , 40, 299-309	5.7	14
188	Unresponsiveness to inhaled antigen is governed by conventional dendritic cells and overridden during infection by monocytes. <i>Science Immunology</i> , 2020 , 5,	28	5
187	Activated PI3KIbreaches multiple B cell tolerance checkpoints and causes autoantibody production. <i>Journal of Experimental Medicine</i> , 2020 , 217,	16.6	18
186	Regulation of the germinal center and humoral immunity by interleukin-21. <i>Journal of Experimental Medicine</i> , 2020 , 217,	16.6	37
185	Dominant-negative mutations in human IL6ST underlie hyper-IgE syndrome. <i>Journal of Experimental Medicine</i> , 2020 , 217,	16.6	36
184	Autoantibodies against type I IFNs in patients with life-threatening COVID-19. Science, 2020, 370,	33.3	1090
183	Primary immunodeficiencies reveal the molecular requirements for effective host defense against EBV infection. <i>Blood</i> , 2020 , 135, 644-655	2.2	40
182	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	8.4	13
181	An essential role for the Zn transporter ZIP7 in B cell development. <i>Nature Immunology</i> , 2019 , 20, 350-3	3 61 5.1	54
180	The FOXP3 isoform supports Treg cell development and protects against severe IPEX syndrome. <i>Journal of Allergy and Clinical Immunology</i> , 2019 , 144, 317-320.e8	11.5	11
179	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	11.5	15

(2018-2019)

178	Human DOCK2 Deficiency: Report of a Novel Mutation and Evidence for Neutrophil Dysfunction. Journal of Clinical Immunology, 2019 , 39, 298-308	5.7	17
177	Activating mutations in PIK3CD disrupt the differentiation and function of human and murine CD4 T cells. <i>Journal of Allergy and Clinical Immunology</i> , 2019 , 144, 236-253	11.5	31
176	Immune Dysregulation and Disease Pathogenesis due to Activating Mutations in PIK3CD-the GoldilocksNEffect. <i>Journal of Clinical Immunology</i> , 2019 , 39, 148-158	5.7	20
175	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	11.5	44
174	A deep intronic splice mutation of underlies hyper IgE syndrome by negative dominance. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 16463-16472	11.5	11
173	Hematopoietic stem cell transplant effectively rescues lymphocyte differentiation and function in DOCK8-deficient patients. <i>JCI Insight</i> , 2019 , 5,	9.9	12
172	Human CD8 T cell cross-reactivity across influenza A, B and C viruses. <i>Nature Immunology</i> , 2019 , 20, 613-	-6251	109
171	Human inborn errors of the actin cytoskeleton affecting immunity: way beyond WAS and WIP. <i>Immunology and Cell Biology</i> , 2019 , 97, 389-402	5	22
170	Chronic mucocutaneous candidiasis and connective tissue disorder in humans with impaired JNK1-dependent responses to IL-17A/F and TGF-\(\Bar{O} \) Science Immunology, 2019 , 4,	28	25
169	Denisovan, modern human and mouse TNFAIP3 alleles tune A20 phosphorylation and immunity. <i>Nature Immunology</i> , 2019 , 20, 1299-1310	19.1	29
168	What can primary immunodeficiencies teach us about Th9 cell differentiation and function?. <i>Immunology and Cell Biology</i> , 2019 , 97, 380-388	5	1
167	STAT3 regulates cytotoxicity of human CD57+ CD4+ T cells in blood and lymphoid follicles. <i>Scientific Reports</i> , 2018 , 8, 3529	4.9	18
166	Combined Immunodeficiency with Ring Chromosome 21. Journal of Clinical Immunology, 2018, 38, 251-2	? 5 67	1
165	Circulating T cells, serological memory, and tissue compartmentalization shape human influenza-specific B cell immunity. <i>Science Translational Medicine</i> , 2018 , 10,	17.5	117
164	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	5.7	1
163	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	11.5	14
162	Is it dead or alive? TLR8 can tell. <i>Nature Immunology</i> , 2018 , 19, 324-326	19.1	2
161	IRF4 haploinsufficiency in a family with WhippleN disease. <i>ELife</i> , 2018 , 7,	8.9	25

160	B cells race the clock to get a second wind. <i>Nature Immunology</i> , 2018 , 19, 791-793	19.1	1
159	Germline-activating mutations in compromise B cell development and function. <i>Journal of Experimental Medicine</i> , 2018 , 215, 2073-2095	16.6	53
158	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	11.5	43
157	Disruption of an antimycobacterial circuit between dendritic and helper T cells in human SPPL2a deficiency. <i>Nature Immunology</i> , 2018 , 19, 973-985	19.1	67
156	Memory B cells are reactivated in subcapsular proliferative foci of lymph nodes. <i>Nature Communications</i> , 2018 , 9, 3372	17.4	50
155	A recessive form of hyper-IgE syndrome by disruption of ZNF341-dependent STAT3 transcription and activity. <i>Science Immunology</i> , 2018 , 3,	28	82
154	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	5.7	510
153	The 2017 IUIS Phenotypic Classification for Primary Immunodeficiencies. <i>Journal of Clinical Immunology</i> , 2018 , 38, 129-143	5.7	345
152	Human IFN-IImmunity to mycobacteria is governed by both IL-12 and IL-23. <i>Science Immunology</i> , 2018 , 3,	28	83
151	Tuberculosis and impaired IL-23-dependent IFN-Immunity in humans homozygous for a common missense variant. <i>Science Immunology</i> , 2018 , 3,	28	88
150	Chronic Aichi Virus Infection in a Patient with X-Linked Agammaglobulinemia. <i>Journal of Clinical Immunology</i> , 2018 , 38, 748-752	5.7	11
149	Human immunity against EBV-lessons from the clinic. <i>Journal of Experimental Medicine</i> , 2017 , 214, 269-	2 88 .6	96
148	The TORC that Gets the GC Cycling. <i>Immunity</i> , 2017 , 46, 974-976	32.3	О
147	Memory B cells: total recall. <i>Current Opinion in Immunology</i> , 2017 , 45, 132-140	7.8	32
146	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	16.6	111
145	Arginine methylation catalyzed by PRMT1 is required for B cell activation and differentiation. <i>Nature Communications</i> , 2017 , 8, 891	17.4	20
144	Cytokine-Mediated Regulation of Human Lymphocyte Development and Function: Insights from Primary Immunodeficiencies. <i>Journal of Immunology</i> , 2017 , 199, 1949-1958	5.3	12
143	DOCK8 Drives Src-Dependent NK Cell Effector Function. <i>Journal of Immunology</i> , 2017 ,	5.3	12

(2016-2017)

142	Defective protein prenylation is a diagnostic biomarker of mevalonate kinase deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2017 , 140, 873-875.e6	11.5	19
141	Low IgE Is Insufficiently Sensitive to Guide Genetic Testing of Gain-of-Function Mutations. <i>Clinical Chemistry</i> , 2017 , 63, 1539-1540	5.5	4
140	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	5.7	5
139	Dedicator of cytokinesis 8-deficient CD4 Titells are biased to a T2 effector fate at the expense of T1 and T17Itells. <i>Journal of Allergy and Clinical Immunology</i> , 2017 , 139, 933-949	11.5	51
138	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	32.3	107
137	Inherited GINS1 deficiency underlies growth retardation along with neutropenia and NK cell deficiency. <i>Journal of Clinical Investigation</i> , 2017 , 127, 1991-2006	15.9	73
136	Immunology: Cytotoxic T cells that escape exhaustion. <i>Nature</i> , 2016 , 537, 312-314	50.4	4
135	IL-27 Directly Enhances Germinal Center B Cell Activity and Potentiates Lupus in Sanroque Mice. <i>Journal of Immunology</i> , 2016 , 197, 3008-3017	5.3	17
134	Dual T cell- and B cell-intrinsic deficiency in humans with biallelic RLTPR mutations. <i>Journal of Experimental Medicine</i> , 2016 , 213, 2413-2435	16.6	75
133	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	11.5	19
132	The Integrin LFA-1 Controls T Follicular Helper Cell Generation and Maintenance. <i>Immunity</i> , 2016 , 45, 831-846	32.3	42
131	The Expanding Spectrum of NFkB1 Deficiency. <i>Journal of Clinical Immunology</i> , 2016 , 36, 531-2	5.7	4
130	Genetic cause of immune dysregulation - one gene or two?. <i>Journal of Clinical Investigation</i> , 2016 , 126, 4065-4067	15.9	4
129	Compartmentalization of Total and Virus-Specific Tissue-Resident Memory CD8+ T Cells in Human Lymphoid Organs. <i>PLoS Pathogens</i> , 2016 , 12, e1005799	7.6	57
128	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, 1589-608	16.6	51
127	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	11.5	10
126	NaWe and memory B cells exhibit distinct biochemical responses following BCR engagement. <i>Immunology and Cell Biology</i> , 2016 , 94, 774-86	5	16
125	Mevalonate kinase deficiency leads to decreased prenylation of Rab GTPases. <i>Immunology and Cell Biology</i> , 2016 , 94, 994-999	5	20

124	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	11.5	126
123	IMMUNODEFICIENCIES. Impairment of immunity to Candida and Mycobacterium in humans with bi-allelic RORC mutations. <i>Science</i> , 2015 , 349, 606-613	33.3	291
122	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	32.3	59
121	STAT3 is a critical cell-intrinsic regulator of human unconventional T cell numbers and function. <i>Journal of Experimental Medicine</i> , 2015 , 212, 855-64	16.6	54
120	Advances in IL-21 biology - enhancing our understanding of human disease. <i>Current Opinion in Immunology</i> , 2015 , 34, 107-15	7.8	54
119	T follicular helper cells have distinct modes of migration and molecular signatures in naive and memory immune responses. <i>Immunity</i> , 2015 , 42, 704-18	32.3	125
118	SnapShot: Interactions between B Cells and T Cells. <i>Cell</i> , 2015 , 162, 926-6.e1	56.2	19
117	Cerebral Vasculitis in X-linked Lymphoproliferative Disease Cured by Matched Unrelated Cord Blood Transplant. <i>Journal of Clinical Immunology</i> , 2015 , 35, 604-9	5.7	13
116	Human TYK2 deficiency: Mycobacterial and viral infections without hyper-IgE syndrome. <i>Journal of Experimental Medicine</i> , 2015 , 212, 1641-62	16.6	209
115	Thucydides and longer-lived plasma cells. <i>Blood</i> , 2015 , 125, 1684-5	2.2	1
114	Dominant-activating germline mutations in the gene encoding the PI(3)K catalytic subunit p110 result in T cell senescence and human immunodeficiency. <i>Nature Immunology</i> , 2014 , 15, 88-97	19.1	453
113	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, 986-90	2.3	3
112	Immune dysregulation in human subjects with heterozygous germline mutations in CTLA4. <i>Science</i> , 2014 , 345, 1623-1627	33.3	563
111	STAT3 is a central regulator of lymphocyte differentiation and function. <i>Current Opinion in Immunology</i> , 2014 , 28, 49-57	7.8	62
110	XLP: clinical features and molecular etiology due to mutations in SH2D1A encoding SAP. <i>Journal of Clinical Immunology</i> , 2014 , 34, 772-9	5.7	84
109	Signaling lymphocytic activation molecule (SLAM)/SLAM-associated protein pathway regulates human B-cell tolerance. <i>Journal of Allergy and Clinical Immunology</i> , 2014 , 133, 1149-61	11.5	31
108	The right "Job" for STAT3 mutant mice!. <i>Blood</i> , 2014 , 123, 2907-9	2.2	1
107	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, 4946-51	11.5	59

106	Cytokine-Mediated Regulation of Plasma Cell Generation: IL-21 Takes Center Stage. <i>Frontiers in Immunology</i> , 2014 , 5, 65	8.4	116
105	Human T follicular helper cells in primary immunodeficiencies. <i>Current Opinion in Pediatrics</i> , 2014 , 26, 720-6	3.2	13
104	T cells require DOCK8 for flexibility and function. <i>Journal of Experimental Medicine</i> , 2014 , 211, 2482-3	16.6	1
103	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, 400-11.e9	11.5	48
102	IL-21 signalling via STAT3 primes human naive B cells to respond to IL-2 to enhance their differentiation into plasmablasts. <i>Blood</i> , 2013 , 122, 3940-50	2.2	84
101	Circulating precursor CCR7(lo)PD-1(hi) CXCR5+ CD4+ T cells indicate Tfh cell activity and promote antibody responses upon antigen reexposure. <i>Immunity</i> , 2013 , 39, 770-81	32.3	449
100	The good, the bad and the ugly - TFH cells in human health and disease. <i>Nature Reviews Immunology</i> , 2013 , 13, 412-26	36.5	402
99	Transitional B cell subsets in human bone marrow. Clinical and Experimental Immunology, 2013, 174, 53-	-%.2	21
98	Inherited human OX40 deficiency underlying classic Kaposi sarcoma of childhood. <i>Journal of Experimental Medicine</i> , 2013 , 210, 1743-59	16.6	99
97	Signal transducer and activator of transcription 3 limits Epstein-Barr virus lytic activation in B lymphocytes. <i>Journal of Virology</i> , 2013 , 87, 11438-46	6.6	37
96	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, 2739-53	16.6	121
95	To B1 or not to B1: that really is still the question!. <i>Blood</i> , 2013 , 121, 5109-10	2.2	36
94	DOCK8 is critical for the survival and function of NKT cells. <i>Blood</i> , 2013 , 122, 2052-61	2.2	60
93	A recurrent dominant negative E47 mutation causes agammaglobulinemia and BCR(-) B cells. <i>Journal of Clinical Investigation</i> , 2013 , 123, 4781-5	15.9	78
92	Functional STAT3 deficiency compromises the generation of human T follicular helper cells. <i>Blood</i> , 2012 , 119, 3997-4008	2.2	230
91	Clinical, molecular, and cellular immunologic findings in patients with SP110-associated veno-occlusive disease with immunodeficiency syndrome. <i>Journal of Allergy and Clinical Immunology</i> , 2012 , 130, 735-742.e6	11.5	41
90	The origins, function, and regulation of T follicular helper cells. <i>Journal of Experimental Medicine</i> , 2012 , 209, 1241-53	16.6	387
89	T cell-B cell interactions in primary immunodeficiencies. <i>Annals of the New York Academy of Sciences</i> , 2012 , 1250, 1-13	6.5	24

88	Expansion of somatically reverted memory CD8+ T cells in patients with X-linked lymphoproliferative disease caused by selective pressure from Epstein-Barr virus. <i>Journal of Experimental Medicine</i> , 2012 , 209, 913-24	16.6	47
87	A new ICB sister journal focuses on clinical and translational immunology. <i>Clinical and Translational Immunology</i> , 2012 , 1, e1	6.8	
86	Human RHOH deficiency causes T cell defects and susceptibility to EV-HPV infections. <i>Journal of Clinical Investigation</i> , 2012 , 122, 3239-47	15.9	109
85	Identification of Bcl-6-dependent follicular helper NKT cells that provide cognate help for B cell responses. <i>Nature Immunology</i> , 2011 , 13, 35-43	19.1	205
84	Molecular pathogenesis of EBV susceptibility in XLP as revealed by analysis of female carriers with heterozygous expression of SAP. <i>PLoS Biology</i> , 2011 , 9, e1001187	9.7	89
83	SLAM family receptors and SAP adaptors in immunity. <i>Annual Review of Immunology</i> , 2011 , 29, 665-705	34.7	350
82	Staying alive: regulation of plasma cell survival. <i>Trends in Immunology</i> , 2011 , 32, 595-602	14.4	100
81	Calcineurin-dependent negative regulation of CD94/NKG2A expression on naive CD8+ T cells. <i>Blood</i> , 2011 , 118, 116-28	2.2	17
80	A subset of interleukin-21+ chemokine receptor CCR9+ T helper cells target accessory organs of the digestive system in autoimmunity. <i>Immunity</i> , 2011 , 34, 602-15	32.3	92
79	Plasmacytoid DCs induce gutsy plasma cells. <i>Immunity</i> , 2011 , 34, 144-6	32.3	2
78	Regulation of T follicular helper cell formation and function by antigen presenting cells. <i>Current Opinion in Immunology</i> , 2011 , 23, 111-8	7.8	58
77	IL-21 is the primary common Ethain-binding cytokine required for human B-cell differentiation in vivo. <i>Blood</i> , 2011 , 118, 6824-35	2.2	115
76	DOCK8 deficiency impairs CD8 T cell survival and function in humans and mice. <i>Journal of Experimental Medicine</i> , 2011 , 208, 2305-20	16.6	140
75	CXCR5 expressing human central memory CD4 T cells and their relevance for humoral immune responses. <i>Journal of Immunology</i> , 2011 , 186, 5556-68	5.3	246
74	Human Th9 cells: inflammatory cytokines modulate IL-9 production through the induction of IL-21. <i>Immunology and Cell Biology</i> , 2010 , 88, 621-3	5	21
73	Comprehensive analysis of the cytokine-rich chromosome 5q31.1 region suggests a role for IL-4 gene variants in prostate cancer risk. <i>Carcinogenesis</i> , 2010 , 31, 1748-54	4.6	34
72	B cell-intrinsic signaling through IL-21 receptor and STAT3 is required for establishing long-lived antibody responses in humans. <i>Journal of Experimental Medicine</i> , 2010 , 207, 155-71	16.6	277
71	IL-27 supports germinal center function by enhancing IL-21 production and the function of T follicular helper cells. <i>Journal of Experimental Medicine</i> , 2010 , 207, 2895-906	16.6	160

(2008-2010)

70	Therapeutic implications of advances in our understanding of transitional B-cell development in humans. <i>Expert Review of Clinical Immunology</i> , 2010 , 6, 765-75	5.1	11
69	Differential expression of CD21 identifies developmentally and functionally distinct subsets of human transitional B cells. <i>Blood</i> , 2010 , 115, 519-29	2.2	89
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