

Raif S Geha

List of Publications by Year in descending order

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Version: 2024-02-01

150
papers

8,434
citations

34016

52
h-index

53109

85
g-index

157
all docs

157
docs citations

157
times ranked

10109
citing authors

#	ARTICLE	IF	CITATIONS
1	Linker-Improved Chimeric Endolysin Selectively Kills <i>Staphylococcus aureus</i> <i>In Vitro</i> , on Reconstituted Human Epidermis, and in a Murine Model of Skin Infection. <i>Antimicrobial Agents and Chemotherapy</i> , 2022, 66, e0227321.	1.4	12
2	Inborn Errors of the Immune System Associated With Atopy. <i>Frontiers in Immunology</i> , 2022, 13, 860821.	2.2	10
3	ITK deficiency presenting as autoimmune lymphoproliferative syndrome. <i>Journal of Allergy and Clinical Immunology</i> , 2021, 147, 743-745.e1.	1.5	8
4	DOCK8 Expression in Regulatory T Cells Maintains their Stability and Limits Contact Hypersensitivity. <i>Journal of Investigative Dermatology</i> , 2021, 141, 1503-1511.e3.	0.3	7
5	Hematopoietic Stem Cell Transplantation Is a Curative Therapy for Transferrin Receptor 1 (TFRC) Deficiency. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2021, 9, 753-759.e2.	2.0	4
6	Mast cell-derived IL-13 downregulates IL-12 production by skin dendritic cells to inhibit the TH1 cell response to cutaneous antigen exposure. <i>Journal of Allergy and Clinical Immunology</i> , 2021, 147, 2305-2315.e3.	1.5	33
7	Efficacy and economics of targeted panel versus whole-exome sequencing in 878 patients with suspected primary immunodeficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2021, 147, 723-726.	1.5	31
8	Combined immunodeficiency due to a mutation in the $\beta 1$ subunit of the coat protein I complex. <i>Journal of Clinical Investigation</i> , 2021, 131, .	3.9	15
9	Multi-kingdom ecological drivers of microbiota assembly in preterm infants. <i>Nature</i> , 2021, 591, 633-638.	13.7	169
10	Macabre TH2 skewing in DOCK8 deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2021, 148, 73-75.	1.5	5
11	Basophil: The cell that itches. <i>Journal of Allergy and Clinical Immunology</i> , 2021, 148, 708-709.	1.5	1
12	Mechanisms underlying genetic susceptibility to multisystem inflammatory syndrome in children (MIS-C). <i>Journal of Allergy and Clinical Immunology</i> , 2021, 148, 732-738.e1.	1.5	84
13	TNFRSF13B genotypes control immune-mediated pathology by regulating the functions of innate B cells. <i>JCI Insight</i> , 2021, 6, .	2.3	4
14	Combined immunodeficiency with autoimmunity caused by a homozygous missense mutation in inhibitor of nuclear factor κ B kinase alpha (IKK α). <i>Science Immunology</i> , 2021, 6, eabf6723.	5.6	6
15	Basophil-derived IL-4 promotes cutaneous <i>Staphylococcus aureus</i> infection. <i>JCI Insight</i> , 2021, 6, .	2.3	15
16	Hypomorphic variants in AK2 reveal the contribution of mitochondrial function to B-cell activation. <i>Journal of Allergy and Clinical Immunology</i> , 2020, 146, 192-202.	1.5	13
17	Dysregulated actin dynamics in activated PI3K δ syndrome. <i>Clinical Immunology</i> , 2020, 210, 108311.	1.4	7
18	Acetaminophen Inhibits the Neutrophil Oxidative Burst: Implications for Diagnostic Testing. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2020, 8, 3543-3548.	2.0	0

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19	APRIL expression is upregulated in atopic dermatitis skin lesions and at sites of antigen driven allergic skin inflammation in mice. <i>Clinical Immunology</i> , 2020, 219, 108556.	1.4	1
20	Immune dysregulation and multisystem inflammatory syndrome in children (MIS-C) in individuals with haploinsufficiency of SOCS1. <i>Journal of Allergy and Clinical Immunology</i> , 2020, 146, 1194-1200.e1.	1.5	92
21	A regulatory T cell Notch4â€“GDF15 axis licenses tissue inflammation in asthma. <i>Nature Immunology</i> , 2020, 21, 1359-1370.	7.0	70
22	Immunoglobulins in the treatment of COVID-19 infection: Proceed with caution!. <i>Clinical Immunology</i> , 2020, 216, 108459.	1.4	91
23	ILC2 activation by keratinocyte-derived IL-25 drives IL-13 production at sites of allergic skin inflammation. <i>Journal of Allergy and Clinical Immunology</i> , 2020, 145, 1606-1614.e4.	1.5	68
24	Genotype and functional correlates of disease phenotype in deficiency of adenosine deaminase 2 (DADA2). <i>Journal of Allergy and Clinical Immunology</i> , 2020, 145, 1664-1672.e10.	1.5	95
25	Severe combined immunodeficiency caused by inositol-trisphosphate 3-kinase B (ITPKB) deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2020, 145, 1696-1699.e6.	1.5	6
26	Cutaneous barrier leakage and gut inflammation drive skin disease in Omenn syndrome. <i>Journal of Allergy and Clinical Immunology</i> , 2020, 146, 1165-1179.e11.	1.5	13
27	Inherited human IFN-Î³ deficiency underlies mycobacterial disease. <i>Journal of Clinical Investigation</i> , 2020, 130, 3158-3171.	3.9	89
28	IL-22 promotes allergic airway inflammation in epicutaneously sensitized mice. <i>Journal of Allergy and Clinical Immunology</i> , 2019, 143, 619-630.e7.	1.5	41
29	A novel truncating mutation in MYD88 in a patient with BCG adenitis, neutropenia and delayed umbilical cord separation. <i>Clinical Immunology</i> , 2019, 207, 40-42.	1.4	9
30	Combined immunodeficiency in a patient with c-Rel deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2019, 144, 606-608.e4.	1.5	32
31	Outcomes and Treatment Strategies for Autoimmunity and Hyperinflammation in Patients with RAG Deficiency. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2019, 7, 1970-1985.e4.	2.0	64
32	Mechanical Skin Injury Promotes Food Anaphylaxis by Driving Intestinal Mast Cell Expansion. <i>Immunity</i> , 2019, 50, 1262-1275.e4.	6.6	158
33	Rethinking newborn screening for severe combined immunodeficiency: Lessons from an international partnership for patients with primary immunodeficiencies in Pakistan. <i>Clinical Immunology</i> , 2019, 202, 29-32.	1.4	2
34	F-BAR domain only protein 1 (FCHO1) deficiency is a novel cause of combined immune deficiency in human subjects. <i>Journal of Allergy and Clinical Immunology</i> , 2019, 143, 2317-2321.e12.	1.5	21
35	Immunodeficiency and EBV-induced lymphoproliferation caused by 4-1BB deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2019, 144, 574-583.e5.	1.5	63
36	Human primary immunodeficiency caused by expression of a kinase-dead p110Î³ mutant. <i>Journal of Allergy and Clinical Immunology</i> , 2019, 143, 797-799.e2.	1.5	33

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37	Hematopoietic Stem Cell Transplantation as Treatment for Patients with DOCK8 Deficiency. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2019, 7, 848-855.	2.0	67
38	Primary immunodeficiencies caused by mutations in actin regulatory proteins. <i>Immunological Reviews</i> , 2019, 287, 121-134.	2.8	40
39	Injury, dysbiosis, and filaggrin deficiency drive skin inflammation through keratinocyte IL-1 β release. <i>Journal of Allergy and Clinical Immunology</i> , 2019, 143, 1426-1443.e6.	1.5	56
40	Atypical phenotype of an old disease or typical phenotype of a new disease: deficiency of adenosine deaminase 2. <i>Turkish Journal of Pediatrics</i> , 2019, 61, 413.	0.3	5
41	ROR γ -expressing T regulatory cells restrain allergic skin inflammation. <i>Science Immunology</i> , 2018, 3, .	5.6	97
42	WASP-mediated regulation of anti-inflammatory macrophages is IL-10 dependent and is critical for intestinal homeostasis. <i>Nature Communications</i> , 2018, 9, 1779.	5.8	40
43	Mechanisms of genotype-phenotype correlation in autosomal dominant anhidrotic ectodermal dysplasia with immune deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2018, 141, 1060-1073.e3.	1.5	22
44	Exaggerated follicular helper T-cell responses in patients with LRBA deficiency caused by failure of CTLA4-mediated regulation. <i>Journal of Allergy and Clinical Immunology</i> , 2018, 141, 1050-1059.e10.	1.5	93
45	DNA recombination defects in Kuwait: Clinical, immunologic and genetic profile. <i>Clinical Immunology</i> , 2018, 187, 68-75.	1.4	11
46	Clinical, immunologic, and genetic spectrum of 696 patients with combined immunodeficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2018, 141, 1450-1458.	1.5	90
47	Novel biallelic TRNT1 mutations resulting in sideroblastic anemia, combined B and T cell defects, hypogammaglobulinemia, recurrent infections, hypertrophic cardiomyopathy and developmental delay. <i>Clinical Immunology</i> , 2018, 188, 20-22.	1.4	24
48	The Lack of WIP Binding to Actin Results in Impaired B Cell Migration and Altered Humoral Immune Responses. <i>Cell Reports</i> , 2018, 24, 619-629.	2.9	17
49	Defective TLR9-driven STAT3 activation in B cells of patients with CVID. <i>Clinical Immunology</i> , 2018, 197, 40-44.	1.4	7
50	MyD88 signaling in T regulatory cells by endogenous ligands dampens skin inflammation in filaggrin deficient mice. <i>Clinical Immunology</i> , 2018, 195, 88-92.	1.4	1
51	Comprehensive Genetic Results for Primary Immunodeficiency Disorders in a Highly Consanguineous Population. <i>Frontiers in Immunology</i> , 2018, 9, 3146.	2.2	37
52	Deficient LRRC8A-dependent volume-regulated anion channel activity is associated with male infertility in mice. <i>JCI Insight</i> , 2018, 3, .	2.3	29
53	Leucine-rich repeat containing 8A (LRRC8A) α -dependent volume-regulated anion channel activity is dispensable for T-cell development and function. <i>Journal of Allergy and Clinical Immunology</i> , 2017, 140, 1651-1659.e1.	1.5	36
54	Human <i>REL1A</i> haploinsufficiency results in autosomal-dominant chronic mucocutaneous ulceration. <i>Journal of Experimental Medicine</i> , 2017, 214, 1937-1947.	4.2	84

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55	Janus kinase 3 deficiency caused by a homozygous synonymous exonic mutation that creates a dominant splice site. <i>Journal of Allergy and Clinical Immunology</i> , 2017, 140, 268-271.e6.	1.5	11
56	DOCK8 Deficiency Presenting as an IPEX-Like Disorder. <i>Journal of Clinical Immunology</i> , 2017, 37, 811-819.	2.0	39
57	Combined immunodeficiency with EBV positive B cell lymphoma and epidermodysplasia verruciformis due to a novel homozygous mutation in RASGRP1. <i>Clinical Immunology</i> , 2017, 183, 142-144.	1.4	43
58	Advances in basic and clinical immunology in 2016. <i>Journal of Allergy and Clinical Immunology</i> , 2017, 140, 959-973.	1.5	5
59	A novel mutation in the JH4 domain of JAK3 causing severe combined immunodeficiency complicated by vertebral osteomyelitis. <i>Clinical Immunology</i> , 2017, 183, 198-200.	1.4	6
60	Cernunnos deficiency associated with BCG adenitis and autoimmunity: First case from the national Iranian registry and review of the literature. <i>Clinical Immunology</i> , 2017, 183, 201-206.	1.4	11
61	<i>Staphylococcus aureus</i> Epicutaneous Exposure Drives Skin Inflammation via IL-36-Mediated T Cell Responses. <i>Cell Host and Microbe</i> , 2017, 22, 653-666.e5.	5.1	170
62	Disseminated <i>Mycobacterium mageritense</i> and <i>Salmonella</i> Infections Associated with a Novel Variant in NFKBIA. <i>Journal of Clinical Immunology</i> , 2017, 37, 415-418.	2.0	13
63	Epidermodysplasia verruciformis as a manifestation of ARTEMIS deficiency in a young adult. <i>Journal of Allergy and Clinical Immunology</i> , 2017, 139, 372-375.e4.	1.5	18
64	Heterozygosity for transmembrane activator and calcium modulator ligand interactor A144E causes haploinsufficiency and pneumococcal susceptibility in mice. <i>Journal of Allergy and Clinical Immunology</i> , 2017, 139, 1293-1301.e4.	1.5	13
65	The LRRC8A Mediated Ca^{2+} Swell Activated Cl^{-} Conductance Is Dispensable for Vacuolar Homeostasis in Neutrophils. <i>Frontiers in Pharmacology</i> , 2017, 8, 262.	1.6	9
66	Uses of Next-Generation Sequencing Technologies for the Diagnosis of Primary Immunodeficiencies. <i>Frontiers in Immunology</i> , 2017, 8, 847.	2.2	95
67	DOCK8 enforces immunological tolerance by promoting IL-2 signaling and immune synapse formation in Tregs. <i>JCI Insight</i> , 2017, 2, .	2.3	31
68	<i>Staphylococcus aureus</i> Exploits Epidermal Barrier Defects in Atopic Dermatitis to Trigger Cytokine Expression. <i>Journal of Investigative Dermatology</i> , 2016, 136, 2192-2200.	0.3	260
69	Chronic mucocutaneous candidiasis associated with an SH2 domain gain-of-function mutation that enhances STAT1 phosphorylation. <i>Journal of Allergy and Clinical Immunology</i> , 2016, 138, 297-299.	1.5	24
70	Hyper IgM Syndrome: a Report from the USIDNET Registry. <i>Journal of Clinical Immunology</i> , 2016, 36, 490-501.	2.0	92
71	Hematopoietic stem cell transplantation outcomes for 11 patients with dedicator of cytokinesis 8 deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2016, 138, 852-859.e3.	1.5	48
72	Combined immunodeficiency due to a homozygous mutation in ORAI1 that deletes the C-terminus that interacts with STIM 1. <i>Clinical Immunology</i> , 2016, 166-167, 100-102.	1.4	11

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73	Rapid molecular diagnostics of severe primary immunodeficiency determined by using targeted next-generation sequencing. <i>Journal of Allergy and Clinical Immunology</i> , 2016, 138, 1142-1151.e2.	1.5	85
74	Mutations in pyrin masquerading as a primary immunodeficiency. <i>Clinical Immunology</i> , 2016, 171, 65-66.	1.4	2
75	IL-22 derived from $\hat{3}\hat{1}$ T cells restricts <i>Staphylococcus aureus</i> infection of mechanically injured skin. <i>Journal of Allergy and Clinical Immunology</i> , 2016, 138, 1098-1107.e3.	1.5	48
76	IL-23 induced in keratinocytes by endogenous TLR4 ligands polarizes dendritic cells to drive IL-22 responses to skin immunization. <i>Journal of Experimental Medicine</i> , 2016, 213, 2147-2166.	4.2	79
77	IL-33 promotes food anaphylaxis in epicutaneously sensitized mice by targeting mast cells. <i>Journal of Allergy and Clinical Immunology</i> , 2016, 138, 1356-1366.	1.5	157
78	Dedicator of cytokinesis 8 regulates signal transducer and activator of transcription 3 activation and promotes TH17 cell differentiation. <i>Journal of Allergy and Clinical Immunology</i> , 2016, 138, 1384-1394.e2.	1.5	70
79	A missense mutation in TFRC, encoding transferrin receptor 1, causes combined immunodeficiency. <i>Nature Genetics</i> , 2016, 48, 74-78.	9.4	219
80	Thymic stromal lymphopoietin and IL-33 promote skin inflammation and vaccinia virus replication in a mouse model of atopic dermatitis. <i>Journal of Allergy and Clinical Immunology</i> , 2016, 138, 283-286.	1.5	22
81	Spectrum of Phenotypes Associated with Mutations in LRBA. <i>Journal of Clinical Immunology</i> , 2016, 36, 33-45.	2.0	180
82	Recurrent viral infections associated with a homozygous CORO1A mutation that disrupts oligomerization and cytoskeletal association. <i>Journal of Allergy and Clinical Immunology</i> , 2016, 137, 879-888.e2.	1.5	41
83	Autoimmune lymphoproliferative syndrome caused by a homozygous FasL mutation that disrupts FasL assembly. <i>Journal of Allergy and Clinical Immunology</i> , 2016, 137, 324-327.e2.	1.5	13
84	Targeted deep sequencing identifies rare loss-of-function variants in IFNGR1 for risk of atopic dermatitis complicated by eczema herpeticum. <i>Journal of Allergy and Clinical Immunology</i> , 2015, 136, 1591-1600.	1.5	42
85	Defective lymphoid organogenesis underlies the immune deficiency caused by a heterozygous S321 mutation in $I\hat{P}B1\hat{2}$. <i>Journal of Experimental Medicine</i> , 2015, 212, 185-202.	4.2	25
86	The microbiota is important for IL-17A expression and neutrophil infiltration in lesional skin of <i>Flgft/ft</i> mice. <i>Clinical Immunology</i> , 2015, 156, 128-130.	1.4	12
87	Filaggrin deficiency promotes the dissemination of cutaneously inoculated vaccinia virus. <i>Journal of Allergy and Clinical Immunology</i> , 2015, 135, 1511-1518.e6.	1.5	15
88	A novel mutation in NCF2 associated with autoimmune disease and a solitary late-onset infection. <i>Clinical Immunology</i> , 2015, 161, 128-130.	1.4	12
89	Wiskott-Aldrich Syndrome Interacting Protein Deficiency Uncovers the Role of the Co-receptor CD19 as a Generic Hub for PI3 Kinase Signaling in B Cells. <i>Immunity</i> , 2015, 43, 660-673.	6.6	68
90	A novel mutation in ORAI1 presenting with combined immunodeficiency and residual T-cell function. <i>Journal of Allergy and Clinical Immunology</i> , 2015, 136, 479-482.e1.	1.5	28

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91	The Rho GTPase Cdc42 Is Essential for the Activation and Function of Mature B Cells. <i>Journal of Immunology</i> , 2015, 194, 4750-4758.	0.4	26
92	Broad spectrum of autoantibodies in patients with Wiskott-Aldrich syndrome and X-linked thrombocytopenia. <i>Journal of Allergy and Clinical Immunology</i> , 2015, 136, 1401-1404.e3.	1.5	25
93	A novel mutation in ICOS presenting as hypogammaglobulinemia with susceptibility to opportunistic pathogens. <i>Journal of Allergy and Clinical Immunology</i> , 2015, 136, 794-797.e1.	1.5	26
94	Regulatory T-cell deficiency and immune dysregulation, polyendocrinopathy, enteropathy, X-linked-like disorder caused by loss-of-function mutations in LRBA. <i>Journal of Allergy and Clinical Immunology</i> , 2015, 135, 217-227.e9.	1.5	223
95	Dedicator of cytokinesis 8-deficient patients have a breakdown in peripheral B-cell tolerance and defective regulatory T cells. <i>Journal of Allergy and Clinical Immunology</i> , 2014, 134, 1365-1374.	1.5	79
96	Flow cytometry diagnosis of dedicator of cytokinesis 8 (DOCK8) deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2014, 134, 221-223.e7.	1.5	62
97	Food allergy: Insights into etiology, prevention, and treatment provided by murine models. <i>Journal of Allergy and Clinical Immunology</i> , 2014, 133, 309-317.	1.5	96
98	Gene hunting in the genomic era: Approaches to diagnostic dilemmas in patients with primary immunodeficiencies. <i>Journal of Allergy and Clinical Immunology</i> , 2014, 134, 262-268.	1.5	34
99	A systematic analysis of recombination activity and genotype-phenotype correlation in human recombination-activating gene 1 deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2014, 133, 1099-1108.e12.	1.5	132
100	Binding of WIP to Actin Is Essential for T Cell Actin Cytoskeleton Integrity and Tissue Homing. <i>Molecular and Cellular Biology</i> , 2014, 34, 4343-4354.	1.1	21
101	The CARD11-BCL10-MALT1 (CBM) signalosome complex: Stepping into the limelight of human primary immunodeficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2014, 134, 276-284.	1.5	133
102	A novel mutation in FOXP1 resulting in SCID: A case report and literature review. <i>Clinical Immunology</i> , 2014, 155, 30-32.	1.4	32
103	Flow cytometry biomarkers distinguish DOCK8 deficiency from severe atopic dermatitis. <i>Clinical Immunology</i> , 2014, 150, 220-224.	1.4	38
104	Lessons in gene hunting: A RAG1 mutation presenting with agammaglobulinemia and absence of B cells. <i>Journal of Allergy and Clinical Immunology</i> , 2014, 134, 983-985.e1.	1.5	22
105	Plasmacytoid dendritic cell depletion in DOCK8 deficiency: Rescue of severe herpetic infections with IFN- γ 2b therapy. <i>Journal of Allergy and Clinical Immunology</i> , 2014, 133, 1753-1755.e3.	1.5	46
106	Presence of hypogammaglobulinemia and abnormal antibody responses in GATA2 deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2014, 134, 223-226.	1.5	25
107	A novel disease-causing CD40L mutation reduces expression of CD40 ligand, but preserves CD40 binding capacity. <i>Clinical Immunology</i> , 2014, 153, 288-291.	1.4	9
108	T-cell receptor ligation causes Wiskott-Aldrich syndrome protein degradation and F-actin assembly downregulation. <i>Journal of Allergy and Clinical Immunology</i> , 2013, 132, 648-655.e1.	1.5	42

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109	C3a receptor promotes viral containment in mice inoculated with vaccinia virus at sites of allergic skin inflammation. <i>Journal of Allergy and Clinical Immunology</i> , 2013, 132, 746-748.e3.	1.5	0
110	Epicutaneous sensitization results in IgE-dependent intestinal mast cell expansion and food-induced anaphylaxis. <i>Journal of Allergy and Clinical Immunology</i> , 2013, 131, 451-460.e6.	1.5	139
111	Mice Lacking Wasp-Interacting Protein Evolve Anti-Platelet IgAs That Impair Platelet Responses Mediated by the Collagen Receptor GPVI.. <i>Blood</i> , 2008, 112, 1231-1231.	0.6	13
112	Epicutaneous antigen exposure induces a Th17 response that drives airway inflammation after inhalation challenge. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2007, 104, 15817-15822.	3.3	179
113	Charles A. Janeway and Fred S. Rosen: The discovery of gamma globulin therapy and primary immunodeficiency diseases at Boston Children's Hospital. <i>Journal of Allergy and Clinical Immunology</i> , 2005, 116, 937-940.	1.5	6
114	The regulation of immunoglobulin E class-switch recombination. <i>Nature Reviews Immunology</i> , 2003, 3, 721-732.	10.6	399
115	An obligate role for T-cell receptor $\hat{1}\hat{2}^+$ T cells but not T-cell receptor $\hat{3}\hat{1}^+$ T cells, B cells, or CD40/CD40L interactions in a mouse model of atopic dermatitis. <i>Journal of Allergy and Clinical Immunology</i> , 2001, 107, 359-366.	1.5	60
116	WIP regulates N-WASP-mediated actin polymerization and filopodium formation. <i>Nature Cell Biology</i> , 2001, 3, 484-491.	4.6	251
117	Allelic Exclusion of the T Cell Receptor $\hat{1}^2$ Locus Requires the Sh2 Domain-Containing Leukocyte Protein (Slp)-76 Adaptor Protein. <i>Journal of Experimental Medicine</i> , 1999, 190, 1093-1102.	4.2	46
118	CD40 ligation and IL-4 use different mechanisms of transcriptional activation of the human lymphotoxin $\hat{1}\hat{2}$ promoter in B cells. <i>European Journal of Immunology</i> , 1998, 28, 901-906.	1.6	32
119	The hyper-IgM (HIM) syndrome. <i>Seminars in Immunopathology</i> , 1998, 19, 383-399.	4.0	26
120	Searching for genes involved in the pathogenesis of primary immunodeficiency diseases: lessons from mouse knockouts. <i>Journal of Clinical Immunology</i> , 1997, 17, 109-126.	2.0	19
121	Affinity maturation without germinal centres in lymphotoxin- $\hat{1}\hat{2}$ -deficient mice. <i>Nature</i> , 1996, 382, 462-466.	13.7	313
122	CD40-mediated lymphotoxin $\hat{1}\hat{2}$ expression in human B cells is tyrosine kinase dependent. <i>European Journal of Immunology</i> , 1995, 25, 2438-2444.	1.6	20
123	Bacterial Superantigens Induce The Proliferation of Resting $\hat{1}^3\hat{1}^7$ Receptor Bearing T Cells. <i>Immunological Investigations</i> , 1995, 24, 713-724.	1.0	12
124	Protein Tyrosine Kinase Activation and Protein Kinase C Translocation Are Functional Components of CD40 Signal Transduction in Resting Human B Cells. <i>Immunological Investigations</i> , 1994, 23, 437-448.	1.0	11
125	Engagement of MHC class II molecules by staphylococcal superantigens activates src-type protein tyrosine kinases. <i>European Journal of Immunology</i> , 1994, 24, 651-658.	1.6	58
126	Decreased expression of the ligand for CD40 in newborn lymphocytes. <i>European Journal of Immunology</i> , 1994, 24, 1925-1928.	1.6	71

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127	Activator protein-1 (AP-1) is stimulated by microbial superantigens in human monocytic cells. <i>European Journal of Immunology</i> , 1993, 23, 2129-2135.	1.6	16
128	Signal Transduction by Microbial Superantigens via MHC class II Molecules. <i>Immunological Reviews</i> , 1993, 131, 43-59.	2.8	59
129	Severe Combined Immunodeficiency with Selective T-Cell Cytokine Genes. <i>Pediatric Research</i> , 1993, 33, S20-S23.	1.1	2
130	Severe Combined Immunodeficiency with Selective T-Cell Cytokine Genes. <i>Pediatric Research</i> , 1993, 33, S20-S23.	1.1	4
131	Novel immune deficiencies: defective transcription of lymphokine genes. <i>Clinical and Experimental Allergy</i> , 1991, 21, 190-194.	1.4	0
132	Binding of toxic shock syndrome toxin-1 to murine major histocompatibility complex class II molecules. <i>European Journal of Immunology</i> , 1990, 20, 1911-1916.	1.6	39
133	Interleukin 4 down-regulates the expression of CD14 in normal human monocytes. <i>European Journal of Immunology</i> , 1990, 20, 2375-2381.	1.6	145
134	Regulation of Human IgE Synthesis. <i>International Reviews of Immunology</i> , 1989, 5, 111-115.	1.5	7
135	Endogenous interleukin 6 plays an obligatory role in interleukin 4-dependent human IgE synthesis. <i>European Journal of Immunology</i> , 1989, 19, 1419-1424.	1.6	208
136	The B-cell binding site on human immunoglobulin E. <i>Nature</i> , 1989, 338, 649-651.	13.7	145
137	Regulatory effects of human IgE-binding factors in the IgE synthesis by human and rat lymphocytes. <i>European Journal of Immunology</i> , 1988, 18, 1663-1670.	1.6	6
138	Regulation of the Human IgE Antibody Response. <i>International Reviews of Immunology</i> , 1987, 2, 75-91.	1.5	52
139	Characterization of human T cell-derived IgE-potentiating factor. <i>European Journal of Immunology</i> , 1986, 16, 985-991.	1.6	20
140	Control of IgE synthesis in man. <i>Journal of Clinical Immunology</i> , 1986, 6, 273-283.	2.0	24
141	Polymerase activity in lymphocyte culture supernatants from patients with Kawasaki disease. <i>Nature</i> , 1986, 323, 814-816.	13.7	104
142	Inhibition of the Prausnitz-K�stner reaction by an immunoglobulin �-chain fragment synthesized in <i>E. coli</i> . <i>Nature</i> , 1985, 315, 577-578.	13.7	69
143	Analysis of antigen uptake and presentation by Epstein-Barr virus-transformed human lymphoblastoid B cells. <i>European Journal of Immunology</i> , 1984, 14, 291-298.	1.6	18
144	Production of IgE-potentiating factor in man by T cell lines bearing Fc receptors for IgE. <i>European Journal of Immunology</i> , 1984, 14, 871-878.	1.6	82

#	ARTICLE	IF	CITATIONS
145	Intestinal Malakoplakia in Childhood: Case Report and Review of Literature. <i>Pediatric Pathology</i> , 1983, 1, 337-343.	0.5	14
146	Nature of the immunogenic moiety recognized by the human T cell proliferating in response to tetanus toxoid antigen. <i>European Journal of Immunology</i> , 1981, 11, 365-371.	1.6	23
147	Macrophage T-cell interaction in man: Handling of tetanus toxoid antigen by human monocytes. <i>Journal of Clinical Immunology</i> , 1981, 1, 21-29.	2.0	14
148	Regulation of Human B Cell Activation. <i>Immunological Reviews</i> , 1979, 45, 275-305.	2.8	44
149	Heterogeneity of "Acquired" or Common Variable Agammaglobulinemia. <i>New England Journal of Medicine</i> , 1974, 291, 1-6.	13.9	211
150	INTERACTION OF HUMAN THYMUS-DERIVED AND NON-THYMUS-DERIVED LYMPHOCYTES IN VITRO. <i>Journal of Experimental Medicine</i> , 1973, 138, 1230-1247.	4.2	123