Sung-Yun Pai

List of Publications by Year
in descending order

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124 papers	9,394 citations	46918 47 h-index	39575 94 g-index
133 all docs	133 docs citations	133 times ranked	10472 citing authors

#	Article	IF	Citations
1	Characteristics and outcomes of autoimmune hemolytic anemia after pediatric allogeneic stem cell transplant. Pediatric Blood and Cancer, 2022, 69, e29410.	0.8	1
2	Intestinal atresias and intestinal failure in patients with TTC7A mutations. Journal of Pediatric Surgery Case Reports, 2022, 80, 102247.	0.1	2
3	A Curative DNA Code for Hematopoietic Defects. Hematology/Oncology Clinics of North America, 2022, 36, 647-665.	0.9	6
4	Infections in Infants with SCID: Isolation, Infection Screening, and Prophylaxis in PIDTC Centers. Journal of Clinical Immunology, 2021, 41, 38-50.	2.0	36
5	Hematopoietic Stem Cell Transplantation Is a Curative Therapy for Transferrin Receptor 1 (TFRC) Deficiency. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 753-759.e2.	2.0	4
6	Post-Transcriptional Genetic Silencing ofBCL11Ato Treat Sickle Cell Disease. New England Journal of Medicine, 2021, 384, 205-215.	13.9	250
7	Novel Compound Heterozygous Mutations in ZAP70 Leading to a SCID Phenotype with Normal Downstream In vitro Signaling. Journal of Clinical Immunology, 2021, 41, 470-472.	2.0	2
8	Ten Years of Newborn Screening for Severe Combined Immunodeficiency (SCID) in Massachusetts. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 2060-2067.e2.	2.0	20
9	Very Early Onset Inflammatory Bowel Disease: A Clinical Approach With a Focus on the Role of Genetics and Underlying Immune Deficiencies. Inflammatory Bowel Diseases, 2020, 26, 820-842.	0.9	100
10	Defining a new immune deficiency syndrome: MAN2B2-CDG. Journal of Allergy and Clinical Immunology, 2020, 145, 1008-1011.	1.5	19
11	Vasculitis as a Major Morbidity Factor in Patients With Partial RAG Deficiency. Frontiers in Immunology, 2020, 11, 574738.	2.2	10
12	Excellent outcomes following hematopoietic cell transplantation for Wiskott-Aldrich syndrome: a PIDTC report. Blood, 2020, 135, 2094-2105.	0.6	87
13	Diagnostic assay to assist clinical decisions for unclassified severe combined immune deficiency. Blood Advances, 2020, 4, 2606-2610.	2.5	28
14	Gene therapy for X-linked severe combined immunodeficiency: Historical outcomes and current status. Journal of Allergy and Clinical Immunology, 2020, 146, 258-261.	1.5	19
15	Hematopoietic Cell Transplantation in Patients With Primary Immune Regulatory Disorders (PIRD): A Primary Immune Deficiency Treatment Consortium (PIDTC) Survey. Frontiers in Immunology, 2020, 11, 239.	2.2	57
16	Transplantation Outcomes for Children with Severe Combined Immune Deficiency (SCID) Have Improved over Time: A 36-Year Summary Report By the Primary Immune Deficiency Treatment Consortium (PIDTC). Biology of Blood and Marrow Transplantation, 2020, 26, S18-S19.	2.0	3
17	Third-Party Virus-Specific T-Cell Infusion for Treatment of Refractory Viral Infections: Interim Results from PBMTC SUP1701. Biology of Blood and Marrow Transplantation, 2020, 26, S89-S90.	2.0	1
18	Lentiviral gene therapy for X-linked chronic granulomatous disease. Nature Medicine, 2020, 26, 200-206.	15.2	175

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19	Pulmonary Complications in Patients with Primary Immunodeficiency Undergoing Hematopoietic Stem Cell Transplantation. Biology of Blood and Marrow Transplantation, 2020, 26, S16-S17.	2.0	1
20	Chronic Granulomatous Disease-Associated IBD Resolves and Does Not Adversely Impact Survival Following Allogeneic HCT. Journal of Clinical Immunology, 2019, 39, 653-667.	2.0	41
21	Vasculitis as a major morbidity factor in patients with hypomorphic RAG mutations. Journal of Allergy and Clinical Immunology, 2019, 143, AB116.	1.5	2
22	174.â€ƒVASCULITIS AS A MAJOR MORBIDITY FACTOR IN PATIENTS WITH HYPOMORPHIC RAG VARIANTS. Rheumatology, 2019, 58, .	0.9	0
23	Universal Newborn Screening for Severe Combined Immunodeficiency (SCID). Frontiers in Pediatrics, 2019, 7, 373.	0.9	82
24	Outcomes and Treatment Strategies for Autoimmunity and Hyperinflammation in Patients with RAG Deficiency. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 1970-1985.e4.	2.0	64
25	T-cell mitochondrial dysfunction and lymphopenia in DOCK2-deficient patients. Journal of Allergy and Clinical Immunology, 2019, 144, 306-309.e2.	1.5	13
26	Treatment of primary immunodeficiency with allogeneic transplant and gene therapy. Hematology American Society of Hematology Education Program, 2019, 2019, 457-465.	0.9	22
27	Hematopoietic Stem Cell Transplantation as Treatment for Patients with DOCK8 Deficiency. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 848-855.	2.0	67
28	Immune Reconstitution Therapy for Immunodeficiency. , 2019, , 1115-1128.e1.		0
29	The genetic landscape of severe combined immunodeficiency in the United States and Canada in the current era (2010-2018). Journal of Allergy and Clinical Immunology, 2019, 143, 405-407.	1.5	64
30	Outcome of Hematopoietic Stem Cell Gene Therapy for Wiskott-Aldrich Syndrome. Blood, 2019, 134, 4629-4629.	0.6	17
31	Inducible Phase Separation of GSK3Î± As a Mechanism for Asparaginase Resistance in Acute Leukemias. Blood, 2019, 134, 169-169.	0.6	0
32	Long-Term Treatment Outcome in IPEX Syndrome Patients: An International Multicenter Retrospective Study. Biology of Blood and Marrow Transplantation, 2018, 24, S86-S87.	2.0	0
33	Patients with CD3G mutations reveal a role for human CD3Î³ in Treg diversity and suppressive function. Blood, 2018, 131, 2335-2344.	0.6	51
34	Long-term follow-up of IPEX syndrome patients after different therapeutic strategies: An international multicenter retrospective study. Journal of Allergy and Clinical Immunology, 2018, 141, 1036-1049.e5.	1.5	233
35	WASP-mediated regulation of anti-inflammatory macrophages is IL-10 dependent and is critical for intestinal homeostasis. Nature Communications, 2018, 9, 1779.	5.8	40
36	Resolution of CGD Related Colitis after Allogeneic Hematopoietic Stem Cell Transplantation in Patients with Chronic Granulomatous Diseaseâ€"Early Results From the 6903 Study of the Primary Immune Deficiency Treatment Consortium (PIDTC). Biology of Blood and Marrow Transplantation, 2018, 24, S80-S81.	2.0	2

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37	Getting Past HSC Security: Cyclosporine H Gives Lentiviruses an Entry Pass. Cell Stem Cell, 2018, 23, 775-776.	5.2	1
38	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
39	SCID genotype and 6-month posttransplant CD4 count predict survival and immune recovery. Blood, 2018, 132, 1737-1749.	0.6	128
40	B-cell differentiation and IL-21 response in IL2RG/JAK3 SCID patients after hematopoietic stem cell transplantation. Blood, 2018, 131, 2967-2977.	0.6	37
41	Disrupted N-linked glycosylation as a disease mechanism in deficiency of ADA2. Journal of Allergy and Clinical Immunology, 2018, 142, 1363-1365.e8.	1.5	28
42	Congenital Disorders of Lymphocyte Function. , 2018, , 710-723.e3.		2
43	Reduced-intensity conditioning for hematopoietic cell transplant for HLH and primary immune deficiencies. Blood, 2018, 132, 1438-1451.	0.6	78
44	Genotype, Phenotype and T Cell Counts at One Year Predict Survival and Long Term Immune Reconstitution after Transplantation in Severe Combined Immune Deficiency (SCID)â€"The Primary Immune Deficiency Treatment Consortium (PIDTC). Biology of Blood and Marrow Transplantation, 2017, 23, S133-S134.	2.0	4
45	Reticular dysgenesis: international survey on clinical presentation, transplantation, and outcome. Blood, 2017, 129, 2928-2938.	0.6	31
46	Immune reconstitution and survival of 100 SCID patients postâ€"hematopoietic cell transplant: a PIDTC natural history study. Blood, 2017, 130, 2718-2727.	0.6	212
47	Pharmacokinetics and Model-Based Dosing to Optimize Fludarabine Therapy in Pediatric Hematopoietic Cell Transplant Recipients. Biology of Blood and Marrow Transplantation, 2017, 23, 1701-1713.	2.0	32
48	Natural Killer Cells from Patients with Recombinase-Activating Gene and Non-Homologous End Joining Gene Defects Comprise a Higher Frequency of CD56bright NKG2A+++ Cells, and Yet Display Increased Degranulation and Higher Perforin Content. Frontiers in Immunology, 2017, 8, 798.	2.2	41
49	Characterization of T and B cell repertoire diversity in patients with RAG deficiency. Science Immunology, 2016, 1, .	5.6	88
50	HSCT for DOCK8 Deficiency - an International Study on 74 Patients. Biology of Blood and Marrow Transplantation, 2016, 22, S103-S104.	2.0	2
51	Plasma and Intracellular Pharmacokinetic (PK) Analysis of Fludarabine in Pediatric Allogeneic Hematopoietic Cell Transplant (alloHCT) Recipients. Biology of Blood and Marrow Transplantation, 2016, 22, S253.	2.0	0
52	O-008â€ƒAberrant Anti-inflammatory Macrophage Function and Differentiation in Wiskott-Aldrich Syndrome Protein-Deficient Mice and Humans. Inflammatory Bowel Diseases, 2016, 22, S3.	0.9	1
53	Hematopoietic stem cell transplantation outcomes for 11 patients with dedicator of cytokinesis 8 deficiency. Journal of Allergy and Clinical Immunology, 2016, 138, 852-859.e3.	1.5	48
54	Stem cell transplantation for tetratricopeptide repeat domain 7A deficiency: long-term follow-up. Blood, 2016, 128, 1306-1308.	0.6	40

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55	Poor T Cell Reconstitution at 100 Days after T Cell-Replete Hematopoietic Cell Transplantation (HCT) for SCID Is Associated with Later Risk of Death or Need for 2nd Transplant in the 6901 Prospective Study of the Pidtc. Biology of Blood and Marrow Transplantation, 2016, 22, S101-S102.	2.0	3
56	A novel mutation in the POLE2 geneÂcausing combined immunodeficiency. Journal of Allergy and Clinical Immunology, 2016, 137, 635-638.e1.	1.5	49
57	Autoimmune lymphoproliferative syndrome caused by a homozygous FasL mutation that disrupts FasL assembly. Journal of Allergy and Clinical Immunology, 2016, 137, 324-327.e2.	1.5	13
58	Preclinical Development of Gene Therapy for X-Linked Severe Combined Immunodeficiency (SCID-X1). Blood, 2016, 128, 4705-4705.	0.6	1
59	Fatal autoimmunity in mice reconstituted with human hematopoietic stem cells encoding defective FOXP3. Blood, 2015, 125, 3886-3895.	0.6	33
60	The syndrome of hemophagocytic lymphohistiocytosis in primary immunodeficiencies: implications for differential diagnosis and pathogenesis. Haematologica, 2015, 100, 978-988.	1.7	161
61	Human HOIP and LUBAC deficiency underlies autoinflammation, immunodeficiency, amylopectinosis, and lymphangiectasia. Journal of Experimental Medicine, 2015, 212, 939-951.	4.2	241
62	A Case of Leaky SCID with Variable Presentation in Two Siblings Identified By Newborn Screening. Journal of Allergy and Clinical Immunology, 2015, 135, AB15.	1.5	1
63	DOCK8 Deficiency: Clinical and Immunological Phenotype and Treatment Options - a Review of 136 Patients. Journal of Clinical Immunology, 2015, 35, 189-198.	2.0	284
64	Early Hematopoietic Cell Transplant (HCT) Outcomes of Children with Severe Combined Immunodeficiency Disease (SCID): The First Seventy Four Patients of the Primary Immune Deficiency Treatment Consortium (PIDTC) Prospective Study 6901. Biology of Blood and Marrow Transplantation, 2015, 21, S289-S291.	2.0	2
65	A novel mutation in ORAI1 presenting with combined immunodeficiency and residual T-cell function. Journal of Allergy and Clinical Immunology, 2015, 136, 479-482.e1.	1.5	28
66	Broad spectrum of autoantibodies in patients with Wiskott-Aldrich syndrome and X-linked thrombocytopenia. Journal of Allergy and Clinical Immunology, 2015, 136, 1401-1404.e3.	1.5	25
67	A novel mutation in ICOS presenting as hypogammaglobulinemia with susceptibility to opportunistic pathogens. Journal of Allergy and Clinical Immunology, 2015, 136, 794-797.e1.	1.5	26
68	Gene Therapy Using a Self-Inactivating Lentiviral Vector Improves Clinical and Laboratory Manifestations of Wiskott-Aldrich Syndrome. Blood, 2015, 126, 260-260.	0.6	12
69	Newborn Screening for Severe Combined Immunodeficiency in 11 Screening Programs in the United States. JAMA - Journal of the American Medical Association, 2014, 312, 729.	3.8	586
70	Stem cell transplantation for primary immunodeficiency diseases. Current Opinion in Allergy and Clinical Immunology, 2014, 14, 521-526.	1.1	36
71	Retrospective Study of 240 Patients with Severe Combined Immunodeficiency Transplanted from 2000-2009: A Report from the Primary Immune Deficiency Treatment Consortium of North America. Biology of Blood and Marrow Transplantation, 2014, 20, S24-S25.	2.0	1
72	Flow cytometry diagnosis of dedicator of cytokinesis 8 (DOCK8) deficiency. Journal of Allergy and Clinical Immunology, 2014, 134, 221-223.e7.	1.5	62

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73	Establishing diagnostic criteria for severe combined immunodeficiency disease (SCID), leaky SCID, and Omenn syndrome: The Primary Immune Deficiency Treatment Consortium experience. Journal of Allergy and Clinical Immunology, 2014, 133, 1092-1098.	1.5	301
74	A systematic analysis of recombination activity andÂgenotype-phenotype correlation in human recombination-activating gene 1 deficiency. Journal of Allergy and Clinical Immunology, 2014, 133, 1099-1108.e12.	1.5	132
75	Primary Immune Deficiency Treatment Consortium (PIDTC) report. Journal of Allergy and Clinical Immunology, 2014, 133, 335-347.e11.	1.5	65
76	Transplantation Outcomes for Severe Combined Immunodeficiency, 2000â€“2009. New England Journal of Medicine, 2014, 371, 434-446.	13.9	594
77	Comparison of outcomes of hematopoietic stem cell transplantation without chemotherapy conditioning by using matched sibling and unrelated donors for treatment ofÂsevere combined immunodeficiency. Journal of Allergy and Clinical Immunology, 2014, 134, 935-943.e15.	1.5	82
78	A Modified Î³-Retrovirus Vector for X-Linked Severe Combined Immunodeficiency. New England Journal of Medicine, 2014, 371, 1407-1417.	13.9	358
79	363 Human Hematopoietic Stem Cells With a Defined Immunodeficiency and Enteropathy Transfer Clinical Phenotype to a Novel Humanized Mouse Strain. Gastroenterology, 2014, 146, S-81.	0.6	2
80	Next Generation Sequencing Reveals Skewing of the T and B Cell Receptor Repertoires in Patients with WiskottÃ¢â‚¬â€œAldrich Syndrome. Frontiers in Immunology, 2014, 5, 340.	2.2	40
81	Itm2a, a Target Gene of GATA-3, Plays a Minimal Role in Regulating the Development and Function of T Cells. PLoS ONE, 2014, 9, e96535.	1.1	26
82	The Natural History of Children with Severe Combined Immunodeficiency: Baseline Features of the First Fifty Patients of the Primary Immune Deficiency Treatment Consortium Prospective Study 6901. Journal of Clinical Immunology, 2013, 33, 1156-1164.	2.0	100
83	Effect of Weight and Maturation on Busulfan Clearance in Infants and Small Children Undergoing Hematopoietic Cell Transplantation. Biology of Blood and Marrow Transplantation, 2013, 19, 1608-1614.	2.0	69
84	Whole-exome sequencing identifies tetratricopeptide repeat domain 7A (TTC7A) mutations for combined immunodeficiency with intestinal atresias. Journal of Allergy and Clinical Immunology, 2013, 132, 656-664.e17.	1.5	140
85	GATA-3 Regulates the Homeostasis and Activation of CD8+ T Cells. Journal of Immunology, 2013, 190, 428-437.	0.4	37
86	First reported case of Omenn syndrome in a patient with reticular dysgenesis. Journal of Allergy and Clinical Immunology, 2013, 131, 1227-1230.e3.	1.5	32
87	Expanding the spectrum of recombination-activating gene 1 deficiency: AÂfamily with early-onset autoimmunity. Journal of Allergy and Clinical Immunology, 2013, 132, 969-971.e2.	1.5	59
88	Gene Therapy Through Autologous Transplantation of Gene-Modified Hematopoietic Stem Cells. Biology of Blood and Marrow Transplantation, 2013, 19, S64-S69.	2.0	23
89	Hypomorphic Janus kinase 3 mutations result in a spectrum of immune defects, including partial maternal T-cell engraftment. Journal of Allergy and Clinical Immunology, 2013, 131, 1136-1145.	1.5	27
90	Multicenter study of banked third-party virus-specific T cells to treat severe viral infections after hematopoietic stem cell transplantation. Blood, 2013, 121, 5113-5123.	0.6	507

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91	P-200â€ƒHuman Hematopoietic Stem Cells with a Defined Immunodeficiency Transfer Clinical Phenotype to Novel Humanized Mouse Strain. Inflammatory Bowel Diseases, 2013, 19, S105.	0.9	0
92	Stem cell transplantation and immune reconstitution in immunodeficiency. , 2013, , 1007-1019.		0
93	Multicenter Study of â€œoff-the-Shelfâ€•Third Party Virus-Specific T Cells (VSTs) to Treat Adenovirus (Adv), Cytomegalovirus (CMV) or Epstein Barr Virus (EBV) Infection After Hemopoietic Stem Cell Transplantation (HSCT). Blood, 2012, 120, 457-457.	0.6	2
94	Reduced thymic output, cell cycle abnormalities, and increased apoptosis of T lymphocytes in patients with cartilage-hair hypoplasia. Journal of Allergy and Clinical Immunology, 2011, 128, 139-146.	1.5	36
95	Most Closely HLA-Matched Allogeneic Virus Specific Cytotoxic T-Lymphocytes (CTL) to Treat Persistent Reactivation or Infection With Adenovirus, CMV and EBV After Hemopoietic Stem Cell Transplantation (HSCT). Biology of Blood and Marrow Transplantation, 2011, 17, S151-S152.	2.0	4
96	A Markov model to analyze cost-effectiveness of screening for severe combined immunodeficiency (SCID). Molecular Genetics and Metabolism, 2011, 104, 383-389.	0.5	93
97	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
98	IL-21 is the primary common Î³ chain-binding cytokine required for human B-cell differentiation in vivo. Blood, 2011, 118, 6824-6835.	0.6	132
99	Somatic Gene Therapy for X-Linked Severe Combined Immunodeficiency Using a Self-Inactivating Modified Gammaretroviral Vector Results in An Improved Preclinical Safety Profile and Early Clinical Efficacy in a Human Patient. Blood, 2011, 118, 164-164.	0.6	3
100	Guidelines for implementation of populationâ€based newborn screening for severe combined immunodeficiency. Journal of Inherited Metabolic Disease, 2010, 33, 273-281.	1.7	60
101	Rac GTPases in Human Diseases. Disease Markers, 2010, 29, 177-187.	0.6	71
102	High-Throughput Multiplexed T-Cellâ€“Receptor Excision Circle Quantitative PCR Assay with Internal Controls for Detection of Severe Combined Immunodeficiency in Population-Based Newborn Screening. Clinical Chemistry, 2010, 56, 1466-1474.	1.5	74
103	p53⁺/mdm2^{â^'} Atypical Lipomatous Tumor/Well-Differentiated Liposarcoma in Young Children: An Early Expression of Li-Fraumeni Syndrome. Pediatric and Developmental Pathology, 2010, 13, 218-224.	0.5	28
104	Hematopoietic Cell Transplantation for Wiskott-Aldrich Syndrome: Advances in Biology and Future Directions for Treatment. Immunology and Allergy Clinics of North America, 2010, 30, 179-194.	0.7	50
105	Successful engraftment of donor marrow after allogeneic hematopoietic cell transplantation in autosomal-recessive hyper-IgE syndrome caused by dedicator of cytokinesis 8 deficiency. Journal of Allergy and Clinical Immunology, 2010, 126, 1304-1305.e3.	1.5	61
106	Identification of an infant with severe combined immunodeficiency by newborn screening. Journal of Allergy and Clinical Immunology, 2010, 126, 1073-1074.	1.5	48
107	Small-molecule screen identifies reactive oxygen species as key regulators of neutrophil chemotaxis. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 3546-3551.	3.3	141
108	Rac GTPases in human diseases. Disease Markers, 2010, 29, 177-87.	0.6	33

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109	CDK Inhibitor p18INK4c Is a Downstream Target of GATA3 and Restrains Mammary Luminal Progenitor Cell Proliferation and Tumorigenesis. Cancer Cell, 2009, 15, 389-401.	7.7	82
110	Allogeneic hematopoietic stem cell transplantation for X-linked ectodermal dysplasia and immunodeficiency: case report and review of outcomes. Immunologic Research, 2009, 44, 89-98.	1.3	48
111	GATA3 and the T-cell lineage: essential functions before and after T-helper-2-cell differentiation. Nature Reviews Immunology, 2009, 9, 125-135.	10.6	344
112	Novel presentation of Omenn syndrome in association with aniridia. Journal of Allergy and Clinical Immunology, 2009, 123, 966-969.	1.5	10
113	GATA-3 Links Tumor Differentiation and Dissemination in a Luminal Breast Cancer Model. Cancer Cell, 2008, 13, 141-152.	7.7	314
114	Allogeneic transplantation successfully corrects immune defects, but not susceptibility to colitis, in a patient with nuclear factor-Î°B essential modulator deficiency. Journal of Allergy and Clinical Immunology, 2008, 122, 1113-1118.e1.	1.5	45
115	Distinct Structural Requirements of GATA-3 for the Regulation of Thymocyte and Th2 Cell Differentiation. Journal of Immunology, 2008, 180, 1050-1059.	0.4	11
116	GATA-3 - not just for Th2 cells anymore. Cellular and Molecular Immunology, 2007, 4, 15-29.	4.8	71
117	Lipid defect underlies selective skin barrier impairment of an epidermal-specific deletion of Gata-3. Journal of Cell Biology, 2006, 175, 661-670.	2.3	80
118	GATA-3 Regulates the Development and Function of Invariant NKT Cells. Journal of Immunology, 2006, 177, 6650-6659.	0.4	108
119	GATA-3 deficiency abrogates the development and maintenance of T helper type 2 cells. Proceedings of the National Academy of Sciences of the United States of America, 2004, 101, 1993-1998.	3.3	299
120	Critical Roles for Transcription Factor GATA-3 in Thymocyte Development. Immunity, 2003, 19, 863-875.	6.6	277
121	c-Rel delivers a one-two punch in Th1 cell differentiation. Journal of Clinical Investigation, 2002, 110, 741-742.	3.9	8
122	c-Rel delivers a one-two punch in Th1 cell differentiation. Journal of Clinical Investigation, 2002, 110, 741-742.	3.9	3
123	Cross-linking CD28 leads to activation of 70-kDa S6 kinase. European Journal of Immunology, 1994, 24, 2364-2368.	1.6	34
124	The current status of treatment for anorexia nervosa and bulimia nervosa. International Journal of Eating Disorders, 1992, 12, 215-220.	2.1	34