Kathleen E Sullivan

List of Publications by Year
in descending order

309	22,716	10389	10445
papers	citations	72 h-index	139 g-index

325	325	325	22255
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Molecular diagnosis of childhood immune dysregulation, polyendocrinopathy, and enteropathy, and implications for clinical management. Journal of Allergy and Clinical Immunology, 2022, 149, 327-339.	2.9	22
2	Distinct immune trajectories in patients with chromosome 22q11.2 deletion syndrome and immune-mediated diseases. Journal of Allergy and Clinical Immunology, 2022, 149, 445-450.	2.9	15
3	Therapeutic options for CTLA-4 insufficiency. Journal of Allergy and Clinical Immunology, 2022, 149, 736-746.	2.9	68
4	Îœ Î½Î¿Ï‚: Power, life force, and purpose in mentoring. Journal of Allergy and Clinical Immunology, 2022, 149, 547-549.	2.9	1
5	Chronic Granulomatous Disease With Inflammatory Bowel Disease: Clinical Presentation, Treatment, and Outcomes From the USIDNET Registry. Journal of Allergy and Clinical Immunology: in Practice, 2022, 10, 1325-1333.e5.	3.8	11
6	Morbidity, Mortality, and Therapeutics in Combined Immunodeficiency: Data from the USIDNET Registry. Journal of Allergy and Clinical Immunology: in Practice, 2022, , .	3.8	0
7	Inborn Errors of Immunity Associated With Type 2 Inflammation in the USIDNET Registry. Frontiers in Immunology, 2022, 13, 831279.	4.8	6
8	Association of Persistent Rubella Virus With Idiopathic Skin Granulomas in Clinically Immunocompetent Adults. JAMA Dermatology, 2022, 158, 626.	4.1	14
9	22q11.2 Deletion and Duplication Syndromes and COVID-19. Journal of Clinical Immunology, 2022, 42, 746-748.	3.8	2
10	Drug Sensitivity of Vaccine-Derived Rubella Viruses and Quasispecies Evolution in Granulomatous Lesions of Two Ataxia-Telangiectasia Patients Treated with Nitazoxanide. Pathogens, 2022, 11, 338.	2.8	5
11	X-Linked Agammaglobulinemia: Infection Frequency and Infection-Related Mortality in the USIDNET Registry. Journal of Clinical Immunology, 2022, 42, 827-836.	3.8	11
12	Ocular Manifestations in Primary Immunodeficiency Disorders: A Report from the United States Immunodeficiency Network (USIDNET) Registry. Journal of Allergy and Clinical Immunology: in Practice, 2022, , .	3.8	2
13	The yin and the yang of early classical pathway complement disorders. Clinical and Experimental Immunology, 2022, 209, 151-160.	2.6	3
14	Human Inborn Errors of Immunity: 2022 Update on the Classification from the International Union of Immunological Societies Expert Committee. Journal of Clinical Immunology, 2022, 42, 1473-1507.	3.8	389
15	Human Adenovirus 7-Associated Hemophagocytic Lymphohistiocytosis-like Illness: Clinical and Virological Characteristics in a Cluster of Five Pediatric Cases. Clinical Infectious Diseases, 2021, 73, e1532-e1538.	5.8	12
16	Colonoids From Patients With Pediatric Inflammatory Bowel Disease Exhibit Decreased Growth Associated With Inflammation Severity and Durable Upregulation of Antigen Presentation Genes. Inflammatory Bowel Diseases, 2021, 27, 256-267.	1.9	7
17	The scary world of variants of uncertain significance (VUS): AÂhitchhikerâ€™s guide to interpretation. Journal of Allergy and Clinical Immunology, 2021, 147, 492-494.	2.9	16
18	Coronavirus disease 2019 in patients with inborn errors of immunity: An international study. Journal of Allergy and Clinical Immunology, 2021, 147, 520-531.	2.9	278

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19	Relationship Between Severity of T Cell Lymphopenia and Immune Dysregulation in Patients with DiGeorge Syndrome (22q11.2 Deletions and/or Related TBX1 Mutations): a USIDNET Study. Journal of Clinical Immunology, 2021, 41, 29-37.	3.8	17
20	Infection Phenotypes Among Patients with Primary Antibody Deficiency Mined from a US Patient Registry. Journal of Clinical Immunology, 2021, 41, 374-381.	3.8	4
21	Infectious Complications Predict Premature CD8+ T-cell Senescence in CD40 Ligand-Deficient Patients. Journal of Clinical Immunology, 2021, 41, 795-806.	3.8	2
22	Palmoplantar keratoderma with deafness phenotypic variability in a patient with an inherited GJB2 frameshift variant and novel missense variant. Molecular Genetics & Genomic Medicine, 2021, 9, e1574.	1.2	0
23	The Ever-Increasing Array of Novel Inborn Errors of Immunity: an Interim Update by the IUIS Committee. Journal of Clinical Immunology, 2021, 41, 666-679.	3.8	165
24	Combined use of emapalumab and ruxolitinib in a patient with refractory hemophagocytic lymphohistiocytosis was safe and effective. Pediatric Blood and Cancer, 2021, 68, e29026.	1.5	11
25	Diagnostic Challenges in Pediatric Hemophagocytic Lymphohistiocytosis. Journal of Clinical Immunology, 2021, 41, 1213-1218.	3.8	10
26	Variants in STXBP3 are Associated with Very Early Onset Inflammatory Bowel Disease, Bilateral Sensorineural Hearing Loss and Immune Dysregulation. Journal of Crohn's and Colitis, 2021, 15, 1908-1919.	1.3	7
27	The culture microenvironment of juvenile idiopathic arthritis synovial fibroblasts is favorable for endochondral bone formation through BMP4 and repressed by chondrocytes. Pediatric Rheumatology, 2021, 19, 72.	2.1	4
28	Constrained chromatin accessibility in PU.1-mutated agammaglobulinemia patients. Journal of Experimental Medicine, 2021, 218, .	8.5	31
29	Heterozygous OAS1 gain-of-function variants cause an autoinflammatory immunodeficiency. Science Immunology, 2021, 6, .	11.9	36
30	Granulomatous Dermatitis Associated With Rubella Virus Infection in an Adult With Immunodeficiency. JAMA Dermatology, 2021, 157, 842.	4.1	14
31	Immune Dysregulation in Human ITCH Deficiency Successfully Treated with Hematopoietic Cell Transplantation. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 2885-2893.e3.	3.8	4
32	Self-Limited COVID-19 in a Patient with Artemis Hypomorphic SCID. Journal of Clinical Immunology, 2021, 41, 1745-1747.	3.8	7
33	Chromatin Modifications in 22q11.2 Deletion Syndrome. Journal of Clinical Immunology, 2021, 41, 1853-1864.	3.8	10
34	The relevance of primary immunodeficiency registries on a global perspective. Journal of Allergy and Clinical Immunology, 2021, 148, 1170-1171.	2.9	5
35	The Role of PF4 Antibodies in Pediatric Sars-Cov-2 Infections. Blood, 2021, 138, 1004-1004.	1.4	0
36	Proteomic profiling of MIS-C patients indicates heterogeneity relating to interferon gamma dysregulation and vascular endothelial dysfunction. Nature Communications, 2021, 12, 7222.	12.8	41

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37	Rubella Virus Infected Macrophages and Neutrophils Define Patterns of Granulomatous Inflammation in Inborn and Acquired Errors of Immunity. Frontiers in Immunology, 2021, 12, 796065.	4.8	19
38	The Unique Disease Course of Children with Very Early onset-Inflammatory Bowel Disease. Inflammatory Bowel Diseases, 2020, 26, 909-918.	1.9	32
39	Prolactin activates IRF1 and leads to altered balance of histone acetylation: Implications for systemic lupus erythematosus. Modern Rheumatology, 2020, 30, 532-543.	1.8	6
40	Diagnostic interpretation of genetic studies in patients with primary immunodeficiency diseases: AÂworking group report of the Primary Immunodeficiency Diseases Committee of the American Academy of Allergy, Asthma & Immunology. Journal of Allergy and Clinical Immunology, 2020, 145, 46-69.	2.9	54
41	Prophylactic Antibiotics Versus Immunoglobulin Replacement in Specific Antibody Deficiency. Journal of Clinical Immunology, 2020, 40, 158-164.	3.8	11
42	North American Society for Pediatric Gastroenterology, Hepatology, and Nutrition Position Paper on the Evaluation and Management for Patients With Very Earlyâ€onset Inflammatory Bowel Disease. Journal of Pediatric Gastroenterology and Nutrition, 2020, 70, 389-403.	1.8	79
43	Juvenile idiopathic arthritis fibroblast-like synoviocytes influence chondrocytes to alter BMP antagonist expression demonstrating an interaction between the two prominent cell types involved in endochondral bone formation. Pediatric Rheumatology, 2020, 18, 89.	2.1	3
44	Global systematic review of primary immunodeficiency registries. Expert Review of Clinical Immunology, 2020, 16, 717-732.	3.0	74
45	Rubella virus-associated chronic inflammation in primary immunodeficiency diseases. Current Opinion in Allergy and Clinical Immunology, 2020, 20, 574-581.	2.3	24
46	Adenosine Deaminase (ADA)â€“Deficient Severe Combined Immune Deficiency (SCID) in the US Immunodeficiency Network (USIDNet) Registry. Journal of Clinical Immunology, 2020, 40, 1124-1131.	3.8	19
47	Convalescent plasma for pediatric patients with SARSâ€CoVâ€2â€associated acute respiratory distress syndrome. Pediatric Blood and Cancer, 2020, 67, e28693.	1.5	37
48	PROMIS-29 survey confirms major impact of fatigue on health-related quality of life in common variable immunodeficiency. Immunologic Research, 2020, 68, 379-388.	2.9	9
49	Excellent outcomes following hematopoietic cell transplantation for Wiskott-Aldrich syndrome: a PIDTC report. Blood, 2020, 135, 2094-2105.	1.4	87
50	Acute Kidney Injury in Children after Hematopoietic Cell Transplantation Is Associated with Elevated Urine CXCL10 and CXCL9. Biology of Blood and Marrow Transplantation, 2020, 26, 1266-1272.	2.0	11
51	Lymphoproliferative Disease in CVID: a Report of Types and Frequencies from a US Patient Registry. Journal of Clinical Immunology, 2020, 40, 524-530.	3.8	34
52	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.	4.8	57
53	The Association of Fetal Thymus Size with Subsequent T Cell Counts in 22q11.2 Deletion Syndrome. Journal of Clinical Immunology, 2020, 40, 783-785.	3.8	4
54	Epigenetic syndromes with immune deficiency. , 2020, , 319-337.		0

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55	European Society for Immunodeficiencies (ESID) and European Reference Network on Rare Primary Immunodeficiency, Autoinflammatory and Autoimmune Diseases (ERN RITA) Complement Guideline: Deficiencies, Diagnosis, and Management. Journal of Clinical Immunology, 2020, 40, 576-591.	3.8	43
56	IL-1 Transcriptional Responses to Lipopolysaccharides Are Regulated by a Complex of RNA Binding Proteins. Journal of Immunology, 2020, 204, 1334-1344.	0.8	12
57	Human Inborn Errors of Immunity: 2019 Update on the Classification from the International Union of Immunological Societies Expert Committee. Journal of Clinical Immunology, 2020, 40, 24-64.	3.8	881
58	Increased Tâ€cell counts in patients with 22q11.2 deletion syndrome who have anxiety. American Journal of Medical Genetics, Part A, 2020, 182, 1815-1818.	1.2	2
59	Human Inborn Errors of Immunity: 2019 Update of the IUIS Phenotypical Classification. Journal of Clinical Immunology, 2020, 40, 66-81.	3.8	525
60	Persons Living With Primary Immunodeficiency Act as Citizen Scientists and Launch Prospective Cohort Body Temperature Study. Journal of Participatory Medicine, 2020, 12, e22297.	1.3	2
61	Understanding of Inheritance and Genetic Variation Assessed through the Use of an Engaging Real-Life Survey. Public Health Genomics, 2020, 23, 246-251.	1.0	0
62	Evidence of Microangiopathy in Children with Sars-Cov-2 Regardless of Clinical Presentation. Blood, 2020, 136, 28-29.	1.4	0
63	Abstract 12622: Pro-inflammatory Urinary CXCL10 and VCAM Are Elevated After Infant Cardiac Surgery-associated Acute Kidney Injury (AKI). Circulation, 2020, 142, .	1.6	0
64	American Academy of Allergy, Asthma and Immunology response to the <scp>EAACI</scp>/<scp>GA</scp>²<scp>LEN</scp>/<scp>EDF</scp>/<scp>WAO</scp> guideline for the definition, classification, diagnosis, and management of Urticaria 2017 revision. Allergy: European Journal of Allergy and Clinical Immunology, 2019, 74, 411-413.	5.7	17
65	Patients with common variable immunodeficiency with autoimmune cytopenias exhibit hyperplastic yet inefficient germinal center responses. Journal of Allergy and Clinical Immunology, 2019, 143, 258-265.	2.9	68
66	Chronic Granulomatous Disease-Associated IBD Resolves and Does Not Adversely Impact Survival Following Allogeneic HCT. Journal of Clinical Immunology, 2019, 39, 653-667.	3.8	41
67	Tollâ€like receptor 2 stimulation augments esophageal barrier integrity. Allergy: European Journal of Allergy and Clinical Immunology, 2019, 74, 2449-2460.	5.7	26
68	Infectious vaccine-derived rubella viruses emerge, persist, and evolve in cutaneous granulomas of children with primary immunodeficiencies. PLoS Pathogens, 2019, 15, e1008080.	4.7	58
69	Immune Biomarkers of Autoimmunity in Chromosome 22q11.2 Deletion Syndrome. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 2377-2378.	3.8	2
70	Common variable immunodeficiencyâ€“associated endotoxemia promotes early commitment to the T follicular lineage. Journal of Allergy and Clinical Immunology, 2019, 144, 1660-1673.	2.9	22
71	Outcomes for Nitazoxanide Treatment in a Case Series of Patients with Primary Immunodeficiencies and Rubella Virus-Associated Granuloma. Journal of Clinical Immunology, 2019, 39, 112-117.	3.8	19
72	Hematopoietic stem cell transplantation for CD40 ligand deficiency: Results from an EBMT/ESID-IEWP-SCETIDE-PIDTC study. Journal of Allergy and Clinical Immunology, 2019, 143, 2238-2253.	2.9	60

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73	A novel human IL2RB mutation results in T and NK cellâ€“driven immune dysregulation. Journal of Experimental Medicine, 2019, 216, 1255-1267.	8.5	64
74	X-linked agammaglobulinemia (XLA): Phenotype, diagnosis, and therapeutic challenges around the world. World Allergy Organization Journal, 2019, 12, 100018.	3.5	83
75	Allogeneic hematopoietic stem cell transplantation in adolescent patients with chronic granulomatous disease. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 1052-1054.e2.	3.8	8
76	Chromosome 22q11.2 deletion syndrome and DiGeorge syndrome. Immunological Reviews, 2019, 287, 186-201.	6.0	81
77	Expansion of the Human Phenotype Ontology (HPO) knowledge base and resources. Nucleic Acids Research, 2019, 47, D1018-D1027.	14.5	539
78	Early-Onset Inflammatory Bowel Disease. Immunology and Allergy Clinics of North America, 2019, 39, 63-79.	1.9	64
79	Rubella Virus-Associated Cutaneous Granulomatous Disease: a Unique Complication in Immune-Deficient Patients, Not Limited to DNA Repair Disorders. Journal of Clinical Immunology, 2019, 39, 81-89.	3.8	56
80	Neutropenia as a sign of immunodeficiency. Journal of Allergy and Clinical Immunology, 2019, 143, 96-100.	2.9	11
81	Hypomorphic caspase activation and recruitment domain 11 (CARD11) mutations associated with diverse immunologic phenotypes with or without atopic disease. Journal of Allergy and Clinical Immunology, 2019, 143, 1482-1495.	2.9	116
82	Rapid induction of expression by LPS is accompanied by favorable chromatin and rapid binding of c-Jun. Molecular Immunology, 2018, 95, 99-106.	2.2	3
83	Phenotypic characterization of patients with rheumatologic manifestations of common variable immunodeficiency. Seminars in Arthritis and Rheumatism, 2018, 48, 318-326.	3.4	26
84	<scp>EMSY</scp> is increased and activates <scp>TSLP</scp> & <scp>CCL</scp>5 expression in eosinophilic esophagitis. Pediatric Allergy and Immunology, 2018, 29, 565-568.	2.6	7
85	Low Serum IgE Is a Sensitive and Specific Marker for Common Variable Immunodeficiency (CVID). Journal of Clinical Immunology, 2018, 38, 225-233.	3.8	48
86	Primary lymphedema and other lymphatic anomalies are associated with 22q11.2 deletion syndrome. European Journal of Medical Genetics, 2018, 61, 411-415.	1.3	5
87	Variable immune deficiency related to deletion size in chromosome 22q11.2 deletion syndrome. American Journal of Medical Genetics, Part A, 2018, 176, 2082-2086.	1.2	53
88	Genomic Circuitry Underlying Immunological Response to Pediatric Acute Respiratory Infection. Cell Reports, 2018, 22, 411-426.	6.4	15
89	Mevalonate kinase deficiency presenting as recurrent rectal abscesses and perianal fistulae. Annals of Allergy, Asthma and Immunology, 2018, 120, 214-215.	1.0	6
90	Cancer in primary immunodeficiency diseases: Cancer incidence in the United States Immune Deficiency Network Registry. Journal of Allergy and Clinical Immunology, 2018, 141, 1028-1035.	2.9	172

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91	Autoimmune Cytopenias and Associated Conditions in CVID: a Report From the USIDNET Registry. Journal of Clinical Immunology, 2018, 38, 28-34.	3.8	79
92	T-cell lymphopenia in 22q11.2 deletion syndrome: Relationship to cardiac disease. Journal of Allergy and Clinical Immunology: in Practice, 2018, 6, 690-691.	3.8	7
93	Autosomal Dominant Hyper-IgE Syndrome in the USIDNET Registry. Journal of Allergy and Clinical Immunology: in Practice, 2018, 6, 996-1001.	3.8	62
94	Nitazoxanide May Modify the Course of Progressive Multifocal Leukoencephalopathy. Journal of Clinical Immunology, 2018, 38, 4-6.	3.8	5
95	International Union of Immunological Societies: 2017 Primary Immunodeficiency Diseases Committee Report on Inborn Errors of Immunity. Journal of Clinical Immunology, 2018, 38, 96-128.	3.8	732
96	The 2017 IUIS Phenotypic Classification for Primary Immunodeficiencies. Journal of Clinical Immunology, 2018, 38, 129-143.	3.8	488
97	Very early-onset inflammatory bowel disease: an integrated approach. Current Opinion in Allergy and Clinical Immunology, 2018, 18, 459-469.	2.3	12
98	Cover Image, Volume 176A, Number 10, October 2018. , 2018, 176, i-i.		0
99	Enhancer RNA and NFÎºB-dependent P300 regulation of ADAMDEC1. Molecular Immunology, 2018, 103, 312-321.	2.2	16
100	What is new with 22q? An update from the 22q and You Center at the Children's Hospital of Philadelphia. American Journal of Medical Genetics, Part A, 2018, 176, 2058-2069.	1.2	106
101	SCID genotype and 6-month posttransplant CD4 count predict survival and immune recovery. Blood, 2018, 132, 1737-1749.	1.4	128
102	Phenotype, penetrance, and treatment of 133 cytotoxic T-lymphocyte antigen 4â€“insufficient subjects. Journal of Allergy and Clinical Immunology, 2018, 142, 1932-1946.	2.9	344
103	Epigenetics of Systemic Lupus Erythematosus. , 2018, , 133-148.		0
104	Prevalence of Granulomas in Patients With Primary Immunodeficiency Disorders, United States: Data From National Health Care Claims and the US Immunodeficiency Network Registry. Journal of Clinical Immunology, 2018, 38, 717-726.	3.8	23
105	Overall Downregulation of mRNAs and Enrichment of H3K4me3 Change Near Genome-Wide Association Study Signals in Systemic Lupus Erythematosus: Cell-Specific Effects. Frontiers in Immunology, 2018, 9, 497.	4.8	24
106	Low Rates of Poliovirus Antibodies in Primary Immunodeficiency Patients on Regular Intravenous Immunoglobulin Treatment. Journal of Clinical Immunology, 2018, 38, 628-634.	3.8	4
107	Renal Disease in Chronic Granulomatous Disease: Data from the USIDNET Registry. Journal of Clinical Immunology, 2018, 38, 556-557.	3.8	1
108	Pathogenesis of Pediatric Rheumatologic Diseases. Pediatric Clinics of North America, 2018, 65, 639-655.	1.8	6

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109	Prior to extension, Transcriptomes of fibroblast-like Synoviocytes from extended and Polyarticular juvenile idiopathic arthritis are indistinguishable. Pediatric Rheumatology, 2018, 16, 3.	2.1	2
110	Complications Associated with Underweight Primary Immunodeficiency Patients: Prevalence and Associations Within the USIDNET Registry. Journal of Clinical Immunology, 2018, 38, 283-293.	3.8	19
111	Cytotoxic T-Lymphocyte-Associated Protein 4 Haploinsufficiency-Associated Inflammation Can Occur Independently of T-Cell Hyperproliferation. Frontiers in Immunology, 2018, 9, 1715.	4.8	13
112	The immune deficiency of chromosome 22q11.2 deletion syndrome. American Journal of Medical Genetics, Part A, 2017, 173, 2366-2372.	1.2	64
113	Frequency of untreated hypogammaglobulinemia in bronchiectasis. Annals of Allergy, Asthma and Immunology, 2017, 119, 83-85.	1.0	4
114	Immunodeficiencies Associated with Abnormal Newborn Screening for T Cell and B Cell Lymphopenia. Journal of Clinical Immunology, 2017, 37, 363-374.	3.8	33
115	C3 nephritic factors: AÂchanging landscape. Journal of Allergy and Clinical Immunology, 2017, 140, 57-59.	2.9	10
116	Secretion of proâ€inflammatory cytokines and chemokines and loss of regulatory signals by fibroblastâ€like synoviocytes in juvenile idiopathic arthritis. Proteomics - Clinical Applications, 2017, 11, 1600088.	1.6	14
117	Monogenic lupus: it's all new!. Current Opinion in Immunology, 2017, 49, 87-95.	5.5	51
118	Inhibition of rubella virus replication by the broad-spectrum drug nitazoxanide in cell culture and in a patient with a primary immune deficiency. Antiviral Research, 2017, 147, 58-66.	4.1	36
119	Immune reconstitution and survival of 100 SCID patients postâ€“hematopoietic cell transplant: a PIDTC natural history study. Blood, 2017, 130, 2718-2727.	1.4	212
120	Two Sides of the Same Coin: Pediatric-Onset and Adult-Onset Common Variable Immune Deficiency. Journal of Clinical Immunology, 2017, 37, 592-602.	3.8	34
121	Inflammatory Bowel Disease in Primary Immunodeficiencies. Current Allergy and Asthma Reports, 2017, 17, 57.	5.3	46
122	Emerging Infections and Pertinent Infections Related to Travel for Patients with Primary Immunodeficiencies. Journal of Clinical Immunology, 2017, 37, 650-692.	3.8	6
123	Pediatric healthcare costs for patients with 22q11.2 deletion syndrome. Molecular Genetics & Genomic Medicine, 2017, 5, 631-638.	1.2	12
124	Identification of 22q11.2 Deletion Syndrome via Newborn Screening for Severe Combined Immunodeficiency. Journal of Clinical Immunology, 2017, 37, 476-485.	3.8	35
125	Lack of Clinical Hypersensitivity to Penicillin Antibiotics in Common Variable Immunodeficiency. Journal of Clinical Immunology, 2017, 37, 22-24.	3.8	11
126	Recurrent and Sustained Viral Infections in Primary Immunodeficiencies. Frontiers in Immunology, 2017, 8, 665.	4.8	37

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127	Common Variable Immunodeficiency Non-Infectious Disease Endotypes Redefined Using Unbiased Network Clustering in Large Electronic Datasets. Frontiers in Immunology, 2017, 8, 1740.	4.8	70
128	SERPINB2 is regulated by dynamic interactions with pause-release proteins and enhancer RNAs. Molecular Immunology, 2017, 88, 20-31.	2.2	37
129	Healthcare utilization in chromosome 22q11.2 deletion patients with cardiac disease and low T cell counts. American Journal of Medical Genetics, Part A, 2016, 170, 1630-1634.	1.2	13
130	Hyper IgM Syndrome: a Report from the USIDNET Registry. Journal of Clinical Immunology, 2016, 36, 490-501.	3.8	92
131	Rubella persistence in epidermal keratinocytes and granuloma M2 macrophages in patients with primary immunodeficiencies. Journal of Allergy and Clinical Immunology, 2016, 138, 1436-1439.e11.	2.9	73
132	Successful unrelated cord blood transplant for complete IFN-Î³ receptor 2 deficiency. Journal of Allergy and Clinical Immunology, 2016, 138, 1489-1491.	2.9	8
133	New insights into the immunopathogenesis of systemic lupus erythematosus. Nature Reviews Rheumatology, 2016, 12, 716-730.	8.0	909
134	Severe, persistent, and fatal Tâ€cell immunodeficiency following therapy for infantile leukemia. Pediatric Blood and Cancer, 2016, 63, 2046-2049.	1.5	12
135	A healthy female with C3 hypocomplementemia and C3 Nephritic Factor. Clinical Immunology, 2016, 169, 14-15.	3.2	9
136	H3K4 tri-methylation breadth at transcription start sites impacts the transcriptome of systemic lupus erythematosus. Clinical Epigenetics, 2016, 8, 14.	4.1	47
137	P-197â€ƒIdentification of a Homozygous Mutation in the ZBTB24 Gene in a Patient with Very Early Onset Inflammatory Bowel Disease. Inflammatory Bowel Diseases, 2016, 22, S68.	1.9	0
138	Risk Factors and Clinical Significance of Lymphopenia in Survivors of the Fontan Procedure for Single-Ventricle Congenital Cardiac Disease. Journal of Allergy and Clinical Immunology: in Practice, 2016, 4, 491-496.	3.8	33
139	Enteroviruses in X-Linked Agammaglobulinemia: Update on Epidemiology and Therapyâ€”. Journal of Allergy and Clinical Immunology: in Practice, 2016, 4, 1059-1065.	3.8	67
140	B cell development in chromosome 22q11.2 deletion syndrome. Clinical Immunology, 2016, 163, 1-9.	3.2	24
141	22q11.2 deletion syndrome. Nature Reviews Disease Primers, 2015, 1, 15071.	30.5	954
142	Antibodies to BK virus in children prior to allogeneic hematopoietic cell transplant. Pediatric Blood and Cancer, 2015, 62, 1670-1673.	1.5	9
143	The Role of MicroRNAs and Human Epidermal Growth Factor Receptor 2 in Proliferative Lupus Nephritis. Arthritis and Rheumatology, 2015, 67, 2415-2426.	5.6	46
144	Rare variants at 16p11.2 are associated with common variable immunodeficiency. Journal of Allergy and Clinical Immunology, 2015, 135, 1569-1577.	2.9	22

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145	A de novo whole gene deletion of XIAP detected by exome sequencing analysis in very early onset inflammatory bowel disease: a case report. BMC Gastroenterology, 2015, 15, 160.	2.0	38
146	Food allergy in patients with primary immunodeficiency diseases: Prevalence within the US Immunodeficiency Network (USIDNET). Journal of Allergy and Clinical Immunology, 2015, 135, 273-275.	2.9	45
147	Interferon regulatory factor 1 and histone H4 acetylation in systemic lupus erythematosus. Epigenetics, 2015, 10, 191-199.	2.7	44
148	Multicenter experience in hematopoietic stem cell transplantation for serious complications of common variable immunodeficiency. Journal of Allergy and Clinical Immunology, 2015, 135, 988-997.e6.	2.9	123
149	Exome Sequencing Analysis Reveals Variants in Primary Immunodeficiency Genes in Patients With Very Early Onset Inflammatory Bowel Disease. Gastroenterology, 2015, 149, 1415-1424.	1.3	99
150	Association of CLEC16A with human common variable immunodeficiency disorder and role in murine B cells. Nature Communications, 2015, 6, 6804.	12.8	63
151	Endotoxin tolerance in monocytes can be mitigated by Â2-interferon. Journal of Leukocyte Biology, 2015, 98, 651-659.	3.3	12
152	Primary Immunodeficiency Diseases: an Update on the Classification from the International Union of Immunological Societies Expert Committee for Primary Immunodeficiency 2015. Journal of Clinical Immunology, 2015, 35, 696-726.	3.8	621
153	Monocyte enhancers are highly altered in systemic lupus erythematosus. Epigenomics, 2015, 7, 921-935.	2.1	27
154	The 2015 IUIS Phenotypic Classification for Primary Immunodeficiencies. Journal of Clinical Immunology, 2015, 35, 727-738.	3.8	199
155	Genetic sharing and heritability of paediatric age of onset autoimmune diseases. Nature Communications, 2015, 6, 8442.	12.8	58
156	Combined immunodeficiency in the United States and Kuwait: Comparison of patients' characteristics and molecular diagnosis. Clinical Immunology, 2015, 161, 170-173.	3.2	22
157	Meta-analysis of shared genetic architecture across ten pediatric autoimmune diseases. Nature Medicine, 2015, 21, 1018-1027.	30.7	212
158	Health care utilization by patients with common variable immune deficiency defined by International Classification of Diseases, Ninth Revision code 279.06. Annals of Allergy, Asthma and Immunology, 2015, 115, 248-250.	1.0	4
159	Interferon Regulatory Factor 1 Marks Activated Genes and Can Induce Target Gene Expression in Systemic Lupus Erythematosus. Arthritis and Rheumatology, 2015, 67, 785-796.	5.6	31
160	Noncoding RNAs and LRRFIP1 Regulate TNF Expression. Journal of Immunology, 2014, 192, 3057-3067.	0.8	23
161	DiGeorge Syndrome: A Serendipitous Discovery. , 2014, , 229-240.		0
162	The intersection of immune deficiency and autoimmunity. Current Opinion in Rheumatology, 2014, 26, 570-578.	4.3	17

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163	Terminal Deletion of 11q with Significant Late-Onset Combined Immune Deficiency. Journal of Clinical Immunology, 2014, 34, 114-118.	3.8	24
164	ICON: The Early Diagnosis of Congenital Immunodeficiencies. Journal of Clinical Immunology, 2014, 34, 398-424.	3.8	34
165	Primary Immune Deficiency Treatment Consortium (PIDTC) report. Journal of Allergy and Clinical Immunology, 2014, 133, 335-347.e11.	2.9	65
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