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	22q11.2 deletion syndrome. Nature Reviews Disease Primers, 2015, 1, 15071.	30.5	954
2	New insights into the immunopathogenesis of systemic lupus erythematosus. Nature Reviews Rheumatology, 2016, 12, 716-730.	8.0	909
3	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
4	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
5	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
6	Transplantation Outcomes for Severe Combined Immunodeficiency, 2000â€“2009. New England Journal of Medicine, 2014, 371, 434-446.	27.0	594
7	A multiinstitutional survey of the Wiskott-Aldrich syndrome. Journal of Pediatrics, 1994, 125, 876-885.	1.8	586
8	Expansion of the Human Phenotype Ontology (HPO) knowledge base and resources. Nucleic Acids Research, 2019, 47, D1018-D1027.	14.5	539
9	Human Inborn Errors of Immunity: 2019 Update of the IUIS Phenotypical Classification. Journal of Clinical Immunology, 2020, 40, 66-81.	3.8	525
10	X-Linked Agammaglobulinemia. Medicine (United States), 2006, 85, 193-202.	1.0	516
11	Velocardiofacial syndrome, DiGeorge syndrome: the chromosome 22q11.2 deletion syndromes. Lancet, The, 2007, 370, 1443-1452.	13.7	513
12	The 2017 IUIS Phenotypic Classification for Primary Immunodeficiencies. Journal of Clinical Immunology, 2018, 38, 129-143.	3.8	488
13	FasGene Mutations in the Canaleâ€“Smith Syndrome, an Inherited Lymphoproliferative Disorder Associated with Autoimmunity. New England Journal of Medicine, 1996, 335, 1643-1649.	27.0	484
14	Practical Guidelines for Managing Patients with 22q11.2 Deletion Syndrome. Journal of Pediatrics, 2011, 159, 332-339.e1.	1.8	481
15	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
16	Chromosome 22q11.2 Deletion Syndrome (DiGeorge Syndrome/Velocardiofacial Syndrome). Medicine (United States), 2011, 90, 1-18.	1.0	381
17	ATM stabilizes DNA double-strand-break complexes during V(D)J recombination. Nature, 2006, 442, 466-470.	27.8	366
18	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

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19	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
20	Immunologic features of chromosome 22q11.2 deletion syndrome (DiGeorge) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 702 Td (syndrome/	1.8	240
21	Meta-analysis of shared genetic architecture across ten pediatric autoimmune diseases. Nature Medicine, 2015, 21, 1018-1027.	30.7	212
22	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
23	The 2015 IUIS Phenotypic Classification for Primary Immunodeficiencies. Journal of Clinical Immunology, 2015, 35, 727-738.	3.8	199
24	Report of 8 Cases and Review of the Literature. Medicine (United States), 1996, 75, 251-261.	1.0	195
25	Epigenetic Regulation of Tumor Necrosis Factor Alpha. Molecular and Cellular Biology, 2007, 27, 5147-5160.	2.3	183
26	Genome-wide association identifies diverse causes of common variable immunodeficiency. Journal of Allergy and Clinical Immunology, 2011, 127, 1360-1367.e6.	2.9	179
27	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
28	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
29	Autosomal dominant â€œOpitzâ€•GBBB syndrome due to a 22q11. 2 deletion. American Journal of Medical Genetics Part A, 1995, 59, 103-113.	2.4	152
30	Mannose-binding protein genetic polymorphisms in black patients with systemic lupus erythematosus. Arthritis and Rheumatism, 1996, 39, 2046-2051.	6.7	152
31	Unmasking Evans syndrome: T-cell phenotype and apoptotic response reveal autoimmune lymphoproliferative syndrome (ALPS). Blood, 2005, 105, 2443-2448.	1.4	151
32	Treatment with sirolimus results in complete responses in patients with autoimmune lymphoproliferative syndrome. British Journal of Haematology, 2009, 145, 101-106.	2.5	151
33	The 22q11.2 Deletion: Screening, Diagnostic Workup, and Outcome of Results; Report on 181 Patients. Genetic Testing and Molecular Biomarkers, 1997, 1, 99-108.	1.7	150
34	Infections in Patients with Inherited Defects in Phagocytic Function. Clinical Microbiology Reviews, 2003, 16, 597-621.	13.6	146
35	A promoter polymorphism of tumor necrosis factor Î± associated with systemic lupus erythematosus in African-Americans. Arthritis and Rheumatism, 1997, 40, 2207-2211.	6.7	145
36	TFc?RIIIA-158F allele is a risk factor for systemic lupus erythematosus. Arthritis and Rheumatism, 1998, 41, 1813-1818.	6.7	133

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37	Recommendations for live viral and bacterial vaccines inÂimmunodeficient patients and their close contacts. Journal of Allergy and Clinical Immunology, 2014, 133, 961-966.	2.9	128
38	SCID genotype and 6-month posttransplant CD4 count predict survival and immune recovery. Blood, 2018, 132, 1737-1749.	1.4	128
39	Poor immunogenicity of the H1N1 2009 vaccine in well controlled HIV-infected individuals. Aids, 2010, 24, 2187-2192.	2.2	123
40	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
41	Chromosome 22q11.2 deletion syndrome (DiGeorge and velocardiofacial syndromes). Current Opinion in Pediatrics, 2002, 14, 678-683.	2.0	122
42	Defective Natural Killer Cell Function in Patients with Hemophagocytic Lymphohistiocytosis and in First Degree Relatives. Pediatric Research, 1998, 44, 465-468.	2.3	120
43	T-cell homeostasis in humans with thymic hypoplasia due to chromosome 22q11.2 deletion syndrome. Blood, 2004, 103, 1020-1025.	1.4	119
44	Association of a Promoter Polymorphism of Tumor Necrosis Factor-Î± with Subacute Cutaneous Lupus Erythematosus and Distinct Photoregulation of Transcription1,21Werth VP, Sullivan K: Strong association of a promoter polymorphism of tumor necrosis factors Î± with a photosensitive form of cutaneous lupus erythematosus. Arthr Rheum 42:S105 1999 (abstr.)2Werth VP, Zhang W, Sullivan K: Role of a promoter polymorphism of tumor necrosis factor-Î± (TNF-Î±) in a photosensitive form of lupus erythematosus (LE): clinical a. Journal of Investigative Dermatology, 2000, 115, 726-730.	0.7	117
45	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
46	Juvenile rheumatoid arthritisâ€like polyarthritis in chromosome 22q11.2 deletion syndrome (digeorge) Tj ETQq0 0 0 rgBT /Overlock 10 T Rheumatism, 1997, 40, 430-436.	6.7	115
47	Defective actin accumulation impairs human natural killer cell function in patients with dedicator of cytokinesis 8 deficiency. Journal of Allergy and Clinical Immunology, 2013, 131, 840-848.	2.9	113
48	Rituximab-Treated Patients Have a Poor Response to Influenza Vaccination. Journal of Clinical Immunology, 2013, 33, 388-396.	3.8	111
49	Lack of Correlation between Impaired T Cell Production, Immunodeficiency, and Other Phenotypic Features in Chromosome 22q11.2 Deletion Syndromes (DiGeorge Syndrome/Velocardiofacial) Tj ETQq1 1 0.784314 rgBT /Overlock 10 T	4.0	110
50	The SLE Transcriptome Exhibits Evidence of Chronic Endotoxin Exposure and Has Widespread Dysregulation of Non-Coding and Coding RNAs. PLoS ONE, 2014, 9, e93846.	2.5	109
51	Immune dysregulation in severe influenza. Journal of Leukocyte Biology, 2009, 85, 1036-1043.	3.3	106
52	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
53	Hemizygous mutations in SNAP29 unmask autosomal recessive conditions and contribute to atypical findings in patients with 22q11.2DS. Journal of Medical Genetics, 2013, 50, 80-90.	3.2	104
54	CHARGE (Coloboma, Heart Defect, Atresia Choanae, Retarded Growth and Development, Genital) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 Comparison of Immunologic and Nonimmunologic Phenotypic Features. Pediatrics, 2009, 123, e871-e877.	2.1	103

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55	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
56	Measurement of Cytokine Secretion, Intracellular Protein Expression, and mRNA in Resting and Stimulated Peripheral Blood Mononuclear Cells. Vaccine Journal, 2000, 7, 920-924.	2.6	98
57	Use of Sirolimus in IPEX and IPEX-Like Children. Journal of Clinical Immunology, 2008, 28, 581-587.	3.8	98
58	Hyper IgM Syndrome: a Report from the USIDNET Registry. Journal of Clinical Immunology, 2016, 36, 490-501.	3.8	92
59	Chromosome 22q11.2 Deletion Syndrome: DiGeorge Syndrome/Velocardiofacial Syndrome. Immunology and Allergy Clinics of North America, 2008, 28, 353-366.	1.9	89
60	Autoimmunity and Inflammation in X-linked Agammaglobulinemia. Journal of Clinical Immunology, 2014, 34, 627-632.	3.8	88
61	Excellent outcomes following hematopoietic cell transplantation for Wiskott-Aldrich syndrome: a PIDTC report. Blood, 2020, 135, 2094-2105.	1.4	87
62	Safety of Live Viral Vaccines in Patients With Chromosome 22q11.2 Deletion Syndrome (DiGeorge) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 5	2.1	86
63	X-linked agammaglobulinemia (XLA): Phenotype, diagnosis, and therapeutic challenges around the world. World Allergy Organization Journal, 2019, 12, 100018.	3.5	83
64	Rapamycin improves lymphoproliferative disease in murine autoimmune lymphoproliferative syndrome (ALPS). Blood, 2006, 108, 1965-1971.	1.4	82
65	The clinical, immunological, and molecular spectrum of chromosome 22q11.2 deletion syndrome and DiGeorge syndrome. Current Opinion in Allergy and Clinical Immunology, 2004, 4, 505-512.	2.3	81
66	Chromosome 22q11.2 deletion syndrome and DiGeorge syndrome. Immunological Reviews, 2019, 287, 186-201.	6.0	81
67	Autoimmune Cytopenias and Associated Conditions in CVID: a Report From the USIDNET Registry. Journal of Clinical Immunology, 2018, 38, 28-34.	3.8	79
68	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
69	Immune abnormalities are a frequent manifestation of Kabuki syndrome. American Journal of Medical Genetics, Part A, 2005, 135A, 278-281.	1.2	78
70	Associations of Tumor Necrosis Factor Î± and HLA Polymorphisms with Adult Dermatomyositis: Implications for a Unique Pathogenesi1. Journal of Investigative Dermatology, 2002, 119, 617-620.	0.7	77
71	Secondary immunologic consequences in chromosome 22q11.2 deletion syndrome (DiGeorge) Tj ETQq1 1 0.784314 rgBT /Overlock	3.2	75
72	Thrombocytopenia in patients with chromosome 22q11.2 deletion syndrome. Journal of Pediatrics, 2003, 143, 277-278.	1.8	74

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73	Global systematic review of primary immunodeficiency registries. Expert Review of Clinical Immunology, 2020, 16, 717-732.	3.0	74
74	Increased Prevalence of Immunoglobulin A Deficiency in Patients with the Chromosome 22q11.2 Deletion Syndrome (DiGeorge Syndrome/Velocardiofacial Syndrome). Vaccine Journal, 1998, 5, 415-417.	2.6	74
75	GENETICS OF SYSTEMIC LUPUS ERYTHEMATOSUS. Rheumatic Disease Clinics of North America, 2000, 26, 229-256.	1.9	73
76	Histone acetylation and chromatin conformation are regulated separately at the TNF-Î± promoter in monocytes and macrophages. Journal of Leukocyte Biology, 2003, 73, 862-871.	3.3	73
77	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
78	Progressive Neurodegeneration in Patients with Primary Immunodeficiency Disease on IVIG Treatment. Clinical Immunology, 2002, 102, 19-24.	3.2	70
79	The TNFÎ± locus is altered in monocytes from patients with systemic lupus erythematosus. Clinical Immunology, 2007, 123, 74-81.	3.2	70
80	Allogeneic hematopoietic cell transplantation for primary immune deficiency diseases: Current status and critical needs. Journal of Allergy and Clinical Immunology, 2008, 122, 1087-1096.	2.9	70
81	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
82	Complement deficiency and autoimmunity. Current Opinion in Pediatrics, 1998, 10, 600-606.	2.0	69
83	Longitudinal Analysis of Lymphocyte Function and Numbers in the First Year of Life in Chromosome 22q11.2 Deletion Syndrome (DiGeorge Syndrome/Velocardiofacial Syndrome). Vaccine Journal, 1999, 6, 906-911.	2.6	68
84	GCF2/LRRFIP1 Represses Tumor Necrosis Factor Alpha Expression. Molecular and Cellular Biology, 2005, 25, 9073-9081.	2.3	68
85	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
86	Therapeutic options for CTLA-4 insufficiency. Journal of Allergy and Clinical Immunology, 2022, 149, 736-746.	2.9	68
87	Enteroviruses in X-Linked Agammaglobulinemia: Update on Epidemiology and Therapyâˆ—. Journal of Allergy and Clinical Immunology: in Practice, 2016, 4, 1059-1065.	3.8	67
88	Geosmithia argillacea: An Emerging Cause of Invasive Mycosis in Human Chronic Granulomatous Disease. Clinical Infectious Diseases, 2011, 52, e136-e143.	5.8	66
89	Primary Immune Deficiency Treatment Consortium (PIDTC) report. Journal of Allergy and Clinical Immunology, 2014, 133, 335-347.e11.	2.9	65
90	The immune deficiency of chromosome 22q11.2 deletion syndrome. American Journal of Medical Genetics, Part A, 2017, 173, 2366-2372.	1.2	64

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91	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
92	Early-Onset Inflammatory Bowel Disease. Immunology and Allergy Clinics of North America, 2019, 39, 63-79.	1.9	64
93	Cytokine and Chemokine Dysregulation in Hyper-IgE Syndrome. Clinical Immunology, 2001, 100, 49-56.	3.2	63
94	Immunoglobulin Deficiencies: The B-Lymphocyte Side of DiGeorge Syndrome. Journal of Pediatrics, 2012, 161, 950-953.e1.	1.8	63
95	Association of CLEC16A with human common variable immunodeficiency disorder and role in murine B cells. Nature Communications, 2015, 6, 6804.	12.8	63
96	Autosomal Dominant Hyper-IgE Syndrome in the USIDNET Registry. Journal of Allergy and Clinical Immunology: in Practice, 2018, 6, 996-1001.	3.8	62
97	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
98	Malignancy in chromosome 22q11.2 deletion syndrome (DiGeorge syndrome/velocardiofacial) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 462	1.2	59
99	Genetic sharing and heritability of paediatric age of onset autoimmune diseases. Nature Communications, 2015, 6, 8442.	12.8	58
100	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
101	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
102	Genetics and Epigenetics of Systemic Lupus Erythematosus. Current Rheumatology Reports, 2013, 15, 369.	4.7	56
103	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
104	Analysis of polymorphisms affecting immune complex handling in systemic lupus erythematosus. British Journal of Rheumatology, 2003, 42, 446-452.	2.3	55
105	Defective antigen-induced lymphocyte proliferation in the X-linked hyper-IgM syndrome. Journal of Pediatrics, 1997, 131, 147-150.	1.8	54
106	Graves' disease in patients with 22q11.2 deletion. Journal of Pediatrics, 2001, 139, 892-895.	1.8	54
107	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
108	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

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109	Monogenic lupus: it's all new!. Current Opinion in Immunology, 2017, 49, 87-95.	5.5	51
110	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
111	H3K4 tri-methylation breadth at transcription start sites impacts the transcriptome of systemic lupus erythematosus. Clinical Epigenetics, 2016, 8, 14.	4.1	47
112	Mannose Binding Lectin (MBL) Polymorphisms Associated with Low MBL Production in Patients with Dermatomyositis1. Journal of Investigative Dermatology, 2002, 119, 1394-1399.	0.7	46
113	Allergies in patients with chromosome 22q11.2 deletion syndrome (DiGeorge) Tj ETQq1 1 0.784314 rgBT /Overlock 10 Tf 50 587 Td (cy Allergy and Immunology, 2005, 16, 226-230.	2.6	46
114	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
115	Inflammatory Bowel Disease in Primary Immunodeficiencies. Current Allergy and Asthma Reports, 2017, 17, 57.	5.3	46
116	The Diverse Clinical Features of Chromosome 22q11.2 Deletion Syndrome (DiGeorge Syndrome). Journal of Allergy and Clinical Immunology: in Practice, 2013, 1, 589-594.	3.8	45
117	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
118	Interferon regulatory factor 1 and histone H4 acetylation in systemic lupus erythematosus. Epigenetics, 2015, 10, 191-199.	2.7	44
119	A Potential Screening Tool for IPEX Syndrome. Pediatric and Developmental Pathology, 2007, 10, 98-105.	1.0	43
120	Pediatric common variable immunodeficiency: Immunologic and phenotypic associations with switched memory B cells. Pediatric Allergy and Immunology, 2010, 21, 852-858.	2.6	43
121	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
122	CD4 + CD25 + T-Cell Production in Healthy Humans and in Patients with Thymic Hypoplasia. Vaccine Journal, 2002, 9, 1129-1131.	3.1	41
123	Genome-wide analysis of interferon regulatory factor I binding in primary human monocytes. Gene, 2011, 487, 21-28.	2.2	41
124	Transfer of influenza vaccineâ€“primed costimulated autologous T cells after stem cell transplantation for multiple myeloma leads to reconstitution of influenza immunity: results of a randomized clinical trial. Blood, 2011, 117, 63-71.	1.4	41
125	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
126	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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127	Adoptive Transfer of Autologous T Cells Improves T-cell Repertoire Diversity and Long-term B-cell Function in Pediatric Patients with Neuroblastoma. Clinical Cancer Research, 2012, 18, 6732-6741.	7.0	39
128	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
129	Recurrent and Sustained Viral Infections in Primary Immunodeficiencies. Frontiers in Immunology, 2017, 8, 665.	4.8	37
130	Convalescent plasma for pediatric patients with SARSâ€CoVâ€2â€associated acute respiratory distress syndrome. Pediatric Blood and Cancer, 2020, 67, e28693.	1.5	37
131	SERPINB2 is regulated by dynamic interactions with pause-release proteins and enhancer RNAs. Molecular Immunology, 2017, 88, 20-31.	2.2	37
132	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
133	Heterozygous OAS1 gain-of-function variants cause an autoinflammatory immunodeficiency. Science Immunology, 2021, 6, .	11.9	36
134	Association of Toxic Shock Syndrome Toxin-Secreting and Exfoliative Toxin-Secreting Staphylococcus aureus with Kawasaki Syndrome Complicated by Coronary Artery Disease. Pediatric Research, 1997, 42, 268-272.	2.3	36
135	Contiguous X-chromosome Deletion Syndrome Encompassing the BTK, TIMM8A, TAF7L, and DRP2 Genes. Journal of Clinical Immunology, 2007, 27, 640-646.	3.8	35
136	Polarization of Primary Human Monocytes by IFN-Î³ Induces Chromatin Changes and Recruits RNA Pol II to the TNF-Î± Promoter. Journal of Immunology, 2008, 180, 5257-5266.	0.8	35
137	A Prospective Study of Influenza Vaccination and a Comparison of Immunologic Parameters in Children and Adults with Chromosome 22q11.2 Deletion Syndrome (DiGeorge) Tj ETQq1 1 0.784314 rgBT /Overlock 10 Tf 50 337 Td (S	3.8	35
138	Identification of 22q11.2 Deletion Syndrome via Newborn Screening for Severe Combined Immunodeficiency. Journal of Clinical Immunology, 2017, 37, 476-485.	3.8	35
139	Cell Cycle Checkpoints and DNA Repair in Nijmegen Breakage Syndrome. Clinical Immunology and Immunopathology, 1997, 82, 43-48.	2.0	34
140	Gamma Interferon and Lipopolysaccharide Interact at the Level of Transcription To Induce Tumor Necrosis Factor Alpha Expression. Infection and Immunity, 2001, 69, 2847-2852.	2.2	34
141	ICON: The Early Diagnosis of Congenital Immunodeficiencies. Journal of Clinical Immunology, 2014, 34, 398-424.	3.8	34
142	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
143	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
144	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

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145	Immunodeficiencies Associated with Abnormal Newborn Screening for T Cell and B Cell Lymphopenia. Journal of Clinical Immunology, 2017, 37, 363-374.	3.8	33
146	The Unique Disease Course of Children with Very Early onset-Inflammatory Bowel Disease. Inflammatory Bowel Diseases, 2020, 26, 909-918.	1.9	32
147	3? polymorphisms of ETS1 are associated with different clinical phenotypes in SLE. Human Mutation, 2000, 16, 49-53.	2.5	31
148	USIDNET: A Strategy to Build a Community of Clinical Immunologists. Journal of Clinical Immunology, 2014, 34, 428-435.	3.8	31
149	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
150	Constrained chromatin accessibility in PU.1-mutated agammaglobulinemia patients. Journal of Experimental Medicine, 2021, 218, .	8.5	31
151	Monocyte enhancers are highly altered in systemic lupus erythematosus. Epigenomics, 2015, 7, 921-935.	2.1	27
152	Phenotypic characterization of patients with rheumatologic manifestations of common variable immunodeficiency. Seminars in Arthritis and Rheumatism, 2018, 48, 318-326.	3.4	26
153	Tollâ€like receptor 2 stimulation augments esophageal barrier integrity. Allergy: European Journal of Allergy and Clinical Immunology, 2019, 74, 2449-2460.	5.7	26
154	Diminished T Cell Numbers in Patients with Chronic Granulomatous Disease. Clinical Immunology, 2002, 105, 273-278.	3.2	24
155	Myeloid differentiation primary response gene 88 (MyD88) deficiency in a large kindred. Journal of Allergy and Clinical Immunology, 2010, 126, 172-175.	2.9	24
156	Protecting Pediatric Oncology Patients From Influenza. Oncologist, 2013, 18, 204-211.	3.7	24
157	Terminal Deletion of 11q with Significant Late-Onset Combined Immune Deficiency. Journal of Clinical Immunology, 2014, 34, 114-118.	3.8	24
158	B cell development in chromosome 22q11.2 deletion syndrome. Clinical Immunology, 2016, 163, 1-9.	3.2	24
159	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
160	Rubella virus-associated chronic inflammation in primary immunodeficiency diseases. Current Opinion in Allergy and Clinical Immunology, 2020, 20, 574-581.	2.3	24
161	Digeorge syndrome/chromosome 22q11.2 deletion syndrome. Current Allergy and Asthma Reports, 2001, 1, 438-444.	5.3	23
162	Regulation of Inflammation. Immunologic Research, 2003, 27, 529-538.	2.9	23

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163	The Efficacy of Influenza Vaccination in a Pediatric Oncology Population. Journal of Pediatric Hematology/Oncology, 2010, 32, e177-e181.	0.6	23
164	Noncoding RNAs and LRRFIP1 Regulate TNF Expression. Journal of Immunology, 2014, 192, 3057-3067.	0.8	23
165	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
166	Characterization of LRRFIP1. Biochemistry and Cell Biology, 2010, 88, 899-906.	2.0	22
167	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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