Charlotte Cunningham-Rundles

List of Publications by Year
in descending order

363 papers	32,993 citations	4388 86 h-index	4774 169 g-index
377 all docs	377 docs citations	377 times ranked	25621 citing authors

#	ARTICLE	IF	CITATIONS
1	Overactive WASp in X-linked neutropenia leads to aberrant B-cell division and accelerated plasma cell generation. Journal of Allergy and Clinical Immunology, 2022, 149, 1069-1084.	2.9	5
2	Rheumatologic diseases in patients with inborn errors of immunity in the USIDNET registry. Clinical Rheumatology, 2022, 41, 2197-2203.	2.2	5
3	Morbidity, Mortality, and Therapeutics in Combined Immunodeficiency: Data from the USIDNET Registry. Journal of Allergy and Clinical Immunology: in Practice, 2022, , .	3.8	0
4	Practical guidance for the diagnosis and management of secondary hypogammaglobulinemia: AÂWork Group Report of the AAAAI Primary Immunodeficiency and Altered Immune Response Committees. Journal of Allergy and Clinical Immunology, 2022, 149, 1525-1560.	2.9	53
5	Seeking Relevant Biomarkers in Common Variable Immunodeficiency. Frontiers in Immunology, 2022, 13, 857050.	4.8	14
6	X-Linked Agammaglobulinemia: Infection Frequency and Infection-Related Mortality in the USIDNET Registry. Journal of Clinical Immunology, 2022, 42, 827-836.	3.8	11
7	eP236: TeleKidSeq: Incorporating telehealth into clinical care of children from diverse backgrounds undergoing clinical genome sequencing. Genetics in Medicine, 2022, 24, S150.	2.4	2
8	Case Series: Convalescent Plasma Therapy for Patients with COVID-19 and Primary Antibody Deficiency. Journal of Clinical Immunology, 2022, 42, 253-265.	3.8	19
9	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
10	Genomic characterization of lymphomas in patients with inborn errors of immunity. Blood Advances, 2022, 6, 5403-5414.	5.2	12
11	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
12	Clinical outcomes and features of COVID-19 in patients with primary immunodeficiencies in New York City. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 490-493.e2.	3.8	72
13	IFN-Î³ receptor 2 deficiency initial mimicry of multisystem inflammatory syndrome in children (MIS-C). Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 989-992.e1.	3.8	6
14	Germline IKAROS dimerization haploinsufficiency causes hematologic cytopenias and malignancies. Blood, 2021, 137, 349-363.	1.4	32
15	LIG1 syndrome mutations remodel a cooperative network of ligand binding interactions to compromise ligation efficiency. Nucleic Acids Research, 2021, 49, 1619-1630.	14.5	14
16	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
17	Interstitial Lung Disease in Common Variable Immunodeficiency. Frontiers in Immunology, 2021, 12, 605945.	4.8	12
18	Clinical disparity of primary antibody deficiency patients at a safety net hospital. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 2923-2925.e1.	3.8	6

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19	International multicenter experience of transjugular intrahepatic portosystemic shunt implantation in patients with common variable immunodeficiency. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 2931-2935.e1.	3.8	4
20	Treatment Strategies for GLILD in Common Variable Immunodeficiency: A Systematic Review. Frontiers in Immunology, 2021, 12, 606099.	4.8	24
21	Convergence of cytokine dysregulation and antibody deficiency in common variable immunodeficiency with inflammatory complications. Journal of Allergy and Clinical Immunology, 2021, , .	2.9	13
22	Lymphoid malignancy in common variable immunodeficiency in a singleâ€center cohort. European Journal of Haematology, 2021, 107, 503-516.	2.2	8
23	Clinical Manifestations and Outcomes of Activated Phosphoinositide 3-Kinase Î´ Syndrome from the USIDNET Cohort. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 4095-4102.	3.8	11
24	Biochemically deleterious human NFKB1 variants underlie an autosomal dominant form of common variable immunodeficiency. Journal of Experimental Medicine, 2021, 218, .	8.5	32
25	On the relevance of immunodeficiency evaluation in haematological cancer. Hematological Oncology, 2021, 39, 721-723.	1.7	3
26	COVID-19 prevalence and outcomes in patients receiving biologic therapies at an infusion center in New York City. Clinical Immunology, 2021, 230, 108803.	3.2	3
27	Crohnâ€™s-like Enteritis in X-Linked Agammaglobulinemia: A Case Series and Systematic Review. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 3466-3478.	3.8	2
28	The NYCKidSeq project: study protocol for a randomized controlled trial incorporating genomics into the clinical care of diverse New York City children. Trials, 2021, 22, 56.	1.6	21
29	Circulating bioactive bacterial DNA is associated with immune activation and complications in common variable immunodeficiency. JCI Insight, 2021, 6, .	5.0	22
30	Serum B-Cell Maturation Antigen (BCMA) Levels Differentiate Primary Antibody Deficiencies. Journal of Allergy and Clinical Immunology: in Practice, 2020, 8, 283-291.e1.	3.8	15
31	Chemical chaperones reverse early suppression of regulatory circuits during unfolded protein response in B cells from common variable immunodeficiency patients. Clinical and Experimental Immunology, 2020, 200, 73-86.	2.6	4
32	M244 ORAL LESIONS IN A PATIENT WITH HYPER IMMUNOGLOBULIN M: DIFFERENTIAL DIAGNOSIS AND MANAGEMENT. Annals of Allergy, Asthma and Immunology, 2020, 125, S82-S83.	1.0	1
33	Three patients with X-linked agammaglobulinemia hospitalized for COVID-19 improved with convalescent plasma. Journal of Allergy and Clinical Immunology: in Practice, 2020, 8, 3594-3596.e3.	3.8	72
34	The Importance of Primary Immune Deficiency Registries. Immunology and Allergy Clinics of North America, 2020, 40, 385-402.	1.9	6
35	Gut T cellâ€“independent IgA responses to commensal bacteria require engagement of the TACI receptor on B cells. Science Immunology, 2020, 5, .	11.9	40
36	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

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37	Targeting FcRn for immunomodulation: Benefits, risks, and practical considerations. Journal of Allergy and Clinical Immunology, 2020, 146, 479-491.e5.	2.9	52
38	Neurologic Conditions and Symptoms Reported Among Common Variable Immunodeficiency Patients in the USIDNET. Journal of Clinical Immunology, 2020, 40, 1181-1183.	3.8	3
39	A Nonsense N â€“Terminus NFKB2 Mutation Leading to Haploinsufficiency in a Patient with a Predominantly Antibody Deficiency. Journal of Clinical Immunology, 2020, 40, 1093-1101.	3.8	7
40	Reticular dysgenesis caused by an intronic pathogenic variant in AK2. Journal of Physical Education and Sports Management, 2020, 6, a005017.	1.2	4
41	A serological assay to detect SARS-CoV-2 seroconversion in humans. Nature Medicine, 2020, 26, 1033-1036.	30.7	1,678
42	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
43	Primary Immunodeficiency Diagnoses seen in Patients with Chronic Lung Disease: Findings from the USIDNET Registry. Journal of Allergy and Clinical Immunology, 2020, 145, AB178.	2.9	0
44	Hypogammaglobulinemia and common variable immune deficiency. , 2020, , 467-497.		4
45	Non-infectious Complications of Common Variable Immunodeficiency: Updated Clinical Spectrum, Sequelae, and Insights to Pathogenesis. Frontiers in Immunology, 2020, 11, 149.	4.8	118
46	Vedolizumab therapy in common variable immune deficiency associated enteropathy: A case series. Clinical Immunology, 2020, 212, 108362.	3.2	12
47	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
48	Characterization of the clinical and immunologic phenotype and management of 157 individuals with 56 distinct heterozygous NFKB1 mutations. Journal of Allergy and Clinical Immunology, 2020, 146, 901-911.	2.9	78
49	CVID-associated intestinal disorders in the USIDNET registry: An analysis of disease phenotypes, functional status, comorbidities, and treatment. Journal of Allergy and Clinical Immunology, 2020, 145, AB80.	2.9	0
50	Current genetic landscape in common variable immune deficiency. Blood, 2020, 135, 656-667.	1.4	109
51	Human Inborn Errors of Immunity: 2019 Update of the IUIS Phenotypical Classification. Journal of Clinical Immunology, 2020, 40, 66-81.	3.8	525
52	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
53	Autoimmunity in common variable immunodeficiency. Annals of Allergy, Asthma and Immunology, 2019, 123, 454-460.	1.0	64
54	Cellular Defects in CVID Patients with Chronic Lung Disease in the USIDNET Registry. Journal of Clinical Immunology, 2019, 39, 569-576.	3.8	12

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55	Respiratory Comorbidities Associated with Bronchiectasis in Patients with Common Variable Immunodeficiency. Journal of Allergy and Clinical Immunology, 2019, 143, AB13.	2.9	0
56	Factors Beyond Lack of Antibody Govern Pulmonary Complications in Primary Antibody Deficiency. Journal of Clinical Immunology, 2019, 39, 440-447.	3.8	29
57	AIRE expression controls the peripheral selection of autoreactive B cells. Science Immunology, 2019, 4, .	11.9	65
58	Common variable immune deficiency: case studies. Hematology American Society of Hematology Education Program, 2019, 2019, 449-456.	2.5	8
59	Common variable immune deficiency: case studies. Blood, 2019, 134, 1787-1795.	1.4	18
60	M292 NUTRITIONAL SUPPLEMENTATION IN PATIENTS WITH COMBINED IMMUNODEFICIENCY SECONDARY TO MTHFD1 DEFICIENCY. Annals of Allergy, Asthma and Immunology, 2019, 123, S122.	1.0	0
61	Common variable immune deficiency: Dissection of the variable. Immunological Reviews, 2019, 287, 145-161.	6.0	59
62	Blacklisting variants common in private cohorts but not in public databases optimizes human exome analysis. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 950-959.	7.1	52
63	Expansion of the Human Phenotype Ontology (HPO) knowledge base and resources. Nucleic Acids Research, 2019, 47, D1018-D1027.	14.5	539
64	Gastrointestinal Manifestations and Complications of Primary Immunodeficiency Disorders. Immunology and Allergy Clinics of North America, 2019, 39, 81-94.	1.9	37
65	Differentiation of Common Variable Immunodeficiency From IgG Deficiency. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 1277-1284.	3.8	43
66	Primary B-cell immunodeficiencies. Human Immunology, 2019, 80, 351-362.	2.4	42
67	BAFF-driven B cell hyperplasia underlies lung disease in common variable immunodeficiency. JCI Insight, 2019, 4, .	5.0	54
68	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
69	Ruxolitinib partially reverses functional natural killer cell deficiency in patients with signal transducer and activator of transcription 1 (STAT1) gain-of-function mutations. Journal of Allergy and Clinical Immunology, 2018, 141, 2142-2155.e5.	2.9	79
70	Ralph Josiah Patrick Wedgwood (1924â€“2017). Journal of Clinical Immunology, 2018, 38, 153-154.	3.8	0
71	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
72	Detection of antiâ€“glutamic acid decarboxylase antibodies in immunoglobulin products. Journal of Allergy and Clinical Immunology: in Practice, 2018, 6, 260-261.	3.8	9

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73	Autoimmune Cytopenias and Associated Conditions in CVID: a Report From the USIDNET Registry. Journal of Clinical Immunology, 2018, 38, 28-34.	3.8	79
74	Autosomal Dominant Hyper-IgE Syndrome in the USIDNET Registry. Journal of Allergy and Clinical Immunology: in Practice, 2018, 6, 996-1001.	3.8	62
75	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
76	The 2017 IUIS Phenotypic Classification for Primary Immunodeficiencies. Journal of Clinical Immunology, 2018, 38, 129-143.	3.8	488
77	2153 The plasma contact system and its role in common variable immunodeficiency (CVID): An explorative study. Journal of Clinical and Translational Science, 2018, 2, 32-32.	0.6	0
78	Evaluation of Lymphoproliferative Disease and Increased Risk of Lymphoma in Activated Phosphoinositide 3 Kinase Delta Syndrome: A Case Report With Discussion. Frontiers in Pediatrics, 2018, 6, 402.	1.9	8
79	BRWD1 orchestrates epigenetic landscape of late B lymphopoiesis. Nature Communications, 2018, 9, 3888.	12.8	24
80	TACI Isoforms Regulate Ligand Binding and Receptor Function. Frontiers in Immunology, 2018, 9, 2125.	4.8	26
81	Jakinibs for the treatment of immune dysregulation in patients with gain-of-function signal transducer and activator of transcription 1 (STAT1) or STAT3 mutations. Journal of Allergy and Clinical Immunology, 2018, 142, 1665-1669.	2.9	196
82	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
83	CDG: An Online Server for Detecting Biologically Closest Disease-Causing Genes and its Application to Primary Immunodeficiency. Frontiers in Immunology, 2018, 9, 1340.	4.8	6
84	Disseminated Cutaneous Warts in X-Linked Hyper IgM Syndrome. Journal of Clinical Immunology, 2018, 38, 454-456.	3.8	3
85	Biallelic mutations in DNA ligase 1 underlie a spectrum of immune deficiencies. Journal of Clinical Investigation, 2018, 128, 5489-5504.	8.2	32
86	Clonal and constricted T cell repertoire in Common Variable Immune Deficiency. Clinical Immunology, 2017, 178, 1-9.	3.2	36
87	Differences in Pulmonary Complications in Common Variable Immunodeficiency and X-Linked Agammaglobulinemia. Journal of Allergy and Clinical Immunology, 2017, 139, AB111.	2.9	0
88	Dysregulation of Innate Lymphoid Cells in Common Variable Immunodeficiency. Current Allergy and Asthma Reports, 2017, 17, 77.	5.3	8
89	Idiopathic CD4 lymphocytopenia. Annals of Allergy, Asthma and Immunology, 2017, 119, 374-378.	1.0	40
90	mTOR intersects antibody-inducing signals from TACI in marginal zone B cells. Nature Communications, 2017, 8, 1462.	12.8	65

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91	Idiopathic T cell lymphopenia identified in New York State Newborn Screening. Clinical Immunology, 2017, 183, 36-40.	3.2	27
92	Fulminant Sepsis Due to Granulibacter bethesdensis in a 4-Year-Old Boy With X-Linked Chronic Granulomatous Disease. Pediatric Infectious Disease Journal, 2017, 36, 1165-1166.	2.0	5
93	Long-term outcomes of 176 patients with X-linked hyper-IgM syndrome treated with or without hematopoietic cell transplantation. Journal of Allergy and Clinical Immunology, 2017, 139, 1282-1292.	2.9	107
94	Lack of Clinical Hypersensitivity to Penicillin Antibiotics in Common Variable Immunodeficiency. Journal of Clinical Immunology, 2017, 37, 22-24.	3.8	11
95	OR064 Ocular manifestations in primary immunodeficiency (PID) patients within the us immunodeficiency network (USIDNET) registry. Annals of Allergy, Asthma and Immunology, 2017, 119, S8-S9.	1.0	0
96	P284 STAT 1 gain of function mutation treated with ruxolitinib. Annals of Allergy, Asthma and Immunology, 2017, 119, S72.	1.0	1
97	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
98	Genetic Diagnosis Using Whole Exome Sequencing in Common Variable Immunodeficiency. Frontiers in Immunology, 2016, 7, 220.	4.8	247
99	Primary Immunodeficiency: New Insights and Practical Clinical Approaches. Journal of Allergy and Clinical Immunology: in Practice, 2016, 4, 1109-1110.	3.8	0
100	P186 Case of signal transducer and activator of transcription (STAT) 3 gain of function mutation. Annals of Allergy, Asthma and Immunology, 2016, 117, S77-S78.	1.0	0
101	Hyper IgM Syndrome: a Report from the USIDNET Registry. Journal of Clinical Immunology, 2016, 36, 490-501.	3.8	92
102	Hemoptysis in a Patient with Elevated Immunoglobulin E. Journal of Allergy and Clinical Immunology: in Practice, 2016, 4, 1054-1058.	3.8	8
103	A healthy female with C3 hypocomplementemia and C3 Nephritic Factor. Clinical Immunology, 2016, 169, 14-15.	3.2	9
104	BK virus encephalopathy and sclerosing vasculopathy in a patient with hypohidrotic ectodermal dysplasia and immunodeficiency. Acta Neuropathologica Communications, 2016, 4, 73.	5.2	13
105	Efficacy, Safety, and Pharmacokinetics of a New 10Â% Liquid Intravenous Immunoglobulin Containing High Titer Neutralizing Antibody to RSV and Other Respiratory Viruses in Subjects with Primary Immunodeficiency Disease. Journal of Clinical Immunology, 2016, 36, 590-599.	3.8	22
106	Eosinophilic esophagitis diagnosed in a patient with common variable immunodeficiency. Journal of Allergy and Clinical Immunology: in Practice, 2016, 4, 995-997.	3.8	10
107	Clinical Experience of CVID Enteropathy. Journal of Allergy and Clinical Immunology, 2016, 137, AB179.	2.9	0
108	International Consensus Document (ICON): Common Variable Immunodeficiency Disorders. Journal of Allergy and Clinical Immunology: in Practice, 2016, 4, 38-59.	3.8	669

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109	Loss of B Cells in Patients with Heterozygous Mutations in IKAROS. New England Journal of Medicine, 2016, 374, 1032-1043.	27.0	217
110	Gastrointestinal Disorders Associated with Common Variable Immune Deficiency (CVID) and Chronic Granulomatous Disease (CGD). Current Gastroenterology Reports, 2016, 18, 17.	2.5	104
111	Microbiota regulate the ability of lung dendritic cells to induce IgA class-switch recombination and generate protective gastrointestinal immune responses. Journal of Experimental Medicine, 2016, 213, 53-73.	8.5	94
112	CD19 controls Toll-like receptor 9 responses in human BÂcells. Journal of Allergy and Clinical Immunology, 2016, 137, 889-898.e6.	2.9	50
113	Expansion of inflammatory innate lymphoid cells in patients with common variable immune deficiency. Journal of Allergy and Clinical Immunology, 2016, 137, 1206-1215.e6.	2.9	69
114	Decreased somatic hypermutation induces an impaired peripheral B cell tolerance checkpoint. Journal of Clinical Investigation, 2016, 126, 4289-4302.	8.2	46
115	Cancer in primary immunodeficiency diseases: An analysis of cancer incidence in the United States immune deficiency network (USIDNET) registry.. Journal of Clinical Oncology, 2016, 34, 1520-1520.	1.6	1
116	Rare variants at 16p11.2 are associated with common variable immunodeficiency. Journal of Allergy and Clinical Immunology, 2015, 135, 1569-1577.	2.9	22
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	TNF receptor superfamily member 13b (TNFRSF13B) hemizygosity reveals transmembrane activator and CAML interactor haploinsufficiency at later stages of B-cell development. Journal of Allergy and Clinical Immunology, 2015, 136, 1315-1325.	2.9	38
119	Tollâ€like receptor signaling in primary immune deficiencies. Annals of the New York Academy of Sciences, 2015, 1356, 1-21.	3.8	71
120	Differential induction of plasma cells by isoforms of human TACI. Blood, 2015, 125, 1749-1758.	1.4	45
121	Antibody Deficiencies. , 2015, , 341-347.		0
122	Infections in the Compromised Host. , 2015, , 435-440.		0
123	Association of CLEC16A with human common variable immunodeficiency disorder and role in murine B cells. Nature Communications, 2015, 6, 6804.	12.8	63
124	TLR7- and TLR9-Responsive Human B Cells Share Phenotypic and Genetic Characteristics. Journal of Immunology, 2015, 194, 3035-3044.	0.8	43
125	A Novel Targeted Screening Tool for Hypogammaglobulinemia: Measurement of Serum Immunoglobulin (IgG, IgM, IgA) Levels from Dried Blood Spots (Ig-DBS Assay). Journal of Clinical Immunology, 2015, 35, 573-582.	3.8	11
126	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

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127	The 2015 IUIS Phenotypic Classification for Primary Immunodeficiencies. Journal of Clinical Immunology, 2015, 35, 727-738.	3.8	199
128	Genetic sharing and heritability of paediatric age of onset autoimmune diseases. Nature Communications, 2015, 6, 8442.	12.8	58
129	Combined immunodeficiency in the United States and Kuwait: Comparison of patients' characteristics and molecular diagnosis. Clinical Immunology, 2015, 161, 170-173.	3.2	22
130	Meta-analysis of shared genetic architecture across ten pediatric autoimmune diseases. Nature Medicine, 2015, 21, 1018-1027.	30.7	212
131	IgH sequences in common variable immune deficiency reveal altered B cell development and selection. Science Translational Medicine, 2015, 7, 302ra135.	12.4	77
132	Progression of Common Variable Immunodeficiency Interstitial Lung Disease Accompanies Distinct Pulmonary and Laboratory Findings. Journal of Allergy and Clinical Immunology: in Practice, 2015, 3, 941-950.	3.8	65
133	High-throughput sequencing reveals an altered T cell repertoire in X-linked agammaglobulinemia. Clinical Immunology, 2015, 161, 190-196.	3.2	9
134	An update on the use of immunoglobulin for the treatment of immunodeficiency disorders. Immunotherapy, 2014, 6, 1113-1126.	2.0	51
135	Hypogammaglobulinemia and Common Variable Immunodeficiency. , 2014, , 347-365.		2
136	Primary Immunodeficiency Diseases: An Update on the Classification from the International Union of Immunological Societies Expert Committee for Primary Immunodeficiency. Frontiers in Immunology, 2014, 5, 162.	4.8	466
137	Newborn Screening for Severe Combined Immunodeficiency in 11 Screening Programs in the United States. JAMA - Journal of the American Medical Association, 2014, 312, 729.	7.4	586
138	Tertiary lymphoid neogenesis is a component of pulmonary lymphoid hyperplasia in patients with common variable immunodeficiency. Journal of Allergy and Clinical Immunology, 2014, 133, 535-542.	2.9	55
139	Expansion Of Circulating T Follicular Helper Cells In CVID Patients With Autoimmune Cytopenias. Journal of Allergy and Clinical Immunology, 2014, 133, AB162.	2.9	3
140	Newborn Screening for SCID in New York State: Experience from the First Two Years. Journal of Clinical Immunology, 2014, 34, 289-303.	3.8	104
141	Phellinus tropicalis Abscesses in a Patient with Chronic Granulomatous Disease. Journal of Clinical Immunology, 2014, 34, 130-133.	3.8	18
142	Burden of copy number variation in common variable immunodeficiency. Clinical and Experimental Immunology, 2014, 177, 269-271.	2.6	20
143	Infants With Idiopathic T Cell Lymphopenia Identified On New York State Newborn Screen: A Follow Up Report. Journal of Allergy and Clinical Immunology, 2014, 133, AB93.	2.9	2
144	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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145	USIDNET: A Strategy to Build a Community of Clinical Immunologists. Journal of Clinical Immunology, 2014, 34, 428-435.	3.8	31
146	Autoimmunity and Inflammation in X-linked Agammaglobulinemia. Journal of Clinical Immunology, 2014, 34, 627-632.	3.8	88
147	Studies On Cohort Of Infants With Di-George Syndrome Detected By New York State Newborn Screening For Severe Combined Immunodeficiency (SCID). Journal of Allergy and Clinical Immunology, 2014, 133, AB96.	2.9	2
148	Signaling lymphocytic activation molecule (SLAM)/SLAM-associated protein pathway regulates human B-cell tolerance. Journal of Allergy and Clinical Immunology, 2014, 133, 1149-1161.	2.9	33
149	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
150	Lloyd Mayer, MD, 1952â€“2013, In Memoriam. Clinical Immunology, 2014, 150, A1-A2.	3.2	0
151	Pulmonary radiologic findings in common variable immunodeficiency: clinical and immunological correlations. Annals of Allergy, Asthma and Immunology, 2014, 113, 452-459.	1.0	86
152	IRAK-4 and MyD88 deficiencies impair IgM responses against T-independent bacterial antigens. Blood, 2014, 124, 3561-3571.	1.4	58
153	Prioritization of Evidence-Based Indications for Intravenous Immunoglobulin. Journal of Clinical Immunology, 2013, 33, 1033-1036.	3.8	20
154	Examining the Use of ICD-9 Diagnosis Codes for Primary Immune Deficiency Diseases in New York State. Journal of Clinical Immunology, 2013, 33, 40-48.	3.8	17
155	Home Care Use of Intravenous and Subcutaneous Immunoglobulin for Primary Immunodeficiency in the United States. Journal of Clinical Immunology, 2013, 33, 49-54.	3.8	14
156	Exploratory laparoscopy for rotational abnormality of the intestine in a child with leukocyte adhesion deficiency type II. Journal of Pediatric Surgery Case Reports, 2013, 1, 244-246.	0.2	0
157	Treatment of hypogammaglobulinemia in adults: A scoring system to guide decisions on immunoglobulin replacement. Journal of Allergy and Clinical Immunology, 2013, 131, 1699-1701.e3.	2.9	36
158	TACI mutations and impaired B-cell function in subjects with CVID and healthy heterozygotes. Journal of Allergy and Clinical Immunology, 2013, 131, 468-476.	2.9	86
159	Treatment of common variable immune deficiency. Expert Opinion on Orphan Drugs, 2013, 1, 157-166.	0.8	1
160	Naturally occurring mutation affecting the <scp>M</scp>y<scp>D</scp>88â€binding site of <scp>TNFRSF</scp>13<scp>B</scp> impairs triggering of class switch recombination. European Journal of Immunology, 2013, 43, 805-814.	2.9	14
161	Interferon Signature in the Blood in Inflammatory Common Variable Immune Deficiency. PLoS ONE, 2013, 8, e74893.	2.5	64
162	CVID-associated TACI mutations affect autoreactive B cell selection and activation. Journal of Clinical Investigation, 2013, 123, 4283-4293.	8.2	153

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163	The many faces of the clinical picture of common variable immune deficiency. Current Opinion in Allergy and Clinical Immunology, 2012, 12, 595-601.	2.3	72
164	Morbidity and mortality in common variable immune deficiency over 4 decades. Blood, 2012, 119, 1650-1657.	1.4	685
165	Human B cell defects in perspective. Immunologic Research, 2012, 54, 227-232.	2.9	28
166	Toll-like receptor 4â€“, 7â€–, and 8â€–activated myeloid cells from patients with X-linked agammaglobulinemia produce enhanced inflammatory cytokines. Journal of Allergy and Clinical Immunology, 2012, 129, 184-190.e4.	2.9	47
167	Common variable immunodeficiency. Journal of Allergy and Clinical Immunology, 2012, 129, 1425-1426.e3.	2.9	62
168	Confirmation and improvement of criteria for clinical phenotyping in common variable immunodeficiency disorders in replicate cohorts. Journal of Allergy and Clinical Immunology, 2012, 130, 1197-1198.e9.	2.9	129
169	TLR-Mediated B Cell Defects and IFN-Î± in Common Variable Immunodeficiency. Journal of Clinical Immunology, 2012, 32, 50-60.	3.8	35
170	The many faces of common variable immunodeficiency. Hematology American Society of Hematology Education Program, 2012, 2012, 301-5.	2.5	122
171	Toll-like receptor function in primary B cell defects. Frontiers in Bioscience - Elite, 2012, E4, 1853.	1.8	7
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