Jolan E Walter

List of Publications by Year
in descending order

131 papers	4,579 citations	101543 36 h-index	110387 64 g-index
136 all docs	136 docs citations	136 times ranked	5980 citing authors

#	Article	IF	Citations
1	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
2	Clinical Evaluation of Youth with Pediatric Acute-Onset Neuropsychiatric Syndrome (PANS): Recommendations from the 2013 PANS Consensus Conference. Journal of Child and Adolescent Psychopharmacology, 2015, 25, 3-13.	1.3	241
3	Association of Immunoglobulin Levels, Infectious Risk, and Mortality With Rituximab and Hypogammaglobulinemia. JAMA Network Open, 2018, 1, e184169.	5.9	210
4	Human RAG mutations: biochemistry and clinical implications. Nature Reviews Immunology, 2016, 16, 234-246.	22.7	200
5	Broad-spectrum antibodies against self-antigens and cytokines in RAG deficiency. Journal of Clinical Investigation, 2015, 125, 4135-4148.	8.2	159
6	IL-21 is the primary common Î³ chain-binding cytokine required for human B-cell differentiation in vivo. Blood, 2011, 118, 6824-6835.	1.4	132
7	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.	2.9	132
8	Astrovirus infection in children. Current Opinion in Infectious Diseases, 2003, 16, 247-253.	3.1	129
9	Activation-induced cytidine deaminase (AID) is required for B-cell tolerance in humans. Proceedings of the National Academy of Sciences of the United States of America, 2011, 108, 11554-11559.	7.1	118
10	B cellâ€“intrinsic deficiency of the Wiskott-Aldrich syndrome protein (WASp) causes severe abnormalities of the peripheral B-cell compartment in mice. Blood, 2012, 119, 2819-2828.	1.4	99
11	Hepatopulmonary Syndrome Is a Frequent Cause of Dyspnea in the Short Telomere Disorders. Chest, 2015, 148, 1019-1026.	0.8	95
12	Expansion of immunoglobulin-secreting cells and defects in B cell tolerance in Rag-dependent immunodeficiency. Journal of Experimental Medicine, 2010, 207, 1541-1554.	8.5	90
13	Characterization of T and B cell repertoire diversity in patients with RAG deficiency. Science Immunology, 2016, 1, .	11.9	88
14	PRKDC mutations associated with immunodeficiency, granuloma, and autoimmune regulatorâ€“dependent autoimmunity. Journal of Allergy and Clinical Immunology, 2015, 135, 1578-1588.e5.	2.9	84
15	Identification of Patients with RAG Mutations Previously Diagnosed with Common Variable Immunodeficiency Disorders. Journal of Clinical Immunology, 2015, 35, 119-124.	3.8	70
16	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
17	Activation-Induced Cytidine Deaminase Expression in Human B Cell Precursors Is Essential for Central B Cell Tolerance. Immunity, 2015, 43, 884-895.	14.3	69
18	Powering the Immune System: Mitochondria in Immune Function and Deficiency. Journal of Immunology Research, 2014, 2014, 1-8.	2.2	68

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19	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.	3.8	64
20	Mechanism-Based Strategies for the Management of Autoimmunity and Immune Dysregulation in Primary Immunodeficiencies. Journal of Allergy and Clinical Immunology: in Practice, 2016, 4, 1089-1100.	3.8	61
21	Loss-of-function mutations in caspase recruitment domain-containing protein 14 (CARD14) are associated with a severe variant of atopic dermatitis. Journal of Allergy and Clinical Immunology, 2019, 143, 173-181.e10.	2.9	60
22	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.	2.9	59
23	Live vaccines after pediatric solid organ transplant: Proceedings of a consensus meeting, 2018. Pediatric Transplantation, 2019, 23, e13571.	1.0	59
24	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
25	Autoimmunity due to RAG deficiency and estimated disease incidence in RAG1/2 mutations. Journal of Allergy and Clinical Immunology, 2014, 133, 880-882.e10.	2.9	54
26	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
27	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
28	Gastrointestinal Manifestations in X-linked Agammaglobulinemia. Journal of Clinical Immunology, 2017, 37, 287-294.	3.8	51
29	Autoimmunity as a continuum in primary immunodeficiency. Current Opinion in Pediatrics, 2019, 31, 851-862.	2.0	46
30	Decreased somatic hypermutation induces an impaired peripheral B cell tolerance checkpoint. Journal of Clinical Investigation, 2016, 126, 4289-4302.	8.2	46
31	Severe eczema and Hyper-IgE in Loeysâ€“Dietz-syndrome â€” Contribution to new findings of immune dysregulation in connective tissue disorders. Clinical Immunology, 2014, 150, 43-50.	3.2	43
32	Role of astroviruses in childhood diarrhea. Current Opinion in Pediatrics, 2000, 12, 275-279.	2.0	42
33	Nextâ€Generation Sequencing Reveals Restriction and Clonotypic Expansion of Treg Cells in Juvenile Idiopathic Arthritis. Arthritis and Rheumatology, 2016, 68, 1758-1768.	5.6	42
34	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.	4.8	41
35	Characterisation of a South African human astrovirus as type 8 by antigenic and genetic analyses. Journal of Medical Virology, 2001, 64, 256-261.	5.0	40
36	Genome Prediction of Putative Genome-Linked Viral Protein (VPg) of Astroviruses. Virus Genes, 2005, 31, 21-30.	1.6	40

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37	Prevalence and clinical challenges among adults with primary immunodeficiency and recombination-activating gene deficiency. Journal of Allergy and Clinical Immunology, 2018, 141, 2303-2306.	2.9	40
38	Homozygous DNA ligase IV R278H mutation in mice leads to leaky SCID and represents a model for human LIG4 syndrome. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 3024-3029.	7.1	39
39	Predisposition to infection and SIRS in mitochondrial disorders: 8 years' experience in an academic center. Journal of Allergy and Clinical Immunology: in Practice, 2014, 2, 465-468.e1.	3.8	39
40	Recombination activity of human recombination-activating gene 2 (RAG2) mutations and correlation with clinical phenotype. Journal of Allergy and Clinical Immunology, 2019, 143, 726-735.	2.9	39
41	Recent Advances in Primary Immunodeficiencies: Identification of Novel Genetic Defects and Unanticipated Phenotypes. Pediatric Research, 2009, 65, 3R-12R.	2.3	38
42	Adult-onset manifestation of idiopathic T-cell lymphopenia due to a heterozygous RAG1 mutation. Journal of Allergy and Clinical Immunology, 2013, 131, 1421-1423.	2.9	37
43	B-cell differentiation and IL-21 response in IL2RG/JAK3 SCID patients after hematopoietic stem cell transplantation. Blood, 2018, 131, 2967-2977.	1.4	37
44	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.	2.9	36
45	Induction of metabolic quiescence defines the transitional to follicular B cell switch. Science Signaling, 2019, 12, .	3.6	35
46	Impaired receptor editing and heterozygous RAG2 mutation in a patient with systemic lupus erythematosus and erosive arthritis. Journal of Allergy and Clinical Immunology, 2015, 135, 272-273.	2.9	30
47	Ligase-4 Deficiency Causes Distinctive Immune Abnormalities in Asymptomatic Individuals. Journal of Clinical Immunology, 2016, 36, 341-353.	3.8	30
48	Sesame allergy: Role of specific IgE and skin-prick testing in predicting food challenge results. Allergy and Asthma Proceedings, 2009, 30, 643-648.	2.2	28
49	Abnormalities of T-cell receptor repertoire in CD4+ regulatory and conventional T cells in patients with RAG mutations: Implications for autoimmunity. Journal of Allergy and Clinical Immunology, 2017, 140, 1739-1743.e7.	2.9	28
50	Quantitation of human astrovirus by real-time reverse-transcription-polymerase chain reaction to examine correlation with clinical illness. Journal of Virological Methods, 2006, 134, 190-196.	2.1	27
51	Generalized Bullous Eruption after Routine Vaccination in a Child with Diffuse Cutaneous Mastocytosis. Journal of Allergy and Clinical Immunology: in Practice, 2013, 1, 94-96.	3.8	27
52	Deletion of WASp and N-WASp in B cells cripples the germinal center response and results in production of IgM autoantibodies. Journal of Autoimmunity, 2015, 62, 81-92.	6.5	25
53	Partial RAG deficiency in a patient withÂvaricella infection, autoimmune cytopenia, and anticytokine antibodies. Journal of Allergy and Clinical Immunology: in Practice, 2018, 6, 1769-1771.e2.	3.8	25
54	Rituximab-induced hypogammaglobulinemia and infection risk in pediatric patients. Journal of Allergy and Clinical Immunology, 2021, 148, 523-532.e8.	2.9	24

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55	Case 41-2015: A Boy with Immune and Liver Abnormalities. New England Journal of Medicine, 2016, 374, 2192-2193.	27.0	23
56	Type I interferon, anti-interferon antibodies, and COVID-19. Lancet Rheumatology, The, 2021, 3, e246-e247.	3.9	23
57	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.	3.8	20
58	Secondary Immune Deficiency and Primary Immune Deficiency Crossovers: Hematological Malignancies and Autoimmune Diseases. Frontiers in Immunology, 0, 13, .	4.8	20
59	Unrelated Hematopoietic Cell Transplantation in a Patient with Combined Immunodeficiency with Granulomatous Disease and Autoimmunity Secondary to RAG Deficiency. Journal of Clinical Immunology, 2016, 36, 725-732.	3.8	19
60	Familial Immune Thrombocytopenia Associated With a Novel Variant in IKZF1. Frontiers in Pediatrics, 2019, 7, 139.	1.9	16
61	The Clinical and Genetic Spectrum of 82 Patients With RAG Deficiency Including a c.256_257delAA Founder Variant in Slavic Countries. Frontiers in Immunology, 2020, 11, 900.	4.8	16
62	Antigen-Specific CD4+ T-Cell Activation in Primary Antibody Deficiency After BNT162b2 mRNA COVID-19 Vaccination. Frontiers in Immunology, 2022, 13, 827048.	4.8	16
63	Estimated disease incidence of RAG1/2 mutations: AÂcase report and querying the Exome Aggregation Consortium. Journal of Allergy and Clinical Immunology, 2017, 139, 690-692.e3.	2.9	13
64	When Screening for Severe Combined Immunodeficiency (SCID) with T Cell Receptor Excision Circles Is Not SCID: a Case-Based Review. Journal of Clinical Immunology, 2021, 41, 294-302.	3.8	13
65	Geographical Distribution, Incidence, Malignancies, and Outcome of 136 Eastern Slavic Patients With Nijmegen Breakage Syndrome and NBN Founder Variant c.657_661del5. Frontiers in Immunology, 2020, 11, 602482.	4.8	13
66	Reduced numbers of circulating group 2 innate lymphoid cells in patients with common variable immunodeficiency. European Journal of Immunology, 2017, 47, 1959-1969.	2.9	12
67	Immunoglobulin A Dysgammaglobulinemia Is Associated with Pediatric-Onset Obsessive-Compulsive Disorder. Journal of Child and Adolescent Psychopharmacology, 2019, 29, 268-275.	1.3	12
68	Rituximab and eculizumab when treating nonmalignant hematologic disorders: infection risk, immunization recommendations, and antimicrobial prophylaxis needs. Hematology American Society of Hematology Education Program, 2020, 2020, 312-318.	2.5	12
69	Primary Immunodeficiency in Children With Autoimmune Cytopenias: Retrospective 154-Patient Cohort. Frontiers in Immunology, 2021, 12, 649182.	4.8	12
70	RAG deficiency with ALPS features successfully treated with TCRÎ±Î²/CD19 cell depleted haploidentical stem cell transplant. Clinical Immunology, 2018, 187, 102-103.	3.2	12
71	Arthritis in Two Patients With Partial Recombination Activating Gene Deficiency. Frontiers in Pediatrics, 2019, 7, 235.	1.9	11
72	Combined Immunodeficiency With Late-Onset Progressive Hypogammaglobulinemia and Normal B Cell Count in a Patient With RAG2 Deficiency. Frontiers in Pediatrics, 2019, 7, 122.	1.9	10

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73	Two Unique Cases of X-linked SCID: A Diagnostic Challenge in the Era of Newborn Screening. Frontiers in Pediatrics, 2019, 7, 55.	1.9	10
74	Vasculitis as a Major Morbidity Factor in Patients With Partial RAG Deficiency. Frontiers in Immunology, 2020, 11, 574738.	4.8	10
75	Diffuse cutaneous mastocytosis with novel somatic <scp>KIT</scp> mutation K509I and association with tuberous sclerosis. Clinical Case Reports (discontinued), 2018, 6, 1834-1840.	0.5	9
76	Severe Facial Herpes Vegetans and Viremia in NFKB2-Deficient Common Variable Immunodeficiency. Frontiers in Pediatrics, 2019, 7, 61.	1.9	9
77	Asymptomatic Infant With Atypical SCID and Novel Hypomorphic RAG Variant Identified by Newborn Screening: A Diagnostic and Treatment Dilemma. Frontiers in Immunology, 2020, 11, 1954.	4.8	9
78	Approaches to patients with variants in RAG genes: from diagnosis to timely treatment. Expert Review of Clinical Immunology, 2019, 15, 1033-1046.	3.0	8
79	Functional Confirmation of DNA Repair Defect in Ataxia Telangiectasia (AT) Infants Identified by Newborn Screening for Severe Combined Immunodeficiency (NBS SCID). Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 723-732.e3.	3.8	8
80	CMV-Seropositive Mothers of SCID: To Breastfeed or Not?. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 2866-2867.	3.8	7
81	Intronic SH2D1A mutation with impaired SAP expression and agammaglobulinemia. Clinical Immunology, 2013, 146, 84-89.	3.2	6
82	Gastrointestinal manifestations in common variable immunodeficiency (CVID) are associated with an altered immunophenotype including B- and T-cell dysregulation. Journal of Allergy and Clinical Immunology: in Practice, 2020, 8, 1436-1438.e1.	3.8	6
83	Bilateral Lung Transplantation in a Patient with Humoral Immune Deficiency: A Case Report with Review of the Literature. Case Reports in Immunology, 2014, 2014, 1-7.	0.4	4
84	Case 41-2015. New England Journal of Medicine, 2015, 373, 2664-2676.	27.0	4
85	TREC Screening for WHIM Syndrome. Journal of Clinical Immunology, 2021, 41, 621-628.	3.8	4
86	Antibody Deficiency, Chronic Lung Disease, and Comorbid Conditions: A Case-Based Approach. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 3899-3908.	3.8	4
87	Recurrent disseminated Mycobacterium avium in a female patient from Thailand with anti-interferon-gamma autoantibodies: dilemma on treatment approach. BMJ Case Reports, 2021, 14, e237909.	0.5	4
88	Lymphoma in Partial DiGeorge Syndrome: Report of 2 Cases. Journal of Pediatric Hematology/Oncology, 2022, 44, e819-e822.	0.6	4
89	Adult-Onset Myopathy in a Patient with Hypomorphic RAG2 Mutations and Combined Immune Deficiency. Journal of Clinical Immunology, 2018, 38, 642-645.	3.8	3
90	Predicting the Occurrence of Variants in RAG1 and RAG2. Journal of Clinical Immunology, 2019, 39, 688-701.	3.8	3

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91	Case Report: A Novel Pathogenic Missense Mutation in FAS: A Multi-Generational Case Series of Autoimmune Lymphoproliferative Syndrome. Frontiers in Pediatrics, 2021, 9, 624116.	1.9	3
92	X-Linked Agammaglobulinemia Presenting as Neutropenia: Case Report and an Overview of Literature. Frontiers in Pediatrics, 2021, 9, 633692.	1.9	3
93	Vasculitis as a major morbidity factor in patients with hypomorphic RAG mutations. Journal of Allergy and Clinical Immunology, 2019, 143, AB116.	2.9	2
94	Hyaluronidase-Facilitated High-Dose Subcutaneous IgG Effectively Controls Parvovirus B19 Infection in a Pediatric Cardiac Transplant Patient With Severe T-Cell Lymphopenia. Open Forum Infectious Diseases, 2020, 7, ofaa076.	0.9	2
95	IN TIME: IMPORTÃ‚NCIA E IMPLICAÃ‡Ã•ES GLOBAIS DATRIAGEM NEONATAL PARA A IMUNODEFICIÃŠNCIA GRAVE COMBINADA. Revista Paulista De Pediatria, 2018, 36, 388-397.	1.0	2
96	The new quest in CTLA-4 insufficiency: How to immune modulate effectively?. Journal of Allergy and Clinical Immunology, 2022, 149, 543-546.	2.9	2
97	NaÃ¯ve B cells are prone to develop into polyreactive autoantibody secreting cells from adult RAG2-deficient patient with combined immunodeficiency. Journal of Allergy and Clinical Immunology, 2018, 141, AB22.	2.9	1
98	The WHIM Syndrome Is No Longer a Whim. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 1578-1579.	3.8	1
99	Introducing a New Epoch in Inborn Errors of Immunity. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 660-662.	3.8	1
100	Management and Outcomes of Immune Cytopenia Following Pediatric Heart Transplantation. Trends in Transplantation, 2021, 14, .	0.2	1
101	The Evolution of Very Early Onset Inflammatory Bowel Disease, Autoimmune Hepatitis, and Primary Sclerosing Cholangitis in a Young Girl. Case Reports in Gastroenterology, 2021, 15, 939-947.	0.6	1
102	Prevalence of Anti-lymphocyte Antibodies in Patients with Good Syndrome and Partial RAG Deficiency. Journal of Allergy and Clinical Immunology, 2022, 149, AB25.	2.9	1
103	HSCT using carrier donors for CD40L deficiency results in excellent immune function and higher CD40L expression in cTfh. Blood Advances, 2022, , .	5.2	1
104	Subcutaneous Nodule in a Young Girl. Clinical Pediatrics, 2006, 45, 661-664.	0.8	0
105	Tolerance of Baked Milk in a Subset of Patients with Cow's Milk-Mediated Eosinophilic Esophagitis. Journal of Allergy and Clinical Immunology, 2013, 131, AB181.	2.9	0
106	Evaluation of a Novel Missense Activation-Induced Deaminase AID Mutation in a Child with Hyper IgM Syndrome: Is it a Pathogenic Mutation?. Journal of Allergy and Clinical Immunology, 2014, 133, AB70.	2.9	0
107	An Atypical Severe Combined Immunodeficiency (SCID) Case Diagnosis Complicated by Alternative Care in the Era of Newborn Screening (NBS) for SCID. Journal of Allergy and Clinical Immunology, 2017, 139, AB18.	2.9	0
108	P508 Increased risk for hematologic and specific solid organ malignancy in common variable immunodeficiency (CVID) patients. Annals of Allergy, Asthma and Immunology, 2017, 119, e4.	1.0	0

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109	Gastrointestinal (GI) Manifestations in Common Variable Immunodeficiency (CVID). Journal of Allergy and Clinical Immunology, 2018, 141, AB82.	2.9	0
110	Rituximab Use and Immunological Monitoring in Pediatric Patients. Journal of Allergy and Clinical Immunology, 2019, 143, AB205.	2.9	0
111	174.â€ƒVASCULITIS AS A MAJOR MORBIDITY FACTOR IN PATIENTS WITH HYPOMORPHIC RAG VARIANTS. Rheumatology, 2019, 58, .	1.9	0
112	Treatment Challenges of Refractory Thrombocytopenia in DiGeorge Syndrome. Journal of Allergy and Clinical Immunology, 2019, 143, AB112.	2.9	0
113	Early Diagnosis of Severe Combined Immunodeficiency. , 2019, , 173-193.		0
114	Editorial: Screening for Primary Immunodeficiency Disorders (PIDDs) in Neonates. Frontiers in Immunology, 2020, 11, 633266.	4.8	0
115	Clinical and Treatment History of Patients with Partial DiGeorge Syndrome and Autoimmune Cytopenia. Journal of Allergy and Clinical Immunology, 2021, 147, AB69.	2.9	0
116	Are we diagnosing too late? RAG deficiency in young adults with end organ damage. Journal of Allergy and Clinical Immunology, 2021, 147, AB69.	2.9	0
117	Diagnostic Dilemma on Novel Pathogenic Variant of Cytotoxic T-Lymphocyte Associated Protein 4 (CTLA-4) in a Family with Chronic ITP and Immune Dysregulation. Journal of Allergy and Clinical Immunology, 2021, 147, AB70.	2.9	0
118	Clinical Characteristics of SARS-CoV2 Infected and Exposed Patients at a Tertiary Care Allergy/Immunology Program in Florida. Journal of Allergy and Clinical Immunology, 2021, 147, AB78.	2.9	0
119	Florida Pediatric Bone Marrow Transplant and Cell Therapy Consortium (FPBCC) Outcomes of Children with Primary Immunodeficiency Disorders Following Allogeneic Hematopoietic Cell Transplantation. Transplantation and Cellular Therapy, 2021, 27, S319-S320.	1.2	0
120	Outcomes of T- and B-Cell Acute Lymphoblastic Leukemias Post-Allogeneic Transplant (alloHCT) Using Enhanced Data Back to Center (eDBtC) Platform: Experience from the Florida Pediatric Bone Marrow Transplant and Cell Therapy Consortium (FPBCC). Transplantation and Cellular Therapy, 2021, 27, S135-S136.	1.2	0
121	Short Telomere Syndromes. , 2019, , 1-3.		0
122	Characterizing Autoimmune Hemolytic Anemia in RAG Deficiency. Blood, 2019, 134, 3508-3508.	1.4	0
123	Identifying Primary Immune Deficiencies in Patients with Autoimmune Cytopenias. Blood, 2019, 134, 2328-2328.	1.4	0
124	Short Telomere Syndromes. , 2020, , 590-592.		0
125	Characterization of a Novel Missense CXCR4 Mutation in a Patient with WHIM-like Syndrome. Blood, 2021, 138, 4309-4309.	1.4	0
126	Diagnostic and Therapeutic Challenges for Infants with Radiosensitive (RS)-SCID Identified by Newborn Screening (NBS). Journal of Allergy and Clinical Immunology, 2022, 149, AB25.	2.9	0

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127	T Cell Responses to SARS-CoV-2 Vaccination and Infection in Antibody Deficiency Diseases. Journal of Allergy and Clinical Immunology, 2022, 149, AB66.	2.9	0
128	The Humoral Immune Response To SARS-CoV-2 Infection And/or Immunization in Immunocompromised Versus Immunocompetent Individuals. Journal of Allergy and Clinical Immunology, 2022, 149, AB21.	2.9	0
129	Effects Of The COVID-19 Pandemic On A Group Of Patients With Pathogenic Variant of Cytotoxic T-Lymphocyte Associated Protein 4 (CTLA-4) in a Tertiary Center in Florida. Journal of Allergy and Clinical Immunology, 2022, 149, AB28.	2.9	0
130	Effects Of Mavorixafor On Functional Impairments Due To A Novel Missense CXCR4 Mutation In A Patient With WHIM-Like Syndrome. Journal of Allergy and Clinical Immunology, 2022, 149, AB20.	2.9	0
131	Autoantibodies in immunodeficiency syndromes: The Janus faces of immune dysregulation. Blood Reviews, 2022, , 100948.	5.7	0