Raz Somech

List of Publications by Year
in descending order

165 papers	4,866 citations	109321 35 h-index	123424 61 g-index
172 all docs	172 docs citations	172 times ranked	7202 citing authors

#	Article	IF	Citations
1	Interleukin-10 Receptor Signaling in Innate Immune Cells Regulates Mucosal Immune Tolerance and Anti-Inflammatory Macrophage Function. Immunity, 2014, 40, 706-719.	14.3	455
2	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
3	The nuclear-envelope protein and transcriptional repressor LAP2Î² interacts with HDAC3 at the nuclear periphery, and induces histone H4 deacetylation. Journal of Cell Science, 2005, 118, 4017-4025.	2.0	189
4	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
5	Timely and spatially regulated maturation of B and T cell repertoire during human fetal development. Science Translational Medicine, 2015, 7, 276ra25.	12.4	148
6	Hematologically important mutations: Leukocyte adhesion deficiency (first update). Blood Cells, Molecules, and Diseases, 2012, 48, 53-61.	1.4	147
7	A Congenital Neutrophil Defect Syndrome Associated with Mutations inVPS45. New England Journal of Medicine, 2013, 369, 54-65.	27.0	122
8	Defining combined immunodeficiency. Journal of Allergy and Clinical Immunology, 2012, 130, 177-183.	2.9	104
9	Mutations in STN1 cause Coats plus syndrome and are associated with genomic and telomere defects. Journal of Experimental Medicine, 2016, 213, 1429-1440.	8.5	100
10	T-cell defects in patients with ARPC1B germline mutations account for combined immunodeficiency. Blood, 2018, 132, 2362-2374.	1.4	99
11	Histone deacetylase inhibitors â€“ a new tool to treat cancer. Cancer Treatment Reviews, 2004, 30, 461-472.	7.7	97
12	Chronic granulomatous disease: Clinical, functional, molecular, and genetic studies. The Israeli experience with 84 patients. American Journal of Hematology, 2017, 92, 28-36.	4.1	93
13	Intestinal Inflammation and Dysregulated Immunity in Patients With Inherited Caspase-8 Deficiency. Gastroenterology, 2019, 156, 275-278.	1.3	92
14	Characterization of T and B cell repertoire diversity in patients with RAG deficiency. Science Immunology, 2016, 1, .	11.9	88
15	Disruption of Thrombocyte and T Lymphocyte Development by a Mutation in ARPC1B. Journal of Immunology, 2017, 199, 4036-4045.	0.8	72
16	First Year of Israeli Newborn Screening for Severe Combined Immunodeficiencyâ€"Clinical Achievements and Insights. Frontiers in Immunology, 2017, 8, 1448.	4.8	67
17	Herpes simplex encephalitis in a patient with a distinctive form of inherited IFNAR1 deficiency. Journal of Clinical Investigation, 2021, 131, .	8.2	64
18	Nuclear Envelopathiesâ€"Raising the Nuclear Veil. Pediatric Research, 2005, 57, 8R-15R.	2.3	60

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19	Leucocyte adhesion deficiencyâ€"A multicentre national experience. European Journal of Clinical Investigation, 2019, 49, e13047.	3.4	54
20	Characterization of Î¶-associated protein, 70 kd (ZAP70)â€"deficient human lymphocytes. Journal of Allergy and Clinical Immunology, 2010, 126, 1226-1233.e1.	2.9	52
21	CD137 deficiency causes immune dysregulation with predisposition to lymphomagenesis. Blood, 2019, 134, 1510-1516.	1.4	52
22	Proteome Analysis of Human Neutrophil Granulocytes From Patients With Monogenic Disease Using Data-independent Acquisition. Molecular and Cellular Proteomics, 2019, 18, 760-772.	3.8	52
23	Reduced central tolerance in Omenn syndrome leads to immature self-reactive oligoclonal T cells. Journal of Allergy and Clinical Immunology, 2009, 124, 793-800.	2.9	51
24	Patients with Primary Immunodeficiencies Are a Reservoir of Poliovirus and a Risk to Polio Eradication. Frontiers in Immunology, 2017, 8, 685.	4.8	50
25	Phase 1/2 study of nilotinib prophylaxis after allogeneic stem cell transplantation in patients with advanced chronic myeloid leukemia or <scp>P</scp>hiladelphia chromosomeâ€"positive acute lymphoblastic leukemia. Cancer, 2015, 121, 863-871.	4.1	48
26	Microarray-based gene expression profiling of hematologic malignancies: basic concepts and clinical applications. Blood Reviews, 2005, 19, 223-234.	5.7	47
27	Changes in Routine Pediatric Practice in Light of Coronavirus 2019 (COVID-19). Journal of Pediatrics, 2020, 224, 190-193.	1.8	46
28	Disturbed B and T cell homeostasis and neogenesis in patients with ataxia telangiectasia. Journal of Clinical Immunology, 2014, 34, 561-572.	3.8	45
29	Newborn screening for severe T and B cell immunodeficiency in Israel: a pilot study. Israel Medical Association Journal, 2013, 15, 404-9.	0.1	45
30	The Kinetics of Early T and B Cell Immune Recovery after Bone Marrow Transplantation in RAG-2-Deficient SCID Patients. PLoS ONE, 2012, 7, e30494.	2.5	44
31	T-cell receptor excision circles in primary immunodeficiencies and other T-cell immune disorders. Current Opinion in Allergy and Clinical Immunology, 2011, 11, 517-524.	2.3	42
32	Novel MALT1 Mutation Linked to Immunodeficiency, Immune Dysregulation, and an Abnormal T Cell Receptor Repertoire. Journal of Clinical Immunology, 2019, 39, 401-413.	3.8	42
33	Thymic function in MHC class IIâ€"deficient patients. Journal of Allergy and Clinical Immunology, 2013, 131, 831-839.	2.9	41
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	Complications of Minocycline Therapy for Acne Vulgaris: Case Reports and Review of theâ€ƒLiterature. Pediatric Dermatology, 1999, 16, 469-472.	0.9	38
36	Matrix Metalloproteinases 2 and 9 Are Markers of Inflammation but Not of the Degree of Fibrosis in Chronic Hepatitis C. Digestion, 2005, 71, 124-130.	2.3	37

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37	Novel Mutations in RASGRP1 are Associated with Immunodeficiency, Immune Dysregulation, and EBV-Induced Lymphoma. Journal of Clinical Immunology, 2018, 38, 699-710.	3.8	37
38	Purine nucleoside phosphorylase deficiency presenting as severe combined immune deficiency. Immunologic Research, 2013, 56, 150-154.	2.9	35
39	Bone marrow transplantation for cartilage-hair-hypoplasia. Bone Marrow Transplantation, 2006, 38, 751-756.	2.4	34
40	Thymic involution, a coâ€morbidity factor in amyotrophic lateral sclerosis. Journal of Cellular and Molecular Medicine, 2010, 14, 2470-2482.	3.6	34
41	Whole exome sequencing in childhood-onset lupus frequently detects single gene etiologies. Pediatric Rheumatology, 2019, 17, 52.	2.1	34
42	Mutation Analysis should be Performed to Rule Out Î³c Deficiency in Children with Functional Severe Combined Immune Deficiency Despite Apparently Normal Immunologic Tests. Journal of Pediatrics, 2005, 147, 555-557.	1.8	33
43	Novel mutations in RAG1/2 and ADA genes in Israeli patients presenting with T-B- SCID or Omenn syndrome. Clinical Immunology, 2011, 140, 284-290.	3.2	32
44	Cysteine and hydrophobic residues in CDR3 serve as distinct T-cell self-reactivity indices. Journal of Allergy and Clinical Immunology, 2019, 144, 333-336.	2.9	31
45	Molecular Assessment of Thymus Capabilities in the Evaluation of T-Cell Immunodeficiency. Pediatric Research, 2010, 67, 211-216.	2.3	29
46	Liver Disease in Pediatric Patients With Ataxia Telangiectasia. Journal of Pediatric Gastroenterology and Nutrition, 2016, 62, 550-555.	1.8	28
47	Highlighting the problematic reliance on CD18 for diagnosing leukocyte adhesion deficiency type 1. Immunologic Research, 2016, 64, 476-482.	2.9	23
48	T and B cell clonal expansion in Ras-associated lymphoproliferative disease (RALD) as revealed by next-generation sequencing. Clinical and Experimental Immunology, 2017, 189, 310-317.	2.6	23
49	Human FCHO1 deficiency reveals role for clathrin-mediated endocytosis in development and function of T cells. Nature Communications, 2020, 11, 1031.	12.8	23
50	Reversible airway obstruction in children with ataxia telangiectasia. Pediatric Pulmonology, 2010, 45, 230-235.	2.0	22
51	Whole exome sequencing (WES) approach for diagnosing primary immunodeficiencies (PIDs) in a highly consanguineous community. Clinical Immunology, 2020, 214, 108376.	3.2	22
52	Enhanced expression of the nuclear envelope LAP2 transcriptional repressors in normal and malignant activated lymphocytes. Annals of Hematology, 2007, 86, 393-401.	1.8	21
53	Matched unrelated bone marrow transplant for T+ combined immunodeficiency. Bone Marrow Transplantation, 2008, 41, 947-952.	2.4	21
54	Evaluation of Immediate Allergic Reactions to Cephalosporins in Non-Penicillin-Allergic Patients. International Archives of Allergy and Immunology, 2009, 150, 205-209.	2.1	21

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55	Polyethylene glycolâ€“modified adenosine deaminase improved lung disease but not liverÂdisease in partial adenosine deaminase deficiency. Journal of Allergy and Clinical Immunology, 2009, 124, 848-850.	2.9	21
56	Survival of the fetus: fetal B and T cell receptor repertoire development. Seminars in Immunopathology, 2017, 39, 577-583.	6.1	21
57	Altered T cell receptor beta repertoire patterns in pediatric ulcerative colitis. Clinical and Experimental Immunology, 2019, 196, 1-11.	2.6	21
58	A Large Cohort of RAG1/2-Deficient SCID Patientsâ€”Clinical, Immunological, and Prognostic Analysis. Journal of Clinical Immunology, 2020, 40, 211-222.	3.8	20
59	Inherited SLP76 deficiency in humans causes severe combined immunodeficiency, neutrophil and platelet defects. Journal of Experimental Medicine, 2021, 218, .	8.5	20
60	Characterizing T Cells in SCID Patients Presenting with Reactive or Residual T Lymphocytes. Clinical and Developmental Immunology, 2012, 2012, 1-9.	3.3	18
61	Thymic and bone marrow output in individuals with 22q11.2 deletion syndrome. Pediatric Research, 2015, 77, 579-585.	2.3	18
62	G23D: Online tool for mapping and visualization of genomic variants on 3D protein structures. BMC Genomics, 2016, 17, 681.	2.8	18
63	Newborn Screening for Severe Combined Immunodeficiency in Israel. International Journal of Neonatal Screening, 2017, 3, 13.	3.2	18
64	Epidemiologic, Socioeconomic, and Clinical Factors Associated with Severity of Respiratory Syncytial Virus Infection in Previously Healthy Infants. Clinical Pediatrics, 2006, 45, 621-627.	0.8	17
65	T-Cell Compartment in Synovial Fluid of Pediatric Patients with JIA Correlates with Disease Phenotype. Journal of Clinical Immunology, 2011, 31, 1021-1028.	3.8	17
66	Severe congenital neutropenia with neurological impairment due to a homozygous VPS45 p.E238K mutation: A case report suggesting a genotypeâ€“phenotype correlation. American Journal of Medical Genetics, Part A, 2015, 167, 3214-3218.	1.2	17
67	Correlation between â€˜<scp>ACKR</scp>1/<scp>DARC</scp> nullâ€™ polymorphism and benign neutropenia in Yemenite Jews. British Journal of Haematology, 2015, 170, 892-895.	2.5	17
68	Treatment options for DOCK8 deficiencyâ€•related severe dermatitis. Journal of Dermatology, 2021, 48, 1386-1393.	1.2	17
69	The Effect of Gentamicin-Induced Readthrough on a Novel Premature Termination Codon of CD18 Leukocyte Adhesion Deficiency Patients. PLoS ONE, 2010, 5, e13659.	2.5	17
70	The role of hematopoietic stem cell transplantation in <scp>SP</scp>110 associated venoâ€•occlusive disease with immunodeficiency syndrome. Pediatric Allergy and Immunology, 2013, 24, 250-256.	2.6	16
71	Impact of Conditioning on Outcome of Hematopoietic Stem Cell Transplantation for Wiskott-Aldrich Syndrome. Journal of Pediatric Hematology/Oncology, 2013, 35, e234-e238.	0.6	16
72	Cytomegalovirus Retinitis in HIV-Negative Patients: A Practical Management Approach. Ophthalmology, 2015, 122, 866-868.e3.	5.2	16

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73	T+ NK+ IL-2 Receptor Î³ Chain Mutation: a Challenging Diagnosis of Atypical Severe Combined Immunodeficiency. Journal of Clinical Immunology, 2018, 38, 527-536.	3.8	16
74	Reduced Function and Diversity of T Cell Repertoire and Distinct Clinical Course in Patients With IL7RA Mutation. Frontiers in Immunology, 2019, 10, 1672.	4.8	16
75	Focal Nodular Hyperplasia in Children. Journal of Pediatric Gastroenterology and Nutrition, 2001, 32, 480-483.	1.8	15
76	Pseudotumor Cerebri After Allogeneic Bone Marrow Transplant Associated With Cyclosporine A Use for Graft-Versus-Host Disease Prophylaxis. Journal of Pediatric Hematology/Oncology, 2007, 29, 66-68.	0.6	15
77	Growth characteristics and endocrine abnormalities in 22q11.2 deletion syndrome. American Journal of Medical Genetics, Part A, 2017, 173, 1301-1308.	1.2	15
78	Elevated IgM levels as a marker for a unique phenotype in patients with Ataxia telangiectasia. BMC Pediatrics, 2018, 18, 185.	1.7	15
79	Lessons Learned From Five Years of Newborn Screening for Severe Combined Immunodeficiency in Israel. Journal of Allergy and Clinical Immunology: in Practice, 2022, 10, 2722-2731.e9.	3.8	15
80	T- and B-cell defects in a novel purine nucleoside phosphorylase mutation. Journal of Allergy and Clinical Immunology, 2012, 130, 539-542.	2.9	14
81	Exome sequencing as a differential diagnosis tool: resolving mild trichohepatoenteric syndrome. Clinical Genetics, 2015, 87, 602-603.	2.0	14
82	Gastrointestinal Hemorrhage: A Manifestation of the Telomere Biology Disorders. Journal of Pediatrics, 2021, 230, 55-61.e4.	1.8	14
83	Cellular and humoral aberrations in a kindred with IL-1 receptorâ€“associated kinase 4 deficiency. Journal of Allergy and Clinical Immunology, 2007, 120, 948-950.	2.9	13
84	Mammalian VPS45 orchestrates trafficking through the endosomal system. Blood, 2021, 137, 1932-1944.	1.4	13
85	Genetic predisposition to infectious pathogens: a review of less familiar variants. Pediatric Infectious Disease Journal, 2003, 22, 457-461.	2.0	12
86	Fatal lung fibrosis associated with immunodeficiency and gonadal dysgenesis in 46XX sistersâ€”A new syndrome. American Journal of Medical Genetics, Part A, 2008, 146A, 8-14.	1.2	12
87	A Call to Include Severe Combined Immunodeficiency in Newborn Screening Program. Rambam Maimonides Medical Journal, 2014, 5, e0001.	1.0	12
88	Novel SMARCAL1 Bi-allelic Mutations Associated with a Chromosomal Breakage Phenotype in a Severe SIOD Patient. Journal of Clinical Immunology, 2014, 34, 76-83.	3.8	12
89	A Novel Mutation in a Critical Region for the Methyl Donor Binding in DNMT3B Causes Immunodeficiency, Centromeric Instability, and Facial Anomalies Syndrome (ICF). Journal of Clinical Immunology, 2016, 36, 801-809.	3.8	12
90	The clinical and laboratory spectrum of dedicator of cytokinesis 8 immunodeficiency syndrome in patients with a unique mutation. Immunologic Research, 2017, 65, 651-657.	2.9	12

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91	Congenital neutropenia with variable clinical presentation in novel mutation of the SRP54 gene. Pediatric Blood and Cancer, 2020, 67, e28237.	1.5	12
92	Clinical Features in a Large Cohort of Patients With 22q11.2 Deletion Syndrome. Journal of Pediatrics, 2021, 238, 215-220.e5.	1.8	12
93	LAP2Î¶ binds BAF and suppresses LAP2Î²-mediated transcriptional repression. European Journal of Cell Biology, 2008, 87, 267-278.	3.6	11
94	Molecular assessment of thymic capacities in patients with Schimke immuno-osseous dysplasia. Clinical Immunology, 2009, 133, 375-381.	3.2	11
95	Thymic functions and gene expression profile distinct double-negative cells from single positive cells in the autoimmune lymphoproliferative syndrome. Autoimmunity Reviews, 2012, 11, 723-730.	5.8	11
96	Genetic and Structural Analysis of a SKIV2L Mutation Causing Tricho-hepato-enteric Syndrome. Digestive Diseases and Sciences, 2018, 63, 1192-1199.	2.3	11
97	Bacille Calmette-Guerin (BCG) complications in children with severe combined immunodeficiency (SCID). Infectious Diseases, 2019, 51, 585-592.	2.8	11
98	Alterations in T and B Cell Receptor Repertoires Patterns in Patients With IL10 Signaling Defects and History of Infantile-Onset IBD. Frontiers in Immunology, 2020, 11, 109.	4.8	11
99	Clinical characteristics of children with 2009 pandemic H1N1 influenza virus infections. Pediatrics International, 2011, 53, 426-430.	0.5	10
100	Quantification of specific T and B cells immunological markers in children with chronic and transient ITP. Pediatric Blood and Cancer, 2017, 64, e26646.	1.5	10
101	Long-term nutritional and gastrointestinal aspects in patients with ataxia telangiectasia. Nutrition, 2018, 46, 48-52.	2.4	10
102	Maturation of the immune system in the fetus and the implications for congenital CMV. Best Practice and Research in Clinical Obstetrics and Gynaecology, 2019, 60, 35-41.	2.8	10
103	An RTEL1 Mutation Links to Infantile-Onset Ulcerative Colitis and Severe Immunodeficiency. Journal of Clinical Immunology, 2020, 40, 1010-1019.	3.8	10
104	High-dose methylprednisolone is effective in the management of acute graft-versus-host disease in severe combined immune deficiency. Journal of Allergy and Clinical Immunology, 2008, 122, 1215-1216.	2.9	9
105	Co-existence of clonal expanded autologous and transplacental-acquired maternal T cells in recombination activating gene-deficient severe combined immunodeficiency. Clinical and Experimental Immunology, 2014, 176, 380-386.	2.6	9
106	Post-childhood Presentation and Diagnosis of DiGeorge Syndrome. Clinical Pediatrics, 2016, 55, 368-373.	0.8	9
107	Late diagnosis of chronic granulomatous disease. Clinical and Experimental Immunology, 2020, 201, 297-305.	2.6	9
108	Procalcitonin correlates with C-reactive protein as an acute-phase reactant in pediatric patients. Israel Medical Association Journal, 2000, 2, 147-50.	0.1	9

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109	Thymus Activity, Vitamin D, and Respiratory Infections in Adolescent Swimmers. Israel Medical Association Journal, 2015, 17, 571-5.	0.1	9
110	Granulomatosis Cheilitis and Crohn Disease. Journal of Pediatric Gastroenterology and Nutrition, 2001, 32, 339-341.	1.8	8
111	Specific self-antigen-driven immune response in pericardial effusion as an isolated GVHD manifestation. Bone Marrow Transplantation, 2010, 45, 1084-1087.	2.4	8
112	Severe Eosinophilia in Children. Journal of Pediatric Hematology/Oncology, 2013, 35, 303-306.	0.6	8
113	MHC II deficient infant identified by newborn screening program for SCID. Immunologic Research, 2018, 66, 537-542.	2.9	8
114	Co-appearance of OPV and BCG vaccine-derived complications in two infants with severe combined immunodeficiency. Immunologic Research, 2018, 66, 437-443.	2.9	8
115	Fetuin-A deficiency is associated with infantile cortical hyperostosis (Caffey disease). Pediatric Research, 2019, 86, 603-607.	2.3	8
116	Neutrophil Functions in Immunodeficiency Due to DOCK8 Deficiency. Immunological Investigations, 2019, 48, 431-439.	2.0	8
117	First report of a persistent oropharyngeal infection of type 2 vaccine-derived poliovirus (iVDPV2) in a primary immune deficient (PID) patient after eradication of wild type 2 poliovirus. International Journal of Infectious Diseases, 2019, 83, 40-43.	3.3	8
118	Chronic demodicosis in patients with immune dysregulation: An unexpected infectious manifestation of Signal transducer and activator of transcription (STAT)1 gainâ€ofâ€function. Clinical and Experimental Immunology, 2021, 206, 56-67.	2.6	8
119	Immunological effects of nilotinib prophylaxis after allogeneic stem cell transplantation in patients with advanced chronic myeloid leukemia or philadelphia chromosome-positive acute lymphoblastic leukemia. Oncotarget, 2017, 8, 418-429.	1.8	8
120	The Clinician Scientist, a Distinct and Disappearing Entity. Journal of Pediatrics, 2019, 212, 252-253.e2.	1.8	7
121	Exogenous interleukin-2 can rescue in-vitro T cell activation and proliferation in patients with a novel capping protein regulator and myosin 1 linker 2 mutation. Clinical and Experimental Immunology, 2020, 200, 215-227.	2.6	7
122	Unusual phenotype in patients with a hypomorphic mutation in the DCLRE1C gene: IgG hypergammaglobulinemia with IgA and IgE deficiency. Clinical Immunology, 2020, 213, 108366.	3.2	7
123	Celiac disease: extraintestinal manifestations, associated diseases, and complications. Advances in Pediatrics, 2002, 49, 191-201.	1.4	7
124	Leptin and C-reactive protein levels correlate during minor infection in children. Israel Medical Association Journal, 2007, 9, 76-8.	0.1	7
125	Insight into normal thymic activity by assessment of peripheral blood samples. Immunologic Research, 2015, 61, 198-205.	2.9	6
126	Monogenic Inflammatory Bowel Disease: It's Never Too Late to Make a Diagnosis. Frontiers in Immunology, 2020, 11, 1775.	4.8	6

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127	Selective clinical and immune response of the oligoclonal autoreactive T cells in Omenn patients after cyclosporin A treatment. Clinical and Experimental Immunology, 2012, 167, 338-345.	2.6	5
128	Hypoparathyroidism and central diabetes insipidus: in search of the link. European Journal of Pediatrics, 2014, 173, 1731-1734.	2.7	5
129	Combined immunodeficiency in a patient with mosaic monosomy 21. Immunologic Research, 2016, 64, 841-847.	2.9	5
130	Foscarnet-related Hypercalcemia During CMV Treatment in an Infant With SCID: A Case Report and Review of Literature. Journal of Pediatric Hematology/Oncology, 2017, 39, e173-e175.	0.6	5
131	Immune reconstitution after HSCT in SCIDâ€"a cohort of conditioned and unconditioned patients. Immunologic Research, 2019, 67, 166-175.	2.9	5
132	New Instrument for the Evaluation of Prodromes and Attacks of Hereditary Angioedema (HAE-EPA). Clinical Reviews in Allergy and Immunology, 2021, 61, 29-39.	6.5	5
133	EVALUATION AND MANAGEMENT OF PEDIATRIC PATIENTS WITH ANAPHYLACTOID REACTIONS TO DEFEROXAMINE MESYLATE. Annals of Allergy, Asthma and Immunology, 2007, 99, 575-576.	1.0	4
134	Testicular failure in a patient with G6PC3 deficiency. Pediatric Research, 2014, 76, 197-201.	2.3	4
135	Analysis of Chronic Granulomatous Disease in the Kavkazi Population in Israel Reveals Phenotypic Heterogeneity in Patients with the Same NCF1 mutation (c.579G>A). Journal of Clinical Immunology, 2018, 38, 193-203.	3.8	4
136	Immune and TRG repertoire signature of the thymus in Down syndrome patients. Pediatric Research, 2021, 89, 102-109.	2.3	4
137	Enhanced Collagen Deposition in the Duodenum of Patients with Hyaline Fibromatosis Syndrome and Protein Losing Enteropathy. International Journal of Molecular Sciences, 2020, 21, 8200.	4.1	3
138	Exploring genetic defects in adults who were clinically diagnosed as severe combined immune deficiency during infancy. Immunologic Research, 2021, 69, 145-152.	2.9	3
139	Kidney and urinary tract findings among patients with Kabuki (make-up) syndrome. Pediatric Nephrology, 2021, 36, 4009-4012.	1.7	3
140	Atypical immune phenotype in severe combined immunodeficiency patients with novel mutations in IL2RG and JAK3. Genes and Immunity, 2020, 21, 326-334.	4.1	2
141	Underperformed and Underreported Testing for Persistent Oropharyngeal Poliovirus Infections in Primary Immune Deficient Patientsâ€"Risk for Reemergence of Polioviruses. Journal of the Pediatric Infectious Diseases Society, 2021, 10, 326-333.	1.3	2
142	Pediatric literature trends: high-level analysis using text-mining. Pediatric Research, 2021, 90, 212-215.	2.3	2
143	Ophthalmic manifestations in Kabuki (make-up) syndrome: A single-center pediatric cohort and systematic review of the literature. European Journal of Medical Genetics, 2021, 64, 104210.	1.3	2
144	Maintenance Therapy with Nilotinib (Tasigna) Post Allogeneic Stem Cell Transplantation (AlloSCT) for Advanced (>CP1) Chronic Myeloid Leukemia (CML) and Ph+ Acute Lymphoblastic Leukemia (ALL). Blood, 2011, 118, 2011-2011.	1.4	2

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145	Self-reactive and transplacental-acquired maternal T cells in SCID patientsâ€"time to update. LymphoSign Journal, 2015, 2, 47-52.	0.2	2
146	B cell repertoire in patients with a novel BTK mutation: expanding the spectrum of atypical X-linked agammaglobulinemia. Immunologic Research, 2022, 70, 216-223.	2.9	2
147	Prodromes as predictors of hereditary angioedema attacks. Allergy: European Journal of Allergy and Clinical Immunology, 2022, 77, 1309-1312.	5.7	2
148	Immunologic Heterogeneity in 2 Cartilage-Hair Hypoplasia Patients With a Distinct Clinical Course. Journal of Investigational Allergology and Clinical Immunology, 2023, 33, 263-270.	1.3	2
149	Eruption of urticaria and angioedema induced by binging and purging in an anorexia nervosa patient. International Journal of Eating Disorders, 2016, 49, 822-825.	4.0	1
150	Trough Concentrations of Specific Antibodies in Primary Immunodeficiency Patients Receiving Intravenous Immunoglobulin Replacement Therapy. Journal of Clinical Medicine, 2021, 10, 592.	2.4	1
151	Nilotinib Treatment Post - Allogeneic Stem Cell Transplantation (alloSCT) in Advanced (>CP1) Chronic Myeloid Leukemia (CML) and Ph+ Acute Lymphoblastic Leukemia (ALL).. Blood, 2009, 114, 1176-1176.	1.4	1
152	Genetic workup as a complementary tool for the diagnosis of primary complement component deficiencies: a multicenter experience. European Journal of Pediatrics, 2022, 181, 1997-2004.	2.7	1
153	Glycogen Storage Disease type IA refractory to cornstarch: Can next generation sequencing offer a solution?. European Journal of Medical Genetics, 2022, , 104518.	1.3	1
154	Sometimes double negative is positive. Arthritis Care and Research, 2013, 65, 161-168.	3.4	0
155	Severe Prolonged Hypothyroidism. Global Pediatric Health, 2015, 2, 2333794X1557467.	0.7	0
156	Autoimmunity and Primary Immunodeficiency. , 2019, , 675-683.		0
157	Reply. Journal of Pediatrics, 2020, 217, 220.	1.8	0
158	A novel zeta-associated protein 70 homozygous mutation causing combined immunodeficiency presenting as neonatal autoimmune hemolytic anemia. Immunologic Research, 2021, 69, 100-106.	2.9	0
159	Assessment of the Effect of Nilotinib (Tasigna) Maintenance Therapy After Allogeneic Stem Cell Transplantation in Patients with Advanced CML and Ph+ ALL On Immune Reconstitution and Lymphocyte Function. Blood, 2012, 120, 4478-4478.	1.4	0
160	Novel Immune Checkpoint Deficiency and Susceptibility to EBV-Associated Lymphoma. Blood, 2019, 134, 2326-2326.	1.4	0
161	First generation cephalosporins as therapy for uncomplicated pyelonephritis in children. A retrospective analysis. Journal of Medicine, 2000, 31, 195-203.	0.1	0
162	Acute encephalopathy preceding Shigella infection. Israel Medical Association Journal, 2001, 3, 384-5.	0.1	0

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163	Uncommon presentation of some common pediatric diseases. Advances in Pediatrics, 2003, 50, 269-304.	1.4	0
164	"The Girl who Grew Horns": Temporal Swelling as an Atypical Presenting Symptom of Epstein-Barr Virus Infection. Israel Medical Association Journal, 2016, 18, 761-762.	0.1	0
165	Combined Gastric and Pancreatic Tissue Inside a Meckel's Diverticulum. Israel Medical Association Journal, 2018, 20, 461-462.	0.1	0