Primary immunodeficiency diseases: An update from th Immunological Societies Primary Immunodeficiency Di

Journal of Allergy and Clinical Immunology

120, 776-794

DOI: 10.1016/j.jaci.2007.08.053

Citation Report

#	Article	IF	Citations
1	Human Primary Immunodeficiency Diseases. Immunity, 2007, 27, 835-845.	14.3	125
2	Is Helicobacter pylori responsible for autoimmune diseases? that is the question. Allergologia Et Immunopathologia, 2007, 35, 221-224.	1.7	0
3	Primary immunodeficiency: Meeting the challenges. Journal of Allergy and Clinical Immunology, 2007, 120, 753-755.	2.9	1
4	Primary Immunodeficiencies Unravel Critical Aspects of the Pathophysiology of Autoimmunity and of the Genetics of Autoimmune Disease. Journal of Clinical Immunology, 2008, 28, 4-10.	3.8	31
5	Autoimmunity in Common Variable Immunodeficiency. Journal of Clinical Immunology, 2008, 28, 46-55.	3.8	39
6	Understanding Systemic Lupus Erythematosus Physiopathology in the Light of Primary Immunodeficiencies. Journal of Clinical Immunology, 2008, 28, 34-41.	3.8	73
7	Leukopenie. Wiener Klinische Wochenschrift Education, 2008, 3, 161-178.	0.0	1
8	Antibody deficiency diseases. European Journal of Immunology, 2008, 38, 327-333.	2.9	63
9	Immunodeficiency-associated lymphomas. Blood Reviews, 2008, 22, 261-281.	5.7	149
10	Forward genetic analysis of TLR-signaling pathways: An evaluationâ˜†. Advanced Drug Delivery Reviews, 2008, 60, 824-829.	13.7	23
11	Abnormal liver function in common variable immunodeficiency disorders due to nodular regenerative hyperplasia. Clinical and Experimental Immunology, 2008, 153, 331-337.	2.6	109
12	Building networks for immunodeficiency diseases and immunology training. Nature Immunology, 2008, 9, 1005-1007.	14.5	10
13	Clinical Immunology Review Series: An approach to the patient with recurrent infections in childhood. Clinical and Experimental Immunology, 2008, 152, 389-396.	2.6	70
14	Longitudinal analysis of immune function in the first 3 years of life in thymectomized neonates during cardiac surgery. Clinical and Experimental Immunology, 2008, 154, 375-383.	2.6	62
15	Revisiting human primary immunodeficiencies. Journal of Internal Medicine, 2008, 264, 115-127.	6.0	59
16	Granulomatous disease: Distinguishing primary antibody disease from sarcoidosis. Clinical Immunology, 2008, 128, 18-22.	3.2	60
17	Advances in the management of primary immunodeficiency. Paediatrics and Child Health (United) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50	0.4	2
18	Toll-like receptors: their roles in bacterial recognition and respiratory infections. Expert Review of Anti-Infective Therapy, 2008, 6, 479-495.	4.4	18

#	Article	IF	Citations
19	Interpretation of flow cytometry in primary immunodeficiency disorders. Annals of Allergy, Asthma and Immunology, 2008, 100, 612-615.	1.0	1
20	Mannose-binding lectin and mannose-binding lectinâ€“associated serine protease 2 in susceptibility, severity, and outcome of pneumonia in adults. Journal of Allergy and Clinical Immunology, 2008, 122, 368-374.e2.	2.9	116
21	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
22	Is it necessary to identify molecular defects in primary immunodeficiency disease?. Journal of Allergy and Clinical Immunology, 2008, 122, 1069-1073.	2.9	26
23	The genetic heterogeneity of mendelian susceptibility to mycobacterial diseases. Journal of Allergy and Clinical Immunology, 2008, 122, 1043-1051.	2.9	214
24	Hypogammaglobulinaemia. Immunology and Allergy Clinics of North America, 2008, 28, 691-713.	1.9	20
25	Phagocytes Defects. , 2008, , 131-166.		6
26	Primary Immunodeficiency Diseases. , 2008, , .		23
27	Common variable immunodeficiency disorders: division into distinct clinical phenotypes. Blood, 2008, 112, 277-286.	1.4	709
28	Combined T and B Cell Immunodeficiencies. , 2008, , 39-95.		3
29	Megadose CD34 + Cell Grafts Improve Recovery of T Cell Engraftment but not B Cell Immunity in Patients with Severe Combined Immunodeficiency Disease Undergoing Haplocompatible Nonmyeloablative Transplantation. Biology of Blood and Marrow Transplantation, 2008, 14, 1125-1133.	2.0	47
31	Genetic Syndromic Immunodeficiencies with Antibody Defects. Immunology and Allergy Clinics of North America, 2008, 28, 715-736.	1.9	10
32	Pierwotne niedobory odpornoÅ›ci. Pediatria Polska, 2008, 83, 697-703.	0.2	1
33	Gene Therapy for Primary Immunodeficiencies. Immunology and Allergy Clinics of North America, 2008, 28, 457-471.	1.9	12
34	The use of commercially available genetic tests in immunodeficiency disorders. Annals of Allergy, Asthma and Immunology, 2008, 101, 212-218.	1.0	1
35	Advances in Hematopoietic Stem Cell Transplantation for Primary Immunodeficiency. Immunology and Allergy Clinics of North America, 2008, 28, 439-456.	1.9	18
36	Genetic Diagnosis of Primary Immune Deficiencies. Immunology and Allergy Clinics of North America, 2008, 28, 387-412.	1.9	7
37	Severe Congenital Neutropenia or Hyper-IgM Syndrome? A Novel Mutation of CD40 Ligand in a Patient with Severe Neutropenia. International Archives of Allergy and Immunology, 2008, 147, 255-259.	2.1	18

#	Article	IF	Citations
38	Predominantly Antibody Deficiencies. , 2008, , 97-130.		17
39	Antibody deficiency: Table 1. Journal of Clinical Pathology, 2008, 61, 994-1000.	2.0	18
40	Progression of Selective IgA Deficiency to Common Variable Immunodeficiency. International Archives of Allergy and Immunology, 2008, 147, 87-92.	2.1	138
41	T cell immunodeficiency. Journal of Clinical Pathology, 2008, 61, 988-993.	2.0	22
42	Serum Bactericidal Antibody Responses to Meningococcal Polysaccharide Vaccination as a Basis for Clinical Classification of Common Variable Immunodeficiency. Vaccine Journal, 2008, 15, 607-611.	3.1	38
43	Renal Disease in Common Variable Immunodeficiencyâ€"Case Report and Literature Review. Pediatric Asthma, Allergy and Immunology, 2008, 21, 35-39.	0.2	3
44	Development of Common Variable Immunodeficiency in an 8-year-old Boy Treated with Rituximab for Idiopathic Thrombocytopenia. Pediatric Asthma, Allergy and Immunology, 2008, 21, 99-104.	0.2	4
45	Mutations causing severe combined immunodeficiency: detection with a custom resequencing microarray. Genetics in Medicine, 2008, 10, 575-585.	2.4	31
46	Recent developments related to the laboratory diagnosis of primary immunodeficiency diseases. Current Opinion in Pediatrics, 2008, 20, 688-697.	2.0	9
48	Recently identified factors predisposing children to infectious diseases. Current Opinion in Infectious Diseases, 2008, 21, 217-222.	3.1	9
50	Applications of flow cytometry for the study of primary immune deficiencies. Current Opinion in Allergy and Clinical Immunology, 2008, 8, 499-509.	2.3	51
51	New approaches to treatment of primary immunodeficiencies: fixing mutations with chemicals. Current Opinion in Allergy and Clinical Immunology, 2008, 8, 540-546.	2.3	11
52	ImunodeficiÃªncias primÃ¡rias: aspectos relevantes para o pneumologista. Jornal Brasileiro De Pneumologia, 2009, 35, 1008-1017.	0.7	9
53	Streptococcus pneumoniae : un pathogÃ¨ne toujours trop prÃ©sent. Annales De Biologie Clinique, 2009, 67, 685-696.	0.1	0
54	Systematic Classification of Primary Immunodeficiencies Based on Clinical, Pathological, and Laboratory Parameters. Journal of Immunology, 2009, 183, 7569-7575.	0.8	28
55	Cytokine Gene Polymorphisms in Common Variable Immunodeficiency. International Archives of Allergy and Immunology, 2009, 150, 1-7.	2.1	21
56	Diagnostisches Vorgehen beim Verdacht auf einen PrimÃren Immundefekt (PID) / Diagnostic approach to suspected primary immunodeficiency. Laboratoriums Medizin, 2009, 33, 179-187.	0.6	3
57	Defects in Jak-STAT-mediated cytokine signals cause hyper-IgE syndrome: lessons from a primary immunodeficiency. International Immunology, 2009, 21, 105-112.	4.0	60

#	Article	IF	Citations
58	Diagnostic approach when suspecting primary immunodeficiency (PID) 1. Laboratoriums Medizin, 2009, 33, -.	0.6	0
59	Hot Topics in Infection and Immunity in Children V. Advances in Experimental Medicine and Biology, 2009, , .	1.6	1
60	Common Variable Immunodeficiency: Etiological and Treatment Issues. International Archives of Allergy and Immunology, 2009, 150, 311-324.	2.1	42
61	Ten years of gene therapy for primary immune deficiencies. Hematology American Society of Hematology Education Program, 2009, 2009, 682-689.	2.5	86
62	STIM1Mutation Associated with a Syndrome of Immunodeficiency and Autoimmunity. New England Journal of Medicine, 2009, 360, 1971-1980.	27.0	459
63	The B-cell Compartment in the Peripheral Blood of Children With Different Types of Primary Humoral Immunodeficiency. Pediatric Research, 2009, 66, 28-34.	2.3	28
64	Delayed Onset of (Severe) Combined Immunodeficiency (S)CID (T-B+NK+): Complete IL-7 Receptor Deficiency in a 22 Months Old Girl. Klinische Padiatrie, 2009, 221, 339-343.	0.6	13
65	Prediction of Candidate Primary Immunodeficiency Disease Genes Using a Support Vector Machine Learning Approach. DNA Research, 2009, 16, 345-351.	3.4	26
66	Primary immunodeficiencies. Pharmaceuticals Policy and Law, 2009, 11, 277-283.	0.1	1
67	RAPID: Resource of Asian Primary Immunodeficiency Diseases. Nucleic Acids Research, 2009, 37, D863-D867.	14.5	37
68	Dissecting the Group of Common Variable Immunodeficiency Disorders. Clinical Infectious Diseases, 2009, 49, 1339-1340.	5.8	2
69	The immunocompromised patient: primary immunodeficiencies. Medicine, 2009, 37, 518-521.	0.4	0
70	Recombinaseâ€activating gene 1 immunodeficiency: different immunological phenotypes in three siblings. Acta Paediatrica, International Journal of Paediatrics, 2009, 98, 1062-1064.	1.5	28
71	Severe combined immunodeficiency (SCID) and attention deficit hyperactivity disorder (ADHD) associated with a coronin-1A mutation and a chromosome 16p11.2 deletion. Clinical Immunology, 2009, 131, 24-30.	3.2	123
72	Granulomatous disease in common variable immunodeficiency. Clinical Immunology, 2009, 133, 198-207.	3.2	178
73	Pharmacokinetics of a new 10% intravenous immunoglobulin in patients receiving replacement therapy for primary immunodeficiency. European Journal of Pharmaceutical Sciences, 2009, 37, 272-278.	4.0	42
74	Clinical heterogeneity can hamper the diagnosis of patients with ZAP70 deficiency. European Journal of Pediatrics, 2009, 168, 87-93.	2.7	103
75	Mannose-binding lectin polymorphisms in common variable immunodeficiency. Clinical and Experimental Medicine, 2009, 9, 285-290.	3.6	7

#	Article	IF	Citations
76	Chinese Patients with Defective IL-12/23-Interferon-Î³ Circuit in Taiwan: Partial Dominant Interferon-Î³ Receptor 1 Mutation Presenting as Cutaneous Granuloma and IL-12 Receptor Î²1 Mutation as Pneumatocele. Journal of Clinical Immunology, 2009, 29, 238-245.	3.8	31
77	Primary Immunodeficiency Diseases in Egyptian Children: A Single-Center Study. Journal of Clinical Immunology, 2009, 29, 343-351.	3.8	75
78	Clinical Characteristics and Outcomes of Primary Immunodeficiencies in Thai Children: An 18-year Experience from a Tertiary Care Center. Journal of Clinical Immunology, 2009, 29, 357-364.	3.8	21
79	Autoimmunity in common variable immunodeficiency. Current Allergy and Asthma Reports, 2009, 9, 347-352.	5.3	165
84	Le Centre de RÃ©fÃ©rence DÃ©ficits Immunitaires HÃ©rÃ©ditaires (CEREDIH). Revue Francophone Des Laboratoires, 2009, 2009, 43-44.	0.0	0
85	Update in understanding common variable immunodeficiency disorders (CVIDs) and the management of patients with these conditions. British Journal of Haematology, 2009, 145, 709-727.	2.5	333
86	Proinflammatory cytokine gene single nucleotide polymorphisms in common variable immunodeficiency. Clinical and Experimental Immunology, 2009, 155, 21-27.	2.6	57
87	Decrease in phenotypic regulatory T cells in subsets of patients with common variable immunodeficiency. Clinical and Experimental Immunology, 2009, 156, 446-454.	2.6	64
88	Immunodeficiencies. Clinical and Experimental Immunology, 2009, 158, 14-22.	2.6	63
89	Economic evaluation of immunoglobulin replacement in patients with primary antibody deficiencies. Clinical and Experimental Immunology, 2009, 160, 240-245.	2.6	66
90	The European internet-based patient and research database for primary immunodeficiencies: results 2006â€“2008. Clinical and Experimental Immunology, 2009, 157, 3-11.	2.6	203
91	Primary Immunodeficiencies: â€œNewâ€•Disease in an Old Country. Cellular and Molecular Immunology, 2009, 6, 397-406.	10.5	17
92	Reticular dysgenesis (aleukocytosis) is caused by mutations in the gene encoding mitochondrial adenylate kinase 2. Nature Genetics, 2009, 41, 101-105.	21.4	205
93	Detection of pulmonary complications in common variable immunodeficiency. Pediatric Allergy and Immunology, 2010, 21, 793-805.	2.6	70
94	Children with frequent infections: A proposal for a stepwise assessment and investigation of the immune system. Pediatric Allergy and Immunology, 2010, 21, 463-473.	2.6	10
96	Invasive fungal infections in the paediatric and neonatal population: diagnostics and management issues. Clinical Microbiology and Infection, 2009, 15, 613-624.	6.0	47
97	Hyper-IgE syndrome. Current Opinion in Immunology, 2009, 21, 487-492.	5.5	97
98	Primary immunodeficiencies: increasing market share. Current Opinion in Immunology, 2009, 21, 461-465.	5.5	19

#	Article	IF	Citations
99	Development of a routine newborn screening protocol for severe combined immunodeficiency. Journal of Allergy and Clinical Immunology, 2009, 124, 522-527.	2.9	173
100	Toll-like receptor 7 and 9 defects in common variable immunodeficiency. Journal of Allergy and Clinical Immunology, 2009, 124, 349-356.e3.	2.9	97
101	ComÃ¨l-Netherton syndrome defined as primary immunodeficiency. Journal of Allergy and Clinical Immunology, 2009, 124, 536-543.	2.9	164
102	ORAI1 deficiency and lack of store-operated Ca2+ entry cause immunodeficiency, myopathy, and ectodermal dysplasia. Journal of Allergy and Clinical Immunology, 2009, 124, 1311-1318.e7.	2.9	289
103	DÃ©ficits immunitaires communs variables (DICV). Signes dâ€™alerte chez lâ€™enfant. Revue Francaise D'allergologie, 2009, 49, 487-489.	0.2	0
104	Common variable immunodeficiency: a multifaceted and puzzling disorder. Expert Review of Clinical Immunology, 2009, 5, 167-180.	3.0	18
105	An update on treatment strategies for common variable immunodeficiency. Expert Review of Clinical Immunology, 2009, 5, 381-390.	3.0	6
106	Recent Advances in Primary Immunodeficiencies: Identification of Novel Genetic Defects and Unanticipated Phenotypes. Pediatric Research, 2009, 65, 3R-12R.	2.3	38
107	Primary immunodeficiency diseases: a practical guide for clinicians. Postgraduate Medical Journal, 2009, 85, 660-666.	1.8	37
108	Lateâ€Onset Combined Immune Deficiency: A Subset of Common Variable Immunodeficiency with Severe T Cell Defect. Clinical Infectious Diseases, 2009, 49, 1329-1338.	5.8	192
109	Disseminated bacille Calmetteâ€“GuÃ©rin in Iranian children with severe combined immunodeficiency. International Journal of Infectious Diseases, 2009, 13, e420-e423.	3.3	42
110	Therapeutic Approaches to Secondary Immune Thrombocytopenic Purpura. Seminars in Hematology, 2009, 46, S44-S58.	3.4	16
112	Immunophenotypic profile of T cells in common variable immunodeficiency: is there an association with different clinical findings?. Allergologia Et Immunopathologia, 2009, 37, 14-20.	1.7	18
113	Neutropenia and Primary Immunodeficiency Diseases. International Reviews of Immunology, 2009, 28, 335-366.	3.3	58
114	DÃ©ficits immunitaires primitifs. , 2009, , 83-112.		1
115	The clinical utility of molecular diagnostic testing for primary immune deficiency disorders. Pathology, 2009, 41, 39.	0.6	0
116	Thymus microenvironment in human primary immunodeficiency diseases. Current Opinion in Allergy and Clinical Immunology, 2009, 9, 489-495.	2.3	25
117	Echocardiographic abnormalities and their correlation with bronchiectasis score in primary antibody deficiencies. Journal of Cardiovascular Medicine, 2010, 11, 244-249.	1.5	4

#	Article	IF	Citations
118	Implementing Routine Testing for Severe Combined Immunodeficiency within Wisconsin's Newborn Screening Program. Public Health Reports, 2010, 125, 88-95.	2.5	65
120	The 752delG26 mutation in the RFXANK gene associated with major histocompatibility complex class II deficiency: evidence for a founder effect in the Moroccan population. European Journal of Pediatrics, 2010, 169, 1069-1074.	2.7	31
121	Molecular- and Flow Cytometry-based Diagnosis of Primary Immunodeficiency Disorders. Current Allergy and Asthma Reports, 2010, 10, 460-467.	5.3	17
122	Selective IgA Deficiency. Journal of Clinical Immunology, 2010, 30, 10-16.	3.8	423
123	Guidelines for implementation of populationâ€based newborn screening for severe combined immunodeficiency. Journal of Inherited Metabolic Disease, 2010, 33, 273-281.	3.6	60
124	A novel radiosensitive SCID patient with a pronounced G2/M sensitivity. DNA Repair, 2010, 9, 365-373.	2.8	3
125	Homologous recombination in human iPS and ES cells for use in gene correction therapy. Drug Discovery Today, 2010, 15, 198-202.	6.4	14
126	Use of intravenous immunoglobulin and adjunctive therapies in the treatment of primary immunodeficiencies. Clinical Immunology, 2010, 135, 255-263.	3.2	86
127	Primary immunodeficiencies of protective immunity to primary infections. Clinical Immunology, 2010, 135, 204-209.	3.2	65
128	The French national registry of primary immunodeficiency diseases. Clinical Immunology, 2010, 135, 264-272.	3.2	137
129	The clinical utility of molecular diagnostic testing for primary immune deficiency disorders: a case based review. Allergy, Asthma and Clinical Immunology, 2010, 6, 12.	2.0	25
130	Primary Immunodeficiency and Autoimmunity: Lessons From Human Diseases. Scandinavian Journal of Immunology, 2010, 71, 317-328.	2.7	101
131	B-cell maturation and antibody responses in individuals carrying a mutated CD19 allele. Genes and Immunity, 2010, 11, 523-530.	4.1	34
132	12.8 Morbus haemolyticus neonatorum. , 2010, , .		0
133	Immunoglobulin G Subclass Deficiency is the Major Phenotype of Primary Immunodeficiency in a Korean Adult Cohort. Journal of Korean Medical Science, 2010, 25, 824.	2.5	28
134	A single center 14 years study of infectious complications leading to hospitalization of patients with primary antibody deficiencies. Brazilian Journal of Infectious Diseases, 2010, 14, 351-355.	0.6	0
135	Mutation@A Glance: An Integrative Web Application for Analysing Mutations from Human Genetic Diseases. DNA Research, 2010, 17, 197-208.	3.4	26
136	Gastrointestinal and hepatic manifestations of primary immune deficiency diseases. Saudi Journal of Gastroenterology, 2010, 16, 66.	1.1	29

#	Article	IF	Citations
137	X-Linked Lymphoproliferative Disease Presenting as Pancytopenia in a 10-Month-Old Boy. Case Reports in Medicine, 2010, 2010, 1-3.	0.7	2
138	Approach to the Child with Recurrent Infections. , 2010, , 81-87.		0
139	Serum Bactericidal Antibody Response 1 Year after Meningococcal Polysaccharide Vaccination of Patients with Common Variable Immunodeficiency. Vaccine Journal, 2010, 17, 524-528.	3.1	27
140	Gene Therapy for Primary Immunodeficiencies. , 2010, , 213-231.		0
141	Molecular Assessment of Thymus Capabilities in the Evaluation of T-Cell Immunodeficiency. Pediatric Research, 2010, 67, 211-216.	2.3	29
142	Gene Therapy for Adenosine Deaminase Deficiency. Immunology and Allergy Clinics of North America, 2010, 30, 249-260.	1.9	16
143	Bone Marrow Transplantation for Primary Immunodeficiency Diseases. Pediatric Clinics of North America, 2010, 57, 207-237.	1.8	31
144	Haploidentical Bone Marrow Transplantation in Primary Immune Deficiency: Stem Cell Selection and Manipulation. Immunology and Allergy Clinics of North America, 2010, 30, 45-62.	1.9	7
145	Primary immunodeficiencies. Journal of Allergy and Clinical Immunology, 2010, 125, S182-S194.	2.9	358
146	Infection outcomes in patients with common variable immunodeficiency disorders: Relationship to immunoglobulin therapy over 22 years. Journal of Allergy and Clinical Immunology, 2010, 125, 1354-1360.e4.	2.9	370
147	Primary immunodeficiencies may reveal potential infectious diseases associated with immune-targeting mAb treatments. Journal of Allergy and Clinical Immunology, 2010, 126, 910-917.	2.9	21
149	Altered functional balance of Gfi-1 and Gfi-1b as an alternative cause of reticular dysgenesis?. Medical Hypotheses, 2010, 74, 445-448.	1.5	5
150	Association of IL-4 and IL-10 gene promoter polymorphisms with common variable immunodeficiency. Immunobiology, 2010, 215, 81-87.	1.9	33
151	DÃ©ficits immunitaires primitifs: approche diagnostique pour les pays Ã©mergents. Immuno-Analyse Et Biologie Specialisee, 2010, 25, 257-265.	0.0	2
154	Therapeutic Use of Immunoglobulins. Advances in Pediatrics, 2010, 57, 185-218.	1.4	25
155	Antibody Deficiency. , 2010, , 88-97.		1
156	Evaluation of Immunoglobulin Levels and Infection Rate in Patients with Common Variable Immunodeficiency After Immunoglobulin Replacement Therapy. Journal of Microbiology, Immunology and Infection, 2010, 43, 11-17.	3.1	9
157	A single center 14 years study of infectious complications leading to hospitalization of patients with primary antibody deficiencies. Brazilian Journal of Infectious Diseases, 2010, 14, 351-355.	0.6	15

#	Article	IF	Citations
158	PIDO: the primary immunodeficiency disease ontology. Bioinformatics, 2011, 27, 3193-3199.	4.1	7
162	Defects in the CD19 complex predispose to glomerulonephritis, as well as IgG1 subclass deficiency. Journal of Allergy and Clinical Immunology, 2011, 127, 538-541.e5.	2.9	31
163	Common variable immunodeficiency presenting as herpes simplex virus encephalitis. Journal of Allergy and Clinical Immunology, 2011, 127, 541-543.	2.9	17
164	Inflammation in common variable immunodeficiency is associated with a distinct CD8+ response to cytomegalovirus. Journal of Allergy and Clinical Immunology, 2011, 127, 1385-1393.e4.	2.9	29
166	Principles of Immunopharmacology. , 2011, , .		14
167	Flow Cytometry Assays in Primary Immunodeficiency Diseases. Methods in Molecular Biology, 2011, 699, 317-335.	0.9	12
168	Haploidentical Bone Marrow Transplantation in Primary Immune Deficiency: Stem Cell Selection and Manipulation. Hematology/Oncology Clinics of North America, 2011, 25, 45-62.	2.2	4
169	C2 Plasma-derived immunoglobulins. , 2011, , 271-301.		1
171	Critical issues and needs in management of primary immunodeficiency diseases in Latin America. Allergologia Et Immunopathologia, 2011, 39, 45-51.	1.7	17
172	Primary immunodeficiency diseases in Latin America: Proceedings of the Second Latin American Society for Immunodeficiencies (LASID) Advisory Board. Allergologia Et Immunopathologia, 2011, 39, 106-110.	1.7	18
173	Diagnosis and treatment of primary immunodeficiency disease: the role of the otolaryngologist. American Journal of Otolaryngology - Head and Neck Medicine and Surgery, 2011, 32, 329-337.	1.3	13
174	Chronic pediatric pulmonary disease and primary humoral antibody based immune disease. Respiratory Medicine, 2011, 105, 511-514.	2.9	13
175	Artemis splice defects cause atypical SCID and can be restored in vitro by an antisense oligonucleotide. Genes and Immunity, 2011, 12, 434-444.	4.1	27
177	HTAs and access to rare diseases therapies: How can clinicians assist in the healthcare assessment of treatments for patients with primary immune deficiencies?. Pharmaceuticals Policy and Law, 2011, 13, 205-212.	0.1	0
178	Allogeneic hematopoietic stem cell transplantation in children with primary immunodeficiencies: Hospital Israelita Albert Einstein experience. Einstein (Sao Paulo, Brazil), 2011, 9, 140-144.	0.7	3
179	Successful Firstâ€line Treatment With Propranolol of Multifocal Infantile Hepatic Hemangioma With Highâ€flow Cardiac Overload. Journal of Pediatric Gastroenterology and Nutrition, 2011, 53, 693-695.	1.8	16
180	Earlyâ€onset Hepatic Fibrosis in Lysinuric Protein Intolerance. Journal of Pediatric Gastroenterology and Nutrition, 2011, 53, 695-698.	1.8	6
181	The Role of the Infectious Disease Specialist in the Diagnosis and Treatment of Primary Immunodeficiency Disease. Infectious Diseases in Clinical Practice, 2011, 19, 316-325.	0.3	2

#	Article	IF	Citations
182	High Prevalence of Intestinal Carriage of Campylobacter coli in Patients With Primary Antibody Deficiencies. Journal of Clinical Gastroenterology, 2011, 45, 474-475.	2.2	6
183	Genotype, phenotype, and outcomes of nine patients with Tâ€B+NK+ SCID. Pediatric Transplantation, 2011, 15, 733-741.	1.0	11
184	Clinical and Molecular Characteristics of Immunodysregulation, Polyendocrinopathy, Enteropathy, Xâ€Linked Syndrome in China. Scandinavian Journal of Immunology, 2011, 74, 304-309.	2.7	21
185	Ageâ€matched Reference Values for Bâ€lymphocyte Subpopulations and CVID Classifications in Children. Scandinavian Journal of Immunology, 2011, 74, 502-510.	2.7	72
186	Primary immune deficiency disorders in the South Pacific: the clinical utility of a customized genetic testing program in New Zealand. Annals of the New York Academy of Sciences, 2011, 1238, 53-64.	3.8	21
187	The demographics of primary immunodeficiency diseases across the unique ethnic groups in Iran, and approaches to diagnosis and treatment. Annals of the New York Academy of Sciences, 2011, 1238, 24-32.	3.8	15
188	Primary immunodeficiencies in highly consanguineous North African populations. Annals of the New York Academy of Sciences, 2011, 1238, 42-52.	3.8	58
189	Potential therapeutic applications of antisense morpholino oligonucleotides in modulation of splicing in primary immunodeficiency diseases. Journal of Immunological Methods, 2011, 365, 1-7.	1.4	20
190	Diagnostic strategy for patients with hypogammaglobulinemia in rheumatology. Joint Bone Spine, 2011, 78, 241-245.	1.6	39
191	Diagnosis and treatment of primary immunodeficiency disease in patients with gastrointestinal symptoms. Clinical Immunology, 2011, 139, 238-248.	3.2	31
192	Distribution and Clinical Features of Primary Immunodeficiency Diseases in Chinese Children (2004â€“2009). Journal of Clinical Immunology, 2011, 31, 297-308.	3.8	23
193	Monozygotic Twin Pair Showing Discordant Phenotype for X-linked Thrombocytopenia and Wiskottâ€“Aldrich Syndrome: a Role for Epigenetics?. Journal of Clinical Immunology, 2011, 31, 773-777.	3.8	30
194	Nationwide Survey of Patients with Primary Immunodeficiency Diseases in Japan. Journal of Clinical Immunology, 2011, 31, 968-976.	3.8	77
195	Study of patients with Hyper-IgM type IV phenotype who recovered spontaneously during late childhood and review of the literature. European Journal of Pediatrics, 2011, 170, 1039-1047.	2.7	9
196	StratÃ©gie diagnostique devant la dÃ©couverte dâ€™une hypogammaglobulinemie en rhumatologie. Revue Du Rhumatisme (Edition Francaise), 2011, 78, 122-127.	0.0	2
197	Primary immunodeficiency. Allergy, Asthma and Clinical Immunology, 2011, 7, S11.	2.0	96
198	The 752delG26 mutation in the RFXANK gene associated with major histocompatibility complex class II deficiency: evidence for a founder effect in the Moroccan population. BMC Proceedings, 2011, 5, .	1.6	1
199	Malignancies in the setting of primary immunodeficiency: Implications for hematologists/oncologists. American Journal of Hematology, 2011, 86, 48-55.	4.1	79

#	Article	IF	Citations
200	Characterization of immunologic defects in patients with common variable immunodeficiency (CVID) with intestinal disease. Inflammatory Bowel Diseases, 2011, 17, 251-259.	1.9	124
202	Parental Consanguinity and the Risk of Primary Immunodeficiency Disorders: Report from the Kuwait National Primary Immunodeficiency Disorders Registry. International Archives of Allergy and Immunology, 2011, 154, 76-80.	2.1	30
203	Clinical Features That Identify Children With Primary Immunodeficiency Diseases. Pediatrics, 2011, 127, 810-816.	2.1	149
204	Primary immune deficiency in the intensive care unit: It is never too late to diagnose and treat. Indian Journal of Critical Care Medicine, 2011, 15, 179-181.	0.9	1
205	Immune Dysregulation, Polyendocrinopathy, Enteropathy, X-Linked Syndrome: A Paradigm of Immunodeficiency with Autoimmunity. Frontiers in Immunology, 2012, 3, 211.	4.8	279
206	Immune Deficiency in Children. InnovAiT, 2012, 5, 266-273.	0.0	2
208	Early-Onset Chronic Inflammatory Disease Associated with Maternal Microchimerism. Case Reports in Pediatrics, 2012, 2012, 1-4.	0.4	4
209	Multiorgan Infiltration by CD8+ T Cells and 1p;16p Translocation in a Patient with Hypogammaglobulinemia and a Reduced Number of B Cells. International Archives of Allergy and Immunology, 2012, 158, 206-210.	2.1	2
210	Primary Immunodeficiency Diseases in Oman: Five Years' Experience at Sultan Qaboos University Hospital. World Allergy Organization Journal, 2012, 5, 52-56.	3.5	26
211	Endocrine complications in primary immunodeficiency diseases in Japan. Clinical Endocrinology, 2012, 77, 628-634.	2.4	6
212	Recurrent Infections. , 2012, , 1-75.		0
213	Rheumatic manifestations of primary immunodeficiencies in children. Indian Journal of Rheumatology, 2012, 7, 52-56.	0.4	2
214	Rac2 expression and its role in neutrophil functions of zebrafish (Danio rerio). Fish and Shellfish Immunology, 2012, 33, 1086-1094.	3.6	25
215	Hematological manifestations of celiac disease. Scandinavian Journal of Gastroenterology, 2012, 47, 1401-1411.	1.5	45
216	BAY 41â€2272, a soluble guanylate cyclase agonist, activates human mononuclear phagocytes. British Journal of Pharmacology, 2012, 166, 1617-1630.	5.4	10
217	Advancing the management of primary immunodeficiency diseases in Latin America: Latin American Society for Immunodeficiencies (LASID) Initiatives. Allergologia Et Immunopathologia, 2012, 40, 187-193.	1.7	14
218	Recombinant human hyaluronidase-facilitated subcutaneous infusion of human immunoglobulins for primary immunodeficiency. Journal of Allergy and Clinical Immunology, 2012, 130, 951-957.e11.	2.9	113
220	Optimal use of ivig in a patient with BehÃ§et syndrome and common variable immunodeficiency. Annals of Allergy, Asthma and Immunology, 2012, 109, 84.	1.0	9

#	Article	IF	Citations
221	Predominantly Antibody Deficiency. , 2012, , 113-192.		0
222	Immunomodulatory and antioxidant effect ofLeptadenia reticulataleaf extract in rodents: possible modulation of cell and humoral immune response. Immunopharmacology and Immunotoxicology, 2012, 34, 1010-1019.	2.4	15
223	Prevalence of Primary Immunodeficiency in Korea. Journal of Korean Medical Science, 2012, 27, 788.	2.5	59
224	Immunoglobulin Treatment of Immunodeficient Patients. , 2012, , .		1
225	Primary Immunodeficiency Diseases. , 2012, , 1615-1622.		0
226	Lessons Learned from Phagocytic Function Studies in a Large Cohort of Patients with Recurrent Infections. Journal of Clinical Immunology, 2012, 32, 454-466.	3.8	21
227	Advances in primary immunodeficiency diseases in Latin America: epidemiology, research, and perspectives. Annals of the New York Academy of Sciences, 2012, 1250, 62-72.	3.8	34
228	The European internet-based patient and research database for primary immunodeficiencies: update 2011. Clinical and Experimental Immunology, 2012, 167, 479-491.	2.6	91
229	The cellular and molecular mechanisms for neutropenia in Barth syndrome. European Journal of Haematology, 2012, 88, 195-209.	2.2	28
230	Bacillus Calmette-GuÃ©rin (BCG) complications associated with primary immunodeficiency diseases. Journal of Infection, 2012, 64, 543-554.	3.3	145
231	Recurrent infection in children: When and how to investigate for primary immunodeficiency?. Journal of Paediatrics and Child Health, 2012, 48, 202-209.	0.8	11
232	Primary Immunodeficiency Diseases Associated with Neurologic Manifestations. Journal of Clinical Immunology, 2012, 32, 1-24.	3.8	42
233	Development of common variable immunodeficiency in IgA- and IgG2-deficient patients with systemic lupus erythematosus. Pediatric Nephrology, 2012, 27, 489-492.	1.7	9
234	The role of mannose-binding lectin in pneumococcal infection. European Respiratory Journal, 2013, 41, 131-139.	6.7	42
235	The Prevalances and Patient Characteristics of Primary Immunodeficiency Diseases in Turkeyâ€"Two Centers Study. Journal of Clinical Immunology, 2013, 33, 74-83.	3.8	66
236	Common variable immunodeficiency unmasked by treatment of immune thrombocytopenic purpura with Rituximab. BMC Blood Disorders, 2013, 13, 4.	0.9	26
237	The immunocompromised patient: primary immunodeficiencies. Medicine, 2013, 41, 619-623.	0.4	1
238	Advances in management of primary immunodeficiency. Paediatrics and Child Health (United Kingdom), 2013, 23, 115-120.	0.4	2

#	Article	IF	Citations
239	Next Generation Sequencing. , 2013, , .		10
240	Asthma and Risk of Selective IgA Deficiency or Common Variable Immunodeficiency: A Population-Based Case-Control Study. Mayo Clinic Proceedings, 2013, 88, 813-821.	3.0	58
241	Rheumatic Disease. , 2013, , 1415-1440.		0
242	Germline Mutations in NFKB2 Implicate the Noncanonical NF-Î²B Pathway in the Pathogenesis of Common Variable Immunodeficiency. American Journal of Human Genetics, 2013, 93, 812-824.	6.2	256
243	Common Variable Immune Deficiency in Childrenâ€"Clinical Characteristics Varies Depending on Defect in Peripheral B Cell Maturation. Journal of Clinical Immunology, 2013, 33, 731-741.	3.8	20
244	Recurrent Lower Respiratory Tract Infections in Children: A Practical Approach to Diagnosis. Paediatric Respiratory Reviews, 2013, 14, 53-60.	1.8	40
245	Primary Immunodeficiency Diseases in Different Age Groups: A Report on 1,008 Cases from a Single Brazilian Reference Center. Journal of Clinical Immunology, 2013, 33, 716-724.	3.8	38
246	When to Think of Immunodeficiency?. Advances in Experimental Medicine and Biology, 2013, 764, 167-177.	1.6	5
247	NGS-Based Clinical Diagnosis of Genetically Heterogeneous Disorders. , 2013, , 115-150.		0
248	Primary Immunodeficiency Diseases at Reference and High-Specialty Hospitals in the State of Guanajuato, Mexico. BioMed Research International, 2013, 2013, 1-6.	1.9	7
249	The Role of the Innate Immune System in Granulomatous Disorders. Frontiers in Immunology, 2013, 4, 120.	4.8	71
250	Evidence-based use of serum protein electrophoresis in laboratory medicine. Clinical Chemistry and Laboratory Medicine, 2013, 51, e113-5.	2.3	5
251	Newborn screening for severe combined immunodeficiency: an opportunity for intervention. Journal of Perinatology, 2013, 33, 657-658.	2.0	4
252	Treatment of comorbid selective mutism and procedural anxiety in a child with pediatric common variable immunodeficiency disease: A case study.. Clinical Practice in Pediatric Psychology, 2013, 1, 129-136.	0.3	1
253	Can Patients with Common Variable Immunodeficiency Have Allergic Rhinitis?. American Journal of Rhinology and Allergy, 2013, 27, 79-83.	2.0	17
255	Common variable immunodeficiency â€“. Central-European Journal of Immunology, 2013, 1, 141-148.	1.2	0
256	Clinical Manifestations of the Human Papillomavirus. , 0, , .		0
257	Primary Immunodeficiency Disorders Â– An Update. Bangladesh Journal of Child Health, 2013, 36, 139-145.	0.1	0

#	Article	IF	Citations
258	Primary immunodeficiency investigation in patients during and after hospitalization in a pediatric Intensive Care Unit. Revista Paulista De Pediatria, 2014, 32, 32-36.	1.0	6
259	Health-related quality of life, anxiety, and depressive symptoms in children with primary immunodeficiencies. Patient Preference and Adherence, 2014, 8, 323.	1.8	37
260	Cost-minimization Analysis of IgPro20, a Subcutaneous Immunoglobulin, in Japanese Patients With Primary Immunodeficiency. Clinical Therapeutics, 2014, 36, 1616-1624.	2.5	22
262	Diseases of the Sinuses. , 2014, , .		15
263	Development of an anti-CD2/CD3/CD28 bead-based T-cell proliferation assay. Bioscience Horizons, 2014, 7, hzu012-hzu012.	0.6	7
264	Long-Lasting Production of New T and B Cells and T-Cell Repertoire Diversity in Patients with Primary Immunodeficiency Who Had Undergone Stem Cell Transplantation: A Single-Centre Experience. Journal of Immunology Research, 2014, 2014, 1-10.	2.2	1
265	ComÃ¨l-Netherton syndrome â€“. Przeglad Dermatologiczny, 2014, 6, 481-486.	0.1	0
267	Synthesis and antimicrobial activities of His(2-aryl)-Arg and Trp-His(2-aryl) classes of dipeptidomimetics. MedChemComm, 2014, 5, 671-676.	3.4	22
268	Laboratory Diagnosis of Primary Immunodeficiencies. Clinical Reviews in Allergy and Immunology, 2014, 46, 154-168.	6.5	53
269	Autoimmunity in Primary Immunodeficiency Disorders. , 2014, , 403-418.		0
270	Response to pneumococcal vaccination in mannose-binding lectin-deficient adults with recurrent respiratory tract infections. Clinical and Experimental Immunology, 2014, 177, 272-279.	2.6	15
271	Pneumococcal Vaccine and Patients with Pulmonary Diseases. American Journal of Medicine, 2014, 127, 886.e1-886.e8.	1.5	28
272	Clinical Course and Outcome Predictors of Critically Ill Infants With Complete DiGeorge Anomaly Following Thymus Transplantation. Pediatric Critical Care Medicine, 2014, 15, e321-e326.	0.5	8
273	Clinical and immunological assessment of 94 patients with primary humoral immunodeficiency: Common variable immunodeficiency, selective IgA deficiency and polysaccharide antibody deficiency syndrome. Journal of Pediatric Infectious Diseases, 2015, 06, 159-166.	0.2	3
274	Genetics of common variable immunodeficiency: role of transmembrane activator and calcium modulator and cyclophilin ligand interactor. International Journal of Immunogenetics, 2015, 42, 239-253.	1.8	14
275	Immunodeficiency Among Children with Recurrent Invasive Pneumococcal Disease. Pediatric Infectious Disease Journal, 2015, 34, 644-651.	2.0	23
276	A Perspective on IL-7RÎ± Deficient Tâ^'B+NK+ Severe Combined Immunodeficiency. Pediatric Infectious Disease Journal, 2015, 34, 301-303.	2.0	0
277	Clinical immunology Recurrent respiratory tract infections in children â€“ analysis of immunological examinations. Central-European Journal of Immunology, 2015, 2, 167-173.	1.2	21

#	Article	IF	Citations
278	CD21 and CD19 deficiency: Two defects in the same complex leading to different disease modalities. Clinical Immunology, 2015, 161, 120-127.	3.2	42
279	Class Switch Recombination Process in Ataxia Telangiectasia Patients with Elevated Serum Levels of IgM. Journal of Immunoassay and Immunochemistry, 2015, 36, 16-26.	1.1	21
280	Que faire devant une monoarthrite chez lâ€™enfantÂ?. Revue Du Rhumatisme (Edition Francaise), 2015, 82, 85-90.	0.0	0
281	Diagnosing primary immunodeficiency: a practical approach for the non-immunologist. Current Medical Research and Opinion, 2015, 31, 697-706.	1.9	23
282	20% subcutaneous immunoglobulin for patients with primary immunodeficiency diseases: A systematic review. International Immunopharmacology, 2015, 25, 457-464.	3.8	5
283	Hematopoietic Stem Cell Transplantation for Severe Combined Immunodeficiency. Current Pediatrics Reports, 2015, 3, 1-10.	4.0	26
284	Primary immunodeficiency diseases. Nurse Practitioner, 2015, 40, 1-7.	0.3	4
285	MBL-associated serine proteases (MASPs) and infectious diseases. Molecular Immunology, 2015, 67, 85-100.	2.2	60
286	Validity of Primary Immunodeficiency Disease Diagnoses in United States Medicaid Data. Journal of Clinical Immunology, 2015, 35, 566-572.	3.8	8
287	Identification of Two Novel Mutations in Patients With X-Linked Primary Immunodeficiencies. Fetal and Pediatric Pathology, 2015, 34, 91-98.	0.7	4
288	Intravenous immunoglobulin replacement therapy to prevent pulmonary infection in a patient with Good's syndrome. Journal of Microbiology, Immunology and Infection, 2015, 48, 229-232.	3.1	17
289	Cancers Related to Immunodeficiencies: Update and Perspectives. Frontiers in Immunology, 2016, 7, 365.	4.8	137
290	Population-Based Screening for Selective Immunoglobulin A (IgA) Deficiency in Lithuanian Children Using a Rapid Antibody-Based Fingertip Test. Medical Science Monitor, 2016, 22, 4773-4778.	1.1	14
291	Long-Term Tolerability, Safety, and Efficacy of Recombinant Human Hyaluronidase-Facilitated Subcutaneous Infusion of Human Immunoglobulin for Primary Immunodeficiency. Journal of Clinical Immunology, 2016, 36, 571-582.	3.8	37
292	Autoimmunity in primary T-cell immunodeficiencies. Expert Review of Clinical Immunology, 2016, 12, 989-1006.	3.0	15
293	Development and Initial Validation of a Questionnaire to Measure Health-Related Quality of Life of Adults with Common Variable Immune Deficiency: The CVID_QoL Questionnaire. Journal of Allergy and Clinical Immunology: in Practice, 2016, 4, 1169-1179.e4.	3.8	29
294	Recombinant human hyaluronidase facilitated subcutaneous immunoglobulin treatment in pediatric patients with primary immunodeficiencies: long-term efficacy, safety and tolerability. Immunotherapy, 2016, 8, 1175-1186.	2.0	30
295	Expanded newborn screening by mass spectrometry: New tests, future perspectives. Mass Spectrometry Reviews, 2016, 35, 71-84.	5.4	107

#	Article	IF	Citations
296	Immunoglobulin A deficiency in celiac disease in the United States. Journal of Gastroenterology and Hepatology (Australia), 2016, 31, 133-137.	2.8	35
297	Development of a Low-Cost Stem-Loop Real-Time Quantification PCR Technique for EBV miRNA Expression Analysis. Molecular Biotechnology, 2016, 58, 540-550.	2.4	3
298	Prevalence of selective immunoglobulin A deficiency in Greek children and adolescents with type 1 diabetes. World Journal of Pediatrics, 2016, 12, 470-476.	1.8	7
299	Immunodeficiencies and the Rheumatic Diseases. , 2016, , 597-608.e5.		1
300	F-actin remodeling defects as revealed in primary immunodeficiency disorders. Clinical Immunology, 2016, 164, 34-42.	3.2	8
301	How effective are the 6 European Society of Immunodeficiency warning signs for primary immunodeficiency disease?. Annals of Allergy, Asthma and Immunology, 2016, 116, 151-155.e1.	1.0	16
302	Immunoglobulin-induced hemolysis, splenomegaly and inflammation in patients with antibody deficiencies. Expert Review of Clinical Immunology, 2016, 12, 725-731.	3.0	0
303	Magnetic Resonance Imaging May Be a Valuable Radiation-Free Technique for Lung Pathologies in Patients with Primary Immunodeficiency. Journal of Clinical Immunology, 2016, 36, 66-72.	3.8	18
304	Identifying Novel Inborn Errors of the Immune System. , 2017, , .		0
305	Pathway-based detection of idiopathic pulmonary fibrosis at an early stage. Molecular Medicine Reports, 2017, 15, 2023-2028.	2.4	0
306	Primary Immunodeficiency Diseases. , 2017, , .		22
307	Synergistic Cytotoxicity of Lenalidomide and Dexamethasone in Mantle Cell Lymphoma via Cereblon-dependent Targeting of the IL-6/STAT3/PI3K Axis. EBioMedicine, 2017, 20, 70-78.	6.1	7
308	Novel RAG1 mutation and the occurrence of mycobacterial and Chromobacterium violaceum infections in a case of leaky SCID. Microbial Pathogenesis, 2017, 109, 114-119.	2.9	10
309	Fungal Infections in Primary and Acquired Immunodeficiencies. , 2017, , 1-34.		0
310	Introduction on Primary Immunodeficiency Diseases. , 2017, , 1-81.		3
311	Predominantly Antibody Deficiencies. , 2017, , 183-244.		2
312	Transcriptome analysis of CD34+ cells from myelodysplastic syndrome patients. Leukemia Research, 2017, 62, 40-50.	0.8	1
313	Approach to the Management of Autoimmunity in Primary Immunodeficiency. Scandinavian Journal of Immunology, 2017, 85, 13-29.	2.7	35

#	Article	IF	Citations
314	Selective IgA Deficiency: Epidemiology, Pathogenesis, Clinical Phenotype, Diagnosis, Prognosis and Management. Scandinavian Journal of Immunology, 2017, 85, 3-12.	2.7	139
315	Evaluation of infectious and non-infectious complications in patients with primary immunodeficiency. Central-European Journal of Immunology, 2017, 42, 336-341.	1.2	45
316	Flow Cytometry Assays in Primary Immunodeficiency Diseases. Methods in Molecular Biology, 2018, 1678, 321-345.	0.9	8
317	Gastrointestinal manifestations of primary immune deficiencies in children. International Reviews of Immunology, 2018, 37, 111-118.	3.3	4
318	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
319	Intra cranial granulomatous disease in common variable immunodeficiency: Case series and review of the literature. Seminars in Arthritis and Rheumatism, 2018, 47, 890-896.	3.4	16
320	Common Variable Immunodeficiency and Liver Involvement. Clinical Reviews in Allergy and Immunology, 2018, 55, 340-351.	6.5	58
321	An Uncommon Cause of Bronchiectasis: A Case Report. Journal of Pulmonary & Respiratory Medicine, 2018, 08, .	0.1	0
322	Cytomegalovirus Retinitis in Primary Immune Deficiency Disease. Case Reports in Ophthalmological Medicine, 2018, 2018, 1-5.	0.5	6
323	Hypogammaglobulinemia Observed One Year after Rituximab Treatment for Idiopathic Thrombocytopenic Purpura. Case Reports in Medicine, 2018, 2018, 1-3.	0.7	2
324	Gastrointestinal Manifestations in Children with Primary Immunodeficiencies: Single Center: 12 Years Experience. Digestive Diseases, 2019, 37, 45-52.	1.9	12
325	Non-parametric Heat Map Representation of Flow Cytometry Data: Identifying Cellular Changes Associated With Genetic Immunodeficiency Disorders. Frontiers in Immunology, 2019, 10, 2134.	4.8	8
326	The olfactory function in patients with common variable immunodeficiency. European Annals of Otorhinolaryngology, Head and Neck Diseases, 2019, 136, 429-434.	0.7	3
328	Autoimmunity in Primary Immunodeficiency Disorders. , 2020, , 513-532.		1
329	The influence of hospitalâ€based intravenous immunoglobulin and homeâ€based selfâ€administrated subcutaneous immunoglobulin therapy in young children with primary immunodeficiency diseases on their parentsâ€™ / caregiversâ€™ satisfaction. Pediatrics International, 2020, 62, 316-318.	0.5	4
330	Trained Immunity Based-Vaccines as a Prophylactic Strategy in Common Variable Immunodeficiency. A Proof of Concept Study. Biomedicines, 2020, 8, 203.	3.2	17
331	Autoimmune Hepatitis in a Patient With Common Variable Immunodeficiency. ACG Case Reports Journal, 2020, 7, e00467.	0.4	5
332	Common variable immunodeficiency with granulomatous-lymphocytic interstitial lung disease and preceding neurological involvement: a case-report. BMC Pulmonary Medicine, 2020, 20, 205.	2.0	5

#	Article	IF	Citations
333	Recurrent Infections and Rash in a 6 Year Old Girl: A Case Report of Congenital C3 Defeciency. Journal of Bangladesh College of Physicians & Surgeons, 2020, 38, 101-104.	0.1	0
334	Variable immunodeficiency study: Evaluation of two European cohorts within a variety of clinical phenotypes. Immunology Letters, 2020, 223, 78-88.	2.5	5
335	Pulmonary Manifestations of Immunodeficiency and Immunosuppressive Diseases Other than Human Immunodeficiency Virus. Pediatric Clinics of North America, 2021, 68, 103-130.	1.8	2
336	Oxidative stress in common variable immunodeficiency. European Journal of Inflammation, 2021, 19, 205873922110024.	0.5	2
337	Predominantly Antibody Deficiencies. , 2021, , .		0
338	Immunodeficiency and thymoma in Good syndrome: Two sides of the same coin. Immunology Letters, 2021, 231, 11-17.	2.5	21
339	Involvement of Variants in the Genes Encoding BRCA1-Associated Genome Surveillance Complex (BASC) in the Development of Human Common Diseases. Molecular Biology, 2021, 55, 278-296.	1.3	5
340	CLINICAL FEATURES, AGE AND SEX DISTRIBUTIONS, RISK FACTORS AND THE TYPE OF BACTERIA ISOLATED IN PERIODONTITIS PATIENTS IN SANA'A, YEMEN. Universal Journal of Pharmaceutical Research, 0, , .	0.1	3
341	Inborn Errors of Immunity and Cancer. Biology, 2021, 10, 313.	2.8	13
342	Common Variable Immunodeficiency and Autoimmune Diseases: A Retrospective Study of 95 Adult Patients in a Single Tertiary Care Center. Frontiers in Immunology, 2021, 12, 652487.	4.8	27
343	Inflammatory bowel disease in primary immunodeficiency disorders is a heterogeneous clinical entity requiring an individualized treatment strategy: A systematic review. Autoimmunity Reviews, 2021, 20, 102872.	5.8	7
344	B-Cell Immunophenotyping to Predict Vaccination Outcome in the Immunocompromised - A Systematic Review. Frontiers in Immunology, 2021, 12, 690328.	4.8	12
345	Severe Combined Immunodeficiency Disorder due to a Novel Mutation in Recombination Activation Gene 2: About 2 Cases. Case Reports in Immunology, 2021, 2021, 1-5.	0.4	3
347	Gastrointestinal Manifestations of Immunodeficiency. , 2021, , 429-450.e7.		0
349	Molecular Diagnostics of Primary Immunodeficiencies: Benefits and Future Challenges. Advances in Experimental Medicine and Biology, 2009, 634, 231-241.	1.6	6
350	Gene Correction of Induced Pluripotent Stem Cells Derived from a Murine Model of X-Linked Chronic Granulomatous Disorder. Methods in Molecular Biology, 2014, 1114, 427-440.	0.9	1
351	An Introduction to Primary Immunodeficiency Diseases. , 2008, , 1-38.		11
352	Other Well-Defined Immunodeficiencies. , 2008, , 251-290.		5

#	Article	IF	Citations
353	Autoimmune Hepatitis Type 2 in a Child With IPEX Syndrome. Journal of Pediatric Gastroenterology and Nutrition, 2011, 53, 690-693.	1.8	29
354	CD81 gene defect in humans disrupts CD19 complex formation and leads to antibody deficiency. Journal of Clinical Investigation, 2010, 120, 1265-1274.	8.2	345
355	Major approaches in early diagnostics of common variable immunodeficiency in adults in Moscow. F1000Research, 2012, 1, 46.	1.6	2
356	Efficacy, safety, and tolerability of IVIG-SN in patients with primary immunodeficiency. LymphoSign Journal, 2015, 2, 21-29.	0.2	3
357	Immunodeficiencies with hypergammaglobulinemia: a review. LymphoSign Journal, 2015, 2, 57-73.	0.2	11
359	Cervical lymph node diseases in children. GMS Current Topics in Otorhinolaryngology, Head and Neck Surgery, 2014, 13, Doc08.	0.8	22
360	Hematopoietic cell transplantation for treatment of primary immune deficiencies. Cellular Therapy and Transplantation, 2010, 2, .	0.3	11
361	Isolation of poliovirus shedding following vaccination in children with antibody deficiency disorders. Journal of Infection in Developing Countries, 2012, 6, 881-885.	1.2	6
362	Evaluation of the Child with Immunodeficiency Disorder. Pediatric Annals, 2011, 40, 205-211.	0.8	2
363	Host Defence against Bacterial Biofilms: â€œMission Impossibleâ€?. ISRN Immunology, 2012, 2012, 1-17.	0.7	31
364	Severe combined immunodeficiency pig as an emerging animal model for human diseases and regenerative medicines. BMB Reports, 2019, 52, 625-634.	2.4	17
367	New Findings in Primary Immunodeficiency. Advances in Experimental Medicine and Biology, 2009, 634, 79-93.	1.6	0
368	å…ç–«ä¸å…¨ã•ã‚ã‚¬ãƫ¼ â€•ç‰¹ã•ï¼Œè‡ªç„¶å…ç–«ç³»ã®å^†åã®ã‚¿ãƳã‚'ã‚¯æ§‹é€ ”Ÿç‰©å¦ çš„è¦–ç‚¹ã•ã‰â€•. Nihon Shon 2009, 23, 203-211.	0.2	0
369	Blutzellen (HÃmatologie). , 2009, , 83-117.		0
370	PRIMARY IMMUNODEFICIENCIES. , 2009, , 1021-1037.		0
372	Immunologie. , 2009, , 299-311.		0
373	Genetic Predispositions for Hematologic and Lymphoid Disorders. Molecular Pathology Library, 2010, , 21-64.	0.1	0
374	White Blood Cell and Immunodeficiency Disorders. Molecular Pathology Library, 2010, , 499-509.	0.1	0

#	Article	IF	Citations
375	Does Advoy.comâ„¢ help patients with primary immunodeficiency diseases experience that their life situation has improved?. Revista Electronica De Comunicacao, Informacao & Inovacao Em Saude: RECIIS, 2010, 4, .	0.2	1
377	IPEX Syndrome: Clinical Profile, Biological Features, and Current Treatment. , 2011, , 129-142.		1
378	IMMUNODEFICIENCIES AND THE RHEUMATIC DISEASES. , 2011, , 627-641.		0
379	Primary Combined Antibody and Cellular Immunodeficiencies. , 2011, , 730-738.e1.		1
380	Frequent infections. , 2011, , 188-193.		0
384	T-Cell Immune Defects. , 2012, , 1297-1306.		0
385	Miscellaneous Conditions Associated with Arthritis. , 2012, , 1615-1622.		0
387	Innate Immune Defects. , 2012, , 1275-1284.		0
388	Immunodeficienze congenite e acquisite. , 2012, , 91-100.		0
389	Clinical Outcome of Children with Post Bacillus Calmette and Guerin Vaccination Complications: A Single Center Experience. World Journal of Vaccines, 2012, 02, 50-54.	0.8	0
390	Primary Immunodeficiency Diseases in Latin America: Epidemiology and Perspectives. , 0, , .		0
392	Immunodeficiency and Sinus Disease. , 2014, , 223-245.		0
393	FcÎ³ receptor polymorphisms in patients with transient hypogammaglobulinemia of infancy presenting mild and severe infections. Asian Pacific Journal of Allergy and Immunology, 2015, 33, 312-9.	0.4	0
394	Immunodysregulation, Polyendocrinopathy, and Enteropathy, X-Linked (IPEX) Syndrome. , 2016, , 444-450.		0
395	Phagocytes Defects. , 2017, , 245-294.		3
396	Combined T- and B-Cell Immunodeficiencies. , 2017, , 83-182.		3
398	Health-related quality of life in patients with common variable immunodeficiency switching from intravenous to subcutaneous immunoglobulin therapy. The Egyptian Journal of Internal Medicine, 2017, 29, 77-82.	0.9	0
399	Immunoglobulin A and its selective deficiency. Kazan Medical Journal, 2017, 98, 809-813.	0.2	4

#	Article	IF	Citations
400	Modern view on a problem Â«children with recurrent infectionsÂ» (frequently ill children) and their immune status. Medical Herald of the South of Russia, 2018, 9, 37-43.	0.4	2
401	Inborn errors of immunity: The missing link in infectious diseases susceptibility. Jammi, 2019, 4, 51-54.	0.5	2
402	The Expanding Spectrum of Primary Immune Defects. Pediatric Annals, 2019, 48, e489-e494.	0.8	0
403	An immune globulin intravenous (human), 10% liquid privigen: for the treatment of primary immunodeficiency diseases. P and T, 2009, 34, 2-21.	0.9	1
404	Approach to the child with recurrent infections. Journal of Family and Community Medicine, 2009, 16, 77-82.	1.1	8
405	History of primary immunodeficiency diseases in iran. Iranian Journal of Pediatrics, 2010, 20, 16-34.	0.3	13
406	Immunocompromised patients: Review of the most common infections happened in 446 hospitalized patients. Journal of Research in Medical Sciences, 2014, 19, S71-3.	0.9	12
407	Primary immunodeficiency in children: report of seven years study. Tanaffos, 2011, 10, 38-43.	0.5	0
408	Costs of Hospital Admission on Primary Immunodeficiency Diseases. Iranian Journal of Public Health, 2017, 46, 342-350.	0.5	2
410	Early diagnostics of severe combined immunodeficiencies. Bukovinian Medical Herald, 2014, 18, .	0.1	0
411	Immunogenetic disorders: treatment with phytomedicines. , 2022, , 89-130.		1
412	Genome Wide Differential Expression Profiles in Nevus Sebaceous Uncovered Low Expression of CDKN2AIP and Construction of a ceRNA Network. Clinical, Cosmetic and Investigational Dermatology, 2022, Volume 15, 519-533.	1.8	0
428	The X in seX-biased immunity and autoimmune rheumatic disease. Journal of Experimental Medicine, 2022, 219, .	8.5	18
429	Immune globulin therapy and kidney disease: Overview and screening, monitoring, and management recommendations. American Journal of Health-System Pharmacy, 2022, 79, 1415-1423.	1.0	1
430	Common variable immunodeficiency in a patient with suspected sarcoidosis. Pneumonologia I Alergologia Polska, 2008, 77, 91-97.	0.6	0
431	Monitoring Cellular Immune Responses after Consumption of Selected Probiotics in Immunocompromised Mice. Food Science of Animal Resources, 2022, 42, 903-914.	4.1	7
432	Biomarkers and their Clinical Applications in Pediatrics. , 2022, , 450-489.		0
433	ImunodeficiÃªncia de imunoglobulinas IgG subclasse 3. ServiÃ§o Social E SaÃºde, 2022, 21, e022005.	0.0	0

#	Article	IF	Citations
434	Performance status and deaths among children registered in Kuwait National Primary Immunodeficiency Disorders Registry. Asian Pacific Journal of Allergy and Immunology, 2010, , .	0.4	5
435	CRISPR/Cas9-mediated gene editing. A promising strategy in hematological disorders. Cytotherapy, 2023, 25, 277-285.	0.7	4
436	The Canadian Society for Immunologyâ€™s 34th Annual Meeting 2022: Symposia Mini Review. Journal of Leukocyte Biology, 0, , .	3.3	0
437	Evaluation of immunological abnormalities in patients with rare syndromes. Central-European Journal of Immunology, 2022, 47, 299-307.	1.2	0
438	Immunityâ€onâ€aâ€Chip: Integration of Immune Components into the Scheme of Organâ€onâ€aâ€Chip Systems. Advanced Biology, 2023, 7, .	2.5	0
439	Reduced numbers of naÃ¯ve CD4Â+ÂT cells and an altered CD4/CD8 balance in depressed common variable immune deficiency (CVID) patients. Is thymosin-Î±1 a possible treatment?. International Immunopharmacology, 2023, 119, 110168.	3.8	1
440	Overview of microbial therapeutics in immunological disorders. , 2023, , 289-353.		1
442	Case Report: Mycosis fungoides as an exclusive manifestation of common variable immunodeficiency in a family with a NFKB2 gene mutation. Frontiers in Oncology, 0, 13, .	2.8	1
443	Norovirus infection among HIV-infected patients in Abuja, Nigeria: impact of combination antiretroviral therapy status. BMC Infectious Diseases, 2023, 23, .	2.9	0
444	Wholeâ€exome sequencing to identify undiagnosed primary immunodeficiency disorders in children with communityâ€acquired sepsis, admitted in the pediatric intensive care unit. Pediatric Allergy and Immunology, 2023, 34, .	2.6	0
445	From variant of uncertain significance to likely pathogenic in two siblings with atypical RAG2 Deficiency: a case report and review of the literature. BMC Pediatrics, 2024, 24, .	1.7	0
446	The conneXion between sex and immune responses. Nature Reviews Immunology, 0, , .	22.7	0