Asghar Aghamohammadi

List of Publications by Year
in descending order

291 papers	8,530 citations	41323 49 h-index	74108 75 g-index
296 all docs	296 docs citations	296 times ranked	6927 citing authors

#	Article	IF	Citations
1	Deleterious Mutations in LRBA Are Associated with a Syndrome of Immune Deficiency and Autoimmunity. American Journal of Human Genetics, 2012, 90, 986-1001.	2.6	452
2	IgA Deficiency: Correlation Between Clinical and Immunological Phenotypes. Journal of Clinical Immunology, 2009, 29, 130-136.	2.0	191
3	Spectrum of Phenotypes Associated with Mutations in LRBA. Journal of Clinical Immunology, 2016, 36, 33-45.	2.0	180
4	Home-Based Subcutaneous Immunoglobulin Versus Hospital-Based Intravenous Immunoglobulin in Treatment of Primary Antibody Deficiencies: Systematic Review and Meta Analysis. Journal of Clinical Immunology, 2012, 32, 1180-1192.	2.0	147
5	Bacillus Calmette-GuÃ©rin (BCG) complications associated with primary immunodeficiency diseases. Journal of Infection, 2012, 64, 543-554.	1.7	145
6	Primary immunodeficiency diseases associated with increased susceptibility to viral infections and malignancies. Journal of Allergy and Clinical Immunology, 2011, 127, 1329-1341.e2.	1.5	140
7	Frequency and Clinical Manifestations of Patients with Primary Immunodeficiency Disorders in Iran: Update from the Iranian Primary Immunodeficiency Registry. Journal of Clinical Immunology, 2006, 26, 519-532.	2.0	139
8	Progression of Selective IgA Deficiency to Common Variable Immunodeficiency. International Archives of Allergy and Immunology, 2008, 147, 87-92.	0.9	138
9	Disease Evolution and Response to Rapamycin in Activated Phosphoinositide 3-Kinase Î´ Syndrome: The European Society for Immunodeficiencies-Activated Phosphoinositide 3-Kinase Î´ Syndrome Registry. Frontiers in Immunology, 2018, 9, 543.	2.2	137
10	Combined immunodeficiency and Epstein-Barr virusâ€“induced B cell malignancy in humans with inherited CD70 deficiency. Journal of Experimental Medicine, 2017, 214, 91-106.	4.2	134
11	Ataxiaâ€telangiectasia: A review of clinical features and molecular pathology. Pediatric Allergy and Immunology, 2019, 30, 277-288.	1.1	121
12	Gastrointestinal Manifestations in Patients with Common Variable Immunodeficiency. Digestive Diseases and Sciences, 2007, 52, 2977-2983.	1.1	113
13	Long-term outcome of LRBA deficiency in 76 patients after various treatment modalities as evaluated by the immune deficiency and dysregulation activity (IDDA) score. Journal of Allergy and Clinical Immunology, 2020, 145, 1452-1463.	1.5	112
14	Novel mutations in TNFRSF7/CD27: Clinical, immunologic, and genetic characterization of human CD27 deficiency. Journal of Allergy and Clinical Immunology, 2015, 136, 703-712.e10.	1.5	109
15	Long-term outcomes of 176 patients with X-linked hyper-IgM syndrome treated with or without hematopoietic cell transplantation. Journal of Allergy and Clinical Immunology, 2017, 139, 1282-1292.	1.5	107
16	Clinical and Immunological Features of 65 Iranian Patients with Common Variable Immunodeficiency. Vaccine Journal, 2005, 12, 825-832.	3.2	106
17	Primary Immunodeficiency Disorders in Iran: Update and New Insights from the Third Report of the National Registry. Journal of Clinical Immunology, 2014, 34, 478-490.	2.0	99
18	Xâ€linked lymphoproliferative syndrome: a genetic condition typified by the triad of infection, immunodeficiency and lymphoma. British Journal of Haematology, 2011, 152, 13-30.	1.2	92

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19	A hypomorphic recombination-activating gene 1 (RAG1) mutation resulting in a phenotype resembling common variable immunodeficiency. Journal of Allergy and Clinical Immunology, 2014, 134, 1375-1380.	1.5	91
20	Clinical, immunologic, and genetic spectrum of 696 patients with combined immunodeficiency. Journal of Allergy and Clinical Immunology, 2018, 141, 1450-1458.	1.5	90
21	RAC2 loss-of-function mutation in 2 siblings with characteristics of common variable immunodeficiency. Journal of Allergy and Clinical Immunology, 2015, 135, 1380-1384.e5.	1.5	89
22	Clinical, Immunologic, and Molecular Spectrum of Patients with LPS-Responsive Beige-Like Anchor Protein Deficiency: A Systematic Review. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 2379-2386.e5.	2.0	88
23	Fourth Update on the Iranian National Registry of Primary Immunodeficiencies: Integration of Molecular Diagnosis. Journal of Clinical Immunology, 2018, 38, 816-832.	2.0	86
24	Clinical implications of systematic phenotyping and exome sequencing in patients with primary antibody deficiency. Genetics in Medicine, 2019, 21, 243-251.	1.1	86
25	Consanguinity in Primary Immunodeficiency Disorders; the Report from Iranian Primary Immunodeficiency Registry. American Journal of Reproductive Immunology, 2006, 56, 145-151.	1.2	85
26	Clinical, Immunological, and Genetic Features in Patients with Activated PI3KÎ´ Syndrome (APDS): a Systematic Review. Clinical Reviews in Allergy and Immunology, 2020, 59, 323-333.	2.9	79
27	Characterization of the clinical and immunologic phenotype and management of 157 individuals with 56 distinct heterozygous NFKB1 mutations. Journal of Allergy and Clinical Immunology, 2020, 146, 901-911.	1.5	78
28	Primary immunodeficiency in Iran: first report of the National Registry of PID in Children and Adults. Journal of Clinical Immunology, 2002, 22, 375-380.	2.0	77
29	Chronic Granulomatous Disease: A Clinical Survey of 41 Patients from the Iranian Primary Immunodeficiency Registry. International Archives of Allergy and Immunology, 2004, 134, 253-259.	0.9	76
30	Initial presenting manifestations in 16,486 patients with inborn errors of immunity include infections and noninfectious manifestations. Journal of Allergy and Clinical Immunology, 2021, 148, 1332-1341.e5.	1.5	75
31	Global systematic review of primary immunodeficiency registries. Expert Review of Clinical Immunology, 2020, 16, 717-732.	1.3	74
32	A review on guidelines for management and treatment of common variable immunodeficiency. Expert Review of Clinical Immunology, 2013, 9, 561-575.	1.3	72
33	Clinical, immunologic, molecular analyses and outcomes of iranian patients with <scp>LRBA</scp> deficiency: A longitudinal study. Pediatric Allergy and Immunology, 2017, 28, 478-484.	1.1	65
34	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.	2.0	64
35	Extended clinical and immunological phenotype and transplant outcome in CD27 and CD70 deficiency. Blood, 2020, 136, 2638-2655.	0.6	64
36	Ataxia-Telangiectasia in Iran: Clinical and Laboratory Features of 104 Patients. Pediatric Neurology, 2007, 37, 21-28.	1.0	62

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37	The hyper IgM syndromes: Epidemiology, pathogenesis, clinical manifestations, diagnosis and management. Clinical Immunology, 2019, 198, 19-30.	1.4	62
38	Lymphoma of mucosa-associated lymphoid tissue in common variable immunodeficiency. Leukemia and Lymphoma, 2006, 47, 343-346.	0.6	60
39	Comparison of pulmonary diseases in common variable immunodeficiency and Xâ€linked agammaglobulinaemia. Respirology, 2010, 15, 289-295.	1.3	60
40	Cytokines in Common Variable Immunodeficiency as Signs of Immune Dysregulation and Potential Therapeutic Targets â€“ A Review of the Current Knowledge. Journal of Clinical Immunology, 2014, 34, 524-543.	2.0	59
41	Efficacy of intravenous immunoglobulin on the prevention of pneumonia in patients with agammaglobulinemia. FEMS Immunology and Medical Microbiology, 2004, 40, 113-118.	2.7	58
42	Neutropenia and Primary Immunodeficiency Diseases. International Reviews of Immunology, 2009, 28, 335-366.	1.5	58
43	Autoimmunity in common variable immunodeficiency: epidemiology, pathophysiology and management. Expert Review of Clinical Immunology, 2017, 13, 101-115.	1.3	55
44	IgA Deficiency and the MHC: Assessment of Relative Risk and Microheterogeneity Within the HLA A1 B8, DR3 (8.1) Haplotype. Journal of Clinical Immunology, 2010, 30, 138-143.	2.0	54
45	Clinical, Immunological and Molecular Characteristics of 37 Iranian Patients with X-Linked Agammaglobulinemia. International Archives of Allergy and Immunology, 2006, 141, 408-414.	0.9	52
46	Chromosomal radiosensitivity in patients with common variable immunodeficiency. Immunobiology, 2008, 213, 447-454.	0.8	52
47	Analysis of Switched Memory B Cells in Patients with IgA Deficiency. International Archives of Allergy and Immunology, 2011, 156, 462-468.	0.9	52
48	Primary Immunodeficiency Diseases and Bacillus Calmette-GuÃ©rin (BCG)-Vaccineâ€“Derived Complications: A Systematic Review. Journal of Allergy and Clinical Immunology: in Practice, 2020, 8, 1371-1386.	2.0	51
49	Patients with Primary Immunodeficiencies Are a Reservoir of Poliovirus and a Risk to Polio Eradication. Frontiers in Immunology, 2017, 8, 685.	2.2	50
50	Effect of regular intravenous immunoglobulin therapy on prevention of pneumonia in patients with common variable immunodeficiency. Journal of Microbiology, Immunology and Infection, 2006, 39, 114-20.	1.5	50
51	Xâ€Linked Agammaglobulinemia: A Survey of 33 Iranian Patients. Immunological Investigations, 2004, 33, 81-93.	1.0	49
52	Clinical and Laboratory Findings in Hyper-IgM Syndrome with Novel CD40L and AICDA Mutations. Journal of Clinical Immunology, 2009, 29, 769-776.	2.0	48
53	Novel Mutations in TACI (TNFRSF13B) Causing Common Variable Immunodeficiency. Journal of Clinical Immunology, 2009, 29, 777-785.	2.0	48
54	Evaluation of CD4+CD25+FOXP3+ regulatory T cells function in patients with common variable immunodeficiency. Cellular Immunology, 2013, 281, 129-133.	1.4	48

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55	Overweight and obesity and their associated factors in adolescents in Tehran, Iran, 2004â€“2005. European Journal of Pediatrics, 2006, 165, 489-493.	1.3	47
56	Clinical and Laboratory Findings in Iranian Patients with Leukocyte Adhesion Deficiency (Study of 15) Tj ETQq0 0 0 rgBT /Overlock 10 Tf	2.0	47
57	Malignancy in common variable immunodeficiency: a systematic review and meta-analysis. Expert Review of Clinical Immunology, 2019, 15, 1105-1113.	1.3	47
58	Indications and safety of intravenous and subcutaneous immunoglobulin therapy. Expert Review of Clinical Immunology, 2011, 7, 301-316.	1.3	46
59	Evaluation of infectious and non-infectious complications in patients with primary immunodeficiency. Central-European Journal of Immunology, 2017, 42, 336-341.	0.4	45
60	Mortality and Morbidity in Common Variable Immunodeficiency. Journal of Tropical Pediatrics, 2006, 53, 32-38.	0.7	44
61	Primary Immunodeficiency Diseases Associated with Neurologic Manifestations. Journal of Clinical Immunology, 2012, 32, 1-24.	2.0	42
62	Severe combined immunodeficiency: A cohort of 40 patients. Pediatric Allergy and Immunology, 2008, 19, 303-306.	1.1	41
63	Vaccine-associated Paralytic Poliomyelitis in Immunodeficient Children, Iran, 1995â€“2008. Emerging Infectious Diseases, 2010, 16, 1133-1136.	2.0	40
64	Autoimmunity in Primary Antibody Deficiencies. International Archives of Allergy and Immunology, 2016, 171, 180-193.	0.9	40
65	Cellular and molecular mechanisms of immune dysregulation and autoimmunity. Cellular Immunology, 2016, 310, 14-26.	1.4	39
66	The Clinical, Immunohematological, and Molecular Study of Iranian Patients with Severe Congenital Neutropenia. Journal of Clinical Immunology, 2007, 27, 525-533.	2.0	38
67	Serum Bactericidal Antibody Responses to Meningococcal Polysaccharide Vaccination as a Basis for Clinical Classification of Common Variable Immunodeficiency. Vaccine Journal, 2008, 15, 607-611.	3.2	38
68	Characterization of 11 New Cases of Leukocyte Adhesion Deficiency Type 1 with Seven Novel Mutations in the ITGB2 Gene. Journal of Clinical Immunology, 2010, 30, 756-760.	2.0	38
69	Autoimmunity in a cohort of 471 patients with primary antibody deficiencies. Expert Review of Clinical Immunology, 2017, 13, 1099-1106.	1.3	38
70	Clinical phenotype classification for selective immunoglobulin A deficiency. Expert Review of Clinical Immunology, 2015, 11, 1245-1254.	1.3	37
71	Inflammation, a significant player of Ataxiaâ€“Telangiectasia pathogenesis?. Inflammation Research, 2018, 67, 559-570.	1.6	37
72	Comparison of Common Monogenic Defects in a Large Predominantly Antibody Deficiency Cohort. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 864-878.e9.	2.0	37

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73	Abnormality of regulatory T cells in common variable immunodeficiency. Cellular Immunology, 2017, 315, 11-17.	1.4	36
74	Ataxia telangiectasia syndrome: moonlighting ATM. Expert Review of Clinical Immunology, 2017, 13, 1155-1172.	1.3	36
75	Adverse reactions of prophylactic intravenous immunoglobulin infusions in Iranian patients with primary immunodeficiency. Annals of Allergy, Asthma and Immunology, 2004, 92, 60-64.	0.5	35
76	Genotype-Phenotype Correlation in Bruton's Tyrosine Kinase Deficiency. Journal of Pediatric Hematology/Oncology, 2008, 30, 679-683.	0.3	35
77	Economic burden of common variable immunodeficiency: annual cost of disease. Expert Review of Clinical Immunology, 2015, 11, 681-688.	1.3	35
78	Role of apoptosis in common variable immunodeficiency and selective immunoglobulin A deficiency. Molecular Immunology, 2016, 71, 1-9.	1.0	35
79	New insights into physiopathology of immunodeficiency-associated vaccine-derived poliovirus infection; systematic review of over 5 decades of data. Vaccine, 2018, 36, 1711-1719.	1.7	35
80	International retrospective study of allogeneic hematopoietic cell transplantation for activated PI3K-delta syndrome. Journal of Allergy and Clinical Immunology, 2022, 149, 410-421.e7.	1.5	34
81	Association of IL-4 and IL-10 gene promoter polymorphisms with common variable immunodeficiency. Immunobiology, 2010, 215, 81-87.	0.8	33
82	Consensus Middle East and North Africa Registry on Inborn Errors of Immunity. Journal of Clinical Immunology, 2021, 41, 1339-1351.	2.0	33
83	Evaluation of Natural Regulatory T Cells in Subjects with Selective IgA Deficiency: From Senior Idea to Novel Opportunities. International Archives of Allergy and Immunology, 2013, 160, 208-214.	0.9	32
84	Bronchiectasis in common variable immunodeficiency: A systematic review and metaâ€analysis. Pediatric Pulmonology, 2020, 55, 292-299.	1.0	32
85	Identification of anSH2D1A mutation in a hypogammaglobulinemic male patient with a diagnosis of common variable immunodeficiency. International Journal of Hematology, 2003, 78, 45-47.	0.7	30
86	Congenital Neutropenia and Primary Immunodeficiency Disorders. Journal of Pediatric Hematology/Oncology, 2005, 27, 351-356.	0.3	30
87	Monogenic mutations associated with IgA deficiency. Expert Review of Clinical Immunology, 2016, 12, 1321-1335.	1.3	30
88	Clinical, Immunological, and Genetic Features in 49 Patients With ZAP-70 Deficiency: A Systematic Review. Frontiers in Immunology, 2020, 11, 831.	2.2	29
89	Serum bactericidal antibody response to serogroup C polysaccharide meningococcal vaccination in children with primary antibody deficiencies. Vaccine, 2007, 25, 5308-5314.	1.7	28
90	Is there a need to redefine the diagnostic criteria for common variable immunodeficiency?. Expert Review of Clinical Immunology, 2014, 10, 1-5.	1.3	28

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91	The use of Immunoglobulin Therapy in Primary Immunodeficiency Diseases. Endocrine, Metabolic and Immune Disorders - Drug Targets, 2016, 16, 80-88.	0.6	28
92	Health-related quality of life in primary antibody deficiency. Iranian Journal of Allergy, Asthma and Immunology, 2011, 10, 47-51.	0.3	28
93	Increased Serum Levels of Soluble CD30 in Patients with Common Variable Immunodeficiency and Its Clinical Implications. Journal of Clinical Immunology, 2008, 28, 78-84.	2.0	27
94	Serum Bactericidal Antibody Response 1 Year after Meningococcal Polysaccharide Vaccination of Patients with Common Variable Immunodeficiency. Vaccine Journal, 2010, 17, 524-528.	3.2	27
95	TNF-alpha single nucleotide polymorphisms in atopic dermatitis. European Cytokine Network, 2012, 23, 163-165.	1.1	27
96	Novel mutation of the activation-induced cytidine deaminase gene in a Tajik family: special review on hyper-immunoglobulin M syndrome. Expert Review of Clinical Immunology, 2012, 8, 539-546.	1.3	27
97	Impaired Akt phosphorylation in B-cells of patients with common variable immunodeficiency. Clinical Immunology, 2017, 175, 124-132.	1.4	27
98	Ataxia-telangiectasia: epidemiology, pathogenesis, clinical phenotype, diagnosis, prognosis and management. Expert Review of Clinical Immunology, 2020, 16, 859-871.	1.3	27
99	Immunologic evaluation of patients with recurrent ear, nose, and throat infections. American Journal of Otolaryngology - Head and Neck Medicine and Surgery, 2008, 29, 385-392.	0.6	26
100	Immunoglobulin class switch recombination deficiency type 1 or CD40 ligand deficiency: from bedside to bench and back again. Expert Review of Clinical Immunology, 2014, 10, 91-105.	1.3	26
101	Important differences in the diagnostic spectrum of primary immunodeficiency in adults versus children. Expert Review of Clinical Immunology, 2015, 11, 289-302.	1.3	26
102	Measurement of Health-Related Quality of Life in Primary Antibody-Deficient Patients. Immunological Investigations, 2017, 46, 329-340.	1.0	26
103	Blood pressure nomograms for school children in Iran. Pediatric Nephrology, 2004, 19, 164-168.	0.9	25
104	Different brands of intravenous immunoglobulin for primary immunodeficiencies: how to choose the best option for the patient?. Expert Review of Clinical Immunology, 2015, 11, 1229-1243.	1.3	25
105	Vaccine-Derived Polioviruses and Children with Primary Immunodeficiency, Iran, 1995â€“2014. Emerging Infectious Diseases, 2016, 22, 1712-1719.	2.0	25
106	Monogenic polyautoimmunity in primary immunodeficiency diseases. Autoimmunity Reviews, 2018, 17, 1028-1039.	2.5	24
107	Alteration in frequency and function of CD4âºCD25âºFOXP3âº regulatory T cells in patients with immune thrombocytopenic purpura. Iranian Journal of Allergy, Asthma and Immunology, 2014, 13, 85-92.	0.3	24
108	Combined immunodeficiency presenting with vaccine-associated paralytic poliomyelitis: a case report and narrative review of literature. Immunological Investigations, 2014, 43, 292-298.	1.0	23

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109	The clinical significance of complete class switching defect in Ataxia telangiectasia patients. Expert Review of Clinical Immunology, 2017, 13, 499-505.	1.3	23
110	The clinical and laboratory survey of Iranian patients with Hyper-IgE syndrome. Scandinavian Journal of Infectious Diseases, 2006, 38, 898-903.	1.5	22
111	B-cellâ€“T-cell activation and interaction in common variable immunodeficiency. Human Immunology, 2010, 71, 355-362.	1.2	22
112	Cohort of Iranian Patients with Congenital Agammaglobulinemia: Mutation Analysis and Novel Gene Defects. Expert Review of Clinical Immunology, 2016, 12, 479-486.	1.3	22
113	Evaluation of Known Defective Signaling-Associated Molecules in Patients Who Primarily Diagnosed as Common Variable Immunodeficiency. International Reviews of Immunology, 2016, 35, 7-24.	1.5	22
114	The imbalance of circulating T helper subsets and regulatory T cells in patients with LRBA deficiency: Correlation with disease severity. Journal of Cellular Physiology, 2018, 233, 8767-8777.	2.0	22
115	Effect of Class Switch Recombination Defect on the Phenotype of Ataxia-Telangiectasia Patients. Immunological Investigations, 2021, 50, 201-215.	1.0	22
116	Antibody response toÂpneumococcal capsular polysaccharide vaccination inÂpatients withÂchronic kidney disease. European Cytokine Network, 2009, 20, 69-74.	1.1	21
117	Cytokine Gene Polymorphisms in Common Variable Immunodeficiency. International Archives of Allergy and Immunology, 2009, 150, 1-7.	0.9	21
118	HLA-DRB1,-DQA1 and -DQB1 Allele and Haplotype Frequencies in Female Patients with Early Onset Breast Cancer. Pathology and Oncology Research, 2012, 18, 49-55.	0.9	21
119	Class Switch Recombination Process in Ataxia Telangiectasia Patients with Elevated Serum Levels of IgM. Journal of Immunoassay and Immunochemistry, 2015, 36, 16-26.	0.5	21
120	Novel Mutation of ZAP-70-related Combined Immunodeficiency: First Case from the National Iranian Registry and Review of the Literature. Immunological Investigations, 2017, 46, 70-79.	1.0	21
121	Polyautoimmunity in Patients with LPS-Responsive Beige-Like Anchor (LRBA) Deficiency. Immunological Investigations, 2018, 47, 457-467.	1.0	21
122	Expanding Clinical Phenotype and Novel Insights into the Pathogenesis of ICOS Deficiency. Journal of Clinical Immunology, 2020, 40, 277-288.	2.0	21
123	Presence of Idiopathic Thrombocytopenic Purpura and autoimmune hemolytic anemia in the patients with common variable immunodeficiency. Iranian Journal of Allergy, Asthma and Immunology, 2008, 7, 169-75.	0.3	21
124	Autoimmunity in common variable immunodeficiency: a systematic review and meta-analysis. Expert Review of Clinical Immunology, 2020, 16, 1227-1235.	1.3	20
125	Health-related quality of life in primary immune deficient patients. Iranian Journal of Allergy, Asthma and Immunology, 2006, 5, 23-7.	0.3	20
126	Common variable immunodeficiency: a heterogeneous group needs further subclassification. Expert Review of Clinical Immunology, 2009, 5, 629-631.	1.3	19

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127	Autoimmunity in X-linked agammaglobulinemia: Kawasaki disease and review of the literature. Expert Review of Clinical Immunology, 2012, 8, 155-159.	1.3	19
128	Long-term evaluation of a historical cohort of Iranian common variable immunodeficiency patients. Expert Review of Clinical Immunology, 2014, 10, 1405-1417.	1.3	19
129	The role of toll-like receptors in B-cell development and immunopathogenesis of common variable immunodeficiency. Expert Review of Clinical Immunology, 2016, 12, 195-207.	1.3	19
130	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.	0.9	18
131	Impact of delayed diagnosis in children with primary antibody deficiencies. Journal of Microbiology, Immunology and Infection, 2011, 44, 229-234.	1.5	18
132	Country Quarantine During COVID-19: Critical or Not?. Disaster Medicine and Public Health Preparedness, 2020, 15, 1-2.	0.7	18
133	Somatic reversion of pathogenic DOCK8 variants alters lymphocyte differentiation and function to effectively cure DOCK8 deficiency. Journal of Clinical Investigation, 2021, 131, .	3.9	18
134	Clinical, immunological, and genetic features in 780 patients with autoimmune lymphoproliferative syndrome (ALPS) and ALPSâ€like diseases: A systematic review. Pediatric Allergy and Immunology, 2021, 32, 1519-1532.	1.1	18
135	Selective immunoglobulin A deficiency in Iranian blood donors: prevalence, laboratory and clinical findings. Iranian Journal of Allergy, Asthma and Immunology, 2008, 7, 157-62.	0.3	18
136	Evaluation of serum IgA levels in Iranian patients with type 1 diabetes mellitus. Acta Diabetologica, 2012, 49, 131-135.	1.2	17
137	Evaluation of class switch recombination in B lymphocytes of patients with common variable immunodeficiency. Journal of Immunological Methods, 2013, 394, 94-99.	0.6	17
138	Managing patients with side effects and adverse events to immunoglobulin therapy. Expert Review of Clinical Pharmacology, 2016, 9, 91-102.	1.3	17
139	IL-10 induces TGF-Î² secretion, TGF-Î² receptor II upregulation, and IgA secretion in B cells. European Cytokine Network, 2019, 30, 107-113.	1.1	17
140	Immunomodulatory Effect of G2013 (Î±-L-Guluronic Acid) on the TLR2 and TLR4 in Human Mononuclear Cells. Current Drug Discovery Technologies, 2018, 15, 123-131.	0.6	17
141	Neutropenia in Iranian patients with primary immunodeficiency disorders. Haematologica, 2005, 90, 554-6.	1.7	17
142	Bilateral basal ganglia involvement in a patient with Griscelli syndrome. European Journal of Paediatric Neurology, 2006, 10, 207-209.	0.7	16
143	The Uncommon Combination of Common Variable Immunodeficiency, Macrophage Activation Syndrome, and Cytomegalovirus Retinitis. Viral Immunology, 2012, 25, 161-165.	0.6	16
144	Autosomal Recessive Agammaglobulinemia: A Novel Non-sense Mutation in CD79a. Journal of Clinical Immunology, 2014, 34, 138-141.	2.0	16

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145	In vitro chromosomal radiosensitivity in patients with common variable immunodeficiency. Central-European Journal of Immunology, 2018, 43, 155-161.	0.4	16
146	Newborn Screening for Presymptomatic Diagnosis of Complement and Phagocyte Deficiencies. Frontiers in Immunology, 2020, 11, 455.	2.2	16
147	Human leukocyte antigens (HLA) associated with selective IgA deficiency in Iran and Sweden. Iranian Journal of Allergy, Asthma and Immunology, 2008, 7, 209-14.	0.3	16
148	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.	0.3	15
149	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.	1.8	15
150	Autoimmunity in primary T-cell immunodeficiencies. Expert Review of Clinical Immunology, 2016, 12, 989-1006.	1.3	15
151	The Heterogeneous Pathogenesis of Selective Immunoglobulin A Deficiency. International Archives of Allergy and Immunology, 2019, 179, 231-246.	0.9	15
152	Variable Abnormalities in T and B Cell Subsets in Ataxia Telangiectasia. Journal of Clinical Immunology, 2021, 41, 76-88.	2.0	15
153	The Clinical and Immunological Features of Patients with Primary Antibody Deficiencies. Endocrine, Metabolic and Immune Disorders - Drug Targets, 2018, 18, 537-545.	0.6	15
154	Diagnostic Approach to the Patients with Suspected Primary Immunodeficiency. Endocrine, Metabolic and Immune Disorders - Drug Targets, 2020, 20, 157-171.	0.6	15
155	Family study of pediatric patients with primary antibody deficiencies. Iranian Journal of Allergy, Asthma and Immunology, 2013, 12, 377-82.	0.3	15
156	Genetic mutations and immunological features of severe combined immunodeficiency patients in Iran. Immunology Letters, 2019, 216, 70-78.	1.1	14
157	Comparison of clinical and immunological features and mortality in common variable immunodeficiency and agammaglobulinemia patients. Immunology Letters, 2019, 210, 55-62.	1.1	14
158	The first cohort of Iranian patients with hyper immunoglobulin E syndrome: A longâ€term followâ€up and genetic analysis. Pediatric Allergy and Immunology, 2019, 30, 469-478.	1.1	14
159	Clinical, immunological and genetic findings in patients with UNC13D deficiency (FHL3): A systematic review. Pediatric Allergy and Immunology, 2021, 32, 186-197.	1.1	14
160	Vitamin D deficiency in chronic idiopathic urticaria. Iranian Journal of Allergy, Asthma and Immunology, 2015, 14, 222-7.	0.3	14
161	Physicians awareness on primary immunodeficiency disorders in Iran. Iranian Journal of Allergy, Asthma and Immunology, 2012, 11, 57-64.	0.3	14
162	Response to polysaccharide vaccination amongst pediatric patients with common variable immunodeficiency correlates with clinical disease. Iranian Journal of Allergy, Asthma and Immunology, 2008, 7, 231-4.	0.3	14

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163	Assessment of Immune Response following Immunization with DTP/Td and MMR Vaccines in Children Treated for Acute Lymphoblastic Leukemia. Pediatric Hematology and Oncology, 2014, 31, 656-663.	0.3	13
164	Adverse reactions toMycobacterium bovisbacille Calmette-GuÃ©rin vaccination against tuberculosis in Iranian children. Clinical and Experimental Vaccine Research, 2015, 4, 195.	1.1	13
165	Monogenic Primary Immunodeficiency Disorder Associated with Common Variable Immunodeficiency and Autoimmunity. International Archives of Allergy and Immunology, 2020, 181, 706-714.	0.9	13
166	History of primary immunodeficiency diseases in iran. Iranian Journal of Pediatrics, 2010, 20, 16-34.	0.1	13
167	LPS-Responsive Beige-Like Anchor Gene Mutation Associated With Possible Bronchiolitis Obliterans Organizing Pneumonia Associated With Hypogammaglobulinemia and Normal IgM Phenotype and Low Number of B Cells. Acta Medica Iranica, 2016, 54, 620-623.	0.8	13
168	Neutropenia in patients with primary antibody deficiency disorders. Iranian Journal of Allergy, Asthma and Immunology, 2004, 3, 77-81.	0.3	13
169	CD40 ligand expression on stimulated T-helper lymphocytes in patients with common variable immunodeficiency. Iranian Journal of Allergy, Asthma and Immunology, 2007, 6, 129-35.	0.3	13
170	Evaluation of humoral immune function in patients with bronchiectasis. Iranian Journal of Allergy, Asthma and Immunology, 2008, 7, 69-77.	0.3	13
171	Neutropenia associated with X-linked Agammaglobulinemia in an Iranian referral center. Iranian Journal of Allergy, Asthma and Immunology, 2009, 8, 43-7.	0.3	13
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