Hassan Abolhassani

List of Publications by Year
in descending order

278 papers	24,154 citations	31902 53 h-index	10708 138 g-index
289 all docs	289 docs citations	289 times ranked	22886 citing authors

#	Article	IF	Citations
1	Global burden of 369 diseases and injuries in 204 countries and territories, 1990â€“2019: a systematic analysis for the Global Burden of Disease Study 2019. Lancet, The, 2020, 396, 1204-1222.	6.3	7,664
2	Global burden of 87 risk factors in 204 countries and territories, 1990â€“2019: a systematic analysis for the Global Burden of Disease Study 2019. Lancet, The, 2020, 396, 1223-1249.	6.3	3,928
3	The global, regional, and national burden of inflammatory bowel disease in 195 countries and territories, 1990â€“2017: a systematic analysis for the Global Burden of Disease Study 2017. The Lancet Gastroenterology and Hepatology, 2020, 5, 17-30.	3.7	1,200
4	Global age-sex-specific fertility, mortality, healthy life expectancy (HALE), and population estimates in 204 countries and territories, 1950â€“2019: a comprehensive demographic analysis for the Global Burden of Disease Study 2019. Lancet, The, 2020, 396, 1160-1203.	6.3	890
5	Cancer Incidence, Mortality, Years of Life Lost, Years Lived With Disability, and Disability-Adjusted Life Years for 29 Cancer Groups From 2010 to 2019. JAMA Oncology, 2022, 8, 420.	3.4	719
6	The global, regional, and national burden of stomach cancer in 195 countries, 1990â€“2017: a systematic analysis for the Global Burden of Disease study 2017. The Lancet Gastroenterology and Hepatology, 2020, 5, 42-54.	3.7	390
7	Five insights from the Global Burden of Disease Study 2019. Lancet, The, 2020, 396, 1135-1159.	6.3	335
8	Measuring universal health coverage based on an index of effective coverage of health services in 204 countries and territories, 1990â€“2019: a systematic analysis for the Global Burden of Disease Study 2019. Lancet, The, 2020, 396, 1250-1284.	6.3	330
9	X-linked recessive TLR7 deficiency in ~1% of men under 60 years old with life-threatening COVID-19. Science Immunology, 2021, 6, .	5.6	267
10	The global, regional, and national burden of oesophageal cancer and its attributable risk factors in 195 countries and territories, 1990â€“2017: a systematic analysis for the Global Burden of Disease Study 2017. The Lancet Gastroenterology and Hepatology, 2020, 5, 582-597.	3.7	241
11	Global, regional, and national progress towards Sustainable Development Goal 3.2 for neonatal and child health: all-cause and cause-specific mortality findings from the Global Burden of Disease Study 2019. Lancet, The, 2021, 398, 870-905.	6.3	229
12	IgA Deficiency: Correlation Between Clinical and Immunological Phenotypes. Journal of Clinical Immunology, 2009, 29, 130-136.	2.0	191
13	Spectrum of Phenotypes Associated with Mutations in LRBA. Journal of Clinical Immunology, 2016, 36, 33-45.	2.0	180
14	Persistence of SARS-CoV-2-specific B and TÂcell responses in convalescent COVID-19 patients 6â€“8Âmonths after the infection. Med, 2021, 2, 281-295.e4.	2.2	153
15	Tuberculosis and impaired IL-23â€“dependent IFN-Î³ immunity in humans homozygous for a common TYK2 missense variant. Science Immunology, 2018, 3, .	5.6	148
16	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
17	Different Aspects of Social Network Analysis. , 2006, , .		144
18	Selective IgA Deficiency: Epidemiology, Pathogenesis, Clinical Phenotype, Diagnosis, Prognosis and Management. Scandinavian Journal of Immunology, 2017, 85, 3-12.	1.3	139

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19	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
20	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
21	Ataxiaâ€telangiectasia: A review of clinical features and molecular pathology. Pediatric Allergy and Immunology, 2019, 30, 277-288.	1.1	121
22	Health system performance in Iran: a systematic analysis for the Global Burden of Disease Study 2019. Lancet, The, 2022, 399, 1625-1645.	6.3	119
23	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
24	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
25	Current genetic landscape in common variable immune deficiency. Blood, 2020, 135, 656-667.	0.6	109
26	Heterologous immunization with inactivated vaccine followed by mRNA-booster elicits strong immunity against SARS-CoV-2 Omicron variant. Nature Communications, 2022, 13, 2670.	5.8	108
27	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
28	Harmony K-means algorithm for document clustering. Data Mining and Knowledge Discovery, 2009, 18, 370-391.	2.4	106
29	SARS-CoV-2â€“related MIS-C: A key to the viral and genetic causes of Kawasaki disease?. Journal of Experimental Medicine, 2021, 218, .	4.2	100
30	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
31	Impact of SARS-CoV-2 Pandemic on Patients with Primary Immunodeficiency. Journal of Clinical Immunology, 2021, 41, 345-355.	2.0	97
32	Measuring routine childhood vaccination coverage in 204 countries and territories, 1980â€“2019: a systematic analysis for the Global Burden of Disease Study 2020, Release 1. Lancet, The, 2021, 398, 503-521.	6.3	93
33	Global, regional, and national mortality among young people aged 10â€“24 years, 1950â€“2019: a systematic analysis for the Global Burden of Disease Study 2019. Lancet, The, 2021, 398, 1593-1618.	6.3	92
34	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
35	Clinical, immunologic, and genetic spectrum of 696 patients with combined immunodeficiency. Journal of Allergy and Clinical Immunology, 2018, 141, 1450-1458.	1.5	90
36	The global burden of adolescent and young adult cancer in 2019: a systematic analysis for the Global Burden of Disease Study 2019. Lancet Oncology, The, 2022, 23, 27-52.	5.1	90

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37	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
38	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
39	Health sector spending and spending on HIV/AIDS, tuberculosis, and malaria, and development assistance for health: progress towards Sustainable Development Goal 3. Lancet, The, 2020, 396, 693-724.	6.3	87
40	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
41	Clinical implications of systematic phenotyping and exome sequencing in patients with primary antibody deficiency. Genetics in Medicine, 2019, 21, 243-251.	1.1	86
42	Global, regional, and national burden of respiratory tract cancers and associated risk factors from 1990 to 2019: a systematic analysis for the Global Burden of Disease Study 2019. Lancet Respiratory Medicine,the, 2021, 9, 1030-1049.	5.2	86
43	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
44	Tracking development assistance for health and for COVID-19: a review of development assistance, government, out-of-pocket, and other private spending on health for 204 countries and territories, 1990â€“2050. Lancet, The, 2021, 398, 1317-1343.	6.3	79
45	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
46	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
47	Global systematic review of primary immunodeficiency registries. Expert Review of Clinical Immunology, 2020, 16, 717-732.	1.3	74
48	A review on guidelines for management and treatment of common variable immunodeficiency. Expert Review of Clinical Immunology, 2013, 9, 561-575.	1.3	72
49	Mapping routine measles vaccination in low- and middle-income countries. Nature, 2021, 589, 415-419.	13.7	71
50	Diabetes mortality and trends before 25 years of age: an analysis of the Global Burden of Disease Study 2019. Lancet Diabetes and Endocrinology,the, 2022, 10, 177-192.	5.5	66
51	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
52	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
53	Extended clinical and immunological phenotype and transplant outcome in CD27 and CD70 deficiency. Blood, 2020, 136, 2638-2655.	0.6	64
54	Next Generation Sequencing Data Analysis in Primary Immunodeficiency Disorders â€“ Future Directions. Journal of Clinical Immunology, 2016, 36, 68-75.	2.0	63

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55	The hyper IgM syndromes: Epidemiology, pathogenesis, clinical manifestations, diagnosis and management. Clinical Immunology, 2019, 198, 19-30.	1.4	62
56	Comparison of pulmonary diseases in common variable immunodeficiency and Xâ€linked agammaglobulinaemia. Respirology, 2010, 15, 289-295.	1.3	60
57	Anemia prevalence in women of reproductive age in low- and middle-income countries between 2000 and 2018. Nature Medicine, 2021, 27, 1761-1782.	15.2	60
58	Recessive inborn errors of type I IFN immunity in children with COVID-19 pneumonia. Journal of Experimental Medicine, 2022, 219, .	4.2	59
59	Immunity to SARS-CoV-2 up to 15Âmonths after infection. IScience, 2022, 25, 103743.	1.9	56
60	Autoimmunity in common variable immunodeficiency: epidemiology, pathophysiology and management. Expert Review of Clinical Immunology, 2017, 13, 101-115.	1.3	55
61	Global, regional, and national sex differences in the global burden of tuberculosis by HIV status, 1990â€“2019: results from the Global Burden of Disease Study 2019. Lancet Infectious Diseases, The, 2022, 22, 222-241.	4.6	53
62	Analysis of Switched Memory B Cells in Patients with IgA Deficiency. International Archives of Allergy and Immunology, 2011, 156, 462-468.	0.9	52
63	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
64	Infectious and Noninfectious Pulmonary Complications in Patients With Primary Immunodeficiency Disorders. Journal of Investigational Allergology and Clinical Immunology, 2017, 27, 213-224.	0.6	50
65	Patients with Primary Immunodeficiencies Are a Reservoir of Poliovirus and a Risk to Polio Eradication. Frontiers in Immunology, 2017, 8, 685.	2.2	50
66	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
67	Evaluation of CD4+CD25+FOXP3+ regulatory T cells function in patients with common variable immunodeficiency. Cellular Immunology, 2013, 281, 129-133.	1.4	48
68	Burden of non-communicable diseases among adolescents aged 10â€“24 years in the EU, 1990â€“2019: a systematic analysis of the Global Burden of Diseases Study 2019. The Lancet Child and Adolescent Health, 2022, 6, 367-383.	2.7	48
69	Malignancy in common variable immunodeficiency: a systematic review and meta-analysis. Expert Review of Clinical Immunology, 2019, 15, 1105-1113.	1.3	47
70	Mapping local patterns of childhood overweight and wasting in low- and middle-income countries between 2000 and 2017. Nature Medicine, 2020, 26, 750-759.	15.2	47
71	Indications and safety of intravenous and subcutaneous immunoglobulin therapy. Expert Review of Clinical Immunology, 2011, 7, 301-316.	1.3	46
72	Evaluation of infectious and non-infectious complications in patients with primary immunodeficiency. Central-European Journal of Immunology, 2017, 42, 336-341.	0.4	45

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73	Clinical, Immunological, and Genetic Features in Patients with Immune Dysregulation, Polyendocrinopathy, Enteropathy, X-linked (IPEX) and IPEX-like Syndrome. Journal of Allergy and Clinical Immunology: in Practice, 2020, 8, 2747-2760.e7.	2.0	45
74	Inherited IFNAR1 Deficiency in a Child with Both Critical COVID-19 Pneumonia and Multisystem Inflammatory Syndrome. Journal of Clinical Immunology, 2022, 42, 471-483.	2.0	44
75	Autoimmunity in Primary Antibody Deficiencies. International Archives of Allergy and Immunology, 2016, 171, 180-193.	0.9	40
76	Frequency and Expression of Inhibitory Markers of <scp>CD</scp>4⁺<scp>CD</scp>25⁺<scp>FOXP</scp>3⁺ Regulatory T Cells in Patients with Common Variable Immunodeficiency. Scandinavian Journal of Immunology, 2013, 77, 405-412.	1.3	39
77	Cellular and molecular mechanisms of immune dysregulation and autoimmunity. Cellular Immunology, 2016, 310, 14-26.	1.4	39
78	Autoimmunity in a cohort of 471 patients with primary antibody deficiencies. Expert Review of Clinical Immunology, 2017, 13, 1099-1106.	1.3	38
79	Clinical phenotype classification for selective immunoglobulin A deficiency. Expert Review of Clinical Immunology, 2015, 11, 1245-1254.	1.3	37
80	Inflammation, a significant player of Ataxiaâ€“Telangiectasia pathogenesis?. Inflammation Research, 2018, 67, 559-570.	1.6	37
81	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
82	Ataxia telangiectasia syndrome: moonlighting ATM. Expert Review of Clinical Immunology, 2017, 13, 1155-1172.	1.3	36
83	Economic burden of common variable immunodeficiency: annual cost of disease. Expert Review of Clinical Immunology, 2015, 11, 681-688.	1.3	35
84	Role of apoptosis in common variable immunodeficiency and selective immunoglobulin A deficiency. Molecular Immunology, 2016, 71, 1-9.	1.0	35
85	Approach to the Management of Autoimmunity in Primary Immunodeficiency. Scandinavian Journal of Immunology, 2017, 85, 13-29.	1.3	35
86	Comparison of various classifications for patients with common variable immunodeficiency (CVID) using measurement of B-cell subsets. Allergologia Et Immunopathologia, 2017, 45, 183-192.	1.0	35
87	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
88	Vaccine breakthrough hypoxemic COVID-19 pneumonia in patients with auto-Abs neutralizing type I IFNs. Science Immunology, 2023, 8, .	5.6	35
89	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
90	X-Linked TLR7 Deficiency Underlies Critical COVID-19 Pneumonia in a Male Patient with Ataxia-Telangiectasia. Journal of Clinical Immunology, 2022, 42, 1-9.	2.0	34

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91	Consensus Middle East and North Africa Registry on Inborn Errors of Immunity. Journal of Clinical Immunology, 2021, 41, 1339-1351.	2.0	33
92	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
93	Epidemiology and pathophysiology of malignancy in common variable immunodeficiency?. Allergologia Et Immunopathologia, 2017, 45, 602-615.	1.0	32
94	Bronchiectasis in common variable immunodeficiency: A systematic review and metaâ€analysis. Pediatric Pulmonology, 2020, 55, 292-299.	1.0	32
95	IL2RG hypomorphic mutation: identification of a novel pathogenic mutation in exon 8 and a review of the literature. Allergy, Asthma and Clinical Immunology, 2019, 15, 2.	0.9	31
96	Monogenic mutations associated with IgA deficiency. Expert Review of Clinical Immunology, 2016, 12, 1321-1335.	1.3	30
97	Clinical, Immunological, and Genetic Features in 49 Patients With ZAP-70 Deficiency: A Systematic Review. Frontiers in Immunology, 2020, 11, 831.	2.2	29
98	The use of Immunoglobulin Therapy in Primary Immunodeficiency Diseases. Endocrine, Metabolic and Immune Disorders - Drug Targets, 2016, 16, 80-88.	0.6	28
99	Health-related quality of life in primary antibody deficiency. Iranian Journal of Allergy, Asthma and Immunology, 2011, 10, 47-51.	0.3	28
100	Impaired Akt phosphorylation in B-cells of patients with common variable immunodeficiency. Clinical Immunology, 2017, 175, 124-132.	1.4	27
101	Autoimmunity and its association with regulatory T cells and B cell subsets in patients with common variable immunodeficiency. Allergologia Et Immunopathologia, 2018, 46, 127-135.	1.0	27
102	Ataxia-telangiectasia: epidemiology, pathogenesis, clinical phenotype, diagnosis, prognosis and management. Expert Review of Clinical Immunology, 2020, 16, 859-871.	1.3	27
103	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
104	Measurement of Health-Related Quality of Life in Primary Antibody-Deficient Patients. Immunological Investigations, 2017, 46, 329-340.	1.0	26
105	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
106	Vaccine-Derived Polioviruses and Children with Primary Immunodeficiency, Iran, 1995â€“2014. Emerging Infectious Diseases, 2016, 22, 1712-1719.	2.0	25
107	Monogenic polyautoimmunity in primary immunodeficiency diseases. Autoimmunity Reviews, 2018, 17, 1028-1039.	2.5	24
108	Development of passive immunity against SARS-CoV-2 for management of immunodeficient patientsâ€”a perspective. Journal of Allergy and Clinical Immunology, 2020, 146, 58-60.	1.5	24

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109	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
110	A survey of complementary and alternative medicine in Iran. Chinese Journal of Integrative Medicine, 2012, 18, 409-416.	0.7	23
111	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
112	The clinical significance of complete class switching defect in Ataxia telangiectasia patients. Expert Review of Clinical Immunology, 2017, 13, 499-505.	1.3	23
113	Impaired respiratory burst contributes to infections in PKCÎ´-deficient patients. Journal of Experimental Medicine, 2021, 218, .	4.2	23
114	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
115	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
116	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
117	Effect of Class Switch Recombination Defect on the Phenotype of Ataxia-Telangiectasia Patients. Immunological Investigations, 2021, 50, 201-215.	1.0	22
118	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
119	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
120	Polyautoimmunity in Patients with LPS-Responsive Beige-Like Anchor (LRBA) Deficiency. Immunological Investigations, 2018, 47, 457-467.	1.0	21
121	Expanding Clinical Phenotype and Novel Insights into the Pathogenesis of ICOS Deficiency. Journal of Clinical Immunology, 2020, 40, 277-288.	2.0	21
122	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
123	Predictive markers for humoral influenza vaccine response in patients with common variable immunodeficiency. Journal of Allergy and Clinical Immunology, 2018, 142, 1922-1931.e2.	1.5	20
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	Autoimmunity in X-linked agammaglobulinemia: Kawasaki disease and review of the literature. Expert Review of Clinical Immunology, 2012, 8, 155-159.	1.3	19
126	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

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127	Impact of delayed diagnosis in children with primary antibody deficiencies. Journal of Microbiology, Immunology and Infection, 2011, 44, 229-234.	1.5	18
128	Review of local herbal compounds found in the Iranian traditional medicine known to optimise male fertility. Andrologia, 2016, 48, 850-859.	1.0	18
129	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
130	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
131	Asthma and Allergic Diseases in a Selected Group of Patients With Common Variable Immunodeficiency. Journal of Investigational Allergology and Clinical Immunology, 2016, 26, 209-211.	0.6	18
132	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
133	Managing patients with side effects and adverse events to immunoglobulin therapy. Expert Review of Clinical Pharmacology, 2016, 9, 91-102.	1.3	17
134	Two Faces of LRBA Deficiency in Siblings: Hypogammaglobulinemia and Normal Immunoglobulin Levels. Journal of Investigational Allergology and Clinical Immunology, 2018, 28, 48-50.	0.6	17
135	Identification of a novel de novo gain-of-function mutation of PIK3CD in a patient with activated phosphoinositide 3-kinase Î´ syndrome. Clinical Immunology, 2018, 197, 60-67.	1.4	17
136	Clinical, immunological, and genetic features in 938 patients with autoimmune polyendocrinopathy candidiasis ectodermal dystrophy (APECED): a systematic review. Expert Review of Clinical Immunology, 2021, 17, 807-817.	1.3	17
137	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
138	The Uncommon Combination of Common Variable Immunodeficiency, Macrophage Activation Syndrome, and Cytomegalovirus Retinitis. Viral Immunology, 2012, 25, 161-165.	0.6	16
139	Autosomal Recessive Agammaglobulinemia: A Novel Non-sense Mutation in CD79a. Journal of Clinical Immunology, 2014, 34, 138-141.	2.0	16
140	In vitro chromosomal radiosensitivity in patients with common variable immunodeficiency. Central-European Journal of Immunology, 2018, 43, 155-161.	0.4	16
141	Newborn Screening for Presymptomatic Diagnosis of Complement and Phagocyte Deficiencies. Frontiers in Immunology, 2020, 11, 455.	2.2	16
142	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
143	Investigation of underlying primary immunodeficiencies in patients with severe atopic dermatitis. Allergologia Et Immunopathologia, 2014, 42, 336-341.	1.0	15
144	Mortality and morbidity in patients with X-linked agammaglobulinaemia. Allergologia Et Immunopathologia, 2015, 43, 62-66.	1.0	15

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145	Autoimmunity in primary T-cell immunodeficiencies. Expert Review of Clinical Immunology, 2016, 12, 989-1006.	1.3	15
146	The Heterogeneous Pathogenesis of Selective Immunoglobulin A Deficiency. International Archives of Allergy and Immunology, 2019, 179, 231-246.	0.9	15
147	Variable Abnormalities in T and B Cell Subsets in Ataxia Telangiectasia. Journal of Clinical Immunology, 2021, 41, 76-88.	2.0	15
148	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
149	Diagnostic Approach to the Patients with Suspected Primary Immunodeficiency. Endocrine, Metabolic and Immune Disorders - Drug Targets, 2020, 20, 157-171.	0.6	15
150	Family study of pediatric patients with primary antibody deficiencies. Iranian Journal of Allergy, Asthma and Immunology, 2013, 12, 377-82.	0.3	15
151	Molecular diagnosis of primary immunodeficiency diseases in a developing country: Iran as an example. Expert Review of Clinical Immunology, 2014, 10, 385-396.	1.3	14
152	Circulating Helper T-Cell Subsets and Regulatory T Cells in Patients With Common Variable Immunodeficiency Without Known Monogenic Disease. Journal of Investigational Allergology and Clinical Immunology, 2018, 28, 172-181.	0.6	14
153	Genetic mutations and immunological features of severe combined immunodeficiency patients in Iran. Immunology Letters, 2019, 216, 70-78.	1.1	14
154	Comparison of clinical and immunological features and mortality in common variable immunodeficiency and agammaglobulinemia patients. Immunology Letters, 2019, 210, 55-62.	1.1	14
155	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
156	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
157	Hallmarks of Cancers: Primary Antibody Deficiency Versus Other Inborn Errors of Immunity. Frontiers in Immunology, 2021, 12, 720025.	2.2	14
158	Physicians awareness on primary immunodeficiency disorders in Iran. Iranian Journal of Allergy, Asthma and Immunology, 2012, 11, 57-64.	0.3	14
159	Monogenic Primary Immunodeficiency Disorder Associated with Common Variable Immunodeficiency and Autoimmunity. International Archives of Allergy and Immunology, 2020, 181, 706-714.	0.9	13
160	The First Purine Nucleoside Phosphorylase Deficiency Patient Resembling IgA Deficiency and a Review of the Literature. Immunological Investigations, 2019, 48, 410-430.	1.0	12
161	The profile of IL-4, IL-5, IL-10 and GATA3 in patients with LRBA deficiency and CVID with no known monogenic disease: Association with disease severity. Allergologia Et Immunopathologia, 2019, 47, 172-178.	1.0	12
162	Comprehensive assessment of respiratory complications in patients with common variable immunodeficiency. Annals of Allergy, Asthma and Immunology, 2020, 124, 505-511.e3.	0.5	12

#	Article	IF	Citations
163	Evaluation of Antibody Response to Polysaccharide Vaccine and Switched Memory B Cells in Pediatric Patients with Inflammatory Bowel Disease. Gut and Liver, 2014, 8, 24-28.	1.4	12
164	Expanding the Clinical and Immunological Phenotypes and Natural History of MALT1 Deficiency. Journal of Clinical Immunology, 2022, 42, 634-652.	2.0	12
165	Cutaneous granulomas in common variable immunodeficiency: case report and review of literature. Acta Dermatovenerologica Croatica, 2010, 18, 107-13.	0.1	12
166	The approach to children with recurrent infections. Iranian Journal of Allergy, Asthma and Immunology, 2012, 11, 89-109.	0.3	12
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