Nima Parvaneh

List of Publications by Citations

79 papers	2,229 citations	27 h-index	45 g-index
81 ext. papers	2,594 ext. citations	4.1 avg, IF	4.12 L-index

#	Paper	IF	Citations
79	IgA deficiency: correlation between clinical and immunological phenotypes. *Journal of Clinical Immunology*, **2009**, 29, 130-6	5.7	159
78	Inherited and acquired immunodeficiencies underlying tuberculosis in childhood. *Immunological Reviews*, **2015**, 264, 103-20	11.3	133
77	Hematologically important mutations: leukocyte adhesion deficiency (first update). *Blood Cells, Molecules, and Diseases*, **2012**, 48, 53-61	2.1	118
76	Progression of selective IgA deficiency to common variable immunodeficiency. *International Archives of Allergy and Immunology*, **2008**, 147, 87-92	3.7	102
75	Gastrointestinal manifestations in patients with common variable immunodeficiency. *Digestive Diseases and Sciences*, **2007**, 52, 2977-83	4	96
74	IL-12R1 deficiency in two of fifty children with severe tuberculosis from Iran, Morocco, and Turkey. *PLoS ONE*, **2011**, 6, e18524	3.7	91
73	Primary immunodeficiency disorders in Iran: update and new insights from the third report of the national registry. *Journal of Clinical Immunology*, **2014**, 34, 478-90	5.7	82
72	Clinical features of Candidiasis in patients with inherited interleukin 12 receptor 1 deficiency. *Clinical Infectious Diseases*, **2014**, 58, 204-13	11.6	81
71	Primary immunodeficiencies predisposed to Epstein-Barr virus-driven haematological diseases. *British Journal of Haematology*, **2013**, 162, 573-86	4.5	76
70	Primary immunodeficiencies: a rapidly evolving story. *Journal of Allergy and Clinical Immunology*, **2013**, 131, 314-23	11.5	68
69	The risk of hemophagocytic lymphohistiocytosis in Hermansky-Pudlak syndrome type 2. *Blood*, **2013**, 121, 2943-51	2.2	60
68	Fourth Update on the Iranian National Registry of Primary Immunodeficiencies: Integration of Molecular Diagnosis. *Journal of Clinical Immunology*, **2018**, 38, 816-832	5.7	57
67	Clinical, immunologic, and genetic spectrum of 696 patients with combined immunodeficiency. *Journal of Allergy and Clinical Immunology*, **2018**, 141, 1450-1458	11.5	56
66	Lymphoma of mucosa-associated lymphoid tissue in common variable immunodeficiency. *Leukemia and Lymphoma*, **2006**, 47, 343-6	1.9	52
65	Comparison of pulmonary diseases in common variable immunodeficiency and X-linked agammaglobulinaemia. *Respirology*, **2010**, 15, 289-95	3.6	49
64	Vitamin D insufficiency among children and adolescents living in Tehran, Iran. *Journal of Tropical Pediatrics*, **2009**, 55, 189-91	1.2	47
63	Clinical, immunological and molecular characteristics of 37 Iranian patients with X-linked agammaglobulinemia. *International Archives of Allergy and Immunology*, **2006**, 141, 408-14	3.7	47

62	Clinical and laboratory findings in hyper-IgM syndrome with novel CD40L and AICDA mutations. *Journal of Clinical Immunology*, **2009**, 29, 769-76	5.7	42
61	Analysis of switched memory B cells in patients with IgA deficiency. *International Archives of Allergy and Immunology*, **2011**, 156, 462-8	3.7	41
60	Disseminated Bacillus Calmette-Guerin infection after BCG vaccination. *Journal of Tropical Pediatrics*, **2008**, 54, 413-6	1.2	38
59	Severe combined immunodeficiency: a cohort of 40 patients. *Pediatric Allergy and Immunology*, **2008**, 19, 303-6	4.2	37
58	Vaccine-associated paralytic poliomyelitis in immunodeficient children, Iran, 1995-2008. *Emerging Infectious Diseases*, **2010**, 16, 1133-6	10.2	36
57	Mortality and morbidity in common variable immunodeficiency. *Journal of Tropical Pediatrics*, **2007**, 53, 32-8	1.2	34
56	Autoimmune phenotype in patients with common variable immunodeficiency. *Journal of Investigational Allergology and Clinical Immunology*, **2013**, 23, 323-9	2.3	31
55	Isolation of a type 3 vaccine-derived poliovirus (VDPV) from an Iranian child with X-linked agammaglobulinemia. *Virus Research*, **2008**, 137, 168-72	6.4	30
54	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, 756-60	5.7	28
53	Treatment of pemphigus vulgaris with mycophenolate mofetil as a steroid-sparing agent. *European Journal of Dermatology*, **2008**, 18, 159-64	0.8	27
52	A Variety of Alu-Mediated Copy Number Variations Can Underlie IL-12Rβ1 Deficiency. *Journal of Clinical Immunology*, **2018**, 38, 617-627	5.7	26
51	Novel BTK mutation presenting with vaccine-associated paralytic poliomyelitis. *European Journal of Pediatrics*, **2008**, 167, 1335-8	4.1	25
50	New therapeutic approach by sirolimus for enteropathy treatment in patients with LRBA deficiency. *European Annals of Allergy and Clinical Immunology*, **2017**, 49, 235-239	1.3	24
49	NCKAP1L defects lead to a novel syndrome combining immunodeficiency, lymphoproliferation, and hyperinflammation. *Journal of Experimental Medicine*, **2020**, 217,	16.6	23
48	Analysis of RAB27A gene in griscelli syndrome type 2: novel mutations including a deletion hotspot. *Journal of Clinical Immunology*, **2008**, 28, 384-9	5.7	22
47	Vaccine-associated paralytic poliomyelitis in a patient with MHC class II deficiency. *Journal of Clinical Virology*, **2007**, 39, 145-8	14.5	22
46	Characterization of six novel mutations in CYBA: the gene causing autosomal recessive chronic granulomatous disease. *British Journal of Haematology*, **2008**, 141, 848-51	4.5	21
45	Progressive multifocal leukoencephalopathy in purine nucleoside phosphorylase deficiency. *Brain and Development*, **2007**, 29, 124-6	2.2	20

44	Inborn errors of metabolism underlying primary immunodeficiencies. *Journal of Clinical Immunology* , **2014**, 34, 753-71	5.7	19
43	Cohort of Iranian Patients with Congenital Agammaglobulinemia: Mutation Analysis and Novel Gene Defects. *Expert Review of Clinical Immunology*, **2016**, 12, 479-86	5.1	18
42	Visceral leishmaniasis in two patients with IL-12p40 and IL-12R deficiencies. *Pediatric Blood and Cancer*, **2017**, 64, e26362	3	18
41	Molecular diagnosis of X-linked chronic granulomatous disease in Iran. *International Journal of Hematology*, **2008**, 87, 398-404	2.3	17
40	Long-term evaluation of a historical cohort of Iranian common variable immunodeficiency patients. *Expert Review of Clinical Immunology*, **2014**, 10, 1405-17	5.1	16
39	Genetic defects in B-cell development and their clinical consequences. *Journal of Investigational Allergology and Clinical Immunology*, **2014**, 24, 6-22; quiz 2 p following 22	2.3	16
38	Evaluation of serum IgA levels in Iranian patients with type 1 diabetes mellitus. *Acta Diabetologica*, **2012**, 49, 131-5	3.9	15
37	Mutation spectra of the ITGB2 gene in Iranian families with leukocyte adhesion deficiency type 1. *Human Immunology*, **2016**, 77, 191-5	2.3	14
36	Human papillomavirus detected in esophageal squamous cell carcinoma in Iran. *European Journal of Internal Medicine*, **2012**, 23, e59-62	3.9	14
35	Impact of delayed diagnosis in children with primary antibody deficiencies. *Journal of Microbiology, Immunology and Infection*, **2011**, 44, 229-34	8.5	14
34	Invasive aspergillosis in chronic granulomatous disease: report of 7 cases. *European Journal of Pediatrics*, **2007**, 166, 83-4	4.1	13
33	Novel UBR1 gene mutation in a patient with typical phenotype of Johanson-Blizzard syndrome. *European Journal of Pediatrics*, **2011**, 170, 233-5	4.1	11
32	Evaluation of antibody response to polysaccharide vaccine and switched memory B cells in pediatric patients with inflammatory bowel disease. *Gut and Liver*, **2014**, 8, 24-8	4.8	11
31	Genetic mutations and immunological features of severe combined immunodeficiency patients in Iran. *Immunology Letters*, **2019**, 216, 70-78	4.1	10
30	Pubertal development in a random sample of 4,020 urban Iranian girls. *Journal of Pediatric Endocrinology and Metabolism*, **2008**, 21, 681-7	1.6	10
29	Intermittent neutropenia as an early feature of mild mevalonate kinase deficiency. *Journal of Clinical Immunology*, **2014**, 34, 123-6	5.7	7
28	Impaired in-vitro responses to IL-12 and IFN- in Iranian patients with Mendelian susceptibility to mycobacterial disease. *Allergologia Et Immunopathologia*, **2015**, 43, 456-60	1.9	7
27	Griscelli Syndrome Type 2 Sine Albinism: Unraveling Differential RAB27A Effector Engagement. *Frontiers in Immunology*, **2020**, 11, 612977	8.4	6

26	Effects of Coronavirus Disease 2019 (COVID-19) on Peripheral Blood Lymphocytes and Their Subsets in Children: Imbalanced CD4/CD8 T Cell Ratio and Disease Severity. *Frontiers in Pediatrics*, **2021**, 9, 643299	3.4	6
25	A Novel STK4 Mutation Presenting with Juvenile Idiopathic Arthritis and Epidermodysplasia Verruciformis. *Journal of Clinical Immunology*, **2019**, 39, 11-14	5.7	6
24	2q34-qter duplication and 4q34.2-qter deletion in a patient with developmental delay. *European Journal of Medical Genetics*, **2012**, 55, 203-10	2.6	5
23	Behavior abnormality following intravenous immunoglobulin treatment in patients with primary antibody deficiencies. *Human Psychopharmacology*, **2010**, 25, 419-22	2.3	5
22	Complement deficiency in pediatric-onset systemic lupus erythematosus. *Journal of Laboratory Physicians*, **2018**, 10, 232-236	1.6	5
21	Vaccine-Derived Poliovirus Infection among Patients with Primary Immunodeficiency and Effect of Patient Screening on Disease Outcomes, Iran. *Emerging Infectious Diseases*, **2019**, 25, 2005-2012	10.2	4
20	A novel deletion mutation in the human hairless (HR) gene in an Iranian family with atrichia and papular lesions. *Clinical and Experimental Dermatology*, **2009**, 34, e498-500	1.8	4
19	NOD2 sequencing in Iranian children with Crohn□s disease. *Zeitschrift Fur Gastroenterologie*, **2011**, 49, 1526-8	1.6	4
18	Different phenotypes of the same XIAP mutation in a family: A case of XIAP deficiency with juvenile idiopathic arthritis. *Pediatric Blood and Cancer*, **2019**, 66, e27593	3	4
17	Diffuse dermal melanocytosis in two patients with Sandhoff disease and mucopolysaccharidosis VI. *International Journal of Dermatology*, **2014**, 53, 736-8	1.7	3
16	Post-mortem Diagnosis of Heme Oxygenase-1 Deficiency by Whole Exome Sequencing in an Iranian Child. *International Journal of Molecular and Cellular Medicine*, **2019**, 8, 300-307	1.2	3
15	DOCK8 deficiency in six Iranian patients. *Clinical Case Reports (discontinued)*, **2016**, 4, 593-600	0.7	3
14	Evaluation of interleukin-12 receptor □1 and interferon gamma receptor 1 deficiency in patients with disseminated BCG infection. *Allergologia Et Immunopathologia*, **2019**, 47, 38-42	1.9	3
13	Impaired respiratory burst contributes to infections in PKC□deficient patients. *Journal of Experimental Medicine*, **2021**, 218,	16.6	3
12	Clinical Manifestations, Immunological Characteristics and Genetic Analysis of Patients with Hyper-Immunoglobulin M Syndrome in Iran. *International Archives of Allergy and Immunology*, **2019**, 180, 52-63	3.7	2
11	Candidate Predisposition Variants in Kaposi Sarcoma as Detected by Whole-Genome Sequencing. *Open Forum Infectious Diseases*, **2019**, 6, ofz337	1	1
10	Retropharyngeal abscess after BCG vaccination. *International Journal of Pediatric Otorhinolaryngology Extra*, **2006**, 1, 279-281		1
9	Associations of Behavioral Disorders with Asthma in Iranian Children. *Iranian Journal of Allergy, Asthma and Immunology*, **2019**, 18, 340-345	1.1	1

8	Delay in Diagnosis of Two Siblings with Severe Ocular Problems and Autoimmune Polyglandular Syndrome. *Iranian Journal of Allergy, Asthma and Immunology*, **2020**, 19, 313-317	1.1	1
7	Invasive Rasamsonia argillacea infection in chronic granulomatous disease: Report of a new case and literature review. *Journal De Mycologie Medicale*, **2021**, 31, 101106	3	1
6	Mendelian susceptibility to mycobacterial disease: Clinical and immunological findings of patients suspected for IL12R1 deficiency. *Allergologia Et Immunopathologia*, **2019**, 47, 491-498	1.9	1
5	Identification of a New Variant in NLRP3 Gene by Whole Exome Sequencing in a Patient with Cryopyrin-Associated Periodic Syndrome. *Case Reports in Immunology*, **2021**, 2021, 2023119	1.9	1
4	The Efficacy of Anti-Tumor Necrosis Factor Therapy in Cryopyrin-Associated Periodic Syndromes: A Report of Two Cases.. *Case Reports in Genetics*, **2022**, 2022, 2898553	0.7	0
3	Ocular Manifestations of Chronic Granulomatous Disease: First Report of Coats Disease and Literature Review. *Journal of Clinical Immunology*, **2020**, 40, 940-947	5.7	
2	Serum sickness-like reactions in Iranian children: a registry-based study in a referral center. *Allergologia Et Immunopathologia*, **2020**, 48, 424-429	1.9	
1	Bacillus Calmette-Guerin (BCG)-associated hemophagocytic lymphohistiocytosis in the setting of IFN-R1 deficiency: A diagnostic dilemma. *EJHaem*, **2020**, 1, 334-337	0.9	