Mohammad Nabavi

List of Publications by Citations

69 papers	1,563 citations	16 h-index	39 g-index
77 ext. papers	1,841 ext. citations	3.2 avg, IF	3.57 L-index

#	Paper	IF	Citations
69	A homozygous CARD9 mutation in a family with susceptibility to fungal infections. *New England Journal of Medicine*, **2009**, 361, 1727-35	59.2	587
68	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, 519-32	5.7	130
67	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
66	Consanguinity in primary immunodeficiency disorders; the report from Iranian Primary Immunodeficiency Registry. *American Journal of Reproductive Immunology*, **2006**, 56, 145-51	3.8	76
65	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
64	Clinical, immunologic, and genetic spectrum of 696 patients with combined immunodeficiency. *Journal of Allergy and Clinical Immunology*, **2018**, 141, 1450-1458	11.5	56
63	Aspirin desensitization for patients with aspirin-exacerbated respiratory disease: A randomized double-blind placebo-controlled trial. *Clinical Immunology*, **2015**, 160, 349-57	9	53
62	Efficacy of intravenous immunoglobulin on the prevention of pneumonia in patients with agammaglobulinemia. *FEMS Immunology and Medical Microbiology*, **2004**, 40, 113-8		51
61	Impact of SARS-CoV-2 Pandemic on Patients with Primary Immunodeficiency. *Journal of Clinical Immunology*, **2021**, 41, 345-355	5.7	51
60	The effects of vitamin D supplementation on airway functions in mild to moderate persistent asthma. *Annals of Allergy, Asthma and Immunology*, **2014**, 113, 404-9	3.2	42
59	Auto-inflammation in a Patient with a Novel Homozygous OTULIN Mutation. *Journal of Clinical Immunology*, **2019**, 39, 138-141	5.7	21
58	Circulating level of CD4+ ́CD25+ ́FOXP3+ T cells in patients with chronic urticaria. *International Journal of Dermatology*, **2014**, 53, e561-6	1.7	21
57	The clinical and laboratory survey of Iranian patients with hyper-IgE syndrome. *Scandinavian Journal of Infectious Diseases*, **2006**, 38, 898-903		20
56	Aspirin hypersensitivity in patients with chronic rhinosinusitis and nasal polyposis: frequency and contributing factors. *American Journal of Rhinology and Allergy*, **2014**, 28, 239-43	2.4	19
55	HLA-DRB and HLA-DQ genetic variability in patients with aspirin-exacerbated respiratory disease. *American Journal of Rhinology and Allergy*, **2015**, 29, e63-9	2.4	18
54	DOCK2 Deficiency in a Patient with Hyper IgM Phenotype. *Journal of Clinical Immunology*, **2018**, 38, 10-13	5.7	17
53	A gain-of-function mutation of STAT1: A novel genetic factor contributing to chronic mucocutaneous candidiasis. *Acta Microbiologica Et Immunologica Hungarica*, **2017**, 64, 191-201	1.8	16

52	Increased level of interleukin-13, but not interleukin-4 and interferon-□n chronic rhinosinusitis with nasal polyps. *Allergologia Et Immunopathologia*, **2014**, 42, 465-71	1.9	16
51	Interleukin-6 and tumor necrosis factor-alpha gene polymorphisms in chronic idiopathic urticaria. *Allergologia Et Immunopathologia*, **2014**, 42, 533-8	1.9	15
50	Common causes of anaphylaxis in children: the first report of anaphylaxis registry in iran. *World Allergy Organization Journal*, **2010**, 3, 9-13	5.2	14
49	Long-term follow-up of ninety eight Iranian patients with primary immune deficiency in a single tertiary centre. *Allergologia Et Immunopathologia*, **2016**, 44, 322-30	1.9	13
48	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		12
47	Clinical, Laboratory, and Molecular Findings for 63 Patients With Severe Combined Immunodeficiency: A Decade□s Experience. *Journal of Investigational Allergology and Clinical Immunology*, **2017**, 27, 299-304	2.3	11
46	Central nervous system involvement in Blau syndrome: a new feature of the syndrome?. *Journal of Rheumatology*, **2007**, 34, 2504-5	4.1	11
45	The persian version of the chronic urticaria quality of life questionnaire: factor analysis, validation, and initial clinical findings. *Iranian Journal of Allergy, Asthma and Immunology*, **2014**, 13, 278-85	1.1	10
44	Pharmacogenetic tests to predict the efficacy of aspirin desensitization in patients with aspirin-exacerbated respiratory diseases; HLA-DQB302. *Expert Review of Respiratory Medicine*, **2015**, 9, 511-8	3.8	9
43	Factors associated with cord blood IgE levels. *Asian Pacific Journal of Allergy and Immunology*, **2013**, 31, 157-62	5.4	9
42	Vitamin D deficiency in chronic idiopathic urticaria. *Iranian Journal of Allergy, Asthma and Immunology*, **2015**, 14, 222-7	1.1	9
41	Novel Patient with Late-Onset Familial Hemophagocytic Lymphohistiocytosis with STXBP2 Mutations Presenting with Autoimmune Hepatitis, Neurological Manifestations and Infections Associated with Hypogammaglobulinemia. *Journal of Clinical Immunology*, **2015**, 35, 22-5	5.7	8
40	Monogenic Primary Immunodeficiency Disorder Associated with Common Variable Immunodeficiency and Autoimmunity. *International Archives of Allergy and Immunology*, **2020**, 181, 706-714	3.7	8
39	Phenotyping and follow up of forty-seven Iranian patients with common variable immunodeficiency. *Allergologia Et Immunopathologia*, **2016**, 44, 226-31	1.9	8
38	Evaluation of a new protocol for wheat desensitization in patients with wheat-induced anaphylaxis. *Immunotherapy*, **2017**, 9, 637-645	3.8	8
37	Association of interleukin 10 and transforming growth factor □gene polymorphisms with chronic idiopathic urticaria. *Acta Dermatovenerologica Croatica*, **2014**, 22, 239-45	0.5	8
36	Successful fludarabine-based hematopoietic stem cell transplantation in a pediatric patient with idiopathic CD4+ lymphocytopenia. *Pediatric Transplantation*, **2013**, 17, E109-11	1.8	7
35	Palatal-Myoclonus as a Presentation of Hashimoto Encephalopathy: an interesting case report. *Iranian Journal of Psychiatry*, **2013**, 8, 149-51	1.9	7

34	Type I and Type II Hereditary Angioedema: Clinical and Laboratory Findings in Iranian Patients. *Archives of Iranian Medicine*, **2015**, 18, 425-9	2.4	7
33	Investigation of ITGB2 gene in 12 new cases of leukocyte adhesion deficiency-type I revealed four novel mutations from Iran. *Archives of Iranian Medicine*, **2015**, 18, 760-4	2.4	7
32	Characteristics, Etiology and Treatment of Pediatric and Adult Anaphylaxis in Iran. *Iranian Journal of Allergy, Asthma and Immunology*, **2017**, 16, 480-487	1.1	6
31	Genetic Analysis of 13 Iranian Families With Leukocyte Adhesion Deficiency Type 1. *Journal of Pediatric Hematology/Oncology*, **2019**, 41, e3-e6	1.2	5
30	Autoimmune manifestations among 461 patients with monogenic inborn errors of immunity. *Pediatric Allergy and Immunology*, **2021**, 32, 1335-1348	4.2	5
29	Frequency and Risk Factors of Penicillin and Amoxicillin Allergy in Suspected Patients with Drug Allergy. *Archives of Iranian Medicine*, **2017**, 20, 34-37	2.4	5
28	Cow's Milk Desensitization in Anaphylactic Patients: A New Personalized-dose Method. *Iranian Journal of Allergy, Asthma and Immunology*, **2017**, 16, 45-52	1.1	4
27	Characterization of 4 New Mutations in the CYBB Gene in 10 Iranian Families With X-linked Chronic Granulomatous Disease. *Journal of Pediatric Hematology/Oncology*, **2018**, 40, e268-e272	1.2	3
26	Persistent papilloma and polyoma virus infection in common variable immunodeficiency with progressive multifocal leukoencephalopathy. *Annals of Allergy, Asthma and Immunology*, **2013**, 110, 119-20	2.2	3
25	DOCK8 deficiency in six Iranian patients. *Clinical Case Reports (discontinued)*, **2016**, 4, 593-600	0.7	3
24	Single nucleotide polymorphisms of IL-2, but not IL-12 and IFN-□are associated with increased susceptibility to chronic spontaneous urticaria. *Allergologia Et Immunopathologia*, **2017**, 45, 333-338	1.9	2
23	AICDA single nucleotide polymorphism in common variable immunodeficiency and selective IgA deficiency. *Allergologia Et Immunopathologia*, **2014**, 42, 422-6	1.9	2
22	Comparison of Diagnostic Tests with Oral Food Challenge in a Clinical Trial for Adult Patients with Sesame Anaphylaxis. *Iranian Journal of Allergy, Asthma and Immunology*, **2020**, 19, 27-34	1.1	2
21	Development of a High-resolution Melting Analysis Method Based on SYBR Green-I for rs7216389 Locus Genotyping in Asthmatic Child Patients. *Avicenna Journal of Medical Biotechnology*, **2014**, 6, 72-80	1.4	2
20	X-linked hyper-IgM syndrome associated with pulmonary manifestations: A very rare case of functional mutation in CD40L gene in Iran. *Current Research in Translational Medicine*, **2019**, 67, 28-30	3.7	2
19	A newly found homozygous mutation in recombination activating gene 1 in a patient with leaky severe combined immunodeficiency disorder. *Molecular Biology Reports*, **2019**, 46, 6571-6575	2.8	1
18	Delayed diagnosis of hereditary angioedema with C1-inhibitor deficiency in iranian children and adolescents. *Pediatric Allergy and Immunology*, **2019**, 30, 395-398	4.2	1
17	Multiple cancers in a patient with common variable immunodeficiency. *Allergologia Et Immunopathologia*, **2014**, 42, 85-7	1.9	1

16	Hypomorphic DOCK8 deletion causes hypereosinophilic syndrome. *Pediatric Blood and Cancer*, **2020**, 67, e28084	3	1
15	Could oral administration of immunoglobulin reduce diarrheal severity in common variable immunodeficiency?. *Allergologia Et Immunopathologia*, **2014**, 42, 371-2	1.9	0
14	A Case of Linear IgA Bullous Dermatosis Induced by Aspirin Therapy. *Iranian Journal of Allergy, Asthma and Immunology*, **2020**, 19, 550-554	1.1	0
13	Genetic Study of Hereditary Angioedema Type I and Type II (First Report from Iranian Patients: Describing Three New Mutations). *Immunological Investigations*, **2020**, 1-12	2.9	0
12	Periodic Severe Angioedema without Exogenous Hormone Exposure. *Iranian Journal of Allergy, Asthma and Immunology*, **2021**, 20, 120-124	1.1	0
11	Adverse reactions in a large cohort of patients with inborn errors of immunity receiving intravenous immunoglobulin. *Clinical Immunology*, **2021**, 230, 108826	9	0
10	Association between p53 codon 72 polymorphism and systemic lupus erythematosus. *Reumatologia*, **2014**, 2, 94-98	1.7	
9	Immunochemical Characterization of Ligustrum Vulgare (Privet) Pollen Allergens: Study of Common Allergenic Plant in Iran.. *Iranian Journal of Allergy, Asthma and Immunology*, **2022**, 21, 55-64	1.1	
8	Chronic Eosinophilic Pneumonia: a Case Report. *Tanaffos*, **2020**, 19, 262-266	0.5	
7	Aspirin Sensitivity in Patients with Moderate to Severe Asthma. *Iranian Journal of Allergy, Asthma and Immunology*, **2020**, 19, 447-451	1.1	
6	Atopy Patch Test in the Diagnosis of Food Allergens in Infants with Allergic Proctocolitis Compared with Elimination/Introduction C. *Iranian Journal of Allergy, Asthma and Immunology*, **2021**, 20, 520-524	1.1	
5	A Rare Case with Quail Egg Allergy without Allergic Reactions to Oral Food Challenge with Henﾀs Egg White. *Iranian Journal of Allergy, Asthma and Immunology*, **2020**, 19, 310-312	1.1	
4	Goodﾀs syndrome (immunodeficiency with thymoma): A separate entity with a broad classification: Report of six cases and review of the literature. *Clinical Case Reports (discontinued)*, **2021**, 9, e04136	0.7	
3	Oral Immunotherapy in Patients with IgE Mediated Reactions to Egg White: A Clinical Trial Study. *Immunological Investigations*, **2021**, 1-14	2.9	
2	Immunodeficiency, Centromeric Region Instability, and Facial Anomalies Syndrome (ICF) in a Boy with Variable Clinical and Immunological Presentations. *Iranian Journal of Allergy, Asthma and Immunology*, **2021**, 20, 249-254	1.1	
1	Atypical Omenn Syndrome Due to RAG2 Gene Mutation, a Case Report. *Iranian Journal of Immunology*, **2019**, 16, 334-338	0.9	