## Seyed Alireza Mahdaviani

List of Publications by Year in descending order

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84 papers

3,653 citations

218592 26 h-index 58 g-index

90 all docs 90 docs citations

90 times ranked 6319 citing authors

#	Article	IF	Citations
1	Human intracellular ISG15 prevents interferon- $\hat{l}\pm\hat{l}^2$ over-amplification and auto-inflammation. Nature, 2015, 517, 89-93.	13.7	432
2	Characterization of Greater Middle Eastern genetic variation for enhanced disease gene discovery. Nature Genetics, 2016, 48, 1071-1076.	9.4	314
3	Human TYK2 deficiency: Mycobacterial and viral infections without hyper-lgE syndrome. Journal of Experimental Medicine, 2015, 212, 1641-1662.	4.2	293
4	Inherited CARD9 deficiency in otherwise healthy children and adults with Candida species–induced meningoencephalitis, colitis, or both. Journal of Allergy and Clinical Immunology, 2015, 135, 1558-1568.e2.	1.5	208
5	Monogenic mutations differentially affect the quantity and quality of T follicular helper cells in patients with human primary immunodeficiencies. Journal of Allergy and Clinical Immunology, 2015, 136, 993-1006.e1.	1.5	181
6	Inherited and acquired immunodeficiencies underlying tuberculosis in childhood. Immunological Reviews, 2015, 264, 103-120.	2.8	180
7	Human IFN- $\hat{I}^3$ immunity to mycobacteria is governed by both IL-12 and IL-23. Science Immunology, 2018, 3, .	<b>5.</b> 6	152
8	Inherited IL-12p40 Deficiency. Medicine (United States), 2013, 92, 109-122.	0.4	151
9	Inherited CARD9 Deficiency in 2 Unrelated Patients With Invasive Exophiala Infection. Journal of Infectious Diseases, 2015, 211, 1241-1250.	1.9	141
10	IL- $12\hat{R}^21$ Deficiency in Two of Fifty Children with Severe Tuberculosis from Iran, Morocco, and Turkey. PLoS ONE, 2011, 6, e18524.	1.1	111
11	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
12	Inheritance Pattern and Clinical Aspects of 93 Iranian Patients with Chronic Granulomatous Disease. Journal of Clinical Immunology, 2011, 31, 792-801.	2.0	94
13	Clinical, immunologic, and genetic spectrum of 696 patients with combined immunodeficiency. Journal of Allergy and Clinical Immunology, 2018, 141, 1450-1458.	1.5	90
14	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
15	Global systematic review of primary immunodeficiency registries. Expert Review of Clinical Immunology, 2020, 16, 717-732.	1.3	74
16	IL-2-Inducible T-Cell Kinase Deficiency with Pulmonary Manifestations due to Disseminated Epstein-Barr Virus Infection. International Archives of Allergy and Immunology, 2012, 158, 418-422.	0.9	71
17	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
18	Proinflammatory Cytokine Gene Polymorphisms among Iranian Patients with Asthma. Journal of Clinical Immunology, 2009, 29, 57-62.	2.0	60

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19	IL-10, TGF-ß, IL-2, IL-12, and IFN-γ Cytokine Gene Polymorphisms in Asthma. Journal of Asthma, 2008, 45, 790-794.	0.9	58
20	Whole-exome sequencing to analyze population structure, parental inbreeding, and familial linkage. Proceedings of the National Academy of Sciences of the United States of America, 2016, 113, 6713-6718.	3.3	53
21	Pulmonary manifestations of chronic granulomatous disease. Expert Review of Clinical Immunology, 2013, 9, 153-160.	1.3	50
22	Classification of Asthma Based on Nonlinear Analysis of Breathing Pattern. PLoS ONE, 2016, 11, e0147976.	1.1	48
23	Inherited deficiency of stress granule ZNFX1 in patients with monocytosis and mycobacterial disease. Proceedings of the National Academy of Sciences of the United States of America, 2021, 118, .	3.3	47
24	A Variety of Alu-Mediated Copy Number Variations Can Underlie IL- $12R\hat{l}^21$ Deficiency. Journal of Clinical Immunology, 2018, 38, 617-627.	2.0	45
25	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
26	Consensus Middle East and North Africa Registry on Inborn Errors of Immunity. Journal of Clinical Immunology, 2021, 41, 1339-1351.	2.0	33
27	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
28	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
29	Skin Prick Test Reactivity to Common Aero and Food Allergens among Children with Allergy. Iranian Journal of Medical Sciences, 2014, 39, 29-35.	0.3	25
30	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
31	Mendelian Susceptibility to Mycobacterial Disease (MSMD): Clinical and Genetic Features of 32 Iranian Patients. Journal of Clinical Immunology, 2020, 40, 872-882.	2.0	22
32	Fungal epidemiology in cystic fibrosis patients with a special focus on Scedosporium species complex. Microbial Pathogenesis, 2019, 129, 168-175.	1.3	19
33	Bronchoalveolar galactomannan in invasive pulmonary aspergillosis: a prospective study in pediatric patients. Medical Mycology, 2015, 53, 709-716.	0.3	18
34	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.	0.6	18
35	Mobile GIS-based monitoring asthma attacks based on environmental factors. Journal of Cleaner Production, 2018, 179, 417-428.	4.6	17
36	Susceptibility to mycobacterial disease due to mutations in IL- $12R\hat{l}^21$ in three Iranian patients. Immunogenetics, 2018, 70, 373-379.	1.2	17

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37	An ensemble learning method for asthma control level detection with leveraging medical knowledge-based classifier and supervised learning. Journal of Medical Systems, 2019, 43, 158.	2.2	15
38	Genetic mutations and immunological features of severe combined immunodeficiency patients in Iran. Immunology Letters, 2019, 216, 70-78.	1.1	14
39	Expression levels of plasma exosomal miR-124, miR-125b, miR-133b, miR-130a and miR-125b-1-3p in severe asthma patients and normal individuals with emphasis on inflammatory factors. Allergy, Asthma and Clinical Immunology, 2021, 17, 51.	0.9	14
40	Paecilomyces formosus Infection in an Adult Patient with Undiagnosed Chronic Granulomatous Disease. Journal of Clinical Immunology, 2017, 37, 342-346.	2.0	13
41	Genetic and molecular findings of 38 Iranian patients with chronic granulomatous disease caused by p47â€ <i>phox</i> defect. Scandinavian Journal of Immunology, 2019, 90, e12767.	1.3	13
42	Monogenic Primary Immunodeficiency Disorder Associated with Common Variable Immunodeficiency and Autoimmunity. International Archives of Allergy and Immunology, 2020, 181, 706-714.	0.9	13
43	Evaluation of a new protocol for wheat desensitization in patients with wheat-induced anaphylaxis. Immunotherapy, 2017, 9, 637-645.	1.0	12
44	Improving the function of neutrophils from chronic granulomatous disease patients using mesenchymal stem cells' exosomes. Human Immunology, 2020, 81, 614-624.	1.2	12
45	Pulmonary computed tomography scan findings in chronic granulomatous disease. Allergologia Et Immunopathologia, 2014, 42, 444-448.	1.0	9
46	Autoimmune manifestations among 461 patients with monogenic inborn errors of immunity. Pediatric Allergy and Immunology, 2021, 32, 1335-1348.	1.1	9
47	Asthma induces psychiatric impairments in association with default mode and salience networks alteration: A resting-state EEG study. Respiratory Physiology and Neurobiology, 2022, 300, 103870.	0.7	9
48	Leukocytoclastic vasculitis in patients with IL12B or IL12RB1 deficiency: case report and review of the literature. Pediatric Rheumatology, 2021, 19, 121.	0.9	8
49	lgG anti-lgA antibodies in paediatric antibody-deficient patients receiving intravenous immunoglobulin. Allergologia Et Immunopathologia, 2015, 43, 403-408.	1.0	7
50	Respiratory Complications in Patients with Hyper IgM Syndrome. Journal of Clinical Immunology, 2019, 39, 557-568.	2.0	7
51	Clinical and Mutation Description of the First Iranian Cohort of Infantile Inflammatory Bowel Disease: The Iranian Primary Immunodeficiency Registry (IPIDR). Immunological Investigations, 2021, 50, 445-459.	1.0	7
52	Effect of Family Empowerment on the Quality of life of School-Aged Children with Asthma. Tanaffos, 2014, 13, 35-42.	0.5	7
53	Good's Syndrome-Association of the Late Onset Combined Immunodeficiency with Thymoma: Review of Literature and Case Report. Iranian Journal of Allergy, Asthma and Immunology, 2018, 17, 85-93.	0.3	7
54	Immunophenotypic and functional analysis of lymphocyte subsets in common variable immunodeficiency patients without monogenic defects. Scandinavian Journal of Immunology, 2022, 96, e13164.	1.3	7

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55	Effective anti-mycobacterial treatment for BCG disease in patients with Mendelian Susceptibility to Mycobacterial Disease (MSMD): a case series. Annals of Clinical Microbiology and Antimicrobials, 2022, 21, 8.	1.7	7
56	The First Iranian Cohort of Pediatric Patients with Activated Phosphoinositide 3-Kinase- $\hat{l}$ (PI3K $\hat{l}$ ) Syndrome (APDS). Immunological Investigations, 2021, , 1-16.	1.0	6
57	The Prevalence of Atopic Manifestations in 313 Iranian Patients with Inborn Errors of Immunity. International Archives of Allergy and Immunology, 2021, 182, 1122-1126.	0.9	6
58	Prevalence of specific immunoglobulin E and G against Aspergillus fumigatus in patients with asthma. Current Medical Mycology, 2018, 4, 7-11.	0.8	6
59	A new ataxia-telangiectasia mutation in an 11-year-old female. Immunogenetics, 2017, 69, 415-419.	1.2	5
60	Atypical Ataxia Presentation in Variant Ataxia Telangiectasia: Iranian Case-Series and Review of the Literature. Frontiers in Immunology, 2021, 12, 779502.	2.2	5
61	Amyloidosis as a Renal Complication of Chronic Granulomatous Disease. Iranian Journal of Kidney Diseases, 2016, 10, 228-32.	0.1	5
62	Association of specific viral infections with childhood asthma exacerbations. Interventional Medicine & Applied Science, 2018, 11, 17-20.	0.2	4
63	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.	0.9	4
64	Evaluation of Expression of LRBA and CTLA-4 Proteins in Common Variable Immunodeficiency Patients. Immunological Investigations, 2022, 51, 381-394.	1.0	4
65	Effect of Family Empowerment on Asthma Control in School-Age Children. Tanaffos, 2018, 17, 47-52.	0.5	4
66	Delay in the Diagnosis of APECED: A Case Report and Review of Literature from Iran. Immunological Investigations, 2020, 49, 299-306.	1.0	3
67	Adverse reactions in a large cohort of patients with inborn errors of immunity receiving intravenous immunoglobulin. Clinical Immunology, 2021, 230, 108826.	1.4	3
68	Pulmonary manifestations in a cohort of patients with inborn errors of immunity: an 8-year follow-up study. Allergologia Et Immunopathologia, 2022, 50, 80-84.	1.0	3
69	Transmission electron microscopy study of suspected primary ciliary dyskinesia patients. Scientific Reports, 2022, 12, 2375.	1.6	3
70	AICDA single nucleotide polymorphism in common variable immunodeficiency and selective IgA deficiency. Allergologia Et Immunopathologia, 2014, 42, 422-426.	1.0	2
71	Association of <i>Mycobacterium</i> infections in patients with Mendelian susceptibility to mycobacterial disease with venous thromboembolism. Microbiology and Immunology, 2016, 60, 678-686.	0.7	2
72	Delayed diagnosis of hereditary angioedema with C1â€inhibitor deficiency in iranian children and adolescents. Pediatric Allergy and Immunology, 2019, 30, 395-398.	1.1	2

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73	Disseminated Mycobacterium simiae Infection in a Patient with Complete IL-12p40 Deficiency. Iranian Journal of Allergy, Asthma and Immunology, 2021, 20, 376-381.	0.3	2
74	Chest CT Manifestations in Children with CVID: A 10-Year Report. Tanaffos, 2012, 11, 56-9.	0.5	2
75	Allergic and nonallergic asthma in children: are they distinct phenotypes?. Iranian Journal of Allergy, Asthma and Immunology, 2014, 13, 370-4.	0.3	2
76	Determination of the Most Common Indoor and Outdoor Allergens in 602 Patients with Allergic Symptoms Using Specific IgE Local Panel. Iranian Journal of Allergy, Asthma and Immunology, 2017, 16, 298-306.	0.3	2
77	Pulmonary complications of predominantly antibody immunodeficiencies in a tertiary lung center. Interventional Medicine & Applied Science, 2018, $11$ , $1$ -7.	0.2	1
78	Use of a Bail-Bearing to Facilitate Goniolens Rotation. Optometry and Vision Science, 1995, 72, 924.	0.6	0
79	Defects in Innate Immunity: Receptors and Signaling Components. , 2012, , 279-307.		O
80	Pulmonary Manifestations of Congenital Defects of Phagocytes. , 2019, , 121-143.		0
81	Relationship between spirometry results and colonisation of Aspergillus species in allergic asthma. Clinical Respiratory Journal, 2020, 14, 748-757.	0.6	0
82	Interleukin 9 serum level and single nucleotide polymorphism in patients with asthma. Acta Biomedica, 2021, 92, e2021206.	0.2	0
83	A fludarabine and melphalan reduced-intensity conditioning regimen for HSCT in fifteen chronic granulomatous disease patients and a literature review. Annals of Hematology, 2022, 101, 869-880.	0.8	O
84	IL-17 Producing T cells as Predictors of Primary Immunodeficiencies in Patients with Candida Infections. Archives of Pediatric Infectious Diseases, 2022, In Press, .	0.1	0