

Seyed Alireza Mahdavian

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

83
papers

2,493
citations

24
h-index

49
g-index

89
ext. papers

3,192
ext. citations

5.1
avg, IF

3.89
L-index

| # | Paper | IF | Citations |
|----|---|------|-----------|
| 83 | Human intracellular ISG15 prevents interferon- γ over-amplification and auto-inflammation. <i>Nature</i> , 2015 , 517, 89-93 | 50.4 | 311 |
| 82 | Human TYK2 deficiency: Mycobacterial and viral infections without hyper-IgE syndrome. <i>Journal of Experimental Medicine</i> , 2015 , 212, 1641-62 | 16.6 | 209 |
| 81 | Characterization of Greater Middle Eastern genetic variation for enhanced disease gene discovery. <i>Nature Genetics</i> , 2016 , 48, 1071-6 | 36.3 | 192 |
| 80 | Inherited CARD9 deficiency in otherwise healthy children and adults with <i>Candida</i> species-induced meningoencephalitis, colitis, or both. <i>Journal of Allergy and Clinical Immunology</i> , 2015 , 135, 1558-68.e2 | 11.5 | 143 |
| 79 | Inherited and acquired immunodeficiencies underlying tuberculosis in childhood. <i>Immunological Reviews</i> , 2015 , 264, 103-20 | 11.3 | 133 |
| 78 | Monogenic mutations differentially affect the quantity and quality of T follicular helper cells in patients with human primary immunodeficiencies. <i>Journal of Allergy and Clinical Immunology</i> , 2015 , 136, 993-1006.e1 | 11.5 | 126 |
| 77 | Inherited IL-12p40 deficiency: genetic, immunologic, and clinical features of 49 patients from 30 kindreds. <i>Medicine (United States)</i> , 2013 , 92, 109-122 | 1.8 | 121 |
| 76 | Inherited CARD9 deficiency in 2 unrelated patients with invasive <i>Exophiala</i> infection. <i>Journal of Infectious Diseases</i> , 2015 , 211, 1241-50 | 7 | 101 |
| 75 | IL-12R β 1 deficiency in two of fifty children with severe tuberculosis from Iran, Morocco, and Turkey. <i>PLoS ONE</i> , 2011 , 6, e18524 | 3.7 | 91 |
| 74 | Human IFN- γ immunity to mycobacteria is governed by both IL-12 and IL-23. <i>Science Immunology</i> , 2018 , 3, | 28 | 83 |
| 73 | Primary immunodeficiency disorders in Iran: update and new insights from the third report of the national registry. <i>Journal of Clinical Immunology</i> , 2014 , 34, 478-90 | 5.7 | 82 |
| 72 | Inheritance pattern and clinical aspects of 93 Iranian patients with chronic granulomatous disease. <i>Journal of Clinical Immunology</i> , 2011 , 31, 792-801 | 5.7 | 73 |
| 71 | IL-2-inducible T-cell kinase deficiency with pulmonary manifestations due to disseminated Epstein-Barr virus infection. <i>International Archives of Allergy and Immunology</i> , 2012 , 158, 418-22 | 3.7 | 65 |
| 70 | Fourth Update on the Iranian National Registry of Primary Immunodeficiencies: Integration of Molecular Diagnosis. <i>Journal of Clinical Immunology</i> , 2018 , 38, 816-832 | 5.7 | 57 |
| 69 | Clinical, immunologic, and genetic spectrum of 696 patients with combined immunodeficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2018 , 141, 1450-1458 | 11.5 | 56 |
| 68 | Proinflammatory cytokine gene polymorphisms among Iranian patients with asthma. <i>Journal of Clinical Immunology</i> , 2009 , 29, 57-62 | 5.7 | 53 |
| 67 | IL-10, TGF-beta, IL-2, IL-12, and IFN-gamma cytokine gene polymorphisms in asthma. <i>Journal of Asthma</i> , 2008 , 45, 790-4 | 1.9 | 53 |

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|----|--|------|----|
| 66 | Clinical, immunologic, molecular analyses and outcomes of Iranian patients with LRBA deficiency: A longitudinal study. <i>Pediatric Allergy and Immunology</i> , 2017 , 28, 478-484 | 4.2 | 48 |
| 65 | Pulmonary manifestations of chronic granulomatous disease. <i>Expert Review of Clinical Immunology</i> , 2013 , 9, 153-60 | 5.1 | 41 |
| 64 | Whole-exome sequencing to analyze population structure, parental inbreeding, and familial linkage. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2016 , 113, 6713-8 | 11.5 | 37 |
| 63 | Classification of Asthma Based on Nonlinear Analysis of Breathing Pattern. <i>PLoS ONE</i> , 2016 , 11, e0147976 | 5.7 | 36 |
| 62 | Global systematic review of primary immunodeficiency registries. <i>Expert Review of Clinical Immunology</i> , 2020 , 16, 717-732 | 5.1 | 29 |
| 61 | Comparison of Common Monogenic Defects in a Large Predominantly Antibody Deficiency Cohort. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2019 , 7, 864-878.e9 | 5.4 | 29 |
| 60 | A Variety of Alu-Mediated Copy Number Variations Can Underlie IL-12R β Deficiency. <i>Journal of Clinical Immunology</i> , 2018 , 38, 617-627 | 5.7 | 26 |
| 59 | Novel mutation of the activation-induced cytidine deaminase gene in a Tajik family: special review on hyper-immunoglobulin M syndrome. <i>Expert Review of Clinical Immunology</i> , 2012 , 8, 539-46 | 5.1 | 23 |
| 58 | Cohort of Iranian Patients with Congenital Agammaglobulinemia: Mutation Analysis and Novel Gene Defects. <i>Expert Review of Clinical Immunology</i> , 2016 , 12, 479-86 | 5.1 | 18 |
| 57 | Skin Prick Test Reactivity to Common Aero and Food Allergens among Children with Allergy. <i>Iranian Journal of Medical Sciences</i> , 2014 , 39, 29-35 | 1.2 | 18 |
| 56 | Autoimmunity and its association with regulatory T cells and B cell subsets in patients with common variable immunodeficiency. <i>Allergologia Et Immunopathologia</i> , 2018 , 46, 127-135 | 1.9 | 18 |
| 55 | Bronchoalveolar galactomannan in invasive pulmonary aspergillosis: a prospective study in pediatric patients. <i>Medical Mycology</i> , 2015 , 53, 709-16 | 3.9 | 16 |
| 54 | Inherited deficiency of stress granule ZNFX1 in patients with monocytosis and mycobacterial disease. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2021 , 118, | 11.5 | 16 |
| 53 | Susceptibility to mycobacterial disease due to mutations in IL-12R β in three Iranian patients. <i>Immunogenetics</i> , 2018 , 70, 373-379 | 3.2 | 13 |
| 52 | Fungal epidemiology in cystic fibrosis patients with a special focus on <i>Scedosporium</i> species complex. <i>Microbial Pathogenesis</i> , 2019 , 129, 168-175 | 3.8 | 11 |
| 51 | Clinical, Laboratory, and Molecular Findings for 63 Patients With Severe Combined Immunodeficiency: A Decade's Experience. <i>Journal of Investigational Allergology and Clinical Immunology</i> , 2017 , 27, 299-304 | 2.3 | 11 |
| 50 | Genetic mutations and immunological features of severe combined immunodeficiency patients in Iran. <i>Immunology Letters</i> , 2019 , 216, 70-78 | 4.1 | 10 |
| 49 | Mendelian Susceptibility to Mycobacterial Disease (MSMD): Clinical and Genetic Features of 32 Iranian Patients. <i>Journal of Clinical Immunology</i> , 2020 , 40, 872-882 | 5.7 | 9 |

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| 48 | Paecilomyces formosus Infection in an Adult Patient with Undiagnosed Chronic Granulomatous Disease. <i>Journal of Clinical Immunology</i> , 2017 , 37, 342-346 | 5.7 | 8 |
| 47 | Genetic and molecular findings of 38 Iranian patients with chronic granulomatous disease caused by p47-phox defect. <i>Scandinavian Journal of Immunology</i> , 2019 , 90, e12767 | 3.4 | 8 |
| 46 | Monogenic Primary Immunodeficiency Disorder Associated with Common Variable Immunodeficiency and Autoimmunity. <i>International Archives of Allergy and Immunology</i> , 2020 , 181, 706-714 | 3.7 | 8 |
| 45 | Evaluation of a new protocol for wheat desensitization in patients with wheat-induced anaphylaxis. <i>Immunotherapy</i> , 2017 , 9, 637-645 | 3.8 | 8 |
| 44 | Mobile GIS-based monitoring asthma attacks based on environmental factors. <i>Journal of Cleaner Production</i> , 2018 , 179, 417-428 | 10.3 | 7 |
| 43 | Pulmonary computed tomography scan findings in chronic granulomatous disease. <i>Allergologia Et Immunopathologia</i> , 2014 , 42, 444-8 | 1.9 | 7 |
| 42 | Effect of Family Empowerment on the Quality of life of School-Aged Children with Asthma. <i>Tanaffos</i> , 2014 , 13, 35-42 | 0.5 | 7 |
| 41 | Good's Syndrome-Association of the Late Onset Combined Immunodeficiency with Thymoma: Review of Literature and Case Report. <i>Iranian Journal of Allergy, Asthma and Immunology</i> , 2018 , 17, 85-93 | 1.1 | 7 |
| 40 | 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. <i>Allergy, Asthma and Clinical Immunology</i> , 2021 , 17, 51 | 3.2 | 6 |
| 39 | Consensus Middle East and North Africa Registry on Inborn Errors of Immunity. <i>Journal of Clinical Immunology</i> , 2021 , 41, 1339-1351 | 5.7 | 6 |
| 38 | An ensemble learning method for asthma control level detection with leveraging medical knowledge-based classifier and supervised learning. <i>Journal of Medical Systems</i> , 2019 , 43, 158 | 5.1 | 5 |
| 37 | IgG anti-IgA antibodies in paediatric antibody-deficient patients receiving intravenous immunoglobulin. <i>Allergologia Et Immunopathologia</i> , 2015 , 43, 403-8 | 1.9 | 5 |
| 36 | Prevalence of specific immunoglobulin E and G against in patients with asthma. <i>Current Medical Mycology</i> , 2018 , 4, 7-11 | 1.1 | 5 |
| 35 | Autoimmune manifestations among 461 patients with monogenic inborn errors of immunity. <i>Pediatric Allergy and Immunology</i> , 2021 , 32, 1335-1348 | 4.2 | 5 |
| 34 | A new ataxia-telangiectasia mutation in an 11-year-old female. <i>Immunogenetics</i> , 2017 , 69, 415-419 | 3.2 | 4 |
| 33 | Amyloidosis as a Renal Complication of Chronic Granulomatous Disease. <i>Iranian Journal of Kidney Diseases</i> , 2016 , 10, 228-32 | 0.9 | 4 |
| 32 | Respiratory Complications in Patients with Hyper IgM Syndrome. <i>Journal of Clinical Immunology</i> , 2019 , 39, 557-568 | 5.7 | 3 |
| 31 | Improving the function of neutrophils from chronic granulomatous disease patients using mesenchymal stem cellsExosomes. <i>Human Immunology</i> , 2020 , 81, 614-624 | 2.3 | 3 |

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| 30 | Clinical and Mutation Description of the First Iranian Cohort of Infantile Inflammatory Bowel Disease: The Iranian Primary Immunodeficiency Registry (IPIDR). <i>Immunological Investigations</i> , 2021 , 50, 445-459 | 2.9 | 3 |
| 29 | Clinical Manifestations, Immunological Characteristics and Genetic Analysis of Patients with Hyper-Immunoglobulin M Syndrome in Iran. <i>International Archives of Allergy and Immunology</i> , 2019 , 180, 52-63 | 3.7 | 2 |
| 28 | AICDA single nucleotide polymorphism in common variable immunodeficiency and selective IgA deficiency. <i>Allergologia Et Immunopathologia</i> , 2014 , 42, 422-6 | 1.9 | 2 |
| 27 | Atypical Ataxia Presentation in Variant Ataxia Telangiectasia: Iranian Case-Series and Review of the Literature.. <i>Frontiers in Immunology</i> , 2021 , 12, 779502 | 8.4 | 2 |
| 26 | Chest CT Manifestations in Children with COVID: A 10-Year Report. <i>Tanaffos</i> , 2012 , 11, 56-9 | 0.5 | 2 |
| 25 | Association of specific viral infections with childhood asthma exacerbations. <i>Interventional Medicine & Applied Science</i> , 2019 , 11, 17-20 | 0.7 | 2 |
| 24 | Leukocytoclastic vasculitis in patients with IL12B or IL12RB1 deficiency: case report and review of the literature. <i>Pediatric Rheumatology</i> , 2021 , 19, 121 | 3.5 | 2 |
| 23 | Allergic and nonallergic asthma in children: are they distinct phenotypes?. <i>Iranian Journal of Allergy, Asthma and Immunology</i> , 2014 , 13, 370-4 | 1.1 | 2 |
| 22 | Determination of the Most Common Indoor and Outdoor Allergens in 602 Patients with Allergic Symptoms Using Specific IgE Local Panel. <i>Iranian Journal of Allergy, Asthma and Immunology</i> , 2017 , 16, 298-306 | 1.1 | 2 |
| 21 | Immunophenotypic and functional analysis of lymphocyte subsets in common variable immunodeficiency patients without monogenic defects.. <i>Scandinavian Journal of Immunology</i> , 2022 , e13164 | 3.4 | 2 |
| 20 | Delayed diagnosis of hereditary angioedema with C1-inhibitor deficiency in Iranian children and adolescents. <i>Pediatric Allergy and Immunology</i> , 2019 , 30, 395-398 | 4.2 | 1 |
| 19 | Effect of Family Empowerment on Asthma Control in School-Age Children. <i>Tanaffos</i> , 2018 , 17, 47-52 | 0.5 | 1 |
| 18 | Evaluation of Expression of LRBA and CTLA-4 Proteins in Common Variable Immunodeficiency Patients. <i>Immunological Investigations</i> , 2020 , 1-14 | 2.9 | 1 |
| 17 | The Prevalence of Atopic Manifestations in 313 Iranian Patients with Inborn Errors of Immunity. <i>International Archives of Allergy and Immunology</i> , 2021 , 182, 1122-1126 | 3.7 | 1 |
| 16 | Association of Mycobacterium infections in patients with Mendelian susceptibility to mycobacterial disease with venous thromboembolism. <i>Microbiology and Immunology</i> , 2016 , 60, 678-686 | 2.7 | 1 |
| 15 | Delay in the Diagnosis of APECED: A Case Report and Review of Literature from Iran. <i>Immunological Investigations</i> , 2020 , 49, 299-306 | 2.9 | 1 |
| 14 | The First Iranian Cohort of Pediatric Patients with Activated Phosphoinositide 3-Kinase- γ (PI3K γ) Syndrome (APDS). <i>Immunological Investigations</i> , 2021 , 1-16 | 2.9 | 1 |
| 13 | Asthma induces psychiatric impairments in association with default mode and salience networks alteration: A resting-state EEG study.. <i>Respiratory Physiology and Neurobiology</i> , 2022 , 300, 103870 | 2.8 | 1 |

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| 12 | Transmission electron microscopy study of suspected primary ciliary dyskinesia patients.. <i>Scientific Reports</i> , 2022 , 12, 2375 | 4.9 | o |
| 11 | Pulmonary manifestations in a cohort of patients with inborn errors of immunity: an 8-year follow-up study.. <i>Allergologia Et Immunopathologia</i> , 2022 , 50, 80-84 | 1.9 | o |
| 10 | Pulmonary complications of predominantly antibody immunodeficiencies in a tertiary lung center. <i>Interventional Medicine & Applied Science</i> , 2019 , 11, 1-7 | 0.7 | o |
| 9 | Adverse reactions in a large cohort of patients with inborn errors of immunity receiving intravenous immunoglobulin. <i>Clinical Immunology</i> , 2021 , 230, 108826 | 9 | o |
| 8 | Effective anti-mycobacterial treatment for BCG disease in patients with Mendelian Susceptibility to Mycobacterial Disease (MSMD): a case series.. <i>Annals of Clinical Microbiology and Antimicrobials</i> , 2022 , 21, 8 | 6.2 | o |
| 7 | Relationship between spirometry results and colonisation of <i>Aspergillus</i> species in allergic asthma. <i>Clinical Respiratory Journal</i> , 2020 , 14, 748 | 1.7 | |
| 6 | Defects in Innate Immunity: Receptors and Signaling Components 2012 , 279-307 | | |
| 5 | Use of a ball-bearing to facilitate goniolens rotation. <i>Optometry and Vision Science</i> , 1995 , 72, 924 | 2.1 | |
| 4 | A fludarabine and melphalan reduced-intensity conditioning regimen for HSCT in fifteen chronic granulomatous disease patients and a literature review.. <i>Annals of Hematology</i> , 2022 , 101, 869 | 3 | |
| 3 | Interleukin 9 serum level and single nucleotide polymorphism in patients with asthma. <i>Acta Biomedica</i> , 2021 , 92, e2021206 | 3.2 | |
| 2 | Disseminated <i>Mycobacterium simiae</i> Infection in a Patient with Complete IL-12p40 Deficiency. <i>Iranian Journal of Allergy, Asthma and Immunology</i> , 2021 , 20, 376-381 | 1.1 | |
| 1 | Pulmonary Manifestations of Congenital Defects of Phagocytes 2019 , 121-143 | | |