Samaneh Delavari

List of Publications by Year
in descending order

24 papers	372 citations	1039406 9 h-index	839053 18 g-index
27 all docs	27 docs citations	27 times ranked	757 citing authors

#	Article	IF	Citations
1	Evaluation of Expression of LRBA and CTLA-4 Proteins in Common Variable Immunodeficiency Patients. Immunological Investigations, 2022, 51, 381-394.	1.0	4
2	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
3	Expanding the Clinical and Immunological Phenotypes and Natural History of MALT1 Deficiency. Journal of Clinical Immunology, 2022, 42, 634-652.	2.0	12
4	Evaluation of Specific Antibody Responses in Patients with Selective IgA Deficiency and Ataxia Telangiectasia. Endocrine, Metabolic and Immune Disorders - Drug Targets, 2022, 22, 640-649.	0.6	1
5	The Effects of Stimulation with PMA/Ionomycin on CD4+ T Cell Proliferation and Surface CD4 Molecule Modulation of Patients with LRBA Deficiency and CVID with the Unsolved Genetic Defect. Endocrine, Metabolic and Immune Disorders - Drug Targets, 2022, 22, 539-544.	0.6	1
6	Evaluation of Radiation Sensitivity in Patients with Hyper IgM Syndrome. Immunological Investigations, 2021, 50, 580-596.	1.0	3
7	Effect of Class Switch Recombination Defect on the Phenotype of Ataxia-Telangiectasia Patients. Immunological Investigations, 2021, 50, 201-215.	1.0	22
8	Variable Abnormalities in T and B Cell Subsets in Ataxia Telangiectasia. Journal of Clinical Immunology, 2021, 41, 76-88.	2.0	15
9	Impact of SARS-CoV-2 Pandemic on Patients with Primary Immunodeficiency. Journal of Clinical Immunology, 2021, 41, 345-355.	2.0	97
10	The First Iranian Cohort of Pediatric Patients with Activated Phosphoinositide 3-Kinase-Î´ (PI3KÎ´) Syndrome (APDS). Immunological Investigations, 2021, , 1-16.	1.0	6
11	The spectrum of ATM gene mutations in Iranian patients with ataxiaâ€telangiectasia. Pediatric Allergy and Immunology, 2021, 32, 1316-1326.	1.1	5
12	Consensus Middle East and North Africa Registry on Inborn Errors of Immunity. Journal of Clinical Immunology, 2021, 41, 1339-1351.	2.0	33
13	Autoimmune manifestations among 461 patients with monogenic inborn errors of immunity. Pediatric Allergy and Immunology, 2021, 32, 1335-1348.	1.1	9
14	Lymphocytes subsets in correlation with clinical profile in CVID patients without monogenic defects. Expert Review of Clinical Immunology, 2021, 17, 1041-1051.	1.3	6
15	Atypical Ataxia Presentation in Variant Ataxia Telangiectasia: Iranian Case-Series and Review of the Literature. Frontiers in Immunology, 2021, 12, 779502.	2.2	5
16	First patient in the Iranian Registry with novel DOCK2 gene mutation, presenting with skeletal tuberculosis, and review of literature. Allergy, Asthma and Clinical Immunology, 2021, 17, 126.	0.9	5
17	Disseminated Intravascular Coagulation Associated with Large Deletion of Immunoglobulin Heavy Chain. Iranian Journal of Allergy, Asthma and Immunology, 2021, 20, 778-783.	0.3	1
18	Global systematic review of primary immunodeficiency registries. Expert Review of Clinical Immunology, 2020, 16, 717-732.	1.3	74

#	Article	IF	Citations
19	Monogenic Primary Immunodeficiency Disorder Associated with Common Variable Immunodeficiency and Autoimmunity. International Archives of Allergy and Immunology, 2020, 181, 706-714.	0.9	13
20	Evaluation of respiratory complications in patients with Xâ€linked and autosomal recessive agammaglobulinemia. Pediatric Allergy and Immunology, 2020, 31, 405-417.	1.1	4
21	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
22	Are asthma and allergic diseases phenotypic markers for patients with common variable immunodeficiency?. Annals of Allergy, Asthma and Immunology, 2020, 124, 636.	0.5	2
23	Graft versus host disease and microchimerism in a JAK3 deficient patient. Allergy, Asthma and Clinical Immunology, 2019, 15, 47.	0.9	3
24	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