

Samaneh Delavari

List of Publications by Year in descending order

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Version: 2024-02-01

24
papers

372
citations

1039406

9
h-index

839053

18
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27
all docs

27
docs citations

27
times ranked

757
citing authors

#	ARTICLE	IF	CITATIONS
1	Impact of SARS-CoV-2 Pandemic on Patients with Primary Immunodeficiency. <i>Journal of Clinical Immunology</i> , 2021, 41, 345-355.	2.0	97
2	Global systematic review of primary immunodeficiency registries. <i>Expert Review of Clinical Immunology</i> , 2020, 16, 717-732.	1.3	74
3	X-Linked TLR7 Deficiency Underlies Critical COVID-19 Pneumonia in a Male Patient with Ataxia-Telangiectasia. <i>Journal of Clinical Immunology</i> , 2022, 42, 1-9.	2.0	34
4	Consensus Middle East and North Africa Registry on Inborn Errors of Immunity. <i>Journal of Clinical Immunology</i> , 2021, 41, 1339-1351.	2.0	33
5	Effect of Class Switch Recombination Defect on the Phenotype of Ataxia-Telangiectasia Patients. <i>Immunological Investigations</i> , 2021, 50, 201-215.	1.0	22
6	Variable Abnormalities in T and B Cell Subsets in Ataxia Telangiectasia. <i>Journal of Clinical Immunology</i> , 2021, 41, 76-88.	2.0	15
7	Monogenic Primary Immunodeficiency Disorder Associated with Common Variable Immunodeficiency and Autoimmunity. <i>International Archives of Allergy and Immunology</i> , 2020, 181, 706-714.	0.9	13
8	Comprehensive assessment of respiratory complications in patients with common variable immunodeficiency. <i>Annals of Allergy, Asthma and Immunology</i> , 2020, 124, 505-511.e3.	0.5	12
9	Expanding the Clinical and Immunological Phenotypes and Natural History of MALT1 Deficiency. <i>Journal of Clinical Immunology</i> , 2022, 42, 634-652.	2.0	12
10	Autoimmune manifestations among 461 patients with monogenic inborn errors of immunity. <i>Pediatric Allergy and Immunology</i> , 2021, 32, 1335-1348.	1.1	9
11	The First Iranian Cohort of Pediatric Patients with Activated Phosphoinositide 3-Kinase- γ (PI3K γ) Syndrome (APDS). <i>Immunological Investigations</i> , 2021, , 1-16.	1.0	6
12	Lymphocytes subsets in correlation with clinical profile in CVID patients without monogenic defects. <i>Expert Review of Clinical Immunology</i> , 2021, 17, 1041-1051.	1.3	6
13	The spectrum of ATM gene mutations in Iranian patients with ataxia-telangiectasia. <i>Pediatric Allergy and Immunology</i> , 2021, 32, 1316-1326.	1.1	5
14	Atypical Ataxia Presentation in Variant Ataxia Telangiectasia: Iranian Case-Series and Review of the Literature. <i>Frontiers in Immunology</i> , 2021, 12, 779502.	2.2	5
15	First patient in the Iranian Registry with novel DOCK2 gene mutation, presenting with skeletal tuberculosis, and review of literature. <i>Allergy, Asthma and Clinical Immunology</i> , 2021, 17, 126.	0.9	5
16	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.	0.9	4
17	Evaluation of Expression of LRBA and CTLA-4 Proteins in Common Variable Immunodeficiency Patients. <i>Immunological Investigations</i> , 2022, 51, 381-394.	1.0	4
18	Evaluation of respiratory complications in patients with X-linked and autosomal recessive agammaglobulinemia. <i>Pediatric Allergy and Immunology</i> , 2020, 31, 405-417.	1.1	4

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19	Graft versus host disease and microchimerism in a JAK3 deficient patient. <i>Allergy, Asthma and Clinical Immunology</i> , 2019, 15, 47.	0.9	3
20	Evaluation of Radiation Sensitivity in Patients with Hyper IgM Syndrome. <i>Immunological Investigations</i> , 2021, 50, 580-596.	1.0	3
21	Are asthma and allergic diseases phenotypic markers for patients with common variable immunodeficiency?. <i>Annals of Allergy, Asthma and Immunology</i> , 2020, 124, 636.	0.5	2
22	Evaluation of Specific Antibody Responses in Patients with Selective IgA Deficiency and Ataxia Telangiectasia. <i>Endocrine, Metabolic and Immune Disorders - Drug Targets</i> , 2022, 22, 640-649.	0.6	1
23	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. <i>Endocrine, Metabolic and Immune Disorders - Drug Targets</i> , 2022, 22, 539-544.	0.6	1
24	Disseminated Intravascular Coagulation Associated with Large Deletion of Immunoglobulin Heavy Chain. <i>Iranian Journal of Allergy, Asthma and Immunology</i> , 2021, 20, 778-783.	0.3	1