

Mehdi Yeganeh

List of Publications by Year in descending order

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Version: 2024-02-01

19
papers

1,458
citations

623734

14
h-index

839539

18
g-index

20
all docs

20
docs citations

20
times ranked

2201
citing authors

#	ARTICLE	IF	CITATIONS
1	Negative regulation of the hepatic fibrogenic response by suppressor of cytokine signaling 1. <i>Cytokine</i> , 2016, 82, 58-69.	3.2	15
2	Tumour-promoting role of SOCS1 in colorectal cancer cells. <i>Scientific Reports</i> , 2015, 5, 14301.	3.3	28
3	The extended clinical phenotype of 64 patients with dedicator of cytokinesis 8 deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2015, 136, 402-412.	2.9	163
4	Debilitating progressive encephalitis in a patient with BTK deficiency. <i>Acta Microbiologica Et Immunologica Hungarica</i> , 2012, 59, 335-342.	0.8	8
5	SOCS1 controls liver regeneration by regulating HGF signaling in hepatocytes. <i>Journal of Hepatology</i> , 2011, 55, 1300-1308.	3.7	50
6	Mutations in STAT3 and diagnostic guidelines for hyper-IgE syndrome. <i>Journal of Allergy and Clinical Immunology</i> , 2010, 125, 424-432.e8.	2.9	247
7	IgA Deficiency: Correlation Between Clinical and Immunological Phenotypes. <i>Journal of Clinical Immunology</i> , 2009, 29, 130-136.	3.8	191
8	Large deletions and point mutations involving the dedicator of cytokinesis 8 (DOCK8) in the autosomal-recessive form of hyper-IgE syndrome. <i>Journal of Allergy and Clinical Immunology</i> , 2009, 124, 1289-1302.e4.	2.9	453
9	Novel mutations of NP in two patients with purine nucleoside phosphorylase deficiency. <i>Clinical Biochemistry</i> , 2008, 41, 350-352.	1.9	14
10	Analysis of RAB27A Gene in Griscelli Syndrome type 2: Novel Mutations Including a Deletion Hotspot. <i>Journal of Clinical Immunology</i> , 2008, 28, 384-389.	3.8	23
11	Novel BTK mutation presenting with vaccine-associated paralytic poliomyelitis. <i>European Journal of Pediatrics</i> , 2008, 167, 1335-1338.	2.7	29
12	Severe combined immunodeficiency: A cohort of 40 patients. <i>Pediatric Allergy and Immunology</i> , 2008, 19, 303-306.	2.6	41
13	Genotype-Phenotype Correlation in Bruton's Tyrosine Kinase Deficiency. <i>Journal of Pediatric Hematology/Oncology</i> , 2008, 30, 679-683.	0.6	35
14	Other Well-Defined Immunodeficiencies. , 2008, , 251-290.		5
15	Evaluation of humoral immune function in patients with bronchiectasis. <i>Iranian Journal of Allergy, Asthma and Immunology</i> , 2008, 7, 69-77.	0.4	13
16	Progressive multifocal leukoencephalopathy in purine nucleoside phosphorylase deficiency. <i>Brain and Development</i> , 2007, 29, 124-126.	1.1	25
17	Mortality and Morbidity in Common Variable Immunodeficiency. <i>Journal of Tropical Pediatrics</i> , 2006, 53, 32-38.	1.5	44
18	The clinical and laboratory survey of Iranian patients with Hyper-IgE syndrome. <i>Scandinavian Journal of Infectious Diseases</i> , 2006, 38, 898-903.	1.5	22

#	ARTICLE	IF	CITATIONS
19	Clinical, Immunological and Molecular Characteristics of 37 Iranian Patients with X-Linked Agammaglobulinemia. International Archives of Allergy and Immunology, 2006, 141, 408-414.	2.1	52