Taher Cheraghi

List of Publications by Year
in descending order

29 papers	786 citations	686830 13 h-index	580395 25 g-index
30 all docs	30 docs citations	30 times ranked	1254 citing authors

#	Article	IF	Citations
1	The First Case of BENTA Disease (B Cell Expansion with NF-Î²B and T Cell Anergy) from Iran. Journal of Clinical Immunology, 2021, 41, 811-813.	2.0	2
2	Immunodeficiencies affecting cellular and humoral immunity. , 2021, , 9-39.		1
3	Consensus Middle East and North Africa Registry on Inborn Errors of Immunity. Journal of Clinical Immunology, 2021, 41, 1339-1351.	2.0	33
4	Autoimmune manifestations among 461 patients with monogenic inborn errors of immunity. Pediatric Allergy and Immunology, 2021, 32, 1335-1348.	1.1	9
5	Adverse reactions in a large cohort of patients with inborn errors of immunity receiving intravenous immunoglobulin. Clinical Immunology, 2021, 230, 108826.	1.4	3
6	Predominantly antibody deficiencies. , 2021, , 93-123.		1
7	Monogenic Primary Immunodeficiency Disorder Associated with Common Variable Immunodeficiency and Autoimmunity. International Archives of Allergy and Immunology, 2020, 181, 706-714.	0.9	13
8	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
9	Mendelian susceptibility to mycobacterial disease: Clinical and immunological findings of patients suspected for IL12RÎ²1 deficiency. Allergologia Et Immunopathologia, 2019, 47, 491-498.	1.0	3
10	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
11	Recurrent Sinusitis and Persistent Giardia Infection. , 2019, , 97-100.		0
12	Recurrent, Non-Pitting Edema of Lips, Extremities, and Genitalia. , 2019, , 647-651.		0
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	Comparison of Pulmonary Function Test in School - Age Children in Clean and Polluted Air in Tehran, Iran. Iranian Journal of Pediatrics, 2018, 28, .	0.1	1
16	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
17	Chronic, Long-Term Social Stress Can Cause Decreased Microtubule Protein Network Activity and Dynamics in Cerebral Cortex of Male Wistar Rats. Journal of Molecular Neuroscience, 2015, 55, 579-586.	1.1	7
18	Long-term evaluation of a historical cohort of Iranian common variable immunodeficiency patients. Expert Review of Clinical Immunology, 2014, 10, 1405-1417.	1.3	19

#	Article	IF	Citations
19	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
20	Prevention and control of infections in patients with severe congenital neutropenia; a follow up study. Iranian Journal of Allergy, Asthma and Immunology, 2012, 11, 51-6.	0.3	7
21	Common Causes of Anaphylaxis in Children. World Allergy Organization Journal, 2010, 3, 9-13.	1.6	23
22	BAK, BAX, and NBK/BIK Proapoptotic Gene Alterations in Iranian Patients with Ataxia Telangiectasia. Journal of Clinical Immunology, 2010, 30, 132-137.	2.0	7
23	Behavior abnormality following intravenous immunoglobulin treatment in patients with primary antibody deficiencies. Human Psychopharmacology, 2010, 25, 419-422.	0.7	5
24	Cutaneous granulomas in common variable immunodeficiency: case report and review of literature. Acta Dermatovenerologica Croatica, 2010, 18, 107-13.	0.1	12
25	IgA Deficiency: Correlation Between Clinical and Immunological Phenotypes. Journal of Clinical Immunology, 2009, 29, 130-136.	2.0	191
26	Characterization of six novel mutations in CYBA: the gene causing autosomal recessive chronic granulomatous disease*. British Journal of Haematology, 2008, 141, 848-851.	1.2	24
27	Chromosomal radiosensitivity in patients with common variable immunodeficiency. Immunobiology, 2008, 213, 447-454.	0.8	52
28	Evaluation of humoral immune function in patients with bronchiectasis. Iranian Journal of Allergy, Asthma and Immunology, 2008, 7, 69-77.	0.3	13
29	Presence of Idiopathic Thrombocytopenic Purpura and autoimmune hemolytic anemia in the patients with common variable immunodeficiency. Iranian Journal of Allergy, Asthma and Immunology, 2008, 7, 169-75.	0.3	21