Ilhan Tezcan

List of Publications by Year
in descending order

55	1,400	361413	345221
papers	citations	20 h-index	36 g-index

58	58	58	2837
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Clinical, Laboratory Features and Clinical Courses of Patients with Wiskott Aldrich Syndrome and Xâ€“linked Thrombocytopeniaâ€“A single center study. Immunological Investigations, 2022, 51, 1272-1283.	2.0	0
2	Nutritional status of children with primary immunodeficiency: A single center experience. Pediatrics International, 2022, 64, .	0.5	0
3	Long Term Follow-Up of the Patients with Severe Combined Immunodeficiency After Hematopoietic Stem Cell Transplantation: A Single-Center Study. Immunological Investigations, 2022, 51, 739-747.	2.0	4
4	Combined immunodeficiency due to purine nucleoside phosphorylase deficiency: Outcome of three patients. European Journal of Medical Genetics, 2022, 65, 104428.	1.3	10
5	Evaluation of Health Status and Quality of Life in Patients Using Intravenous and Subcutaneous Forms of Immunoglobulin Replacement. Journal of Ankara University Faculty of Medicine, 2022, 75, 77-83.	0.1	0
6	First allogeneic hematopoietic stem cell transplantation in RASGRP1 deficiency: long-term follow-up. Bone Marrow Transplantation, 2022, , .	2.4	0
7	GIMAP6 regulates autophagy, immune competence, and inflammation in mice and humans. Journal of Experimental Medicine, 2022, 219, .	8.5	4
8	Frequency of HLA Class I and Class II Alleles in Patients with CVID from Turkey. Immunological Investigations, 2021, 50, 363-371.	2.0	0
9	Evaluation of periodontal status and cytokine/chemokine profile of GCF in patients with severe congenital neutropenia. Odontology / the Society of the Nippon Dental University, 2021, 109, 474-482.	1.9	7
10	A Patient With AIRE Mutation Who Presented With Severe Diarrhea and Lung Abscess. Pediatric Infectious Disease Journal, 2021, 40, 66-69.	2.0	3
11	Clinical and Immunological Characteristics of 63 Patients with Chronic Granulomatous Disease: Hacettepe Experience. Journal of Clinical Immunology, 2021, 41, 992-1003.	3.8	8
12	Selective IgM deficiency: Followâ€up and outcome. Pediatric Allergy and Immunology, 2021, 32, 1327-1334.	2.6	8
13	Genomic Spectrum and Phenotypic Heterogeneity of Human IL-21 Receptor Deficiency. Journal of Clinical Immunology, 2021, 41, 1272-1290.	3.8	25
14	Expression of HLA class I and class II genes in patients with multiple skin warts. Experimental Dermatology, 2021, 30, 1642-1649.	2.9	1
15	Diversity in Serine/Threonine Protein Kinase-4 Deficiency and Review of the Literature. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 3752-3766.e4.	3.8	13
16	Differential diagnosis of primary immunodeficiency in patients with BCGitis and BCGosis: A singleâ€centre study. Scandinavian Journal of Immunology, 2021, 94, e13084.	2.7	8
17	COVID-19 in Patients with Primary Immunodeficiency. Journal of Clinical Immunology, 2021, 41, 1515-1522.	3.8	38
18	Clinical and laboratory findings in patients with leukocyte adhesion deficiency type I: A multicenter study in Turkey. Clinical and Experimental Immunology, 2021, 206, 47-55.	2.6	10

#	Article	IF	Citations
19	In case of recurrent wheezing and bronchiolitis: Think again, it may be a primary immunodeficiency. Asian Pacific Journal of Allergy and Immunology, 2021, , .	0.4	1
20	Cytokine profile in serum and gingival crevicular fluid of children with inflammatory bowel disease: A caseâ€control study. Journal of Periodontology, 2021, , .	3.4	2
21	A Monogenic Disease with a Variety of Phenotypes: Deficiency of Adenosine Deaminase 2. Journal of Rheumatology, 2020, 47, 117-125.	2.0	65
22	Recurrent Demyelinating Episodes as Sole Manifestation of Inherited CD59 Deficiency. Neuropediatrics, 2020, 51, 206-210.	0.6	6
23	Adenosine Deaminase Type II Deficiency: Severe Chronic Neutropenia, Lymphoid Infiltration in Bone Marrow, and Inflammatory Features. Immunological Investigations, 2020, , 1-9.	2.0	3
24	Recurrent skin abscesses in a female Xâ€linked chronic granulomatous disease carrier. Journal of Cosmetic Dermatology, 2020, 19, 1810-1812.	1.6	1
25	Elevated Interleukinâ€17A expression in amlodipineâ€induced gingival overgrowth. Journal of Periodontal Research, 2020, 55, 613-621.	2.7	3
26	Impact of mannoseâ€binding lectin 2 gene polymorphisms on disease severity in noncystic fibrosis bronchiectasis in children. Pediatric Pulmonology, 2020, 55, 1190-1198.	2.0	5
27	Lymphocyte Subgroups and KREC Numbers in Common Variable Immunodeficiency: A Single Center Study. Journal of Clinical Immunology, 2020, 40, 494-502.	3.8	2
28	Levels of pro- and anti-inflammatory cytokines in cystic fibrosis patients with or without gingivitis. Cytokine, 2020, 127, 154987.	3.2	7
29	Retroperitoneal Abscess in Severe Combined Immunodeficiency Probably Due to BCG Vaccine. Asim, Allerji, Immunoloji, 2020, 18, 98-101.	0.0	1
30	Three patients with glucose-6 phosphatase catalytic subunit 3 deficiency. Journal of Pediatric Endocrinology and Metabolism, 2020, 33, 957-961.	0.9	0
31	A Spectrum of Clinical Findings from ALPS to CVID: Several Novel LRBA Defects. Journal of Clinical Immunology, 2019, 39, 726-738.	3.8	45
32	Clinical Features and HSCT Outcome for SCID in Turkey. Journal of Clinical Immunology, 2019, 39, 316-323.	3.8	17
33	Effects of oral prophylaxis including tongue cleaning on halitosis and gingival inflammation in gingivitis patientsâ€”a randomized controlled clinical trial. Clinical Oral Investigations, 2019, 23, 1829-1836.	3.0	14
34	Selective loss of function variants in IL6ST cause Hyper-IgE syndrome with distinct impairments of T-cell phenotype and function. Haematologica, 2019, 104, 609-621.	3.5	74
35	Primary Immunodeficiencies Associated with Atopic Dermatitis. Asim, Allerji, Immunoloji, 2019, 17, 70-77.	0.0	1
36	Neurologic Involvement in Primary Immunodeficiency Disorders. Journal of Child Neurology, 2018, 33, 320-328.	1.4	12

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37	Successful hematopoietic stem cell transplantation after myeloablative conditioning in three patients with dedicator of cytokinesis 8 deficiency (DOCK8) related Hyper IgE syndrome. Bone Marrow Transplantation, 2018, 53, 339-343.	2.4	8
38	ADA Deficiency: Evaluation of the Clinical and Laboratory Features and the Outcome. Journal of Clinical Immunology, 2018, 38, 484-493.	3.8	26
39	A novel mutation in TAP1 gene leading to MHC class I deficiency: Report of two cases and review of the literature. Clinical Immunology, 2017, 178, 74-78.	3.2	31
40	Acute myeloid leukemia in a child with dedicator of cytokinesis 8 (DOCK8) deficiency. Pediatric Blood and Cancer, 2017, 64, e26695.	1.5	6
41	ISG15 deficiency and increased viral resistance in humans but not mice. Nature Communications, 2016, 7, 11496.	12.8	156
42	RASGRP1 deficiency causes immunodeficiency with impaired cytoskeletal dynamics. Nature Immunology, 2016, 17, 1352-1360.	14.5	115
43	Statins and IL-1Î², IL-10, and MPO Levels in Gingival Crevicular Fluid: Preliminary Results. Inflammation, 2016, 39, 1547-1557.	3.8	31
44	Recurrent viral infections associated with a homozygous CORO1A mutation that disrupts oligomerization and cytoskeletal association. Journal of Allergy and Clinical Immunology, 2016, 137, 879-888.e2.	2.9	41
45	The extended clinical phenotype of 64 patients with dedicator of cytokinesis 8 deficiency. Journal of Allergy and Clinical Immunology, 2015, 136, 402-412.	2.9	163
46	Combined immunodeficiency with CD4 lymphopenia and sclerosing cholangitis caused by a novel loss-of-function mutation affecting IL21R. Haematologica, 2015, 100, e216-e219.	3.5	46
47	Analysis of TNF-Î± (-308) polymorphism and gingival crevicular fluid TNF-Î± levels in aggressive and chronic periodontitis: A preliminary report. Cytokine, 2015, 72, 173-177.	3.2	25
48	Functional analysis of naturally occurring DCLRE1C mutations and correlation with the clinical phenotype of ARTEMIS deficiency. Journal of Allergy and Clinical Immunology, 2015, 136, 140-150.e7.	2.9	63
49	STK4 (MST1) deficiency in two siblings with autoimmune cytopenias: A novel mutation. Clinical Immunology, 2015, 161, 316-323.	3.2	73
50	Identification of ITK deficiency as a novel genetic cause of idiopathic CD4+ T-cell lymphopenia. Blood, 2014, 124, 655-657.	1.4	51
51	Additional Diverse Findings Expand the Clinical Presentation of DOCK8 Deficiency. Journal of Clinical Immunology, 2012, 32, 698-708.	3.8	84
52	Thirty years of primary immunodeficiencies in Turkey. Annals of the New York Academy of Sciences, 2011, 1238, 15-23.	3.8	25
53	Genetic analysis of patients with leukocyte adhesion deficiency. Experimental Hematology, 2002, 30, 252-261.	0.4	41
54	Bruton tyrosine kinase gene mutations in Turkish patients with presumed X-linked agammaglobulinemia. Human Mutation, 2001, 18, 356-356.	2.5	15

#	Article	IF	Citations
55	Flow Cytometry is a Reliable Tool in the Diagnosis of STK4 Deficiency. Asim, Allerji, Immunoloji, 0, , .	0.0	2