

# Ilhan Tezcan

## List of Publications by Year in descending order

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Version: 2024-02-01

55  
papers

1,400  
citations

361413

20  
h-index

345221

36  
g-index

58  
all docs

58  
docs citations

58  
times ranked

2837  
citing authors

#	ARTICLE	IF	CITATIONS
1	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
2	ISG15 deficiency and increased viral resistance in humans but not mice. <i>Nature Communications</i> , 2016, 7, 11496.	12.8	156
3	RASGRP1 deficiency causes immunodeficiency with impaired cytoskeletal dynamics. <i>Nature Immunology</i> , 2016, 17, 1352-1360.	14.5	115
4	Additional Diverse Findings Expand the Clinical Presentation of DOCK8 Deficiency. <i>Journal of Clinical Immunology</i> , 2012, 32, 698-708.	3.8	84
5	Selective loss of function variants in <i>IL6ST</i> cause Hyper-IgE syndrome with distinct impairments of T-cell phenotype and function. <i>Haematologica</i> , 2019, 104, 609-621.	3.5	74
6	STK4 (MST1) deficiency in two siblings with autoimmune cytopenias: A novel mutation. <i>Clinical Immunology</i> , 2015, 161, 316-323.	3.2	73
7	A Monogenic Disease with a Variety of Phenotypes: Deficiency of Adenosine Deaminase 2. <i>Journal of Rheumatology</i> , 2020, 47, 117-125.	2.0	65
8	Functional analysis of naturally occurring DCLRE1C mutations and correlation with the clinical phenotype of ARTEMIS deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2015, 136, 140-150.e7.	2.9	63
9	Identification of ITK deficiency as a novel genetic cause of idiopathic CD4+ T-cell lymphopenia. <i>Blood</i> , 2014, 124, 655-657.	1.4	51
10	Combined immunodeficiency with CD4 lymphopenia and sclerosing cholangitis caused by a novel loss-of-function mutation affecting IL21R. <i>Haematologica</i> , 2015, 100, e216-e219.	3.5	46
11	A Spectrum of Clinical Findings from ALPS to CVID: Several Novel LRBA Defects. <i>Journal of Clinical Immunology</i> , 2019, 39, 726-738.	3.8	45
12	Genetic analysis of patients with leukocyte adhesion deficiency. <i>Experimental Hematology</i> , 2002, 30, 252-261.	0.4	41
13	Recurrent viral infections associated with a homozygous CORO1A mutation that disrupts oligomerization and cytoskeletal association. <i>Journal of Allergy and Clinical Immunology</i> , 2016, 137, 879-888.e2.	2.9	41
14	COVID-19 in Patients with Primary Immunodeficiency. <i>Journal of Clinical Immunology</i> , 2021, 41, 1515-1522.	3.8	38
15	Statins and IL-1 $\beta$ , IL-10, and MPO Levels in Gingival Crevicular Fluid: Preliminary Results. <i>Inflammation</i> , 2016, 39, 1547-1557.	3.8	31
16	A novel mutation in TAP1 gene leading to MHC class I deficiency: Report of two cases and review of the literature. <i>Clinical Immunology</i> , 2017, 178, 74-78.	3.2	31
17	ADA Deficiency: Evaluation of the Clinical and Laboratory Features and the Outcome. <i>Journal of Clinical Immunology</i> , 2018, 38, 484-493.	3.8	26
18	Thirty years of primary immunodeficiencies in Turkey. <i>Annals of the New York Academy of Sciences</i> , 2011, 1238, 15-23.	3.8	25

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19	Analysis of TNF- $\alpha$ (-308) polymorphism and gingival crevicular fluid TNF- $\alpha$ levels in aggressive and chronic periodontitis: A preliminary report. <i>Cytokine</i> , 2015, 72, 173-177.	3.2	25
20	Genomic Spectrum and Phenotypic Heterogeneity of Human IL-21 Receptor Deficiency. <i>Journal of Clinical Immunology</i> , 2021, 41, 1272-1290.	3.8	25
21	Clinical Features and HSCT Outcome for SCID in Turkey. <i>Journal of Clinical Immunology</i> , 2019, 39, 316-323.	3.8	17
22	Bruton tyrosine kinase gene mutations in Turkish patients with presumed X-linked agammaglobulinemia. <i>Human Mutation</i> , 2001, 18, 356-356.	2.5	15
23	Effects of oral prophylaxis including tongue cleaning on halitosis and gingival inflammation in gingivitis patients: a randomized controlled clinical trial. <i>Clinical Oral Investigations</i> , 2019, 23, 1829-1836.	3.0	14
24	Diversity in Serine/Threonine Protein Kinase-4 Deficiency and Review of the Literature. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2021, 9, 3752-3766.e4.	3.8	13
25	Neurologic Involvement in Primary Immunodeficiency Disorders. <i>Journal of Child Neurology</i> , 2018, 33, 320-328.	1.4	12
26	Clinical and laboratory findings in patients with leukocyte adhesion deficiency type I: A multicenter study in Turkey. <i>Clinical and Experimental Immunology</i> , 2021, 206, 47-55.	2.6	10
27	Combined immunodeficiency due to purine nucleoside phosphorylase deficiency: Outcome of three patients. <i>European Journal of Medical Genetics</i> , 2022, 65, 104428.	1.3	10
28	Successful hematopoietic stem cell transplantation after myeloablative conditioning in three patients with dedicator of cytokinesis 8 deficiency (DOCK8) related Hyper IgE syndrome. <i>Bone Marrow Transplantation</i> , 2018, 53, 339-343.	2.4	8
29	Clinical and Immunological Characteristics of 63 Patients with Chronic Granulomatous Disease: Hacettepe Experience. <i>Journal of Clinical Immunology</i> , 2021, 41, 992-1003.	3.8	8
30	Selective IgM deficiency: Follow-up and outcome. <i>Pediatric Allergy and Immunology</i> , 2021, 32, 1327-1334.	2.6	8
31	Differential diagnosis of primary immunodeficiency in patients with BCGitis and BCGosis: A single-centre study. <i>Scandinavian Journal of Immunology</i> , 2021, 94, e13084.	2.7	8
32	Levels of pro- and anti-inflammatory cytokines in cystic fibrosis patients with or without gingivitis. <i>Cytokine</i> , 2020, 127, 154987.	3.2	7
33	Evaluation of periodontal status and cytokine/chemokine profile of GCF in patients with severe congenital neutropenia. <i>Odontology / the Society of the Nippon Dental University</i> , 2021, 109, 474-482.	1.9	7
34	Acute myeloid leukemia in a child with dedicator of cytokinesis 8 (DOCK8) deficiency. <i>Pediatric Blood and Cancer</i> , 2017, 64, e26695.	1.5	6
35	Recurrent Demyelinating Episodes as Sole Manifestation of Inherited CD59 Deficiency. <i>Neuropediatrics</i> , 2020, 51, 206-210.	0.6	6
36	Impact of mannose-binding lectin 2 gene polymorphisms on disease severity in noncystic fibrosis bronchiectasis in children. <i>Pediatric Pulmonology</i> , 2020, 55, 1190-1198.	2.0	5

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37	Long Term Follow-Up of the Patients with Severe Combined Immunodeficiency After Hematopoietic Stem Cell Transplantation: A Single-Center Study. <i>Immunological Investigations</i> , 2022, 51, 739-747.	2.0	4
38	GIMAP6 regulates autophagy, immune competence, and inflammation in mice and humans. <i>Journal of Experimental Medicine</i> , 2022, 219, .	8.5	4
39	Adenosine Deaminase Type II Deficiency: Severe Chronic Neutropenia, Lymphoid Infiltration in Bone Marrow, and Inflammatory Features. <i>Immunological Investigations</i> , 2020, , 1-9.	2.0	3
40	Elevated Interleukin-17A expression in amlodipine-induced gingival overgrowth. <i>Journal of Periodontal Research</i> , 2020, 55, 613-621.	2.7	3
41	A Patient With AIRE Mutation Who Presented With Severe Diarrhea and Lung Abscess. <i>Pediatric Infectious Disease Journal</i> , 2021, 40, 66-69.	2.0	3
42	Lymphocyte Subgroups and KREC Numbers in Common Variable Immunodeficiency: A Single Center Study. <i>Journal of Clinical Immunology</i> , 2020, 40, 494-502.	3.8	2
43	Flow Cytometry is a Reliable Tool in the Diagnosis of STK4 Deficiency. <i>Asim, Allerji, Immunoloji</i> , 0, , .	0.0	2
44	Cytokine profile in serum and gingival crevicular fluid of children with inflammatory bowel disease: A case-control study. <i>Journal of Periodontology</i> , 2021, , .	3.4	2
45	Recurrent skin abscesses in a female X-linked chronic granulomatous disease carrier. <i>Journal of Cosmetic Dermatology</i> , 2020, 19, 1810-1812.	1.6	1
46	Expression of HLA class I and class II genes in patients with multiple skin warts. <i>Experimental Dermatology</i> , 2021, 30, 1642-1649.	2.9	1
47	Retroperitoneal Abscess in Severe Combined Immunodeficiency Probably Due to BCG Vaccine. <i>Asim, Allerji, Immunoloji</i> , 2020, 18, 98-101.	0.0	1
48	Primary Immunodeficiencies Associated with Atopic Dermatitis. <i>Asim, Allerji, Immunoloji</i> , 2019, 17, 70-77.	0.0	1
49	In case of recurrent wheezing and bronchiolitis: Think again, it may be a primary immunodeficiency. <i>Asian Pacific Journal of Allergy and Immunology</i> , 2021, , .	0.4	1
50	Frequency of HLA Class I and Class II Alleles in Patients with CVID from Turkey. <i>Immunological Investigations</i> , 2021, 50, 363-371.	2.0	0
51	Clinical, Laboratory Features and Clinical Courses of Patients with Wiskott Aldrich Syndrome and X-linked Thrombocytopenia: A single center study. <i>Immunological Investigations</i> , 2022, 51, 1272-1283.	2.0	0
52	Nutritional status of children with primary immunodeficiency: A single center experience. <i>Pediatrics International</i> , 2022, 64, .	0.5	0
53	Three patients with glucose-6 phosphatase catalytic subunit 3 deficiency. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2020, 33, 957-961.	0.9	0
54	Evaluation of Health Status and Quality of Life in Patients Using Intravenous and Subcutaneous Forms of Immunoglobulin Replacement. <i>Journal of Ankara University Faculty of Medicine</i> , 2022, 75, 77-83.	0.1	0

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55	First allogeneic hematopoietic stem cell transplantation in RASGRP1 deficiency: long-term follow-up. Bone Marrow Transplantation, 2022, , .	2.4	0