Ä°smail Reisli

List of Publications by Year
in descending order

33	1,167	516710 16	454955 30
papers	citations	h-index	g-index

36	36	36	2580
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	STK4 deficiency and EBV-associated lymphoproliferative disorders, emphasis on histomorphology, and review of literature. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2022, 480, 393-401.	2.8	5
2	Long-Term Experience of Subcutaneous Immunoglobulin Therapy in Pediatric Primary Immunodeficient Patients with Low and Normal Body Weight. Journal of Clinical Immunology, 2022, 42, 64-71.	3.8	2
3	Comparing the levels of CTLAâ€4â€dependent biological defects in patients with LRBA deficiency and CTLAâ€4 insufficiency. Allergy: European Journal of Allergy and Clinical Immunology, 2022, 77, 3108-3123.	5.7	7
4	A family screening of CD19 gene mutation by PCR-RFLP. European Journal of Clinical and Experimental Medicine, 2022, 20, 141-145.	0.1	1
5	Protein functionality as a potential bottleneck for somatic revertant variants. Journal of Allergy and Clinical Immunology, 2021, 147, 391-393.e8.	2.9	3
6	Consensus Middle East and North Africa Registry on Inborn Errors of Immunity. Journal of Clinical Immunology, 2021, 41, 1339-1351.	3.8	33
7	Simulation Based Endotracheal Intubation Education for Residents of Pediatrics. Eurasian Journal of Emergency Medicine, 2021, 20, 91-94.	0.2	0
8	Simple Measurement of IgA Predicts Immunity and Mortality in Ataxia-Telangiectasia. Journal of Clinical Immunology, 2021, 41, 1878-1892.	3.8	9
9	An association between immune status and chest CT scores in COVIDâ€19 patients. International Journal of Clinical Practice, 2021, 75, e14767.	1.7	3
10	Primary antibody deficiencies in Turkey: molecular and clinical aspects. Immunologic Research, 2021, , 1.	2.9	2
11	Combined immunodeficiency caused by a loss-of-function mutation in DNA polymerase delta 1. Journal of Allergy and Clinical Immunology, 2020, 145, 391-401.e8.	2.9	28
12	ILC3 deficiency and generalized ILC abnormalities in DOCK8â€deficient patients. Allergy: European Journal of Allergy and Clinical Immunology, 2020, 75, 921-932.	5.7	17
13	P0088THE INFILTRATION OF T AND B LYMPHOCYTES AND NK CELLS IN KIDNEY BIOPSIES OF PATIENTS WITH FABRY DISESE UNDER ENZYME REPLACEMENT THERAPHY. Nephrology Dialysis Transplantation, 2020, 35, .	0.7	0
14	Mutational landscape of severe combined immunodeficiency patients from Turkey. International Journal of Immunogenetics, 2020, 47, 529-538.	1.8	14
15	Abatacept as a Long-Term Targeted Therapy for LRBA Deficiency. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 2790-2800.e15.	3.8	112
16	Inherited IFNAR1 deficiency in otherwise healthy patients with adverse reaction to measles and yellow fever live vaccines. Journal of Experimental Medicine, 2019, 216, 2057-2070.	8.5	127
17	Spatiotemporal Gradient of Cortical Neuron Death Contributes to Microcephaly in Knock-In Mouse Model of Ligase 4 Syndrome. American Journal of Pathology, 2019, 189, 2440-2449.	3.8	2
18	B cellâ€“intrinsic requirement for STK4 in humoral immunity in mice and human subjects. Journal of Allergy and Clinical Immunology, 2019, 143, 2302-2305.	2.9	21

#	ARTICLE	IF	CITATIONS
19	F-BAR domain only protein 1 (FCHO1) deficiency is a novel cause of combined immune deficiency in human subjects. Journal of Allergy and Clinical Immunology, 2019, 143, 2317-2321.e12.	2.9	21
20	Intrauterine detection of DCLRE1C (Artemis) mutation by restriction fragment length polymorphism. Pediatric Allergy and Immunology, 2019, 30, 668-671.	2.6	4
21	Patients with CD3G mutations reveal a role for human CD3Î³ in Treg diversity and suppressive function. Blood, 2018, 131, 2335-2344.	1.4	51
22	Type I IFNâ€“related NETosis in ataxia telangiectasia and Artemis deficiency. Journal of Allergy and Clinical Immunology, 2018, 142, 246-257.	2.9	47
23	Tuberculosis and impaired IL-23â€“dependent IFN-Î³ immunity in humans homozygous for a common TYK2 missense variant. Science Immunology, 2018, 3, .	11.9	148
24	Evaluation of Clinical and Immunological Characteristics of Children with Common Variable Immunodeficiency. International Journal of Pediatrics (United Kingdom), 2018, 2018, 1-8.	0.8	12
25	Hematopoietic stem cell transplantation from unrelated donors in children with <scp>DOCK</scp>8 deficiency. Pediatric Transplantation, 2017, 21, e13015.	1.0	12
26	Natural Killer Cells from Patients with Recombinase-Activating Gene and Non-Homologous End Joining Gene Defects Comprise a Higher Frequency of CD56bright NKG2A+++ Cells, and Yet Display Increased Degranulation and Higher Perforin Content. Frontiers in Immunology, 2017, 8, 798.	4.8	41
27	Common variable immunodeficiency in adults requires reserved protocols for long-term follow-up. Turkish Journal of Medical Sciences, 2016, 46, 430-436.	0.9	7
28	Dedicator of cytokinesis 8 regulates signal transducer and activator of transcription 3 activation and promotes TH17Âcell differentiation. Journal of Allergy and Clinical Immunology, 2016, 138, 1384-1394.e2.	2.9	70
29	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
30	Mutations in CDCA7 and HELLS cause immunodeficiencyâ€“centromeric instabilityâ€“facial anomalies syndrome. Nature Communications, 2015, 6, 7870.	12.8	148
31	Human CD19 and CD40L deficiencies impair antibody selection and differentially affect somatic hypermutation. Journal of Allergy and Clinical Immunology, 2014, 134, 135-144.e7.	2.9	71
32	Plasmacytoid dendritic cell depletion in DOCK8 deficiency: Rescue of severe herpetic infections with IFN-Î± 2b therapy. Journal of Allergy and Clinical Immunology, 2014, 133, 1753-1755.e3.	2.9	46
33	Acute rheumatic fever in Konya, Turkey. Pediatrics International, 2000, 42, 71-75.	0.5	28