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	Mutations in CDCA7 and HELLS cause immunodeficiencyâ€“centromeric instabilityâ€“facial anomalies syndrome. Nature Communications, 2015, 6, 7870.	12.8	148
2	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
3	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
4	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
5	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
6	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
7	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
8	Patients with CD3G mutations reveal a role for human CD3Î³ in Treg diversity and suppressive function. Blood, 2018, 131, 2335-2344.	1.4	51
9	Type I IFNâ€“related NETosis in ataxia telangiectasia and Artemis deficiency. Journal of Allergy and Clinical Immunology, 2018, 142, 246-257.	2.9	47
10	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
11	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
12	Consensus Middle East and North Africa Registry on Inborn Errors of Immunity. Journal of Clinical Immunology, 2021, 41, 1339-1351.	3.8	33
13	Acute rheumatic fever in Konya, Turkey. Pediatrics International, 2000, 42, 71-75.	0.5	28
14	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
15	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
16	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
17	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
18	Mutational landscape of severe combined immunodeficiency patients from Turkey. International Journal of Immunogenetics, 2020, 47, 529-538.	1.8	14

#	ARTICLE	IF	CITATIONS
19	Hematopoietic stem cell transplantation from unrelated donors in children with <scp>DOCK</scp>8 deficiency. Pediatric Transplantation, 2017, 21, e13015.	1.0	12
20	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
21	Simple Measurement of IgA Predicts Immunity and Mortality in Ataxia-Telangiectasia. Journal of Clinical Immunology, 2021, 41, 1878-1892.	3.8	9
22	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
23	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
24	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
25	Intrauterine detection of DCLRE1C (Artemis) mutation by restriction fragment length polymorphism. Pediatric Allergy and Immunology, 2019, 30, 668-671.	2.6	4
26	Protein functionality as a potential bottleneck for somatic revertant variants. Journal of Allergy and Clinical Immunology, 2021, 147, 391-393.e8.	2.9	3
27	An association between immune status and chest CT scores in COVIDâ€19 patients. International Journal of Clinical Practice, 2021, 75, e14767.	1.7	3
28	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
29	Primary antibody deficiencies in Turkey: molecular and clinical aspects. Immunologic Research, 2021, , 1.	2.9	2
30	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
31	A family screening of CD19 gene mutation by PCR-RFLP. European Journal of Clinical and Experimental Medicine, 2022, 20, 141-145.	0.1	1
32	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
33	Simulation Based Endotracheal Intubation Education for Residents of Pediatrics. Eurasian Journal of Emergency Medicine, 2021, 20, 91-94.	0.2	0