Meri Kaustio

List of Publications by Year
in descending order

14	524	1163117 8	1372567 10
papers	citations	h-index	g-index

15	15	15	1324
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Loss of DIAPH1 causes SCBMS, combined immunodeficiency, and mitochondrial dysfunction. Journal of Allergy and Clinical Immunology, 2021, 148, 599-611.	2.9	23
2	Loss-of-function mutation in IKZF2 leads to immunodeficiency with dysregulated germinal center reactions and reduction of MAIT cells. Science Immunology, 2021, 6, eabe3454.	11.9	30
3	Identification of novel regulators of STAT3 activity. PLoS ONE, 2020, 15, e0230819.	2.5	12
4	Identification of novel regulators of STAT3 activity. , 2020, 15, e0230819.		0
5	Identification of novel regulators of STAT3 activity. , 2020, 15, e0230819.		0
6	Identification of novel regulators of STAT3 activity. , 2020, 15, e0230819.		0
7	Identification of novel regulators of STAT3 activity. , 2020, 15, e0230819.		0
8	Primary Immunodeficiency, a Possible Cause of Neutrophilic Necrotizing Dermatosis. JAMA Dermatology, 2019, 155, 863.	4.1	5
9	ADA2 deficiency: Clonal lymphoproliferation in a subset of patients. Journal of Allergy and Clinical Immunology, 2018, 141, 1534-1537.e8.	2.9	71
10	Damaging heterozygous mutations in NFKB1 lead to diverse immunologic phenotypes. Journal of Allergy and Clinical Immunology, 2017, 140, 782-796.	2.9	113
11	Constant B cell lymphocytosis since early age in a patient with CARD11 mutation: A 20-year follow-up. Clinical Immunology, 2016, 165, 19-20.	3.2	17
12	Enrichment of rare variants in population isolates: single AICDA mutation responsible for hyper-IgM syndrome type 2 in Finland. European Journal of Human Genetics, 2016, 24, 1473-1478.	2.8	22
13	Autoimmunity, hypogammaglobulinemia, lymphoproliferation, and mycobacterial disease in patients with activating mutations in STAT3. Blood, 2015, 125, 639-648.	1.4	229
14	Dominant NFKB1 Mutations Cause Antibody Deficiency and Autoinflammatory Episodes. Blood, 2015, 126, 206-206.	1.4	1