Maximilian Heeg

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/6986040/publications.pdf

Version: 2024-02-01

22 papers	777 citations	623734 14 h-index	888059 17 g-index
23 all docs	23 docs citations	23 times ranked	1238 citing authors

#	Article	IF	Citations
1	Disease Evolution and Response to Rapamycin in Activated Phosphoinositide 3-Kinase Î´ Syndrome: The European Society for Immunodeficiencies-Activated Phosphoinositide 3-Kinase Î´ Syndrome Registry. Frontiers in Immunology, 2018, 9, 543.	4.8	137
2	Clinical and Immunological Phenotype of Patients With Primary Immunodeficiency Due to Damaging Mutations in NFKB2. Frontiers in Immunology, 2019, 10, 297.	4.8	117
3	Tissue-resident memory CD8+ T cells possess unique transcriptional, epigenetic and functional adaptations to different tissue environments. Nature Immunology, 2022, 23, 1121-1131.	14.5	84
4	Early-onset autoimmunity associated with SOCS1 haploinsufficiency. Nature Communications, 2020, 11, 5341.	12.8	74
5	The German National Registry of Primary Immunodeficiencies (2012â€“2017). Frontiers in Immunology, 2019, 10, 1272.	4.8	71
6	Primary and secondary hemophagocytic lymphohistiocytosis have different patterns of Tâ€cell activation, differentiation and repertoire. European Journal of Immunology, 2017, 47, 364-373.	2.9	69
7	Distinct molecular response patterns of activating STAT3 mutations associate with penetrance of lymphoproliferation and autoimmunity. Clinical Immunology, 2020, 210, 108316.	3.2	40
8	Neuroinflammatory Disease as an Isolated Manifestation of Hemophagocytic Lymphohistiocytosis. Journal of Clinical Immunology, 2020, 40, 901-916.	3.8	33
9	Is an infectious trigger always required for primary hemophagocytic lymphohistiocytosis? Lessons from in utero and neonatal disease. Pediatric Blood and Cancer, 2018, 65, e27344.	1.5	26
10	Germline STAT3 gain-of-function mutations in primary immunodeficiency: Impact on the cellular and clinical phenotype. Biomedical Journal, 2021, 44, 412-421.	3.1	25
11	A distinct CD38+CD45RA+ population of CD4+, CD8+, and double-negative T cells is controlled by FAS. Journal of Experimental Medicine, 2021, 218, .	8.5	25
12	Hemophagocytic lymphohistiocytosis as presenting manifestation of profound combined immunodeficiency due to an ORAI1 mutation. Journal of Allergy and Clinical Immunology, 2017, 140, 1721-1724.	2.9	23
13	Seletalisib for Activated PI3KÎ´ Syndromes: Open-Label Phase 1b and Extension Studies. Journal of Immunology, 2020, 205, 2979-2987.	0.8	21
14	Profound immunodeficiency with severe skin disease explained by concomitant novel CARMIL2 and PLEC1 loss-of-function mutations. Clinical Immunology, 2019, 208, 108228.	3.2	20
15	Functional flow cytometry of monocytes for routine diagnosis of innate primary immunodeficiencies. Journal of Allergy and Clinical Immunology, 2020, 145, 434-437.e4.	2.9	5
16	International multicenter experience of transjugular intrahepatic portosystemic shunt implantation in patients with common variable immunodeficiency. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 2931-2935.e1.	3.8	4
17	Therapeutic approaches to pediatric COVID-19: an online survey of pediatric rheumatologists. Rheumatology International, 2021, 41, 911-920.	3.0	3
18	Title is missing!. , 2020, 17, e1003076.		0

#	Article	IF	Citations
19	Title is missing!. , 2020, 17, e1003076.		0
20	Title is missing!. , 2020, 17, e1003076.		0
21	Title is missing!. , 2020, 17, e1003076.		0
22	Title is missing!. , 2020, 17, e1003076.		0