Maximilian Heeg

List of Publications by Year
in descending order

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

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22	777	623734	888059
papers	citations	14 h-index	17 g-index

23	23	23	1238
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	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
2	Therapeutic approaches to pediatric COVID-19: an online survey of pediatric rheumatologists. Rheumatology International, 2021, 41, 911-920.	3.0	3
3	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
4	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
5	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
6	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
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	Early-onset autoimmunity associated with SOCS1 haploinsufficiency. Nature Communications, 2020, 11, 5341.	12.8	74
9	Seletalisib for Activated PI3KÎ´ Syndromes: Open-Label Phase 1b and Extension Studies. Journal of Immunology, 2020, 205, 2979-2987.	0.8	21
10	Neuroinflammatory Disease as an Isolated Manifestation of Hemophagocytic Lymphohistiocytosis. Journal of Clinical Immunology, 2020, 40, 901-916.	3.8	33
11	Title is missing!. , 2020, 17, e1003076.		0
12	Title is missing!. , 2020, 17, e1003076.		0
13	Title is missing!. , 2020, 17, e1003076.		0
14	Title is missing!. , 2020, 17, e1003076.		0
15	Title is missing!. , 2020, 17, e1003076.		0
16	The German National Registry of Primary Immunodeficiencies (2012â€“2017). Frontiers in Immunology, 2019, 10, 1272.	4.8	71
17	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
18	Clinical and Immunological Phenotype of Patients With Primary Immunodeficiency Due to Damaging Mutations in NFKB2. Frontiers in Immunology, 2019, 10, 297.	4.8	117

#	Article	IF	Citations
19	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
20	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
21	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
22	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