

HÃ©lÃ¨ne Decaluwe

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/773807/publications.pdf>

Version: 2024-02-01

24
papers

765
citations

516561

16
h-index

642610

23
g-index

24
all docs

24
docs citations

24
times ranked

1587
citing authors

#	ARTICLE	IF	CITATIONS
1	The mutational landscape of SARS-CoV-2 variants diversifies TÃ©cell targets in an HLA-supertype-dependent manner. <i>Cell Systems</i> , 2022, 13, 143-157.e3.	2.9	22
2	Successful management of familial hemophagocytic lymphohistiocytosis by the JAK 1/2 inhibitor ruxolitinib. <i>Pediatric Blood and Cancer</i> , 2021, 68, e28954.	0.8	6
3	Immune profiling of a patient with relapsed HIV-Related and EBV-positive diffuse large B-Cell lymphoma treated with pembrolizumab. <i>Leukemia and Lymphoma</i> , 2021, , 1-5.	0.6	0
4	Association of outcomes in acute flaccid myelitis with identification of enterovirus at presentation: a Canadian, nationwide, longitudinal study. <i>The Lancet Child and Adolescent Health</i> , 2020, 4, 828-836.	2.7	9
5	Common gamma chain cytokines and CD8 T cells in cancer. <i>Seminars in Immunology</i> , 2019, 42, 101307.	2.7	25
6	¹⁸ F-Fluorodeoxyglucose positron emission tomography with computed tomography (FDG PET/CT) findings in children with encephalitis and comparison to conventional imaging. <i>European Journal of Nuclear Medicine and Molecular Imaging</i> , 2019, 46, 1309-1324.	3.3	24
7	SCID genotype and 6-month posttransplant CD4 count predict survival and immune recovery. <i>Blood</i> , 2018, 132, 1737-1749.	0.6	128
8	B-cell differentiation and IL-21 response in IL2RG/JAK3 SCID patients after hematopoietic stem cell transplantation. <i>Blood</i> , 2018, 131, 2967-2977.	0.6	37
9	A Novel <i>PGM3</i> Mutation Is Associated With a Severe Phenotype of Bone Marrow Failure, Severe Combined Immunodeficiency, Skeletal Dysplasia, and Congenital Malformations. <i>Journal of Bone and Mineral Research</i> , 2017, 32, 1853-1859.	3.1	28
10	Very Early-Onset Inflammatory Manifestations of X-Linked Chronic Granulomatous Disease. <i>Frontiers in Immunology</i> , 2017, 8, 1167.	2.2	23
11	Primary Immune Deficiency Treatment Consortium (PIDTC) update. <i>Journal of Allergy and Clinical Immunology</i> , 2016, 138, 375-385.	1.5	33
12	IL2R β -dependent signals drive terminal exhaustion and suppress memory development during chronic viral infection. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2016, 113, E5444-53.	3.3	45
13	Cytokines and persistent viral infections. <i>Cytokine</i> , 2016, 82, 4-15.	1.4	33
14	IL β and IL α 15 regulate CD8 ⁺ memory T α cell differentiation but are dispensable for protective recall responses. <i>European Journal of Immunology</i> , 2015, 45, 3324-3338.	1.6	27
15	Multiple Intestinal Atresia With Combined Immune Deficiency Related to TTC7A Defect Is a Multiorgan Pathology. <i>Medicine (United States)</i> , 2014, 93, e327.	0.4	35
16	SCIg vs. IVIg: let α ™s give patients the choice!. <i>Allergy, Asthma and Clinical Immunology</i> , 2014, 10, A13.	0.9	1
17	Exome sequencing identifies mutations in the gene <i>TTC7A</i> in French-Canadian cases with hereditary multiple intestinal atresia. <i>Journal of Medical Genetics</i> , 2013, 50, 324-329.	1.5	119
18	Ataxia-Telangiectasia Presenting With a Novel Immunodeficiency. <i>Pediatric Neurology</i> , 2012, 46, 322-324.	1.0	11

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19	Î³_c deficiency precludes CD8⁺ T cell memory despite formation of potent T cell effectors. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 9311-9316.	3.3	28
20	IFN-Î³ Mediates the Rejection of Haematopoietic Stem Cells in IFN-Î³R1-Deficient Hosts. PLoS Medicine, 2008, 5, e26.	3.9	67
21	Î³_c, cytokines provide multiple homeostatic signals to naive CD4⁺ T cells. European Journal of Immunology, 2007, 37, 2606-2616.	1.6	28
22	Procalcitonin in Children with Escherichia coli O157:H7 Associated Hemolytic Uremic Syndrome. Pediatric Research, 2006, 59, 579-583.	1.1	14
23	Rabies in a Nine-Year-Old Child: The Myth of the Bite. Canadian Journal of Infectious Diseases & Medical Microbiology, 2002, 13, 121-125.	0.3	9
24	Molecular characterization of a pediatric pheochromocytoma with suspected bilateral disease. Journal of Pediatrics, 2001, 138, 269-273.	0.9	13