

Anne Rensing-Ehl

List of Publications by Year in descending order

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Version: 2024-02-01

18
papers

1,596
citations

567281

15
h-index

794594

19
g-index

19
all docs

19
docs citations

19
times ranked

3216
citing authors

#	ARTICLE	IF	CITATIONS
1	Autosomal dominant immune dysregulation syndrome in humans with CTLA4 mutations. <i>Nature Medicine</i> , 2014, 20, 1410-1416.	30.7	723
2	Deficiency of Innate and Acquired Immunity Caused by an <i>IKBKB</i> Mutation. <i>New England Journal of Medicine</i> , 2013, 369, 2504-2514.	27.0	161
3	Autoimmune lymphoproliferative syndrome-like disease in patients with LRBA mutation. <i>Clinical Immunology</i> , 2015, 159, 84-92.	3.2	96
4	The expansion of human T-bet ^{high} CD21 ^{low} B cells is T cell dependent. <i>Science Immunology</i> , 2021, 6, eabh0891.	11.9	82
5	Hyperactive mTOR pathway promotes lymphoproliferation and abnormal differentiation in autoimmune lymphoproliferative syndrome. <i>Blood</i> , 2016, 128, 227-238.	1.4	77
6	Primary and secondary hemophagocytic lymphohistiocytosis have different patterns of T _H cell activation, differentiation and repertoire. <i>European Journal of Immunology</i> , 2017, 47, 364-373.	2.9	69
7	Germline TET2 loss of function causes childhood immunodeficiency and lymphoma. <i>Blood</i> , 2020, 136, 1055-1066.	1.4	58
8	Abnormally differentiated CD4 ⁺ or CD8 ⁺ T cells with phenotypic and genetic features of double negative T cells in human Fas deficiency. <i>Blood</i> , 2014, 124, 851-860.	1.4	54
9	Evolution of disease activity and biomarkers on and off rapamycin in 28 patients with autoimmune lymphoproliferative syndrome. <i>Haematologica</i> , 2017, 102, e52-e56.	3.5	49
10	Activated $\text{PI}3\text{K}\gamma$ syndrome type 2: Two patients, a novel mutation, and review of the literature. <i>Pediatric Allergy and Immunology</i> , 2016, 27, 640-644.	2.6	46
11	Distinct molecular response patterns of activating STAT3 mutations associate with penetrance of lymphoproliferation and autoimmunity. <i>Clinical Immunology</i> , 2020, 210, 108316.	3.2	40
12	SYK expression endows human ZAP70-deficient CD8 T cells with residual TCR signaling. <i>Clinical Immunology</i> , 2015, 161, 103-109.	3.2	38
13	Clinical and Molecular Heterogeneity of RTEL1 Deficiency. <i>Frontiers in Immunology</i> , 2017, 8, 449.	4.8	35
14	Frequency, function and CLA expression of CD4 ⁺ CD25 ⁺ FOXP3 ⁺ regulatory T cells in bullous pemphigoid. <i>Experimental Dermatology</i> , 2007, 16, 13-21.	2.9	33
15	A distinct CD38 ⁺ CD45RA ⁺ population of CD4 ⁺ , CD8 ⁺ , and double-negative T cells is controlled by FAS. <i>Journal of Experimental Medicine</i> , 2021, 218, .	8.5	25
16	IgG4-related disease in autoimmune lymphoproliferative syndrome. <i>Clinical Immunology</i> , 2017, 180, 97-99.	3.2	5
17	Complete CD95/FAS deficiency due to complex homozygous germline TNFRSF6 mutations in an adult patient with mild autoimmune lymphoproliferative syndrome (ALPS). <i>Clinical Immunology</i> , 2021, 228, 108757.	3.2	3
18	Hyperactive mTOR Pathway Promotes Lymphoproliferation and Abnormal Differentiation in Human Autoimmune Lymphoproliferative Syndrome. <i>Blood</i> , 2015, 126, 1020-1020.	1.4	1