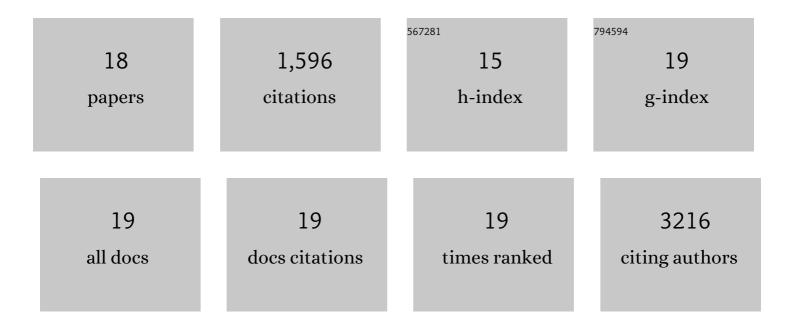
## Anne Rensing-Ehl

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/5760658/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Complete CD95/FAS deficiency due to complex homozygous germline TNFRSF6 mutations in an adult patient with mild autoimmune lymphoproliferative syndrome (ALPS). Clinical Immunology, 2021, 228, 108757.	3.2	3
2	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
3	The expansion of human T-bet <sup>high</sup> CD21 <sup>low</sup> B cells is T cell dependent. Science Immunology, 2021, 6, eabh0891.	11.9	82
4	Distinct molecular response patterns of activating STAT3 mutations associate with penetrance of lymphoproliferation and autoimmunity. Clinical Immunology, 2020, 210, 108316.	3.2	40
5	Germline TET2 loss of function causes childhood immunodeficiency and lymphoma. Blood, 2020, 136, 1055-1066.	1.4	58
6	Evolution of disease activity and biomarkers on and off rapamycin in 28 patients with autoimmune lymphoproliferative syndrome. Haematologica, 2017, 102, e52-e56.	3.5	49
7	lgG4-related disease in autoimmune lymphoproliferative syndrome. Clinical Immunology, 2017, 180, 97-99.	3.2	5
8	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
9	Clinical and Molecular Heterogeneity of RTEL1 Deficiency. Frontiers in Immunology, 2017, 8, 449.	4.8	35
10	Hyperactive mTOR pathway promotes lymphoproliferation and abnormal differentiation in autoimmune lymphoproliferative syndrome. Blood, 2016, 128, 227-238.	1.4	77
11	Activated <scp>PI</scp> 3Kl̂´syndrome type 2: Two patients, a novel mutation, and review of the literature. Pediatric Allergy and Immunology, 2016, 27, 640-644.	2.6	46
12	SYK expression endows human ZAP70-deficient CD8 T cells with residual TCR signaling. Clinical Immunology, 2015, 161, 103-109.	3.2	38
13	Autoimmune lymphoproliferative syndrome-like disease in patients with LRBA mutation. Clinical Immunology, 2015, 159, 84-92.	3.2	96
14	Hyperactive mTOR Pathway Promotes Lymphoproliferation and Abnormal Differentiation in Human Autoimmune Lymphoproliferative Syndrome. Blood, 2015, 126, 1020-1020.	1.4	1
15	Abnormally differentiated CD4+ or CD8+ T cells with phenotypic and genetic features of double negative T cells in human Fas deficiency. Blood, 2014, 124, 851-860.	1.4	54
16	Autosomal dominant immune dysregulation syndrome in humans with CTLA4 mutations. Nature Medicine, 2014, 20, 1410-1416.	30.7	723
17	Deficiency of Innate and Acquired Immunity Caused by an <i>IKBKB</i> Mutation. New England Journal of Medicine, 2013, 369, 2504-2514.	27.0	161
18	Frequency, function and CLA expression of CD4+CD25+FOXP3+ regulatory T cells in bullous pemphigoid. Experimental Dermatology, 2007, 16, 13-21.	2.9	33