

Elisabeth Salzer

List of Publications by Year in descending order

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Version: 2024-02-01

17
papers

1,151
citations

567281

15
h-index

940533

16
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all docs

17
docs citations

17
times ranked

2209
citing authors

#	ARTICLE	IF	CITATIONS
1	Extended clinical and immunological phenotype and transplant outcome in CD27 and CD70 deficiency. <i>Blood</i> , 2020, 136, 2638-2655.	1.4	64
2	The cytoskeletal regulator HEM1 governs B cell development and prevents autoimmunity. <i>Science Immunology</i> , 2020, 5, .	11.9	37
3	Human DEF6 deficiency underlies an immunodeficiency syndrome with systemic autoimmunity and aberrant CTLA-4 homeostasis. <i>Nature Communications</i> , 2019, 10, 3106.	12.8	48
4	PD-1 and PD1 expression in post-transplantation lymphoproliferative disease (PTLD) of childhood and adolescence: An inter- and intra-individual descriptive study covering the whole spectrum of PTLD categories. <i>Cancer Medicine</i> , 2019, 8, 4656-4668.	2.8	14
5	Polymerase δ deficiency causes syndromic immunodeficiency with replicative stress. <i>Journal of Clinical Investigation</i> , 2019, 129, 4194-4206.	8.2	41
6	Identifying Novel Inborn Errors of the Immune System. , 2017, , .		0
7	WIP deficiency severely affects human lymphocyte architecture during migration and synapse assembly. <i>Blood</i> , 2017, 130, 1949-1953.	1.4	25
8	Protein Kinase C δ : a Gatekeeper of Immune Homeostasis. <i>Journal of Clinical Immunology</i> , 2016, 36, 631-640.	3.8	69
9	RASGRP1 deficiency causes immunodeficiency with impaired cytoskeletal dynamics. <i>Nature Immunology</i> , 2016, 17, 1352-1360.	14.5	115
10	Combined immunodeficiency with CD4 lymphopenia and sclerosing cholangitis caused by a novel loss-of-function mutation affecting IL21R. <i>Haematologica</i> , 2015, 100, e216-e219.	3.5	46
11	Potentially Beneficial Effect of Hydroxychloroquine in a Patient with a Novel Mutation in Protein Kinase C δ Deficiency. <i>Journal of Clinical Immunology</i> , 2015, 35, 523-526.	3.8	37
12	Combined Immunodeficiency Evolving into Predominant CD4+ Lymphopenia Caused by Somatic Chimerism in JAK3. <i>Journal of Clinical Immunology</i> , 2014, 34, 941-953.	3.8	25
13	Early-onset inflammatory bowel disease and common variable immunodeficiency-like disease caused by IL-21 deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2014, 133, 1651-1659.e12.	2.9	124
14	Biallelic loss-of-function mutation in NIK causes a primary immunodeficiency with multifaceted aberrant lymphoid immunity. <i>Nature Communications</i> , 2014, 5, 5360.	12.8	116
15	JAGN1 deficiency causes aberrant myeloid cell homeostasis and congenital neutropenia. <i>Nature Genetics</i> , 2014, 46, 1021-1027.	21.4	119
16	B-cell deficiency and severe autoimmunity caused by deficiency of protein kinase C δ . <i>Blood</i> , 2013, 121, 3112-3116.	1.4	118
17	Combined immunodeficiency with life-threatening EBV-associated lymphoproliferative disorder in patients lacking functional CD27. <i>Haematologica</i> , 2013, 98, 473-478.	3.5	153