Jana Pachlopnik Schmid

List of Publications by Year
in descending order

58	4,122	186265	155660
papers	citations	28 h-index	55 g-index

58	58	58	5343
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Therapeutic options for CTLA-4 insufficiency. Journal of Allergy and Clinical Immunology, 2022, 149, 736-746.	2.9	68
2	Wenn hinter einer Immunthrombozytopenie, einer vermeintlichen Sarkoidose, rezidivierendem Fieber oder einer Lymphadenopathie ein angeborener Immundefekt steckt. Paediatrica, 2022, 33, .	0.1	0
3	Lorsque derriÃ¨re une thrombocytopÃ©nie, une sarcoÃ¯dose prÃ©sumÃ©e, une fiÃ¨vre rÃ©cidivante ou une lymphadÃ©nopathie se cache un dÃ©ficit immunitaire congÃ©nital. Paediatrica, 2022, 33, .	0.0	0
4	Sequence-Specific Features of Short Double-Strand, Blunt-End RNAs Have RIG-I- and Type 1 Interferon-Dependent or -Independent Anti-Viral Effects. Viruses, 2022, 14, 1407.	3.3	1
5	Patterns of Immune Dysregulation in Primary Immunodeficiencies: A Systematic Review. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 792-802.e10.	3.8	16
6	Best Practice Recommendations for the Diagnosis and Management of Children With Pediatric Inflammatory Multisystem Syndrome Temporally Associated With SARS-CoV-2 (PIMS-TS; Multisystem) Tj ETQq0 0 0 rgBT /Overlock 10 T	1.9	81
7	X-Linked Lymphoproliferative Disease Mimicking Multisystem Inflammatory Syndrome in Childrenâ€”A Case Report. Frontiers in Pediatrics, 2021, 9, 691024.	1.9	6
8	Thrombotic Microangiopathy Associated with Macrophage Activation Syndrome: A Multinational Study of 23 Patients. Journal of Pediatrics, 2021, 235, 196-202.	1.8	7
9	Multisystem inflammation and susceptibility to viral infections in human ZNFX1 deficiency. Journal of Allergy and Clinical Immunology, 2021, 148, 381-393.	2.9	40
10	Erythropoiesis defect observed in STAT3 GOF patients with severe anemia. Journal of Allergy and Clinical Immunology, 2020, 145, 1297-1301.	2.9	18
11	Distinct molecular response patterns of activating STAT3 mutations associate with penetrance of lymphoproliferation and autoimmunity. Clinical Immunology, 2020, 210, 108316.	3.2	40
12	Maturation of the Human Immunoglobulin Heavy Chain Repertoire With Age. Frontiers in Immunology, 2020, 11, 1734.	4.8	46
13	Targeted busulfan-based reduced-intensity conditioning and HLA-matched HSCT cure hemophagocytic lymphohistiocytosis. Blood Advances, 2020, 4, 1998-2010.	5.2	30
14	Obesity-Induced Increase in Cystatin C Alleviates Tissue Inflammation. Diabetes, 2020, 69, 1927-1935.	0.6	14
15	Case Report: Case Series of Children With Multisystem Inflammatory Syndrome Following SARS-CoV-2 Infection in Switzerland. Frontiers in Pediatrics, 2020, 8, 594127.	1.9	24
16	Swiss newborn screening for severe T and B cell deficiency with a combined TREC/KREC assay â€“ management recommendations. Swiss Medical Weekly, 2020, 150, w20254.	1.6	17
17	Epithelial proliferation in inflammatory skin disease is regulated by tetratricopeptide repeat domain 7 (Ttc7) in fibroblasts and lymphocytes. Journal of Allergy and Clinical Immunology, 2019, 143, 292-304.e8.	2.9	4
18	Recurrent inflammatory disease caused by a heterozygous mutation in CD48. Journal of Allergy and Clinical Immunology, 2019, 144, 1441-1445.e17.	2.9	9

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19	MHC-II Deficiency Among Egyptians: Novel Mutations and Unique Phenotypes. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 856-863.	3.8	19
20	Murine Models of Familial Cytokine Storm Syndromes. , 2019, , 467-489.		0
21	Aberrant tRNA processing causes an autoinflammatory syndrome responsive to TNF inhibitors. Annals of the Rheumatic Diseases, 2018, 77, 612-619.	0.9	49
22	Life-Threatening Primary Varicella Zoster Virus Infection With Hemophagocytic Lymphohistiocytosis-Like Disease in GATA2 Haploinsufficiency Accompanied by Expansion of Double Negative T-Lymphocytes. Frontiers in Immunology, 2018, 9, 2766.	4.8	10
23	Phenotype, penetrance, and treatment of 133 cytotoxic T-lymphocyte antigen 4â€“insufficient subjects. Journal of Allergy and Clinical Immunology, 2018, 142, 1932-1946.	2.9	344
24	Allogeneic hematopoietic cell transplantation in patients with GATA2 deficiencyâ€”a case report and comprehensive review of the literature. Annals of Hematology, 2018, 97, 1961-1973.	1.8	24
25	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
26	Persistent mammalian orthoreovirus, coxsackievirus and adenovirus co-infection in a child with a primary immunodeficiency detected by metagenomic sequencing: a case report. BMC Infectious Diseases, 2018, 18, 33.	2.9	16
27	Screening of 181 Patients With Antibody Deficiency for Deficiency of Adenosine Deaminase 2 Sheds New Light on the Disease in Adulthood. Arthritis and Rheumatology, 2017, 69, 1689-1700.	5.6	103
28	Causes of low neonatal T-cell receptor excision circles: A systematic review. Journal of Allergy and Clinical Immunology: in Practice, 2017, 5, 1457-1460.e22.	3.8	29
29	Hematopoietic stem cell transplantation in 29 patients hemizygous for hypomorphic IKBKG/NEMO mutations. Blood, 2017, 130, 1456-1467.	1.4	95
30	Unusual dermatological presentation and immune phenotype in <scp>SCID</scp> due to an <scp>IL</scp>7R mutation: the value of wholeâ€exome sequencing and the potential benefit of newborn screening. Journal of the European Academy of Dermatology and Venereology, 2017, 31, e147-e148.	2.4	2
31	Lymphadenopathy driven by TCR-Vβ8Vβ1 T-cell expansion in FAS-related autoimmune lymphoproliferative syndrome. Blood Advances, 2017, 1, 1101-1106.	5.2	3
32	The Phenotype and Genotype of Mevalonate Kinase Deficiency: A Series of 114 Cases From the Eurofever Registry. Arthritis and Rheumatology, 2016, 68, 2795-2805.	5.6	168
33	2016 Classification Criteria for Macrophage Activation Syndrome Complicating Systemic Juvenile Idiopathic Arthritis: A European League Against Rheumatism/American College of Rheumatology/Paediatric Rheumatology International Trials Organisation Collaborative Initiative. Arthritis and Rheumatology, 2016, 68, 566-576.	5.6	427
34	Expert consensus on dynamics of laboratory tests for diagnosis of macrophage activation syndrome complicating systemic juvenile idiopathic arthritis. RMD Open, 2016, 2, e000161.	3.8	57
35	2016 Classification Criteria for Macrophage Activation Syndrome Complicating Systemic Juvenile Idiopathic Arthritis. Annals of the Rheumatic Diseases, 2016, 75, 481-489.	0.9	338
36	The syndrome of hemophagocytic lymphohistiocytosis in primary immunodeficiencies: implications for differential diagnosis and pathogenesis. Haematologica, 2015, 100, 978-988.	3.5	161

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37	Immune deficiencyâ€“related enteropathy-lymphocytopenia-alopecia syndrome results from tetratricopeptide repeat domain 7A deficiency. Journal of Allergy and Clinical Immunology, 2014, 134, 1354-1364.e6.	2.9	66
38	Modern management of primary Tâ€cell immunodeficiencies. Pediatric Allergy and Immunology, 2014, 25, 300-313.	2.6	29
39	Distinct severity of HLH in both human and murine mutants with complete loss of cytotoxic effector PRF1, RAB27A, and STX11. Blood, 2013, 121, 595-603.	1.4	60
40	Polymerase Îµ1 mutation in a human syndrome with facial dysmorphism, immunodeficiency, livedo, and short stature (â€œFILS syndromeâ€). Journal of Experimental Medicine, 2012, 209, 2323-2330.	8.5	83
41	MST1 mutations in autosomal recessive primary immunodeficiency characterized by defective naive T-cell survival. Blood, 2012, 119, 3458-3468.	1.4	244
42	X-linked lymphoproliferative disease due to SAP/SH2D1A deficiency: a multicenter study on the manifestations, management and outcome of the disease. Blood, 2011, 117, 53-62.	1.4	268
43	Clinical similarities and differences of patients with X-linked lymphoproliferative syndrome type 1 (XLP-1/SAP deficiency) versus type 2 (XLP-2/XIAP deficiency). Blood, 2011, 117, 1522-1529.	1.4	320
44	Eosinophilia during Psoriasis Treatment with TNF Antagonists. Dermatology, 2011, 223, 311-315.	2.1	32
45	Premature Birth, Respiratory Distress, Intracerebral Hemorrhage, and Silvery-gray Hair. Journal of Pediatric Hematology/Oncology, 2010, 32, 494-496.	0.6	12
46	Inherited defects in lymphocyte cytotoxic activity. Immunological Reviews, 2010, 235, 10-23.	6.0	143
47	Neutralization of IFNÎ³ defeats haemophagocytosis in LCMVâ€infected perforinâ€and Rab27aâ€deficient mice. EMBO Molecular Medicine, 2009, 1, 112-124.	6.9	165
48	Hematopoietic stem cell transplantation in Griscelli syndrome type 2: a single-center report on 10 patients. Blood, 2009, 114, 211-218.	1.4	53
49	A Griscelli syndrome type 2 murine model of hemophagocytic lymphohistiocytosis (HLH). European Journal of Immunology, 2008, 38, 3219-3225.	2.9	54
50	Maternal Tobacco Smoking and Decreased Leukocytes, Including Dendritic Cells, in Neonates. Pediatric Research, 2007, 61, 462-466.	2.3	27
51	Non-lethal fetal toxicity of the angiotensin receptor blocker candesartan. Pediatric Nephrology, 2006, 21, 1329-1330.	1.7	23
52	Transient Hemophagocytosis With Deficient Cellular Cytotoxicity, Monoclonal Immunoglobulin M Gammopathy, Increased T-Cell Numbers, and Hypomorphic NEMO Mutation. Pediatrics, 2006, 117, e1049-e1056.	2.1	50
53	Natural anti-FcÎµRIÎ± autoantibodies may interfere with diagnostic tests for autoimmune urticaria. Journal of Autoimmunity, 2004, 22, 43-51.	6.5	37
54	Hemolytic uremic syndrome linked to infectious mononucleosis. Pediatric Nephrology, 2003, 18, 1193-1194.	1.7	10

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55	Allergic Manifestations as the Results of a Conditional Autoimmune Response. International Archives of Allergy and Immunology, 2001, 124, 411-413.	2.1	7
56	Conditional autoimmunity mediated by human natural antiâ€FcÎµRIÎ± autoantibodies?. FASEB Journal, 2001, 15, 2268-2274.	0.5	52
57	Interaction of Human IgE with Fc Epsilon RI Alpha Exposes Hidden Epitopes on IgE. International Archives of Allergy and Immunology, 1999, 120, 295-302.	2.1	7
58	Antigen Interaction and Heat Inactivation Expose New Epitopes on Human IgE. International Archives of Allergy and Immunology, 1998, 117, 231-238.	2.1	7