

# Janine Reichenbach

## List of Publications by Year in descending order

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87  
papers

8,686  
citations

66234

42  
h-index

56606

83  
g-index

88  
all docs

88  
docs citations

88  
times ranked

11429  
citing authors

#	ARTICLE	IF	CITATIONS
1	CRISPR-Directed Therapeutic Correction at the NCF1 Locus Is Challenged by Frequent Incidence of Chromosomal Deletions. <i>Molecular Therapy - Methods and Clinical Development</i> , 2020, 17, 936-943.	1.8	8
2	Membrane Dynamics and Organization of the Phagocyte NADPH Oxidase in PLB-985 Cells. <i>Frontiers in Cell and Developmental Biology</i> , 2020, 8, 608600.	1.8	7
3	Swiss newborn screening for severe T and B cell deficiency with a combined TREC/KREC assay – management recommendations. <i>Swiss Medical Weekly</i> , 2020, 150, w20254.	0.8	17
4	High Levels of IL-18 and IFN- $\gamma$ in Chronically Inflamed Tissue in Chronic Granulomatous Disease. <i>Frontiers in Immunology</i> , 2019, 10, 2236.	2.2	15
5	Human DOCK2 Deficiency: Report of a Novel Mutation and Evidence for Neutrophil Dysfunction. <i>Journal of Clinical Immunology</i> , 2019, 39, 298-308.	2.0	31
6	Novel Diagnostic Tool for p47-Deficient Chronic Granulomatous Disease Patient and Carrier Detection. <i>Molecular Therapy - Methods and Clinical Development</i> , 2019, 13, 274-278.	1.8	3
7	Ikaros family zinc finger 1 regulates dendritic cell development and function in humans. <i>Nature Communications</i> , 2018, 9, 1239.	5.8	62
8	Disease Evolution and Response to Rapamycin in Activated Phosphoinositide 3-Kinase $\gamma$ Syndrome: The European Society for Immunodeficiencies-Activated Phosphoinositide 3-Kinase $\gamma$ Syndrome Registry. <i>Frontiers in Immunology</i> , 2018, 9, 543.	2.2	137
9	Persistent mammalian orthoreovirus, coxsackievirus and adenovirus co-infection in a child with a primary immunodeficiency detected by metagenomic sequencing: a case report. <i>BMC Infectious Diseases</i> , 2018, 18, 33.	1.3	16
10	Lentiviral gene therapy vector with UCOE stably restores function in iPSC-derived neutrophils of a CDG patient. <i>Matters</i> , 2018, 2018, .	1.0	5
11	Development of a pCCLChim Lentiviral Vector for Gene Therapy of Patients with Chronic Granulomatous Disease (CGD) due to p47-phox Deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2017, 139, AB186.	1.5	0
12	CRISPR/Cas9-generated p47phox-deficient cell line for Chronic Granulomatous Disease gene therapy vector development. <i>Scientific Reports</i> , 2017, 7, 44187.	1.6	15
13	Non-invasive near-infrared fluorescence imaging of the neutrophil response in a mouse model of transient cerebral ischaemia. <i>Journal of Cerebral Blood Flow and Metabolism</i> , 2017, 37, 2833-2847.	2.4	18
14	Unusual dermatological presentation and immune phenotype in SCID due to an IL7R $\alpha$ mutation: the value of whole-exome sequencing and the potential benefit of newborn screening. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2017, 31, e147-e148.	1.3	2
15	Modern management of phagocyte defects. <i>Pediatric Allergy and Immunology</i> , 2017, 28, 124-134.	1.1	9
16	Severe glucose-6-phosphate dehydrogenase deficiency leads to susceptibility to infection and absent NETosis. <i>Journal of Allergy and Clinical Immunology</i> , 2017, 139, 212-219.e3.	1.5	56
17	Heterozygous STAT1 gain-of-function mutations underlie an unexpectedly broad clinical phenotype. <i>Blood</i> , 2016, 127, 3154-3164.	0.6	465
18	Clinical and immunologic phenotype associated with activated phosphoinositide 3-kinase $\gamma$ syndrome 2: A cohort study. <i>Journal of Allergy and Clinical Immunology</i> , 2016, 138, 210-218.e9.	1.5	215

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19	Loss of B Cells in Patients with Heterozygous Mutations in IKAROS. <i>New England Journal of Medicine</i> , 2016, 374, 1032-1043.	13.9	217
20	Hyperinflammation in patients with chronic granulomatous disease leads to impairment of hematopoietic stem cell functions. <i>Journal of Allergy and Clinical Immunology</i> , 2016, 138, 219-228.e9.	1.5	74
21	Neutrophil oxidative burst activates ATM to regulate cytokine production and apoptosis. <i>Blood</i> , 2015, 126, 2842-2851.	0.6	58
22	The Swiss National Registry for Primary Immunodeficiencies: report on the first 6 yearsâ€™ activity from 2008 to 2014. <i>Clinical and Experimental Immunology</i> , 2015, 182, 45-50.	1.1	46
23	Newborn Screening for Primary Immunodeficiencies: Focus on Severe Combined Immunodeficiency (SCID) and Other Severe T-Cell Lymphopenias. <i>International Journal of Neonatal Screening</i> , 2015, 1, 89-100.	1.2	4
24	Preliminary Evidence for a Compromised T-Cell Compartment in Maltreated Children with Depression and Posttraumatic Stress Disorder. <i>NeuroImmunoModulation</i> , 2015, 22, 303-310.	0.9	1
25	Defective nuclear entry of hydrolases prevents neutrophil extracellular trap formation in patients with chronic granulomatous disease. <i>Journal of Allergy and Clinical Immunology</i> , 2015, 136, 1703-1706.e5.	1.5	14
26	Gene Therapy for X-Linked Chronic Granulomatous Disease (Net4CGD). <i>Human Gene Therapy Clinical Development</i> , 2015, 26, 88-90.	3.2	1
27	TALEN-mediated functional correction of X-linked chronic granulomatous disease in patient-derived induced pluripotent stem cells. <i>Biomaterials</i> , 2015, 69, 191-200.	5.7	76
28	Successful Combination of Sequential Gene Therapy and Rescue Allo-HSCT in Two Children with X-CGD - Importance of Timing. <i>Current Gene Therapy</i> , 2015, 15, 416-427.	0.9	61
29	The European Society for Immunodeficiencies (ESID) registry 2014. <i>Clinical and Experimental Immunology</i> , 2014, 178, 18-20.	1.1	43
30	Gene Therapy for Chronic Granulomatous Disease: Current Status and Future Perspectives. <i>Current Gene Therapy</i> , 2014, 14, 447-460.	0.9	21
31	Human miR223 Promoter as a Novel Myelo-Specific Promoter for Chronic Granulomatous Disease Gene Therapy. <i>Human Gene Therapy Methods</i> , 2013, 24, 151-159.	2.1	18
32	Autophagy proteins stabilize pathogen-containing phagosomes for prolonged MHC II antigen processing. <i>Journal of Cell Biology</i> , 2013, 203, 757-766.	2.3	172
33	Autophagy proteins stabilize pathogen-containing phagosomes for prolonged MHC II antigen processing. <i>Journal of Experimental Medicine</i> , 2013, 210, 21013OIA64.	4.2	0
34	Immunoglobulin Deficiencies: The B-Lymphocyte Side of DiGeorge Syndrome. <i>Journal of Pediatrics</i> , 2012, 161, 950-953.e1.	0.9	63
35	Increased number of activated T cells in lymphocyte subsets of maltreated children: Data from a pilot study. <i>Journal of Psychosomatic Research</i> , 2012, 73, 313-318.	1.2	12
36	Heterozygous signal transducer and activator of transcription 3 mutations in hyper-IgE syndrome result in altered B-cell maturation. <i>Journal of Allergy and Clinical Immunology</i> , 2012, 129, 559-562.e2.	1.5	41

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37	Clinical, molecular, and cellular immunologic findings in patients with SP110-associated veno-occlusive disease with immunodeficiency syndrome. <i>Journal of Allergy and Clinical Immunology</i> , 2012, 130, 735-742.e6.	1.5	49
38	Derivation and Functional Analysis of Patient-Specific Induced Pluripotent Stem Cells as an In Vitro Model of Chronic Granulomatous Disease. <i>Stem Cells</i> , 2012, 30, 599-611.	1.4	69
39	Restoration of anti-Aspergillus defense by neutrophil extracellular traps in human chronic granulomatous disease after gene therapy is calprotectin-dependent. <i>Journal of Allergy and Clinical Immunology</i> , 2011, 127, 1243-1252.e7.	1.5	221
40	Quantification of $\lambda$ -deleting recombination excision circles in Guthrie cards for the identification of early B-cell maturation defects. <i>Journal of Allergy and Clinical Immunology</i> , 2011, 128, 223-225.e2.	1.5	91
41	NEMO is a key component of NF- $\kappa$ B and IRF-3 dependent TLR3-mediated immunity to herpes simplex virus. <i>Journal of Allergy and Clinical Immunology</i> , 2011, 128, 610-617.e4.	1.5	66
42	Modern management of primary B-cell immunodeficiencies. <i>Pediatric Allergy and Immunology</i> , 2011, 22, 758-769.	1.1	23
43	Gene Therapy of Chronic Granulomatous Disease: The Engraftment Dilemma. <i>Molecular Therapy</i> , 2011, 19, 28-35.	3.7	147
44	Gain-of-function human STAT1 mutations impair IL-17 immunity and underlie chronic mucocutaneous candidiasis. <i>Journal of Experimental Medicine</i> , 2011, 208, 1635-1648.	4.2	739
45	Revisiting Human IL-12R $\beta$ 1 Deficiency. <i>Medicine (United States)</i> , 2010, 89, 381-402.	0.4	367
46	Inflammasome activation in NADPH oxidase defective mononuclear phagocytes from patients with chronic granulomatous disease. <i>Blood</i> , 2010, 116, 1570-1573.	0.6	249
47	Extracerebellar MRI Lesions in Ataxia Telangiectasia Go Along with Deficiency of the GH/IGF-1 Axis, Markedly Reduced Body Weight, High Ataxia Scores and Advanced Age. <i>Cerebellum</i> , 2010, 9, 190-197.	1.4	35
48	Intact indoleamine 2,3-dioxygenase activity in human chronic granulomatous disease. <i>Clinical Immunology</i> , 2010, 137, 1-4.	1.4	19
49	Signed outside: a surface marker system for transgenic cytoplasmic proteins. <i>Gene Therapy</i> , 2010, 17, 1193-1199.	2.3	5
50	The transmembrane activator TACI triggers immunoglobulin class switching by activating B cells through the adaptor MyD88. <i>Nature Immunology</i> , 2010, 11, 836-845.	7.0	295
51	Reply to Agger and Kowalski. <i>Clinical Infectious Diseases</i> , 2010, 50, 1325-1326.	2.9	0
52	Reactive Oxygen Species Abrogate the Anticarcinogenic Effect of Eicosapentaenoic Acid in Atm-Deficient Mice. <i>Nutrition and Cancer</i> , 2010, 62, 584-592.	0.9	4
53	Diagnostic approach to the hyper-IgE syndromes: Immunologic and clinical key findings to differentiate hyper-IgE syndromes from atopic dermatitis. <i>Journal of Allergy and Clinical Immunology</i> , 2010, 126, 611-617.e1.	1.5	140
54	Clinical Features and Outcome of Patients With IRAK-4 and MyD88 Deficiency. <i>Medicine (United States)</i> , 2010, 89, 403-425.	0.4	366

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55	Actinomyces in Chronic Granulomatous Disease: An Emerging and Unanticipated Pathogen. <i>Clinical Infectious Diseases</i> , 2009, 49, 1703-1710.	2.9	74
56	Identification of Severe Combined Immunodeficiency by T-Cell Receptor Excision Circles Quantification Using Neonatal Guthrie Cards. <i>Journal of Pediatrics</i> , 2009, 155, 829-833.	0.9	108
57	S.7. IL-17 Signaling Defects in Patients with <i>Candida Albicans</i> and/or <i>Staphylococcus Aureus</i> Infections. <i>Clinical Immunology</i> , 2009, 131, S135.	1.4	0
58	Restoration of NET formation by gene therapy in CGD controls aspergillosis. <i>Blood</i> , 2009, 114, 2619-2622.	0.6	500
59	Response: Protecting against <i>Aspergillus</i> infection in CGD. <i>Blood</i> , 2009, 114, 3498-3498.	0.6	2
60	Catapult-like release of mitochondrial DNA by eosinophils contributes to antibacterial defense. <i>Nature Medicine</i> , 2008, 14, 949-953.	15.2	836
61	Novel signal transducer and activator of transcription 3 (STAT3) mutations, reduced TH17 cell numbers, and variably defective STAT3 phosphorylation in hyper-IgE syndrome. <i>Journal of Allergy and Clinical Immunology</i> , 2008, 122, 181-187.	1.5	290
62	First Successful Bone Marrow Transplantation for X-linked Chronic Granulomatous Disease by Using Preimplantation Female Gender Typing and HLA Matching. <i>Pediatrics</i> , 2008, 122, e778-e782.	1.0	26
63	Mutations in <i>STAT3</i> and <i>IL12RB1</i> impair the development of human IL-17 <sup>+</sup> producing T cells. <i>Journal of Experimental Medicine</i> , 2008, 205, 1543-1550.	4.2	406
64	Effectiveness of budesonide nebulising suspension compared to disodium cromoglycate in early childhood asthma. <i>Current Medical Research and Opinion</i> , 2006, 22, 367-373.	0.9	7
65	The NEMO Mutation Creating the Most-Upstream Premature Stop Codon Is Hypomorphic Because of a Reinitiation of Translation. <i>American Journal of Human Genetics</i> , 2006, 78, 691-701.	2.6	89
66	Immunoglobulins and Inflammatory Cytokines in Nasal Secretions in Humoral Immunodeficiencies. <i>Laryngoscope</i> , 2006, 116, 239-244.	1.1	7
67	Impaired interferon- $\gamma$ production in response to live bacteria and Toll-like receptor agonists in patients with ataxia telangiectasia. <i>Clinical and Experimental Immunology</i> , 2006, 146, 381-389.	1.1	14
68	Treatment of HCV infection with interferon alpha-2b and ribavirin in a patient with X-linked lymphoproliferative syndrome. <i>European Journal of Pediatrics</i> , 2006, 165, 348-350.	1.3	4
69	Fatal Neonatal-Onset Mitochondrial Respiratory Chain Disease with T Cell Immunodeficiency. <i>Pediatric Research</i> , 2006, 60, 321-326.	1.1	30
70	Autosomal-dominant primary immunodeficiencies. <i>Current Opinion in Hematology</i> , 2005, 12, 22-30.	1.2	20
71	Growth factor deficiency in patients with ataxia telangiectasia. <i>Clinical and Experimental Immunology</i> , 2005, 140, 517-519.	1.1	50
72	From idiopathic infectious diseases to novel primary immunodeficiencies. <i>Journal of Allergy and Clinical Immunology</i> , 2005, 116, 426-430.	1.5	57

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73	Bacillus Calmette GuÃ©rin triggers the IL-12/IFN-Î³ axis by an IRAK-4- and NEMO-dependent, non-cognate interaction between monocytes, NK, and Tâ€š lymphocytes. <i>European Journal of Immunology</i> , 2004, 34, 3276-3284.	1.6	133
74	Nuclear factor Î²B essential modulatorâ€œ deficient child with immunodeficiency yet without anhidrotic ectodermal dysplasia. <i>Journal of Allergy and Clinical Immunology</i> , 2004, 114, 1456-1462.	1.5	122
75	IMMUNOGENICITY OF THE SEVEN VALENT PNEUMOCOCCAL CONJUGATE VACCINE IN PATIENTS WITH ATAXIA-TELANGIECTASIA. <i>Pediatric Infectious Disease Journal</i> , 2004, 23, 269-270.	1.1	29
76	Safety and Tolerability of Methacholine Challenge in Infants with Recurrent Wheeze. <i>Journal of Asthma</i> , 2003, 40, 795-802.	0.9	9
77	A hypermorphic Î²B mutation is associated with autosomal dominant anhidrotic ectodermal dysplasia and T cell immunodeficiency. <i>Journal of Clinical Investigation</i> , 2003, 112, 1108-1115.	3.9	325
78	Elevated Oxidative Stress in Patients with Ataxia Telangiectasia. <i>Antioxidants and Redox Signaling</i> , 2002, 4, 465-469.	2.5	152
79	Serum ECP levels and methacholine challenge in infants with recurrent wheezing. <i>Annals of Allergy, Asthma and Immunology</i> , 2002, 89, 498-502.	0.5	5
80	Deficiencies in CD4 + and CD8 + T cell subsets in ataxia telangiectasia. <i>Clinical and Experimental Immunology</i> , 2002, 129, 125-132.	1.1	54
81	Mycobacterial diseases in primary immunodeficiencies. <i>Current Opinion in Allergy and Clinical Immunology</i> , 2001, 1, 503-511.	1.1	94
82	Impaired Interferon Gamma-Mediated Immunity and Susceptibility to Mycobacterial Infection in Childhood. <i>Pediatric Research</i> , 2001, 50, 8-13.	1.1	97
83	Interleukinâ€œ12 Receptor Î²1 Deficiency in a Patient with Abdominal Tuberculosis. <i>Journal of Infectious Diseases</i> , 2001, 184, 231-236.	1.9	159
84	Spontaneous and oxidative stress-induced programmed cell death in lymphocytes from patients with ataxia telangiectasia (AT). <i>Clinical and Experimental Immunology</i> , 2000, 119, 140-147.	1.1	13
85	Antioxidative capacity in patients with common variable immunodeficiency. <i>Journal of Clinical Immunology</i> , 2000, 20, 221-226.	2.0	5
86	Immunogenicity and Tolerance of a 7-Valent Pneumococcal Conjugate Vaccine in Nonresponders to the 23-Valent Pneumococcal Vaccine. <i>Infection and Immunity</i> , 2000, 68, 1435-1440.	1.0	72
87	Severe adenovirus bronchiolitis in children. <i>Acta Paediatrica, International Journal of Paediatrics</i> , 2000, 89, 1387-9.	0.7	9