

Dominik Hartl

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/2265160/publications.pdf>

Version: 2024-02-01

95
papers

6,710
citations

71061

41
h-index

64755

79
g-index

96
all docs

96
docs citations

96
times ranked

10965
citing authors

#	ARTICLE	IF	CITATIONS
1	Neutrophils: Between Host Defence, Immune Modulation, and Tissue Injury. <i>PLoS Pathogens</i> , 2015, 11, e1004651.	2.1	532
2	Role of breast regression protein 39 (BRP-39)/chitinase 3-like-1 in Th2 and IL-13-induced tissue responses and apoptosis. <i>Journal of Experimental Medicine</i> , 2009, 206, 1149-1166.	4.2	376
3	Inflammation in cystic fibrosis lung disease: Pathogenesis and therapy. <i>Journal of Cystic Fibrosis</i> , 2015, 14, 419-430.	0.3	371
4	Quantitative and functional impairment of pulmonary CD4+CD25hi regulatory T cells in pediatric asthma. <i>Journal of Allergy and Clinical Immunology</i> , 2007, 119, 1258-1266.	1.5	366
5	Chitin regulation of immune responses: an old molecule with new roles. <i>Current Opinion in Immunology</i> , 2008, 20, 684-689.	2.4	315
6	Cleavage of CXCR1 on neutrophils disables bacterial killing in cystic fibrosis lung disease. <i>Nature Medicine</i> , 2007, 13, 1423-1430.	15.2	291
7	Immune Mechanisms in Pulmonary Fibrosis. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2016, 55, 309-322.	1.4	245
8	CFTR: cystic fibrosis and beyond. <i>European Respiratory Journal</i> , 2014, 44, 1042-1054.	3.1	207
9	Infiltrated Neutrophils Acquire Novel Chemokine Receptor Expression and Chemokine Responsiveness in Chronic Inflammatory Lung Diseases. <i>Journal of Immunology</i> , 2008, 181, 8053-8067.	0.4	199
10	Ultrastructural characterization of cystic fibrosis sputum using atomic force and scanning electron microscopy. <i>Journal of Cystic Fibrosis</i> , 2012, 11, 84-92.	0.3	199
11	Pulmonary TH2 response in <i>Pseudomonas aeruginosa</i> -infected patients with cystic fibrosis. <i>Journal of Allergy and Clinical Immunology</i> , 2006, 117, 204-211.	1.5	172
12	Neutrophil extracellular trap-associated RNA and LL37 enable self-amplifying inflammation in psoriasis. <i>Nature Communications</i> , 2020, 11, 105.	5.8	146
13	Extrapulmonary <i>Aspergillus</i> infection in patients with CARD9 deficiency. <i>JCI Insight</i> , 2016, 1, e89890.	2.3	141
14	Flagellin Induces Myeloid-Derived Suppressor Cells: Implications for <i>Pseudomonas aeruginosa</i> Infection in Cystic Fibrosis Lung Disease. <i>Journal of Immunology</i> , 2013, 190, 1276-1284.	0.4	118
15	In vivo genome editing using nuclease-encoding mRNA corrects SP-B deficiency. <i>Nature Biotechnology</i> , 2015, 33, 584-586.	9.4	113
16	Innate Immunity of the Lung: From Basic Mechanisms to Translational Medicine. <i>Journal of Innate Immunity</i> , 2018, 10, 487-501.	1.8	101
17	Free DNA in Cystic Fibrosis Airway Fluids Correlates with Airflow Obstruction. <i>Mediators of Inflammation</i> , 2015, 2015, 1-11.	1.4	100
18	Pathogenic Fungi Regulate Immunity by Inducing Neutrophilic Myeloid-Derived Suppressor Cells. <i>Cell Host and Microbe</i> , 2015, 17, 507-514.	5.1	99

#	ARTICLE	IF	CITATIONS
19	Myeloid-Derived Suppressor Cells in Bacterial Infections. <i>Frontiers in Cellular and Infection Microbiology</i> , 2016, 6, 37.	1.8	99
20	Granulocytic Myeloid-Derived Suppressor Cells Accumulate in Human Placenta and Polarize toward a Th2 Phenotype. <i>Journal of Immunology</i> , 2016, 196, 1132-1145.	0.4	88
21	TLR Expression on Neutrophils at the Pulmonary Site of Infection: TLR1/TLR2-Mediated Up-Regulation of TLR5 Expression in Cystic Fibrosis Lung Disease. <i>Journal of Immunology</i> , 2008, 181, 2753-2763.	0.4	86
22	Chemokines Indicate Allergic Bronchopulmonary Aspergillosis in Patients with Cystic Fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2006, 173, 1370-1376.	2.5	83
23	Expression of checkpoint molecules on myeloid-derived suppressor cells. <i>Immunology Letters</i> , 2017, 192, 1-6.	1.1	82
24	Regulatory T-Cell Impairment in Cystic Fibrosis Patients with Chronic <i>Pseudomonas</i> Infection. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2015, 191, 914-923.	2.5	77
25	Airway Mucus Obstruction Triggers Macrophage Activation and Matrix Metalloproteinase 12-Dependent Emphysema. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2014, 51, 709-720.	1.4	76
26	Myeloid-Derived Suppressor Cells in Infection: A General Overview. <i>Journal of Innate Immunity</i> , 2018, 10, 407-413.	1.8	76
27	The fungal ligand chitin directly binds TLR 2 and triggers inflammation dependent on oligomer size. <i>EMBO Reports</i> , 2018, 19, .	2.0	75
28	Inhalation Treatment with Glutathione in Patients with Cystic Fibrosis. A Randomized Clinical Trial. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2013, 188, 83-89.	2.5	73
29	Pulmonary chemokines and their receptors differentiate children with asthma and chronic cough. <i>Journal of Allergy and Clinical Immunology</i> , 2005, 115, 728-736.	1.5	70
30	Role of Breast Regression Protein 39 in the Pathogenesis of Cigarette Smoke-Induced Inflammation and Emphysema. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2011, 44, 777-786.	1.4	67
31	Neutrophils in cystic fibrosis. <i>Biological Chemistry</i> , 2016, 397, 485-496.	1.2	64
32	Microbial colonization and lung function in adolescents with cystic fibrosis. <i>Journal of Cystic Fibrosis</i> , 2016, 15, 340-349.	0.3	63
33	Myeloid-derived suppressor cells modulate B-cell responses. <i>Immunology Letters</i> , 2017, 188, 108-115.	1.1	59
34	Current concepts: host-pathogen interactions in cystic fibrosis airways disease. <i>European Respiratory Review</i> , 2014, 23, 320-332.	3.0	55
35	Peripheral blood myeloid-derived suppressor cells reflect disease status in idiopathic pulmonary fibrosis. <i>European Respiratory Journal</i> , 2016, 48, 1171-1183.	3.1	55
36	Novel biomarkers in asthma: chemokines and chitinase-like proteins. <i>Current Opinion in Allergy and Clinical Immunology</i> , 2009, 9, 60-66.	1.1	54

#	ARTICLE	IF	CITATIONS
37	The role of chitin, chitinases, and chitinase-like proteins in pediatric lung diseases. <i>Molecular and Cellular Pediatrics</i> , 2015, 2, 3.	1.0	52
38	Prospective multicenter German study on pulmonary colonization with <i>Scenedosporium</i> / <i>Lomentospora</i> species in cystic fibrosis: Epidemiology and new association factors. <i>PLoS ONE</i> , 2017, 12, e0171485.	1.1	47
39	The emerging role of myeloid-derived suppressor cells in lung diseases. <i>European Respiratory Journal</i> , 2016, 47, 967-977.	3.1	46
40	A role for MCP-1/CCR2 in interstitial lung disease in children. <i>Respiratory Research</i> , 2005, 6, 93.	1.4	44
41	The Chitinase-Like Protein YKL-40 Modulates Cystic Fibrosis Lung Disease. <i>PLoS ONE</i> , 2011, 6, e24399.	1.1	44
42	Acidic Mammalian Chitinase Regulates Epithelial Cell Apoptosis via a Chitinolytic-Independent Mechanism. <i>Journal of Immunology</i> , 2009, 182, 5098-5106.	0.4	43
43	Progress in Definition, Prevention and Treatment of Fungal Infections in Cystic Fibrosis. <i>Mycopathologia</i> , 2018, 183, 21-32.	1.3	43
44	Pathogenesis, imaging and clinical characteristics of CF and non-CF bronchiectasis. <i>BMC Pulmonary Medicine</i> , 2018, 18, 79.	0.8	43
45	Differential neutrophil activation in viral infections: Enhanced TLR7/8-mediated CXCL8 release in asthma. <i>Respirology</i> , 2016, 21, 172-179.	1.3	42
46	RNA and Imidazoquinolines Are Sensed by Distinct TLR7/8 Ectodomain Sites Resulting in Functionally Disparate Signaling Events. <i>Journal of Immunology</i> , 2014, 192, 5963-5973.	0.4	38
47	Acidic Mammalian Chitinase Is Secreted via an ADAM17/Epidermal Growth Factor Receptor-dependent Pathway and Stimulates Chemokine Production by Pulmonary Epithelial Cells. <i>Journal of Biological Chemistry</i> , 2008, 283, 33472-33482.	1.6	37
48	Chemokines in Allergic Aspergillosis - From Animal Models to Human Lung Diseases. <i>Inflammation and Allergy: Drug Targets</i> , 2006, 5, 219-228.	1.8	36
49	Mechanisms and disease relevance of neutrophil extracellular trap formation. <i>European Journal of Clinical Investigation</i> , 2018, 48, e12919.	1.7	36
50	Oxidative stress in cystic fibrosis lung disease: an early event, but worth targeting?. <i>European Respiratory Journal</i> , 2014, 44, 17-19.	3.1	35
51	Regulatory Immune Cells in Idiopathic Pulmonary Fibrosis: Friends or Foes?. <i>Frontiers in Immunology</i> , 2021, 12, 663203.	2.2	33
52	A functional inflammasome activation assay differentiates patients with pathogenic NLRP3 mutations and symptomatic patients with low penetrance variants. <i>Clinical Immunology</i> , 2015, 157, 56-64.	1.4	32
53	Visualization and quantification of <i>in vivo</i> homing kinetics of myeloid-derived suppressor cells in primary and metastatic cancer. <i>Theranostics</i> , 2019, 9, 5869-5885.	4.6	31
54	Immunological mechanisms behind the cystic fibrosis-ABPA link. <i>Medical Mycology</i> , 2009, 47, S183-S191.	0.3	28

#	ARTICLE	IF	CITATIONS
55	Chitinase activation in patients with fungus-associated cystic fibrosis lung disease. <i>Journal of Allergy and Clinical Immunology</i> , 2016, 138, 1183-1189.e4.	1.5	28
56	<i>CXCR1</i> and <i>CXCR2</i> haplotypes synergistically modulate cystic fibrosis lung disease. <i>European Respiratory Journal</i> , 2012, 39, 1385-1390.	3.1	27
57	Human monocytic myeloid-derived suppressor cells impair B cell phenotype and function in vitro. <i>European Journal of Immunology</i> , 2020, 50, 33-47.	1.6	26
58	Influenza A(H1N1)pdm09 and Cystic Fibrosis Lung Disease: A Systematic Meta-Analysis. <i>PLoS ONE</i> , 2014, 9, e78583.	1.1	25
59	Airway, but not serum or urinary, levels of YKL-40 reflect inflammation in early cystic fibrosis lung disease. <i>BMC Pulmonary Medicine</i> , 2014, 14, 28.	0.8	25
60	Differential Regulation of Myeloid-Derived Suppressor Cells by <i>Candida</i> Species. <i>Frontiers in Microbiology</i> , 2016, 7, 1624.	1.5	25
61	The Initial Inflammatory Response to Bioactive Implants Is Characterized by NETosis. <i>PLoS ONE</i> , 2015, 10, e0121359.	1.1	25
62	Janus-Faced Neutrophil Extracellular Traps in Periodontitis. <i>Frontiers in Immunology</i> , 2017, 8, 1404.	2.2	24
63	Surfactant proteins in pediatric interstitial lung disease. <i>Pediatric Research</i> , 2016, 79, 34-41.	1.1	23
64	Transcriptomic profile of cystic fibrosis patients identifies type I interferon response and ribosomal stalk proteins as potential modifiers of disease severity. <i>PLoS ONE</i> , 2017, 12, e0183526.	1.1	23
65	<i>Pseudomonas aeruginosa</i> Airway Infection Recruits and Modulates Neutrophilic Myeloid-Derived Suppressor Cells. <i>Frontiers in Cellular and Infection Microbiology</i> , 2016, 6, 167.	1.8	22
66	mRNA-Mediated Gene Supplementation of Toll-Like Receptors as Treatment Strategy for Asthma In Vivo. <i>PLoS ONE</i> , 2016, 11, e0154001.	1.1	20
67	Immune Response, Diagnosis and Treatment of Allergic Bronchopulmonary Aspergillosis in Cystic Fibrosis Lung Disease. <i>Current Pharmaceutical Design</i> , 2013, 19, 3669-3678.	0.9	20
68	In Vivo Hypoxia PET Imaging Quantifies the Severity of Arthritic Joint Inflammation in Line with Overexpression of Hypoxia-Inducible Factor and Enhanced Reactive Oxygen Species Generation. <i>Journal of Nuclear Medicine</i> , 2017, 58, 853-860.	2.8	19
69	Recurrent pericarditis in children: elevated cardiac autoantibodies. <i>Clinical Research in Cardiology</i> , 2007, 96, 168-175.	1.5	18
70	Characterization of rapid neutrophil extracellular trap formation and its cooperation with phagocytosis in human neutrophils. <i>Discoveries</i> , 2014, 2, e19.	1.5	18
71	Staphylococcal Enterotoxins Dose-Dependently Modulate the Generation of Myeloid-Derived Suppressor Cells. <i>Frontiers in Cellular and Infection Microbiology</i> , 2018, 8, 321.	1.8	17
72	The chemokine CCL18 characterises <i>Pseudomonas</i> infections in cystic fibrosis lung disease. <i>European Respiratory Journal</i> , 2014, 44, 1608-1615.	3.1	16

#	ARTICLE	IF	CITATIONS
73	Increased CCL17 serum levels are associated with improved survival in advanced melanoma. <i>Cancer Immunology, Immunotherapy</i> , 2015, 64, 1075-1082.	2.0	16
74	A semiquantitative MRI-Score can predict loss of lung function in patients with cystic fibrosis: Preliminary results. <i>European Radiology</i> , 2018, 28, 74-84.	2.3	16
75	Allergic bronchopulmonary aspergillosis: the hunt for a diagnostic serological marker in cystic fibrosis patients. <i>Expert Review of Molecular Diagnostics</i> , 2009, 9, 157-164.	1.5	14
76	Induction of Myeloid-Derived Suppressor Cells in Cryopyrin-Associated Periodic Syndromes. <i>Journal of Innate Immunity</i> , 2016, 8, 493-506.	1.8	14
77	Myeloid-Derived Suppressor Cells Dampen Airway Inflammation Through Prostaglandin E2 Receptor 4. <i>Frontiers in Immunology</i> , 2021, 12, 695933.	2.2	13
78	Expression and Regulation of Interferon-Related Development Regulator 1 in Cystic Fibrosis Neutrophils. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2013, 48, 71-77.	1.4	12
79	Update on host-pathogen interactions in cystic fibrosis lung disease. <i>Molecular and Cellular Pediatrics</i> , 2016, 3, 12.	1.0	12
80	Cystic fibrosis in Europe: patients live longer but are we ready?. <i>European Respiratory Journal</i> , 2015, 46, 11-12.	3.1	11
81	CXCR4 ⁺ granulocytes reflect fungal cystic fibrosis lung disease. <i>European Respiratory Journal</i> , 2015, 46, 395-404.	3.1	10
82	CHI3L1 polymorphisms, cord blood YKL-40 levels and later asthma development. <i>BMC Pulmonary Medicine</i> , 2016, 16, 81.	0.8	10
83	Suspicion of respiratory tract infection with multidrug-resistant Enterobacteriaceae: epidemiology and risk factors from a Paediatric Intensive Care Unit. <i>BMC Infectious Diseases</i> , 2017, 17, 163.	1.3	10
84	Anti-inflammatory role of CD11b ⁺ Ly6G ⁺ neutrophilic cells in allergic airway inflammation in mice. <i>Immunology Letters</i> , 2018, 204, 67-74.	1.1	10
85	Fungi in Cystic Fibrosis: Recent Findings and Unresolved Questions. <i>Current Fungal Infection Reports</i> , 2015, 9, 1-5.	0.9	8
86	An informative intragenic microsatellite marker suggests the IL-1 receptor as a genetic modifier in cystic fibrosis. <i>European Respiratory Journal</i> , 2017, 50, 1700426.	3.1	8
87	Human T cells modulate myeloid-derived suppressor cells through a TNF- α -mediated mechanism. <i>Immunology Letters</i> , 2018, 202, 31-37.	1.1	8
88	Temporal Dynamics of Reactive Oxygen and Nitrogen Species and NF- κ B Activation During Acute and Chronic T Cell-Driven Inflammation. <i>Molecular Imaging and Biology</i> , 2020, 22, 504-514.	1.3	8
89	Is osseointegration inflammation-triggered?. <i>Medical Hypotheses</i> , 2016, 93, 1-4.	0.8	7
90	Cystic fibrosis – From basic science to clinical benefit: A review series. <i>Journal of Cystic Fibrosis</i> , 2015, 14, 415-416.	0.3	5

#	ARTICLE	IF	CITATIONS
91	Enhanced IgG 1 mediated antibody response towards thymus dependent immunization in CXCR1 deficient mice. <i>Immunity, Inflammation and Disease</i> , 2021, 9, 210-222.	1.3	4
92	Arginase 1 ⁺ IL10 ⁺ polymorphonuclear myeloid derived suppressor cells are elevated in patients with active pemphigus and correlate with an increased Th2/Th1 response. <i>Experimental Dermatology</i> , 2021, 30, 782-791.	1.4	4
93	Developmental control of CFTR: from bioinformatics to novel therapeutic approaches. <i>European Respiratory Journal</i> , 2015, 45, 18-20.	3.1	3
94	Macrophages and platelets join forces to release kidney-damaging DNA traps. <i>Nature Medicine</i> , 2018, 24, 128-129.	15.2	2
95	JCF 2014 and beyond. <i>Journal of Cystic Fibrosis</i> , 2014, 13, 610-611.	0.3	0