Carsten Schwarz

List of Publications by Year
in descending order

44	1,005	567281 15	454955 30
papers	citations	h-index	g-index

47	47	47	1782
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Clinician variability in the diagnosis and treatment of aspergillus fumigatus-related conditions in cystic fibrosis: An international survey. Journal of Cystic Fibrosis, 2022, 21, 136-142.	0.7	10
2	Tezacaftor/ivacaftor in people with cystic fibrosis who stopped lumacaftor/ivacaftor due to respiratory adverse events. Journal of Cystic Fibrosis, 2021, 20, 228-233.	0.7	21
3	Genetic diversification of persistent Mycobacterium abscessus within cystic fibrosis patients. Virulence, 2021, 12, 2415-2429.	4.4	14
4	Biomarkers for the Diagnosis of Allergic Bronchopulmonary Aspergillosis in Cystic Fibrosis: A Systematic Review and Meta-Analysis. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 1909-1930.e4.	3.8	15
5	Clinical impact of levofloxacin inhalation solution in cystic fibrosis patients in a real-world setting. Journal of Cystic Fibrosis, 2021, 20, 1035-1039.	0.7	7
6	CFTR Modulator Therapy and Its Impact on Lung Transplantation in Cystic Fibrosis. Pulmonary Therapy, 2021, 7, 377-393.	2.2	11
7	Deposition of Inhaled Levofloxacin in Cystic Fibrosis Lungs Assessed by Functional Respiratory Imaging. Pharmaceutics, 2021, 13, 2051.	4.5	3
8	Antisense oligonucleotide eluforsen is safe and improves respiratory symptoms in F508DEL cystic fibrosis. Journal of Cystic Fibrosis, 2020, 19, 99-107.	0.7	46
9	GLPG2737 in lumacaftor/ivacaftor-treated CF subjects homozygous for the F508del mutation: A randomized phase 2A trial (PELICAN). Journal of Cystic Fibrosis, 2020, 19, 292-298.	0.7	11
10	Urban Life as Risk Factor for Aspergillosis. Frontiers in Cellular and Infection Microbiology, 2020, 10, 601834.	3.9	6
11	Risk factors for respiratory Aspergillus fumigatus in German Cystic Fibrosis patients and impact on lung function. Scientific Reports, 2020, 10, 18999.	3.3	30
12	Mutation on lysX from Mycobacterium avium hominissuis impacts the hostâ€“pathogen interaction and virulence phenotype. Virulence, 2020, 11, 132-144.	4.4	11
13	Prospective Evaluation of Aspergillus fumigatus-Specific IgG in Patients With Cystic Fibrosis. Frontiers in Cellular and Infection Microbiology, 2020, 10, 602836.	3.9	8
14	Frequent Pet Contact as Risk Factor for Allergic Bronchopulmonary Aspergillosis in Cystic Fibrosis. Frontiers in Cellular and Infection Microbiology, 2020, 10, 601821.	3.9	5
15	Biatrial Remodeling in Patients with Cystic Fibrosis. Journal of Clinical Medicine, 2019, 8, 1141.	2.4	3
16	Clinical manifestations and risk factors of arthropathy in cystic fibrosis. Respiratory Medicine, 2019, 147, 66-71.	2.9	11
17	Type I interferon induced by TLR2-TLR4-MyD88-TRIF-IRF3 controls Mycobacterium abscessus subsp. abscessus persistence in murine macrophages via nitric oxide. International Journal of Medical Microbiology, 2019, 309, 307-318.	3.6	16
18	Long needed guidelines: Referral to transplant for cystic fibrosis patients. Journal of Cystic Fibrosis, 2019, 18, 305-306.	0.7	0

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19	Human Anti-fungal Th17 Immunity and Pathology Rely on Cross-Reactivity against Candida albicans. Cell, 2019, 176, 1340-1355.e15.	28.9	321
20	Risk of piperacillinâ€induced hemolytic anemia in patients with cystic fibrosis and antipseudomonal treatment: a prospective observational study. Transfusion, 2019, 59, 3746-3754.	1.6	9
21	Combined antifungal therapy is superior to monotherapy in pulmonary scedosporiosis in cystic fibrosis. Journal of Cystic Fibrosis, 2019, 18, 227-232.	0.7	26
22	Environmental Mycobacterium avium subsp. hominissuis have a higher probability to act as a recipient in conjugation than clinical strains. Plasmid, 2018, 95, 28-35.	1.4	7
23	Fungal Respiratory Infections in Cystic Fibrosis (CF): Recent Progress and Future Research Agenda. Mycopathologia, 2018, 183, 1-5.	3.1	9
24	Fungus-Specific CD4 T Cells as Specific Sensors for Identification of Pulmonary Fungal Infections. Mycopathologia, 2018, 183, 213-226.	3.1	17
25	Developing collaborative works for faster progress on fungal respiratory infections in cystic fibrosis. Medical Mycology, 2018, 56, S42-S59.	0.7	27
26	Organization of Patient Management and Fungal Epidemiology in Cystic Fibrosis. Mycopathologia, 2018, 183, 7-19.	3.1	32
27	Mepolizumabâ€"a novel option for the treatment of hypereosinophilic syndrome in childhood. Pediatric Allergy and Immunology, 2018, 29, 28-33.	2.6	13
28	Invasive Pulmonary Fungal Infections in Cystic Fibrosis. Mycopathologia, 2018, 183, 33-43.	3.1	27
29	Aspergillus Bronchitis in Patients with Cystic Fibrosis. Mycopathologia, 2018, 183, 61-69.	3.1	65
30	Progress in Definition, Prevention and Treatment of Fungal Infections in Cystic Fibrosis. Mycopathologia, 2018, 183, 21-32.	3.1	43
31	Prospective multicenter German study on pulmonary colonization with Scedosporium /Lomentospora species in cystic fibrosis: Epidemiology and new association factors. PLoS ONE, 2017, 12, e0171485.	2.5	47
32	Allergic bronchopulmonary aspergillosis is associated with pet ownership in cystic fibrosis. Pediatric Allergy and Immunology, 2016, 27, 597-603.	2.6	15
33	Piperacillin-induced mild haemolytic anaemia in a 44-year-old patient with cystic fibrosis. BMJ Case Reports, 2016, 2016, bcr2016216937.	0.5	8
34	Impact of Long-Term Tiotropium Bromide Therapy on Annual Lung Function Decline in Adult Patients with Cystic Fibrosis. PLoS ONE, 2016, 11, e0158193.	2.5	2
35	Cystic fibrosis in Europe: patients live longer but are we ready?. European Respiratory Journal, 2015, 46, 11-12.	6.7	11
36	ColobreatheÂ® for the Treatment of Cystic Fibrosis-Associated Pulmonary Infections. Pulmonary Therapy, 2015, 1, 19-30.	2.2	13

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37	Different approaches to evaluate patient experience and satisfaction in CF centres. Journal of Cystic Fibrosis, 2015, 14, E19-E20.	0.7	0
38	Patient experience in cystic fibrosis care: Development of a disease-specific questionnaire. Chronic Illness, 2015, 11, 108-125.	1.5	10
39	Arxula adeninivorans causing invasive pulmonary mycosis and fungaemia in cystic fibrosis. Lancet, The, 2015, 385, 1476.	13.7	10
40	Cystic fibrosis (CF) care through the patients' eyes â€“ A nationwide survey on experience and satisfaction with services using a disease-specific questionnaire. Respiratory Medicine, 2015, 109, 79-87.	2.9	7
41	Scedosporium apiospermum: a fungal pathogen causing pneumonia in a patient with cystic fibrosis. JMM Case Reports, 2015, 2, .	1.3	10
42	Hypersensitivity to antibiotics in patients with cystic fibrosis. Journal of Cystic Fibrosis, 2014, 13, 205-211.	0.7	45
43	Hypersensitivity to parenteral antibiotics in patients with cystic fibrosis. Clinical and Translational Allergy, 2014, 4, P66.	3.2	0
44	Zystische Fibrose: Mukoviszidose ist lÃngst keine Kinderkrankheit mehr. , 0, , .		0