

Caroline Kannengiesser

List of Publications by Year in descending order

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Version: 2024-02-01

33
papers

1,943
citations

361413

20
h-index

454955

30
g-index

34
all docs

34
docs citations

34
times ranked

2024
citing authors

#	ARTICLE	IF	CITATIONS
1	<i>MUC5B</i> Promoter Variant and Rheumatoid Arthritis with Interstitial Lung Disease. <i>New England Journal of Medicine</i> , 2018, 379, 2209-2219.	27.0	326
2	Shared genetic predisposition in rheumatoid arthritis-interstitial lung disease and familial pulmonary fibrosis. <i>European Respiratory Journal</i> , 2017, 49, 1602314.	6.7	154
3	The <i>MUC5B</i> Variant Is Associated with Idiopathic Pulmonary Fibrosis but Not with Systemic Sclerosis Interstitial Lung Disease in the European Caucasian Population. <i>PLoS ONE</i> , 2013, 8, e70621.	2.5	142
4	Prevalence and characteristics of <i>TERT</i> and <i>TERC</i> mutations in suspected genetic pulmonary fibrosis. <i>European Respiratory Journal</i> , 2016, 48, 1721-1731.	6.7	136
5	Heterozygous <i>RTEL1</i> mutations are associated with familial pulmonary fibrosis. <i>European Respiratory Journal</i> , 2015, 46, 474-485.	6.7	135
6	Germline <i>SFTPA1</i> mutation in familial idiopathic interstitial pneumonia and lung cancer. <i>Human Molecular Genetics</i> , 2016, 25, 1457-1467.	2.9	119
7	Methotrexate and rheumatoid arthritis associated interstitial lung disease. <i>European Respiratory Journal</i> , 2021, 57, 2000337.	6.7	114
8	Severe Pulmonary Fibrosis as the First Manifestation of Interferonopathy (TMEM173 Mutation). <i>Chest</i> , 2016, 150, e65-e71.	0.8	112
9	Severe hematologic complications after lung transplantation in patients with telomerase complex mutations. <i>Journal of Heart and Lung Transplantation</i> , 2015, 34, 538-546.	0.6	109
10	Resequencing Study Confirms That Host Defense and Cell Senescence Gene Variants Contribute to the Risk of Idiopathic Pulmonary Fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2019, 200, 199-208.	5.6	90
11	Is Telomeropathy the Explanation for Combined Pulmonary Fibrosis and Emphysema Syndrome?: Report of a Family with <i>TERT</i> Mutation. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2014, 189, 753-754.	5.6	57
12	Management of suspected monogenic lung fibrosis in a specialised centre. <i>European Respiratory Review</i> , 2017, 26, 160122.	7.1	54
13	Regulator of telomere length 1 (<i>RTEL1</i>) mutations are associated with heterogeneous pulmonary and extra-pulmonary phenotypes. <i>European Respiratory Journal</i> , 2019, 53, 1800508.	6.7	45
14	The genetics of interstitial lung diseases. <i>European Respiratory Review</i> , 2019, 28, 190053.	7.1	41
15	Genetic landscape of adult Langerhans cell histiocytosis with lung involvement. <i>European Respiratory Journal</i> , 2020, 55, 1901190.	6.7	38
16	Safety and efficacy of pirfenidone and nintedanib in patients with idiopathic pulmonary fibrosis and carrying a telomere-related gene mutation. <i>European Respiratory Journal</i> , 2021, 57, 2003198.	6.7	36
17	Safety and efficacy of pirfenidone in patients carrying telomerase complex mutation. <i>European Respiratory Journal</i> , 2018, 51, 1701875.	6.7	34
18	Pleuroparenchymal fibroelastosis associated with telomerase reverse transcriptase mutations. <i>European Respiratory Journal</i> , 2017, 49, 1602022.	6.7	32

#	ARTICLE	IF	CITATIONS
19	Pilot experience of multidisciplinary team discussion dedicated to inherited pulmonary fibrosis. Orphanet Journal of Rare Diseases, 2019, 14, 280.	2.7	24
20	Functional assessment and phenotypic heterogeneity of <i>SFTPA1</i> and <i>SFTPA2</i> mutations in interstitial lung diseases and lung cancer. European Respiratory Journal, 2020, 56, 2002806.	6.7	23
21	The <i>MUC5B</i> promoter risk allele for idiopathic pulmonary fibrosis predisposes to asbestosis. European Respiratory Journal, 2020, 55, 1902361.	6.7	23
22	Bi-allelic missense <i>ABCA3</i> mutations in a patient with childhood ILD who reached adulthood. ERJ Open Research, 2019, 5, 00066-2019.	2.6	22
23	First heterozygous <i>NOP10</i> mutation in familial pulmonary fibrosis. European Respiratory Journal, 2020, 55, 1902465.	6.7	13
24	Pneumocystosis revealing immunodeficiency secondary to <i>TERC</i> mutation. European Respiratory Journal, 2017, 50, 1701443.	6.7	12
25	Determinants of survival after lung transplantation in telomerase-related gene mutation carriers: A retrospective cohort. American Journal of Transplantation, 2022, 22, 1236-1244.	4.7	11
26	Genetic testing in interstitial lung disease: An international survey. Respirology, 0, , .	2.3	10
27	Impact of genetic factors on fibrosing interstitial lung diseases. Incidence and clinical presentation in adults. Presse Medicale, 2020, 49, 104024.	1.9	9
28	Pulmonary fibrosis: Genetic analysis of telomere-related genes, telomere length measurement or both?. Respirology, 2019, 24, 97-98.	2.3	8
29	The Genetic Diagnosis of Interstitial Lung Disease: A Need for an International Consensus. American Journal of Respiratory and Critical Care Medicine, 2017, 195, 1538-1539.	5.6	7
30	Interstitial lung diseases associated with mutations of poly(A)-specific ribonuclease: A multicentre retrospective study. Respirology, 2022, 27, 226-235.	2.3	6
31	Aspergillus-induced pneumonia in adult without obvious immunodeficiency: test the burst!. European Respiratory Journal, 2018, 51, 1702711.	6.7	1
32	The Genetics of Interstitial Lung Diseases. , 2022, , 96-113.		0
33	Telomere syndrome and the lung. , 2019, , 391-403.		0