Caroline Kannengiesser

List of Publications by Year
in descending order

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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	The Genetics of Interstitial Lung Diseases. , 2022, , 96-113.		0
2	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
3	Interstitial lung diseases associated with mutations of poly(A)â€specific ribonuclease: A multicentre retrospective study. Respirology, 2022, 27, 226-235.	2.3	6
4	Methotrexate and rheumatoid arthritis associated interstitial lung disease. European Respiratory Journal, 2021, 57, 2000337.	6.7	114
5	Safety and efficacy of pirfenidone and nintedanib in patients with idiopathic pulmonary fibrosis and carrying a telomere-related gene mutation. European Respiratory Journal, 2021, 57, 2003198.	6.7	36
6	Genetic landscape of adult Langerhans cell histiocytosis with lung involvement. European Respiratory Journal, 2020, 55, 1901190.	6.7	38
7	Functional assessment and phenotypic heterogeneity of SFTPA1 and SFTPA2 mutations in interstitial lung diseases and lung cancer. European Respiratory Journal, 2020, 56, 2002806.	6.7	23
8	The MUC5B promoter risk allele for idiopathic pulmonary fibrosis predisposes to asbestosis. European Respiratory Journal, 2020, 55, 1902361.	6.7	23
9	First heterozygous NOP10 mutation in familial pulmonary fibrosis. European Respiratory Journal, 2020, 55, 1902465.	6.7	13
10	Impact of genetic factors on fibrosing interstitial lung diseases. Incidence and clinical presentation in adults. Presse Medicale, 2020, 49, 104024.	1.9	9
11	Resequencing Study Confirms That Host Defense and Cell Senescence Gene Variants Contribute to the Risk of Idiopathic Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2019, 200, 199-208.	5.6	90
12	Bi-allelic missense ABCA3 mutations in a patient with childhood ILD who reached adulthood. ERJ Open Research, 2019, 5, 00066-2019.	2.6	22
13	The genetics of interstitial lung diseases. European Respiratory Review, 2019, 28, 190053.	7.1	41
14	Pilot experience of multidisciplinary team discussion dedicated to inherited pulmonary fibrosis. Orphanet Journal of Rare Diseases, 2019, 14, 280.	2.7	24
15	Pulmonary fibrosis: Genetic analysis of telomereâ€related genes, telomere length measurementâ€or both?. Respirology, 2019, 24, 97-98.	2.3	8
16	Regulator of telomere length 1 (RTEL1) mutations are associated with heterogeneous pulmonary and extra-pulmonary phenotypes. European Respiratory Journal, 2019, 53, 1800508.	6.7	45
17	Telomere syndrome and the lung. , 2019, , 391-403.		0
18	Aspergillus-induced pneumonia in adult without obvious immunodeficiency: test the burst!. European Respiratory Journal, 2018, 51, 1702711.	6.7	1

#	Article	IF	Citations
19	Safety and efficacy of pirfenidone in patients carrying telomerase complex mutation. European Respiratory Journal, 2018, 51, 1701875.	6.7	34
20	MUC5B Promoter Variant and Rheumatoid Arthritis with Interstitial Lung Disease. New England Journal of Medicine, 2018, 379, 2209-2219.	27.0	326
21	Shared genetic predisposition in rheumatoid arthritis-interstitial lung disease and familial pulmonary fibrosis. European Respiratory Journal, 2017, 49, 1602314.	6.7	154
22	Pleuroparenchymal fibroelastosis associated with telomerase reverse transcriptase mutations. European Respiratory Journal, 2017, 49, 1602022.	6.7	32
23	Management of suspected monogenic lung fibrosis in a specialised centre. European Respiratory Review, 2017, 26, 160122.	7.1	54
24	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
25	Pneumocystosis revealing immunodeficiency secondary to TERC mutation. European Respiratory Journal, 2017, 50, 1701443.	6.7	12
26	Prevalence and characteristics ofTERTandTERCmutations in suspected genetic pulmonary fibrosis. European Respiratory Journal, 2016, 48, 1721-1731.	6.7	136
27	Severe Pulmonary Fibrosis as the First Manifestation of Interferonopathy (TMEM173 Mutation). Chest, 2016, 150, e65-e71.	0.8	112
28	GermlineSFTPA1mutation in familial idiopathic interstitial pneumonia and lung cancer. Human Molecular Genetics, 2016, 25, 1457-1467.	2.9	119
29	Severe hematologic complications after lung transplantation in patients with telomerase complex mutations. Journal of Heart and Lung Transplantation, 2015, 34, 538-546.	0.6	109
30	HeterozygousRTEL1mutations are associated with familial pulmonary fibrosis. European Respiratory Journal, 2015, 46, 474-485.	6.7	135
31	Is Telomeropathy the Explanation for Combined Pulmonary Fibrosis and Emphysema Syndrome?: Report of a Family with TERT Mutation. American Journal of Respiratory and Critical Care Medicine, 2014, 189, 753-754.	5.6	57
32	The MUC5B Variant Is Associated with Idiopathic Pulmonary Fibrosis but Not with Systemic Sclerosis Interstitial Lung Disease in the European Caucasian Population. PLoS ONE, 2013, 8, e70621.	2.5	142
33	Genetic testing in interstitial lung disease: An international survey. Respirology, 0, , .	2.3	10