Manuela Funke-Chambour

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Pulmonary function and radiological features 4 months after COVID-19: first results from the national prospective observational Swiss COVID-19 lung study. European Respiratory Journal, 2021, 57, 2003690.	3.1	291
2	Medium throughput breathing human primary cell alveolus-on-chip model. Scientific Reports, 2018, 8, 14359.	1.6	132
3	Acute exacerbation of idiopathic pulmonary fibrosis: international survey and call for harmonisation. European Respiratory Journal, 2020, 55, 1901760.	3.1	61
4	Serum metabolic profiling identified a distinct metabolic signature in patients with idiopathic pulmonary fibrosis – a potential biomarker role for LysoPC. Respiratory Research, 2018, 19, 7.	1.4	58
5	Swiss Recommendations for the Follow-Up and Treatment of Pulmonary Long COVID. Respiration, 2021, 100, 826-841.	1.2	41
6	Surfactant replacement therapy reduces acute lung injury and collapse induration-related lung remodeling in the bleomycin model. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2017, 313, L313-L327.	1.3	39
7	Human microvasculature-on-aÂchip: anti-neovasculogenic effect of nintedanib in vitro. Angiogenesis, 2018, 21, 861-871.	3.7	36
8	Human-Based Advanced in vitro Approaches to Investigate Lung Fibrosis and Pulmonary Effects of COVID-19. Frontiers in Medicine, 2021, 8, 644678.	1.2	31
9	Idiopathic Pulmonary Fibrosis in Switzerland: Diagnosis and Treatment. Respiration, 2017, 93, 363-378.	1.2	26
10	Azithromycin has enhanced effects on lung fibroblasts from idiopathic pulmonary fibrosis (IPF) patients compared to controls. Respiratory Research, 2020, 21, 25.	1.4	26
11	Lysophosphatidic Acid Signaling through the Lysophosphatidic Acid-1 Receptor Is Required for Alveolarization. American Journal of Respiratory Cell and Molecular Biology, 2016, 55, 105-116.	1.4	24
12	Incidence and Prognostic Significance of Hypoxemia in Fibrotic Interstitial Lung Disease. Chest, 2021, 160, 994-1005.	0.4	20
13	Azithromycin for the Treatment of Chronic Cough in Idiopathic Pulmonary Fibrosis: A Randomized Controlled Crossover Trial. Annals of the American Thoracic Society, 2021, 18, 2018-2026.	1.5	19
14	Idiopathic pulmonary fibrosis: the turning point is now!. Swiss Medical Weekly, 2015, 145, w14139.	0.8	18
15	Imaging in the aftermath of COVID-19: what to expect. European Radiology, 2021, 31, 4390-4392.	2.3	17
16	Multiple breath washout: A new and promising lung function test for patients with idiopathic pulmonary fibrosis. Respirology, 2018, 23, 764-770.	1.3	14
17	Novel insights in cough and breathing patterns of patients with idiopathic pulmonary fibrosis performing repeated 24-hour-respiratory polygraphies. Respiratory Research, 2017, 18, 190.	1.4	13
18	Serum calprotectin as new biomarker for disease severity in idiopathic pulmonary fibrosis: a cross-sectional study in two independent cohorts. BMI Open Respiratory Research, 2021, 8, e000827	1.2	13

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19	Frailty assessment for COVID-19 follow-up: a prospective cohort study. BMJ Open Respiratory Research, 2022, 9, e001227.	1.2	12
20	MRI Shows Lung Perfusion Changes after Vaping and Smoking. Radiology, 2022, 304, 195-204.	3.6	9
21	Management of Acute Exacerbation of Idiopathic Pulmonary Fibrosis in Specialised and Non-specialised ILD Centres Around the World. Frontiers in Medicine, 2021, 8, 699644.	1.2	8
22	Idiopathic pulmonary fibrosis in a Swiss interstitial lung disease reference centre. Swiss Medical Weekly, 2018, 148, w14577.	0.8	8
23	Fatigue in Post-COVID-19 Syndrome: Clinical Phenomenology, Comorbidities and Association With Initial Course of COVID-19. Journal of Central Nervous System Disease, 2022, 14, 117957352211027.	0.7	6
24	Perioperative Lung Function Monitoring for Anatomic Lung Resections. Annals of Thoracic Surgery, 2017, 104, 1725-1732.	0.7	5
25	Multidisciplinary discussion for diagnosis of interstitial lung disease in real life. Swiss Medical Weekly, 2016, 146, w14318.	0.8	5
26	New radiological diagnostic criteria: impact on idiopathic pulmonary fibrosis diagnosis. European Respiratory Journal, 2019, 54, 1900905.	3.1	4
27	Performance of a diagnostic algorithm for fibrotic hypersensitivity pneumonitis. A case–control study. Respiratory Research, 2021, 22, 120.	1.4	4
28	Pulmonary hypertension associated with chronic lung diseases. Swiss Medical Weekly, 2016, 146, w14363.	0.8	4
29	Evaluation of a Novel Ear Pulse Oximeter: Towards Automated Oxygen Titration in Eyeglass Frames. Sensors, 2020, 20, 3301.	2.1	2
30	Suggestions for improving clinical utility of future guidelines for diagnosis and management of idiopathic pulmonary fibrosis: results of a Delphi survey. European Respiratory Journal, 2021, 57, 2004219.	3.1	2
31	Registries for Idiopathic Pulmonary Fibrosis: When Is It Time to Go Global?. Annals of the American Thoracic Society, 2020, 17, 1378-1379.	1.5	2
32	The Octopus Sign—A New HRCT Sign in Pulmonary Langerhans Cell Histiocytosis. Diagnostics, 2022, 12, 937.	1.3	2
33	Influence of ventilation inhomogeneity on diffusing capacity of carbon monoxide in smokers without COPD. ERJ Open Research, 2021, 7, 00706-2020.	1.1	1
34	P054 <break></break> Exhaled breath condensate- a potential biomarker tool for patients with idiopathic pulmonary fibrosis?. QJM - Monthly Journal of the Association of Physicians, 2016, , .	0.2	0
35	Antifibrotics: Shrinking the Box of Therapeutic Uncertainty. Respiration, 2019, 97, 202-204.	1.2	0