

# Xinlun Tian

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/9244011/publications.pdf>

Version: 2024-02-01

63  
papers

1,137  
citations

394421

19  
h-index

454955

30  
g-index

72  
all docs

72  
docs citations

72  
times ranked

3673  
citing authors

#	ARTICLE	IF	CITATIONS
1	Inhibition of Autophagy Ameliorates Acute Lung Injury Caused by Avian Influenza A H5N1 Infection. <i>Science Signaling</i> , 2012, 5, ra16.	3.6	140
2	mTOR Overactivation and Compromised Autophagy in the Pathogenesis of Pulmonary Fibrosis. <i>PLoS ONE</i> , 2015, 10, e0138625.	2.5	77
3	Lung function and systemic inflammation associated with short-term air pollution exposure in chronic obstructive pulmonary disease patients in Beijing, China. <i>Environmental Health</i> , 2020, 19, 12.	4.0	58
4	The role of vascular endothelial growth factor-D in diagnosis of lymphangi leiomyomatosis (LAM). <i>Respiratory Medicine</i> , 2013, 107, 263-268.	2.9	54
5	A paper-based skin patch for the diagnostic screening of cystic fibrosis. <i>Chemical Communications</i> , 2015, 51, 6365-6368.	4.1	45
6	Lymphocytic Interstitial Pneumonia and Other Benign Lymphoid Disorders. <i>Seminars in Respiratory and Critical Care Medicine</i> , 2012, 33, 450-461.	2.1	43
7	&lt;p&gt;Short-term effects of ambient air pollution on chronic obstructive pulmonary disease admissions in Beijing, China (2013&ndash;2017)&lt;/p&gt;. <i>International Journal of COPD</i> , 2019, Volume 14, 297-309.	2.3	40
8	Aetiology of severe community acquired pneumonia in adults identified by combined detection methods: a multi-centre prospective study in China. <i>Emerging Microbes and Infections</i> , 2022, 11, 556-566.	6.5	40
9	Characterization of gene mutations and phenotypes of cystic fibrosis in <sc>C</sc>hinese patients. <i>Respirology</i> , 2015, 20, 312-318.	2.3	34
10	p.G970D is the most frequent CFTR mutation in Chinese patients with cystic fibrosis. <i>Human Genome Variation</i> , 2016, 3, 15063.	0.7	34
11	Diffuse cystic lung diseases. <i>Frontiers of Medicine</i> , 2013, 7, 316-327.	3.4	33
12	Pneumocystis Pneumonia in Patients with Autoimmune Diseases: A Retrospective Study Focused on Clinical Characteristics and Prognostic Factors Related to Death. <i>PLoS ONE</i> , 2015, 10, e0139144.	2.5	32
13	Clinical and genetic characteristics of cystic fibrosis in CHINESE patients: a systemic review of reported cases. <i>Orphanet Journal of Rare Diseases</i> , 2018, 13, 224.	2.7	32
14	Clinical and genetic characteristics of chinese patients with Birt-Hogg-DubÃ© syndrome. <i>Orphanet Journal of Rare Diseases</i> , 2017, 12, 104.	2.7	29
15	Long-term efficacy and safety of sirolimus therapy in patients with lymphangi leiomyomatosis. <i>Orphanet Journal of Rare Diseases</i> , 2019, 14, 206.	2.7	28
16	Lymphangi leiomyomatosis. <i>Seminars in Respiratory and Critical Care Medicine</i> , 2020, 41, 256-268.	2.1	25
17	Polymorphisms of &lt;i&gt;PHF11&lt;/i&gt; and &lt;i&gt;DPP10 &lt;/i&gt; Are Associated with Asthma and Related Traits in a Chinese Population. <i>Respiration</i> , 2010, 79, 17-24.	2.6	24
18	Tsc1 deficiency-mediated mTOR hyperactivation in vascular endothelial cells causes angiogenesis defects and embryonic lethality. <i>Human Molecular Genetics</i> , 2014, 23, 693-705.	2.9	24

#	ARTICLE	IF	CITATIONS
19	Diffuse Cystic Lung Diseases: Diagnostic Considerations. <i>Seminars in Respiratory and Critical Care Medicine</i> , 2016, 37, 457-467.	2.1	24
20	Rapamycin for lymphangioleiomyomatosis: optimal timing and optimal dosage. <i>Thorax</i> , 2018, 73, 308-310.	5.6	16
21	Functional improvements in patients with lymphangioleiomyomatosis after sirolimus: an observational study. <i>Orphanet Journal of Rare Diseases</i> , 2018, 13, 34.	2.7	16
22	SPC-Cre-ERT2 Transgenic Mouse for Temporal Gene Deletion in Alveolar Epithelial Cells. <i>PLoS ONE</i> , 2012, 7, e46076.	2.5	15
23	Secondary pulmonary alveolar proteinosis: a single-center retrospective study (a case series and) <i>Tj ETQq1 1 0.784314 rgBT /Overlock 15</i>	2.0	15
24	Pregnancy after the diagnosis of lymphangioleiomyomatosis (LAM). <i>Orphanet Journal of Rare Diseases</i> , 2021, 16, 133.	2.7	14
25	Genotypic characteristics of Chinese patients with BHD syndrome and functional analysis of FLCN variants. <i>Orphanet Journal of Rare Diseases</i> , 2019, 14, 223.	2.7	13
26	Recurrent pulmonary mucormycosis after lobectomy in a non-smoking patient without predisposing risk factors. <i>Brazilian Journal of Infectious Diseases</i> , 2012, 16, 590-593.	0.6	11
27	Clinical characteristics in lymphangioleiomyomatosis-related pulmonary hypertension: an observation on 50 patients. <i>Frontiers of Medicine</i> , 2019, 13, 259-266.	3.4	11
28	Inhaled granulocyte-macrophage colony stimulating factor for mild-to-moderate autoimmune pulmonary alveolar proteinosis - a six month phase II randomized study with 24 months of follow-up. <i>Orphanet Journal of Rare Diseases</i> , 2020, 15, 174.	2.7	11
29	Clinical characteristics and genetic spectrum of 26 individuals of Chinese origin with primary ciliary dyskinesia. <i>Orphanet Journal of Rare Diseases</i> , 2021, 16, 293.	2.7	11
30	Clinicopathological findings of focal organizing pneumonia: a retrospective study of 37 cases. <i>International Journal of Clinical and Experimental Pathology</i> , 2015, 8, 511-6.	0.5	11
31	A novel homozygous complex deletion in CFTR caused cystic fibrosis in a Chinese patient. <i>Molecular Genetics and Genomics</i> , 2017, 292, 1083-1089.	2.1	10
32	Lymphangioleiomyomatosis Association with Underlying Genotype in Patients with Tuberous Sclerosis Complex. <i>Annals of the American Thoracic Society</i> , 2021, 18, 815-819.	3.2	10
33	Quantitative assessment of Pulmonary Alveolar Proteinosis (PAP) with ultra-dose CT and correlation with Pulmonary Function Tests (PFTs). <i>PLoS ONE</i> , 2017, 12, e0172958.	2.5	10
34	Characterization of clinical and genetic spectrum of Chinese patients with cystic fibrosis. <i>Orphanet Journal of Rare Diseases</i> , 2020, 15, 150.	2.7	9
35	Impaired lipid metabolism in idiopathic pulmonary alveolar proteinosis. <i>Lipids in Health and Disease</i> , 2011, 10, 54.	3.0	8
36	Global lung function initiative 2012 reference values for spirometry in Asian Americans. <i>BMC Pulmonary Medicine</i> , 2018, 18, 95.	2.0	8

#	ARTICLE	IF	CITATIONS
37	The efficacy and adverse events of mTOR inhibitors in lymphangioleiomyomatosis: systematic review and meta-analysis. <i>Orphanet Journal of Rare Diseases</i> , 2018, 13, 134.	2.7	8
38	Characterization of CT scans of patients with Birt-Hogg-Dub� syndrome compared with those of Chinese patients with non-BHD diffuse cyst lung diseases. <i>Orphanet Journal of Rare Diseases</i> , 2020, 15, 176.	2.7	8
39	Recent advances in the management of lymphangioleiomyomatosis. <i>F1000Research</i> , 2018, 7, 758.	1.6	8
40	Thymic neoplasms patients complicated with bronchiectasis: Case series in a Chinese hospital and literature review. <i>Thoracic Cancer</i> , 2019, 10, 791-798.	1.9	7
41	The etiology of diffuse cystic lung diseases: an analysis of 1010 consecutive cases in a LAM clinic. <i>Orphanet Journal of Rare Diseases</i> , 2021, 16, 273.	2.7	7
42	Dynamic Observation of Autophagy and Transcriptome Profiles in a Mouse Model of Bleomycin-Induced Pulmonary Fibrosis. <i>Frontiers in Molecular Biosciences</i> , 2021, 8, 664913.	3.5	7
43	Exacerbation of Pneumomediastinum After Air Travel in a Patient with Dermatomyositis. <i>Aviation, Space, and Environmental Medicine</i> , 2011, 82, 734-736.	0.5	6
44	Fatal myositis and spontaneous haematoma induced by combined immune checkpoint inhibitor treatment in a patient with pancreatic adenocarcinoma. <i>BMC Cancer</i> , 2019, 19, 1193.	2.6	6
45	Cystic fibrosis: a rare disease emerging in China. <i>Science China Life Sciences</i> , 2020, 63, 1082-1084.	4.9	6
46	Pulmonary manifestations of Erdheim�Chester disease: clinical characteristics, outcomes and comparison with Langerhans cell histiocytosis. <i>British Journal of Haematology</i> , 2021, 194, 1024-1033.	2.5	6
47	Pulmonary hypertension associated with combined fibrosing mediastinitis and bronchial anthracofibrosis: A retrospective analysis in a single Chinese hospital. <i>Clinical Respiratory Journal</i> , 2018, 12, 1134-1140.	1.6	5
48	Pulmonary high-resolution computed tomography findings in patients with synovitis, acne, pustulosis, hyperostosis and osteitis syndrome. <i>PLoS ONE</i> , 2018, 13, e0206858.	2.5	5
49	Treatment of steroid�resistant checkpoint inhibitor pneumonitis with pirfenidone: A case report. <i>Thoracic Cancer</i> , 2021, 12, 2214-2216.	1.9	5
50	Clinical and Genetic Comparison of Birt�Hogg�Dub� Syndrome (Hornstein�Knickenberg Syndrome) in Chinese: A Systemic Review of Reported Cases. <i>International Journal of General Medicine</i> , 0, Volume 15, 5111-5121.	1.8	5
51	Sirolimus reduces the risk of pneumothorax recurrence in patients with lymphangioleiomyomatosis: a historical prospective self-controlled study. <i>Orphanet Journal of Rare Diseases</i> , 2022, 17, .	2.7	5
52	Airway-invasion-associated pulmonary computed tomography presentations characteristic of invasive pulmonary Aspergillosis in non-immunocompromised adults: a National Multicenter Retrospective Survey in China. <i>Respiratory Research</i> , 2020, 21, 173.	3.6	4
53	Humoral response to inactivated SARS-CoV-2 vaccines in patients on sirolimus alone. <i>Science China Life Sciences</i> , 2022, 65, 2118-2120.	4.9	4
54	The value of transbronchial lung biopsy in the diagnosis of lymphangioleiomyomatosis. <i>BMC Pulmonary Medicine</i> , 2021, 21, 146.	2.0	3

#	ARTICLE	IF	CITATIONS
55	Novel mutation c.1210-3C > G in cis with a poly-T tract of 5T affects CFTR mRNA splicing in a Chinese patient with cystic fibrosis. <i>Frontiers of Medicine</i> , 2022, 16, 150-155.	3.4	3
56	Expiratory Lymphatic Sacs. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2011, 184, 1085-1085.	5.6	2
57	Cushing's disease with pulmonary <i>Cryptococcus neoformans</i> infection in a single center in Beijing, China: A retrospective study and literature review. <i>Journal of the Formosan Medical Association</i> , 2019, 118, 285-290.	1.7	2
58	Case report of neurofibromatosis type 1 combined with primary ciliary dyskinesia. <i>Frontiers of Medicine</i> , 2021, , 1.	3.4	2
59	Tunnel Sign. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2019, 199, 795-796.	5.6	1
60	Expression profiles and potential functions of long noncoding RNAs and mRNAs in autoimmune pulmonary alveolar proteinosis patients. <i>Aging</i> , 2021, 13, 10535-10554.	3.1	1
61	Ga <sup>68</sup> EDTA aerosols in evaluation of inhaled <sup>68</sup> particle deposition and clearance of obstructive pulmonary diseases: A pilot prospective study compared with Galligas. <i>European Journal of Clinical Investigation</i> , 2021, 51, e13620.	3.4	1
62	Lung function and air pollution exposure in adults with asthma in Beijing: a 2-year longitudinal panel study. <i>Frontiers of Medicine</i> , 2022, , 1.	3.4	1
63	Vasculitis Secondary to Pulmonary Bacterial Infection: A Case Report. <i>Diagnostics</i> , 2022, 12, 772.	2.6	0