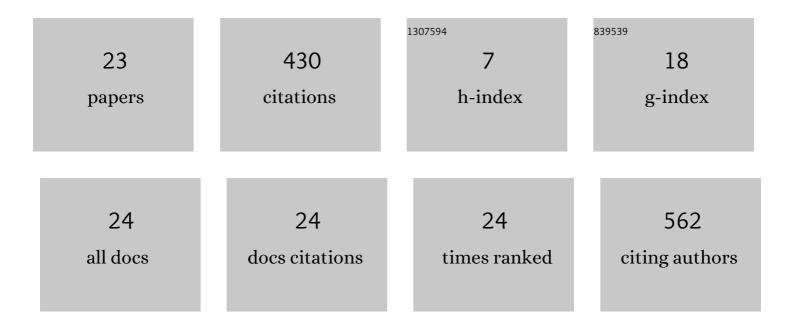
Cheng Lei

List of Publications by Year in descending order

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CHENCLE

#	Article	IF	CITATIONS
1	Nuclear functions of mammalian MicroRNAs in gene regulation, immunity and cancer. Molecular Cancer, 2018, 17, 64.	19.2	257
2	Blood and Bronchoalveolar Lavage Fluid Metagenomic Next-Generation Sequencing in Pneumonia. Canadian Journal of Infectious Diseases and Medical Microbiology, 2020, 2020, 1-9.	1.9	45
3	Bi-allelic BRWD1 variants cause male infertility with asthenoteratozoospermia and likely primary ciliary dyskinesia. Human Genetics, 2021, 140, 761-773.	3.8	23
4	DRC1 deficiency caused primary ciliary dyskinesia and MMAF in a Chinese patient. Journal of Human Genetics, 2022, 67, 197-201.	2.3	16
5	PGS@B–N: an efficient flame retardant to improve simultaneously the interfacial interaction and the flame retardancy of EVA. RSC Advances, 2016, 6, 65921-65929.	3.6	14
6	Construction and validation of a bronchoalveolar lavage cell-associated gene signature for prognosis prediction in idiopathic pulmonary fibrosis. International Immunopharmacology, 2021, 92, 107369.	3.8	9
7	Identification of key modules and hub genes associated with lung function in idiopathic pulmonary fibrosis. PeerJ, 2020, 8, e9848.	2.0	8
8	Case Report: Community-Acquired Legionella gormanii Pneumonia in an Immunocompetent Patient Detected by Metagenomic Next-Generation Sequencing. Frontiers in Medicine, 2022, 9, 819425.	2.6	8
9	Wholeâ€exome sequencing identified a novel homozygous <i>ASPH</i> frameshift variant causing Traboulsi syndrome in a Chinese family. Molecular Genetics & Genomic Medicine, 2021, 9, e1553.	1.2	7
10	Novel Compound Heterozygous Variants in CCDC40 Associated with Primary Ciliary Dyskinesia and Multiple Morphological Abnormalities of the Sperm Flagella. Pharmacogenomics and Personalized Medicine, 2022, Volume 15, 341-350.	0.7	7
11	Mutant CARD10 in a family with progressive immunodeficiency and autoimmunity. Cellular and Molecular Immunology, 2020, 17, 782-784.	10.5	6
12	Case Report: Identification of a Novel ODAD3 Variant in a Patient With Primary Ciliary Dyskinesia. Frontiers in Genetics, 2021, 12, 652381.	2.3	6
13	Identification of Two Novel DNAAF2 Variants in Two Consanguineous Families with Primary Ciliary Dyskinesia. Pharmacogenomics and Personalized Medicine, 2021, Volume 14, 1415-1423.	0.7	6
14	Association between PTCH1 gene polymorphisms and chronic obstructive pulmonary disease susceptibility in a Chinese Han population: a case-control study. Chinese Medical Journal, 2020, 133, 2071-2077.	2.3	4
15	Preparation and Catalytic Properties of Carbon Carrier-Supported Ruthenium Catalysts for Acetalization/Ketalization Reactions. ChemistrySelect, 2017, 2, 9377-9386.	1.5	2
16	A novel switch beam design method with extending switching radio-frequency bandwidth. Microsystem Technologies, 2021, 27, 315-324.	2.0	2
17	The contribution of <i>Pseudomonas aeruginosa</i> infection to clinical outcomes in bronchiectasis: a prospective cohort study. Annals of Medicine, 2021, 53, 459-469.	3.8	2
18	Whole-Exome Sequencing Identified CFTR Variants in Two Consanguineous Families in China. Frontiers in Genetics, 2021, 12, 631221.	2.3	2

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#	Article	IF	CITATIONS
19	Novel RSPH4A Variants Associated With Primary Ciliary Dyskinesia–Related Infertility in Three Chinese Families. Frontiers in Genetics, 0, 13, .	2.3	2
20	Identification of a Novel OFD1 Variant in a Patient with Primary Ciliary Dyskinesia. Pharmacogenomics and Personalized Medicine, 0, Volume 15, 697-704.	0.7	2
21	Case Report: DNAAF4 Variants Cause Primary Ciliary Dyskinesia and Infertility in Two Han Chinese Families. Frontiers in Genetics, 0, 13, .	2.3	2
22	Research on 355 nm all-solid-state ultraviolet laser processing through silicon holes. Journal of Laser Applications, 2019, 31, 022003.	1.7	0
23	Clinical phenotypes of primary ciliary dyskinesia Journal of Central South University (Medical) Tj ETQq1 1 0.784	314 rgBT /	Overlock 10