

# Deepika Polineni

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/1701649/publications.pdf>

Version: 2024-02-01

17  
papers

562  
citations

1163117

8  
h-index

1199594

12  
g-index

18  
all docs

18  
docs citations

18  
times ranked

717  
citing authors

#	ARTICLE	IF	CITATIONS
1	A pilot study of cystic fibrosis exacerbation response phenotypes reveals contrasting serum and sputum iron trends. <i>Scientific Reports</i> , 2021, 11, 4897.	3.3	3
2	Cystic fibrosis. <i>Lancet</i> , The, 2021, 397, 2195-2211.	13.7	316
3	Losartan Rescues Inflammation-related Mucociliary Dysfunction in Relevant Models of Cystic Fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2020, 201, 313-324.	5.6	34
4	Gender Equity. <i>Journal of Cystic Fibrosis</i> , 2020, 19, 328-329.	0.7	0
5	Mining GWAS and eQTL data for CF lung disease modifiers by gene expression imputation. <i>PLoS ONE</i> , 2020, 15, e0239189.	2.5	9
6	Mining GWAS and eQTL data for CF lung disease modifiers by gene expression imputation. , 2020, 15, e0239189.		0
7	Mining GWAS and eQTL data for CF lung disease modifiers by gene expression imputation. , 2020, 15, e0239189.		0
8	Mining GWAS and eQTL data for CF lung disease modifiers by gene expression imputation. , 2020, 15, e0239189.		0
9	Mining GWAS and eQTL data for CF lung disease modifiers by gene expression imputation. , 2020, 15, e0239189.		0
10	Aztreonam Lysine Inhalation Solution in Cystic Fibrosis. <i>Clinical Medicine Insights: Circulatory, Respiratory and Pulmonary Medicine</i> , 2019, 13, 117954841984282.	0.9	10
11	Perspectives on anemia and iron deficiency from the cystic fibrosis care community. <i>Pediatric Pulmonology</i> , 2019, 54, 939-940.	2.0	6
12	Unraveling the CFTR Functionâ€™Phenotype Connection for Precision Treatment in Cystic Fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2019, 199, 1053-1054.	5.6	3
13	Genetic association and transcriptome integration identify contributing genes and tissues at cystic fibrosis modifier loci. <i>PLoS Genetics</i> , 2019, 15, e1008007.	3.5	56
14	Airway Mucosal Host Defense Is Key to Genomic Regulation of Cystic Fibrosis Lung Disease Severity. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2018, 197, 79-93.	5.6	46
15	Analysis of a large cohort of cystic fibrosis patients with severe liver disease indicates lung function decline does not significantly differ from that of the general cystic fibrosis population. <i>PLoS ONE</i> , 2018, 13, e0205257.	2.5	16
16	Accuracy of Nasal Nitric Oxide Measurement as a Diagnostic Test for Primary Ciliary Dyskinesia: A Systematic Review and Meta-Analysis. <i>Annals of the American Thoracic Society</i> , 2017, 14, 1184-1196.	3.2	41
17	Effect of extracorporeal photopheresis on lung function decline for severe bronchiolitis obliterans syndrome following allogeneic stem cell transplantation. <i>Journal of Clinical Apheresis</i> , 2016, 31, 347-352.	1.3	22