Harriet Corvol

List of Publications by Year
in descending order

144 papers	4,837 citations	81839 39 h-index	123376 61 g-index
166 all docs	166 docs citations	166 times ranked	6328 citing authors

#	Article	IF	Citations
1	Genome-wide association meta-analysis identifies five modifier loci of lung disease severity in cystic fibrosis. Nature Communications, 2015, 6, 8382.	5.8	242
2	Alveolar epithelial cells: Master regulators of lung homeostasis. International Journal of Biochemistry and Cell Biology, 2013, 45, 2568-2573.	1.2	187
3	Multiple apical plasma membrane constituents are associated with susceptibility to meconium ileus in individuals with cystic fibrosis. Nature Genetics, 2012, 44, 562-569.	9.4	177
4	Rapid Improvement after Starting Elexacaftorâ€“Tezacaftorâ€“Ivacaftor in Patients with Cystic Fibrosis and Advanced Pulmonary Disease. American Journal of Respiratory and Critical Care Medicine, 2021, 204, 64-73.	2.5	139
5	Influence of Interleukinâ€10 onAspergillus fumigatusInfection in Patients with Cystic Fibrosis. Journal of Infectious Diseases, 2005, 191, 1988-1991.	1.9	115
6	A multinational report to characterise SARS-CoV-2 infection in people with cystic fibrosis. Journal of Cystic Fibrosis, 2020, 19, 355-358.	0.3	113
7	Interstitial lung diseases in children. Orphanet Journal of Rare Diseases, 2010, 5, 22.	1.2	112
8	Distinct cytokine production by lung and blood neutrophils from children with cystic fibrosis. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2003, 284, L997-L1003.	1.3	108
9	The very low penetrance of cystic fibrosis for the R117H mutation: a reappraisal for genetic counselling and newborn screening. Journal of Medical Genetics, 2009, 46, 752-758.	1.5	106
10	New insights into pediatric idiopathic pulmonary hemosiderosis: the French RespiRareÂ® cohort. Orphanet Journal of Rare Diseases, 2013, 8, 161.	1.2	95
11	Real-Life Safety and Effectiveness of Lumacaftorâ€“Ivacaftor in Patients with Cystic Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2020, 201, 188-197.	2.5	95
12	Features of Severe Liver Disease With Portal Hypertension inÂPatients With Cystic Fibrosis. Clinical Gastroenterology and Hepatology, 2016, 14, 1207-1215.e3.	2.4	94
13	Cystic Fibrosis Liver Disease: Outcomes and Risk Factors in a Large Cohort of French Patients. Hepatology, 2019, 69, 1648-1656.	3.6	93
14	RSPH3 Mutations Cause Primary Ciliary Dyskinesia with Central-Complex Defects and a Near Absence of Radial Spokes. American Journal of Human Genetics, 2015, 97, 153-162.	2.6	88
15	Cystic fibrosis gene modifierSLC26A9modulates airway response to CFTR-directed therapeutics. Human Molecular Genetics, 2016, 25, ddw290.	1.4	81
16	Multiple-Locus Variable-Number Tandem-Repeat Analysis for Longitudinal Survey of Sources of Pseudomonas aeruginosa Infection in Cystic Fibrosis Patients. Journal of Clinical Microbiology, 2007, 45, 3175-3183.	1.8	79
17	A national internet-linked based database for pediatric interstitial lung diseases: the French network. Orphanet Journal of Rare Diseases, 2012, 7, 40.	1.2	79
18	Atypical presentation of COVID-19 in young infants. Lancet, The, 2020, 395, 1481.	6.3	78

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19	The global impact of SARS-CoV-2 in 181 people with cystic fibrosis. Journal of Cystic Fibrosis, 2020, 19, 868-871.	0.3	74
20	Neutrophils in Cystic Fibrosis Display a Distinct Gene Expression Pattern. Molecular Medicine, 2008, 14, 36-44.	1.9	69
21	Lung disease modifier genes in cystic fibrosis. International Journal of Biochemistry and Cell Biology, 2014, 52, 83-93.	1.2	66
22	Sources of Variation in Sweat Chloride Measurements in Cystic Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2016, 194, 1375-1382.	2.5	62
23	Clinical characteristics of SARS-CoV-2 infection in children with cystic fibrosis: An international observational study. Journal of Cystic Fibrosis, 2021, 20, 25-30.	0.3	62
24	MicroRNA-9 downregulates the ANO1 chloride channel and contributes to cystic fibrosis lung pathology. Nature Communications, 2017, 8, 710.	5.8	56
25	Genetic association and transcriptome integration identify contributing genes and tissues at cystic fibrosis modifier loci. PLoS Genetics, 2019, 15, e1008007.	1.5	56
26	Dramatic improvement after tocilizumab of severe <scp>COVID</scp>â€19 in a child with sickle cell disease and acute chest syndrome. American Journal of Hematology, 2020, 95, E192-E194.	2.0	56
27	Translating the genetics of cystic fibrosis to personalized medicine. Translational Research, 2016, 168, 40-49.	2.2	54
28	HRCT and MRI of the lung in children with cystic fibrosis: Comparison of different scoring systems. Journal of Cystic Fibrosis, 2014, 13, 198-204.	0.3	51
29	Adherence of airway neutrophils and inflammatory response are increased in CF airway epithelial cell-neutrophil interactions. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2006, 290, L588-L596.	1.3	50
30	Lung alveolar epithelium and interstitial lung disease. International Journal of Biochemistry and Cell Biology, 2009, 41, 1643-1651.	1.2	50
31	Surfactant protein A: A key player in lung homeostasis. International Journal of Biochemistry and Cell Biology, 2016, 81, 151-155.	1.2	50
32	Lung sarcoidosis in children: update on disease expression and management. Thorax, 2015, 70, 537-542.	2.7	49
33	A Joint Location-Scale Test Improves Power to Detect Associated SNPs, Gene Sets, and Pathways. American Journal of Human Genetics, 2015, 97, 125-138.	2.6	48
34	Circulating and airway neutrophils in cystic fibrosis display different TLR expression and responsiveness to interleukin-10. Cytokine, 2008, 41, 54-60.	1.4	47
35	Airway Mucosal Host Defense Is Key to Genomic Regulation of Cystic Fibrosis Lung Disease Severity. American Journal of Respiratory and Critical Care Medicine, 2018, 197, 79-93.	2.5	46
36	Variants in Solute Carrier SLC26A9 Modify Prenatal Exocrine Pancreatic Damage in Cystic Fibrosis. Journal of Pediatrics, 2015, 166, 1152-1157.e6.	0.9	45

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37	FAM13A is a modifier gene of cystic fibrosis lung phenotype regulating rhoa activity, actin cytoskeleton dynamics and epithelial-mesenchymal transition. Journal of Cystic Fibrosis, 2018, 17, 190-203.	0.3	45
38	Azithromycin analogue <scp>CSY</scp>0073 attenuates lung inflammation induced by <scp>LPS</scp> challenge. British Journal of Pharmacology, 2014, 171, 1783-1794.	2.7	44
39	Genetic ancestry modifies pharmacogenetic geneâ€“gene interaction for asthma. Pharmacogenetics and Genomics, 2009, 19, 489-496.	0.7	42
40	Long-term effects of azithromycin in patients with cystic fibrosis. Respiratory Medicine, 2016, 117, 1-6.	1.3	42
41	Genetic variations in inflammatory mediators influence lung disease progression in cystic fibrosis. Pediatric Pulmonology, 2008, 43, 1224-1232.	1.0	41
42	Anoctamin 1 dysregulation alters bronchial epithelial repair in cystic fibrosis. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2013, 1832, 2340-2351.	1.8	40
43	Phenotype of children with inconclusive cystic fibrosis diagnosis after newborn screening. Pediatric Pulmonology, 2020, 55, 918-928.	1.0	39
44	New Insights about miRNAs in Cystic Fibrosis. American Journal of Pathology, 2015, 185, 897-908.	1.9	37
45	Stenotrophomonas maltophilia: A marker of lung disease severity. Pediatric Pulmonology, 2018, 53, 426-430.	1.0	37
46	AGER -429T/C Is Associated with an Increased Lung Disease Severity in Cystic Fibrosis. PLoS ONE, 2012, 7, e41913.	1.1	36
47	Human Bronchial Epithelial Cells Inhibit Aspergillus fumigatus Germination of Extracellular Conidia via FleA Recognition. Scientific Reports, 2018, 8, 15699.	1.6	35
48	Small RNA and transcriptome sequencing reveal the role of miRâ€199aâ€3p in inflammatory processes in cystic fibrosis airways. Journal of Pathology, 2018, 245, 410-420.	2.1	35
49	Genetic Modifiers of Cystic Fibrosis-Related Diabetes Have Extensive Overlap With Type 2 Diabetes and Related Traits. Journal of Clinical Endocrinology and Metabolism, 2020, 105, 1401-1415.	1.8	34
50	Incidence of SARS-CoV-2 in people with cystic fibrosis in Europe between February and June 2020. Journal of Cystic Fibrosis, 2021, 20, 566-577.	0.3	34
51	First Wave of COVID-19 in French Patients with Cystic Fibrosis. Journal of Clinical Medicine, 2020, 9, 3624.	1.0	33
52	Pulmonary hemosiderosis in children with Down syndrome: a national experience. Orphanet Journal of Rare Diseases, 2018, 13, 60.	1.2	32
53	Modifier genes in cystic fibrosis-related liver disease. Current Opinion in Gastroenterology, 2019, 35, 88-92.	1.0	32
54	The role of LTA4H and ALOX5AP genes in the risk for asthma in Latinos. Clinical and Experimental Allergy, 2010, 40, 582-589.	1.4	31

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55	CFTR p.Arg117His associated with CBAVD and other CFTR-related disorders. Journal of Medical Genetics, 2013, 50, 220-227.	1.5	31
56	Moving beyond genetics: isFAM13Aa major biological contributor in lung physiology and chronic lung diseases?. Journal of Medical Genetics, 2014, 51, 646-649.	1.5	31
57	Bronchial Epithelial Cells from Cystic Fibrosis Patients Express a Specific Long Non-coding RNA Signature upon Pseudomonas aeruginosa Infection. Frontiers in Cellular and Infection Microbiology, 2017, 7, 218.	1.8	31
58	Novel Anti-Inflammatory Approaches for Cystic Fibrosis Lung Disease: Identification of Molecular Targets and Design of Innovative Therapies. Frontiers in Pharmacology, 2020, 11, 1096.	1.6	30
59	DNA methylation at modifier genes of lung disease severity is altered in cystic fibrosis. Clinical Epigenetics, 2017, 9, 19.	1.8	29
60	Emerging microRNA Therapeutic Approaches for Cystic Fibrosis. Frontiers in Pharmacology, 2018, 9, 1113.	1.6	29
61	SLC26A9 Gene Is Associated With Lung Function Response to Ivacaftor in Patients With Cystic Fibrosis. Frontiers in Pharmacology, 2018, 9, 828.	1.6	29
62	Glucocorticoid receptor gene polymorphisms associated with progression of lung disease in young patients with cystic fibrosis. Respiratory Research, 2007, 8, 88.	1.4	28
63	Omalizumab treatment for allergic bronchopulmonary aspergillosis in young patients with cystic fibrosis. Respiratory Medicine, 2017, 133, 12-15.	1.3	28
64	The Wide Spectrum of COVID-19 Clinical Presentation in Children. Journal of Clinical Medicine, 2020, 9, 2950.	1.0	28
65	Whole exome sequencing in three families segregating a pediatric case of sarcoidosis. BMC Medical Genomics, 2018, 11, 23.	0.7	26
66	Interstitial lung disease: Physiopathology in the context of lung growth. Paediatric Respiratory Reviews, 2011, 12, 216-222.	1.2	25
67	Glucocorticoids reduce inflammation in cystic fibrosis bronchial epithelial cells. Cellular Signalling, 2012, 24, 1093-1099.	1.7	25
68	CHAC1 Is Differentially Expressed in Normal and Cystic Fibrosis Bronchial Epithelial Cells and Regulates the Inflammatory Response Induced by Pseudomonas aeruginosa. Frontiers in Immunology, 2018, 9, 2823.	2.2	25
69	SERPINA1 Z allele is associated with cystic fibrosis liver disease. Genetics in Medicine, 2019, 21, 2151-2155.	1.1	25
70	Infant bronchiolitis dramatically reduced during the second French COVIDâ€19 outbreak. Acta Paediatrica, International Journal of Paediatrics, 2021, 110, 1297-1299.	0.7	25
71	Genetic variation in CFTR and modifier loci may modulate cystic fibrosis disease severity. Journal of Cystic Fibrosis, 2020, 19, S10-S14.	0.3	24
72	Clinical response to lumacaftor-ivacaftor in patients with cystic fibrosis according to baseline lung function. Journal of Cystic Fibrosis, 2021, 20, 220-227.	0.3	24

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73	Ancestral haplotype 8.1 and lung disease severity in European cystic fibrosis patients. Journal of Cystic Fibrosis, 2012, 11, 63-67.	0.3	22
74	Normal and Cystic Fibrosis Human Bronchial Epithelial Cells Infected with Pseudomonas aeruginosa Exhibit Distinct Gene Activation Patterns. PLoS ONE, 2015, 10, e0140979.	1.1	22
75	Pharmacogenetic response to albuterol among asthmatics. Pharmacogenomics, 2008, 9, 505-510.	0.6	21
76	Macrolides: New therapeutic perspectives in lung diseases. International Journal of Biochemistry and Cell Biology, 2011, 43, 1241-1246.	1.2	21
77	Reference percentiles for FEV1 and BMI in European children and adults with cystic fibrosis. Orphanet Journal of Rare Diseases, 2012, 7, 64.	1.2	21
78	Lessons from a French collaborative caseâ€“control study in cystic fibrosis patients during the 2009 A/H1N1 influenza pandemy. BMC Infectious Diseases, 2015, 16, 55.	1.3	21
79	Biomarkers in Interstitial lung diseases. Paediatric Respiratory Reviews, 2015, 16, 219-224.	1.2	20
80	Longitudinal survey of Staphylococcus aureus in cystic fibrosis patients using a multiple-locus variable-number of tandem-repeats analysis method. BMC Microbiology, 2010, 10, 24.	1.3	19
81	ALOX5AP and LTA4H polymorphisms modify augmentation of bronchodilator responsiveness by leukotriene modifiers in Latinos. Journal of Allergy and Clinical Immunology, 2010, 126, 853-858.	1.5	19
82	Bronchial Epithelial Cells on the Front Line to Fight Lung Infection-Causing Aspergillus fumigatus. Frontiers in Immunology, 2020, 11, 1041.	2.2	19
83	Factors for severe outcomes following SARS-CoV-2 infection in people with cystic fibrosis in Europe. ERJ Open Research, 2021, 7, 00411-2021.	1.1	19
84	Update on SLC6A14 in lung and gastrointestinal physiology and physiopathology: focus on cystic fibrosis. Cellular and Molecular Life Sciences, 2020, 77, 3311-3323.	2.4	18
85	Respiratory Epithelial Cells Can Remember Infection: A Proof of Concept Study. Journal of Infectious Diseases, 2019, 221, 1000-1005.	1.9	17
86	Cystic fibrosisâ€“related diabetes onset can be predicted using biomarkers measured at birth. Genetics in Medicine, 2021, 23, 927-933.	1.1	17
87	Nocturnal hypoventilation in Down syndrome children with or without sleep apnea. Pediatric Pulmonology, 2020, 55, 1246-1253.	1.0	16
88	AGTR2 absence or antagonism prevents cystic fibrosis pulmonary manifestations. Journal of Cystic Fibrosis, 2019, 18, 127-134.	0.3	15
89	Achromobacter xylosoxidans airway infection is associated with lung disease severity in children with cystic fibrosis. ERJ Open Research, 2021, 7, 00076-2021.	1.1	15
90	Factors associated with clinical progression to severe COVID-19 in people with cystic fibrosis: A global observational study. Journal of Cystic Fibrosis, 2022, 21, e221-e231.	0.3	15

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91	Mild cystic fibrosis revealed by persistent hyponatremia during the French 2003 heat wave, associated with the S1455X C-terminus CFTR mutation. Clinical Genetics, 2005, 68, 552-553.	1.0	14
92	SARS-CoV-2 B.1.1.529 (Omicron) Variant Causes an Unprecedented Surge in Children Hospitalizations and Distinct Clinical Presentation Compared to the SARS-CoV-2 B.1.617.2 (Delta) Variant. Frontiers in Pediatrics, 0, 10, .	0.9	14
93	Vaccine coverage in CF children: A French multicenter study. Journal of Cystic Fibrosis, 2015, 14, 615-620.	0.3	12
94	miR-636: A Newly-Identified Actor for the Regulation of Pulmonary Inflammation in Cystic Fibrosis. Frontiers in Immunology, 2019, 10, 2643.	2.2	11
95	Variant classifications, databases and genotype-phenotype correlations. Archives De Pediatrie, 2020, 27, eS13-eS18.	0.4	10
96	Factors associated with humoral immune response to pandemic A/H1N1(v) 2009 influenza vaccine in cystic fibrosis. Vaccine, 2014, 32, 4515-4521.	1.7	9
97	Flagellin concentrations in expectorations from cystic fibrosis patients. BMC Pulmonary Medicine, 2014, 14, 100.	0.8	9
98	Novel variation at chr11p13 associated with cystic fibrosis lung disease severity. Human Genome Variation, 2016, 3, 16020.	0.4	9
99	Improving imputation in disease-relevant regions: lessons from cystic fibrosis. Npj Genomic Medicine, 2018, 3, 8.	1.7	9
100	Mining GWAS and eQTL data for CF lung disease modifiers by gene expression imputation. PLoS ONE, 2020, 15, e0239189.	1.1	9
101	Flagellin From Pseudomonas aeruginosa Modulates SARS-CoV-2 Infectivity in Cystic Fibrosis Airway Epithelial Cells by Increasing TMPRSS2 Expression. Frontiers in Immunology, 2021, 12, 714027.	2.2	9
102	Cumulative Incidence and Risk Factors for Severe Coronavirus Disease 2019 in French People With Cystic Fibrosis. Clinical Infectious Diseases, 2022, 75, 2135-2144.	2.9	9
103	Risk factors for Pseudomonas aeruginosa airway infection and lung function decline in children with cystic fibrosis. Journal of Cystic Fibrosis, 2022, 21, 45-51.	0.3	8
104	Sweat test practice in pediatric pulmonology after introduction of cystic fibrosis newborn screening. European Journal of Pediatrics, 2015, 174, 1613-1620.	1.3	6
105	Two-hybrid screening of FAM13A protein partners in lung epithelial cells. BMC Research Notes, 2019, 12, 804.	0.6	6
106	Severe Acute Respiratory Syndrome Coronavirus 2 Variant Delta Infects All 6 Siblings but Spares Comirnaty (BNT162b2, BioNTech/Pfizer)-Vaccinated Parents. Journal of Infectious Diseases, 2021, 224, 1984-1986.	1.9	6
107	A rare CFTR intronic mutation related to a mild CF disease in a 12-year-old girl. BMJ Case Reports, 2012, 2012, bcr2012006918-bcr2012006918.	0.2	6
108	TMEM16A/ANO1: Current Strategies and Novel Drug Approaches for Cystic Fibrosis. Cells, 2021, 10, 2867.	1.8	6

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109	Leveraging TOPMed imputation server and constructing a cohort-specific imputation reference panel to enhance genotype imputation among cystic fibrosis patients. Human Genetics and Genomics Advances, 2022, 3, 100090.	1.0	6
110	More on BNT162b2 Covid-19 Vaccine in Children 5 to 11 Years of Age. New England Journal of Medicine, 2022, 386, 1191-1193.	13.9	6
111	Evaluation of the trachea and intrathoracic airways by the acoustic reflection method in children with cystic fibrosis. Respiratory Physiology and Neurobiology, 2012, 181, 74-78.	0.7	5
112	Endobronchial avium mycobacteria infection in an immunocompetent child. BMJ Case Reports, 2013, 2013, bcr2013200776-bcr2013200776.	0.2	5
113	COVID-19 vaccine prioritisation for people with cystic fibrosis. Journal of Cystic Fibrosis, 2021, 20, 715-716.	0.3	5
114	Respiratory management of spinal muscular atrophy type 1 patients treated with Nusinersen. Pediatric Pulmonology, 2022, 57, 1505-1512.	1.0	5
115	Gene Therapy: A Possible Alternative to CFTR Modulators?. Frontiers in Pharmacology, 2021, 12, 648203.	1.6	4
116	Severe central apnea secondary to cerebellar dysplasia in a child: look past Joubert syndrome. Journal of Clinical Sleep Medicine, 2020, 16, 2113-2116.	1.4	4
117	An idiopathic congenital chylothorax: surgery or conservative management?. BMJ Case Reports, 2014, 2014, bcr2014204147-bcr2014204147.	0.2	3
118	Ultrasonography and Computed Tomographic Manifestations of Abdominal Sarcoidosis in Children. Journal of Pediatric Gastroenterology and Nutrition, 2016, 63, 195-199.	0.9	3
119	Gas exchanges in children with cystic fibrosis or primary ciliary dyskinesia: A retrospective study. Respiratory Physiology and Neurobiology, 2018, 251, 1-7.	0.7	3
120	Chest physiotherapy enhances detection of Pseudomonas aeruginosa in nonexpectorating children with cystic fibrosis. ERJ Open Research, 2021, 7, 00513-2020.	1.1	3
121	Atypical Severe Organizing Pneumonia Following Coronavirus Disease 2019 in an Immunocompromised Teenager. Clinical Infectious Diseases, 2022, 74, 938-939.	2.9	3
122	Factors Predisposing the Response to Lumacaftor/Ivacaftor in People with Cystic Fibrosis. Journal of Personalized Medicine, 2022, 12, 252.	1.1	3
123	SLC6A14 Impacts Cystic Fibrosis Lung Disease Severity via mTOR and Epithelial Repair Modulation. Frontiers in Molecular Biosciences, 2022, 9, 850261.	1.6	3
124	Opposite Expression of Hepatic and Pulmonary Corticosteroid-Binding Globulin in Cystic Fibrosis Patients. Frontiers in Pharmacology, 2018, 9, 545.	1.6	2
125	Pulmonary sarcoid-like granulomatous disease in an 11-month-old girl. BMJ Case Reports, 2013, 2013, bcr2012008024-bcr2012008024.	0.2	2
126	Lung ultrasound in children with interstitial lung disease: a pilot study. , 2020, , .		2

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127	Benefits and risks of bronchoalveolar lavage in severe asthma in children. ERJ Open Research, 2021, 7, 00332-2021.	1.1	2
128	Hordeum murinum aspiration revealed by a pneumopleurocutaneous fistula in a 15-month-old infant. BMC Pediatrics, 2021, 21, 550.	0.7	2
129	Inorganic exposome in pediatric sarcoidosis: The PEDIASARC study. , 2015, , .		1
130	French translation and linguistic validation of the QOL-PCD, a quality of life questionnaire for patients with primary ciliary dyskinesia. , 2016, , .		1
131	Down syndrome and pulmonary hemosiderosis: an under-recognized association. , 2018, , .		1
132	Prophylactic azithromycin in patients with primary ciliary dyskinesia.. , 2018, , .		1
133	Retour dâ€™expÃ©rience dâ€™un hÃ´pital pÃ©diatrique pendant la crise Covid-19 en ÃŽle-de-France. Annales Francaises De Medecine D'Urgence, 2020, 10, 261-265.	0.0	1
134	Dramatic Improvement by Macrolides in Surfactant Deficiency with ABCA3 Mutation.. , 2009, , .		0
135	Azithromycin long term effects in children with cystic fibrosis. Journal of Cystic Fibrosis, 2009, 8, S66.	0.3	0
136	Azithromycin In Interstitial Lung Disease Associated With Surfactant Metabolism Disorders. , 2010, , .		0
137	149* 2009 H1N1 influenza A in cystic fibrosis patients. A French collaborative study. Journal of Cystic Fibrosis, 2011, 10, S38.	0.3	0
138	Search For Interstitial Lung Disease Etiology In Children: A Step By Step Approach. , 2011, , .		0
139	Are CF carriers predisposed to asthma?. Journal of Cystic Fibrosis, 2016, 15, 555-556.	0.3	0
140	Pulmonary Hemorrhage Revealing Multiple Vascular Malformations in a Child with KCNT1 Developmental Epileptic Encephalopathy. Journal of Pediatrics, 2021, 237, 311-312.	0.9	0
141	642: SLC6A14 is associated with lung function in patients with cystic fibrosis, regulates epithelial repair and mTOR signaling in bronchial epithelial cells. Journal of Cystic Fibrosis, 2021, 20, S305.	0.3	0
142	Rare localization of NUT midline carcinoma revealed by pneumonia and pleural effusion in a 13 years old patient: A case report. , 2015, , .		0
143	Gaz exchange, nasal nitric oxyde levels and lung function tests in children with primary ciliary dyskinesia. , 2015, , .		0
144	Creation and validation of a questionnaire assessing quality of lifeÂof parents of children with interstitial lung disease (chILD). , 2021, , .		0