Hayriye Ugur Ozcelik

List of Publications by Year
in descending order

173 papers	2,728 citations	236925 25 h-index	223800 46 g-index
175 all docs	175 docs citations	175 times ranked	3656 citing authors

#	Article	IF	Citations
1	Mycobacterial Disease and Impaired IFN-Î³ Immunity in Humans with Inherited ISG15 Deficiency. Science, 2012, 337, 1684-1688.	12.6	455
2	Mutations in SLC34A2 Cause Pulmonary Alveolar Microlithiasis and Are Possibly Associated with Testicular Microlithiasis. American Journal of Human Genetics, 2006, 79, 650-656.	6.2	226
3	Congenital lobar emphysema: Evaluation and long-term follow-up of thirty cases at a single center. Pediatric Pulmonology, 2003, 35, 384-391.	2.0	101
4	High resolution CT in children with cystic fibrosis: correlation with pulmonary functions and radiographic scores. European Journal of Radiology, 2001, 37, 54-59.	2.6	68
5	Treatment of Hydatid Disease. Paediatric Drugs, 2001, 3, 123-135.	3.1	68
6	Postinfectious Bronchiolitis obliterans in Children: Clinical and Radiological Profile and Prognostic Factors. Respiration, 2003, 70, 371-375.	2.6	64
7	Medical treatment of pulmonary hydatid disease: for which child?. Parasitology International, 2005, 54, 135-138.	1.3	57
8	Homozygous and Compound-heterozygous Type I Plasminogen Deficiency Is a Common Cause of Ligneous Conjunctivitis. Thrombosis and Haemostasis, 2001, 85, 1004-1010.	3.4	56
9	Longâ€term results of disodium etidronate treatment in pulmonary alveolar microlithiasis. Pediatric Pulmonology, 2010, 45, 514-517.	2.0	52
10	Conservative Treatment of Empyema in Children. Respiration, 1993, 60, 182-185.	2.6	50
11	Gorhamâ€“Stout Syndrome with chylothorax: Successful remission by interferon alphaâ€2b. Pediatric Pulmonology, 2009, 44, 613-615.	2.0	47
12	A comparison of respiratory and peripheral muscle strength, functional exercise capacity, activities of daily living and physical fitness in patients with cystic fibrosis and healthy subjects. Research in Developmental Disabilities, 2015, 45-46, 147-156.	2.2	47
13	Hydatid disease in childhood: A retrospective analysis of 376 cases. , 1998, 26, 190-196.		46
14	Genotype and phenotype evaluation of patients with primary ciliary dyskinesia: First results from Turkey. Pediatric Pulmonology, 2020, 55, 383-393.	2.0	46
15	Different features of lung involvement in Niemann-Pick disease and Gaucher disease. Respiratory Medicine, 2012, 106, 1278-1285.	2.9	45
16	Clinical features and treatment approaches in cystic fibrosis with pseudo-Bartter syndrome. Annals of Tropical Paediatrics, 2005, 25, 119-124.	1.0	39
17	Bronchiectasis: the Consequence of Late Diagnosis in Chronic Respiratory Symptoms. Journal of Tropical Pediatrics, 2005, 51, 362-365.	1.5	39
18	Psychiatric and general health effects of COVIDâ€19 pandemic on children with chronic lung disease and parents' coping styles. Pediatric Pulmonology, 2020, 55, 3579-3586.	2.0	39

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19	Tracheomalacia and Bronchomalacia in 34 Children: Clinical and Radiologic Profiles and Associations with Other Diseases. Clinical Pediatrics, 2005, 44, 777-781.	0.8	37
20	Pulmonary exacerbations in patients with primary ciliary dyskinesia: an expert consensus definition for use in clinical trials. ERJ Open Research, 2019, 5, 00147-2018.	2.6	37
21	Tracheobronchopathia Osteochondroplastica in a 9-year-old Girl. Pediatric Pulmonology, 2006, 41, 95-97.	2.0	32
22	Sodium chloride deficiency in cystic fibrosis patients. European Journal of Pediatrics, 1994, 153, 829-831.	2.7	31
23	Effects of chest physiotherapy and aerobic exercise training on physical fitness in young children with cystic fibrosis. Italian Journal of Pediatrics, 2012, 38, 2.	2.6	29
24	Hepatobiliary manifestations of cystic fibrosis in children: correlation of CT and US findings. European Journal of Radiology, 2002, 41, 26-33.	2.6	28
25	Omalizumab Treatment for Allergic Bronchopulmonary Aspergillosis in Cystic Fibrosis. Annals of Pharmacotherapy, 2016, 50, 188-193.	1.9	27
26	Cystic fibrosis in Turkey: First data from the national registry. Pediatric Pulmonology, 2020, 55, 541-548.	2.0	27
27	Pulmonary and extrapulmonary features in bronchopulmonary dysplasia: a comparison with healthy children. Journal of Physical Therapy Science, 2015, 27, 1761-1765.	0.6	26
28	An unusual pattern of arthritis in a child with Kawasaki syndrome. Clinical Rheumatology, 2004, 23, 73-75.	2.2	25
29	Prevalence and course of disease after lung resection in primary ciliary dyskinesia: a cohort & nested case-control study. Respiratory Research, 2019, 20, 212.	3.6	23
30	Current Approach in the Diagnosis and Management of Allergic Bronchopulmonary Aspergillosis in Children With Cystic Fibrosis. Frontiers in Pediatrics, 2020, 8, 582964.	1.9	23
31	Childhood Parapneumonic Effusions. Chest, 2005, 128, 1436-1441.	0.8	22
32	An epidemic of pseudo-Bartter syndrome in cystic fibrosis patients. European Journal of Pediatrics, 2007, 167, 115-116.	2.7	22
33	Clinical Strains ofChryseobacteriumandElizabethkingiaspp. Isolated from Pediatric Patients in a University Hospital: Performance of MALDI-TOF MS-Based Identification, Antimicrobial Susceptibilities, and Baseline Patient Characteristics. Microbial Drug Resistance, 2018, 24, 816-821.	2.0	22
34	Treatment and follow-up of pulmonary alveolar microlithiasis with disodium editronate: radiological demonstration. Pediatric Radiology, 2002, 32, 380-383.	2.0	20
35	Childhood diffuse panbronchiolitis: A case report. Pediatric Pulmonology, 2005, 40, 354-357.	2.0	20
36	Preventing tuberculosis in children receiving anti-tnf treatment. Clinical Rheumatology, 2010, 29, 389-392.	2.2	20

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37	A review of the etiology and clinical presentation of non-cystic fibrosis bronchiectasis: A tertiary care experience. Respiratory Medicine, 2018, 137, 35-39.	2.9	20
38	Pulmonary involvement in a child with ligneous conjunctivitis and homozygous type I plasminogen deficiency. Pediatric Pulmonology, 2001, 32, 179-183.	2.0	19
39	Xanthoma disseminatum: A child with respiratory system involvement and bronchiectasis. Pediatric Pulmonology, 2005, 39, 84-87.	2.0	18
40	The role of TAP1 and TAP2 gene polymorphism in idiopathic bronchiectasis in children. Pediatric Pulmonology, 2007, 42, 237-241.	2.0	18
41	Study protocol: the earâ€“noseâ€“throat (ENT) prospective international cohort of patients with primary ciliary dyskinesia (EPIC-PCD). BMJ Open, 2021, 11, e051433.	1.9	18
42	Pneumomediastinum, pneumorrhachis and subcutaneous emphysema associated with viral infections: Report of three cases. Pediatrics International, 2015, 57, 1038-1040.	0.5	17
43	Horseshoe lung associated with left-lung hypoplasia, left pulmonary artery sling and bilateral agenesis of upper lobe bronchi. Pediatric Radiology, 2009, 39, 1002-1005.	2.0	16
44	Relation of bone mineral density with clinical and laboratory parameters in preâ€pubertal children with cystic fibrosis. Pediatric Pulmonology, 2009, 44, 706-712.	2.0	16
45	The role of human leucocyte antigens in children with hydatid disease: their association with clinical condition and prognosis. Parasitology Research, 2010, 106, 795-800.	1.6	16
46	The success of the different eradication therapy regimens for Pseudomonas aeruginosa in cystic fibrosis. Journal of Clinical Pharmacy and Therapeutics, 2016, 41, 419-423.	1.5	16
47	Surgical Management of Infants with Congenital Lobar Emphysema and Concomitant Congenital Heart Disease. Heart Surgery Forum, 2004, 7, E644-E649.	0.5	16
48	Hypersensitivity pneumonitis in children: pigeon breeders disease. Annals of Tropical Paediatrics, 2004, 24, 349-355.	1.0	15
49	Successful unilateral partial lung lavage in a child with pulmonary alveolar proteinosis. Journal of Clinical Anesthesia, 2009, 21, 127-130.	1.6	15
50	Reduced anaerobic and aerobic performance in children with primary ciliary dyskinesia. European Journal of Pediatrics, 2018, 177, 765-773.	2.7	15
51	BALF Nitrite as an Indicator of Inflammation in Children with Cystic Fibrosis. Respiration, 2004, 71, 625-629.	2.6	14
52	Pleural Fluid PCR Method for Detection of Staphylococcus aureus, Streptococcus pneumoniae and Haemophilus influenzae in Pediatric Parapneumonic Effusions. Respiration, 2008, 75, 437-442.	2.6	14
53	TAP1 and TAP2 gene polymorphisms in childhood cystic echinococcosis. Parasitology International, 2010, 59, 283-285.	1.3	14
54	Effects of blood transfusion on cytokine profile and pulmonary function in patients with thalassemia major. Clinical Respiratory Journal, 2016, 10, 153-162.	1.6	14

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55	Environmental Tobacco Smoke Exposure and Respiratory Morbidity in Children. Inhalation Toxicology, 2007, 19, 779-785.	1.6	12
56	Congenital Tuberculosis after in-vitro Fertilization in a Woman Previously Undiagnosed with Tuberculosis Salpingitis. Pediatrics and Neonatology, 2016, 57, 539-540.	0.9	12
57	Developmental and behavioral problems in preschool-aged primary ciliary dyskinesia patients. European Journal of Pediatrics, 2019, 178, 995-1003.	2.7	12
58	Clinical features and accompanying findings of Pseudoâ€Bartter Syndrome in cystic fibrosis. Pediatric Pulmonology, 2020, 55, 2011-2016.	2.0	12
59	Multicentric analysis of childhood tuberculosis in Turkey. Turkish Journal of Pediatrics, 2013, 55, 121-9.	0.6	12
60	Tuberculin skin test positivity in pediatric allogeneic BMT recipients and donors in Turkey. Pediatric Transplantation, 2007, 11, 414-418.	1.0	11
61	Auditory nerve-brainstem responses in cystic fibrosis patients. International Journal of Pediatric Otorhinolaryngology, 1996, 35, 165-169.	1.0	10
62	Diagnosis and treatment of pulmonary alveolar microlithiasis. Pediatrics International, 2016, 58, 805-807.	0.5	10
63	Neutrophil chemotaxis in acutely infected and clinically stable cystic fibrosis patients*. Pediatrics International, 1999, 41, 514-518.	0.5	9
64	Gastric organo-axial malrotation coexisting respiratory symptoms. European Journal of Pediatrics, 2009, 168, 491-494.	2.7	9
65	Telephone surveillance during 2019 novel coronavirus disease: Is it a helpful diagnostic tool for detecting acute pulmonary exacerbations in children with chronic lung disease?. Journal of Telemedicine and Telecare, 2020, , 1357633X2097200.	2.7	9
66	Cardiac hydatid cyst and pulmonary hydatidosis in a child. Pediatric Infectious Disease Journal, 2002, 21, 1178-1180.	2.0	8
67	Pigeon-breeder's disease in a child with selective IgA deficiency. Pediatrics International, 2003, 45, 216-218.	0.5	8
68	Clinical status, ocular surface changes and tear ferning in patients with cystic fibrosis. Acta Ophthalmologica, 1996, 74, 563-565.	0.3	8
69	Does Defective Apoptosis Play A Role in Cystic Fibrosis Lung Disease?. Archives of Medical Research, 2009, 40, 561-564.	3.3	8
70	Successful treatment of pulmonary hemangioma with propranolol. Pediatric Pulmonology, 2014, 49, 829-833.	2.0	8
71	Variation in the bombesin staining of pulmonary neuroendocrine cells in pediatric pulmonary disordersâ€"A useful marker for airway maturity. Pediatric Pulmonology, 2020, 55, 2383-2388.	2.0	8
72	Does cystic fibrosis make susceptible to celiac disease?. European Journal of Pediatrics, 2021, 180, 2807-2813.	2.7	8

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73	Pulmonary complications following hematopoietic stem cell transplantation in children. Turkish Journal of Pediatrics, 2019, 61, 59.	0.6	8
74	Foreign-body aspiration mimicking congenital lobar emphysema in a forty-eight-day-old girl. Pediatric Pulmonology, 2005, 39, 189-191.	2.0	7
75	A 4-month-old boy with acrodermatitis enteropathica-like symptoms. European Journal of Pediatrics, 2009, 168, 119-121.	2.7	7
76	DYNC2H1 mutation causes Jeune syndrome and recurrent lung infections associated with ciliopathy. Clinical Respiratory Journal, 2018, 12, 1017-1020.	1.6	7
77	Impact of Surgery on Growth, Pulmonary Functions, and Acute Pulmonary Exacerbations in Children with Non-Cystic Fibrosis Bronchiectasis. Thoracic and Cardiovascular Surgeon, 2019, 67, 058-066.	1.0	7
78	Deviations of body functions and structure, activity limitations, and participation restrictions of the International Classification of Functioning, Disability, and Health model in children with cystic fibrosis and nonâ€“cystic fibrosis bronchiectasis. Pediatric Pulmonology, 2020, 55, 1207-1216.	2.0	7
79	Nonmyeloablative hematopoietic stem cell transplantation in a patient with hereditary pulmonary alveolar proteinosis. Pediatric Pulmonology, 2021, 56, 341-343.	2.0	7
80	Motor repertoire is age-inadequate in infants with cystic fibrosis. Pediatric Research, 2021, 89, 1291-1296.	2.3	7
81	Prenatal period to adolescence: the variable presentations of congenital cystic adenomatoid malformation. Pediatrics International, 2006, 48, 626-630.	0.5	6
82	PulmonaryMycobacterium abscessusInfection in a Patient with Triple A Syndrome. Journal of Tropical Pediatrics, 2016, 62, 324-327.	1.5	6
83	Hospitalization of Children with Cystic Fibrosis Adversely Affects Mothersâ€™ Physical Activity, Sleep Quality, and Psychological Status. Journal of Child and Family Studies, 2017, 26, 800-809.	1.3	6
84	Childhood diffuse parenchymal lung diseases: We need a new classification. Clinical Respiratory Journal, 2020, 14, 102-108.	1.6	6
85	Nasal nitric oxide levels in primary ciliary dyskinesia, cystic fibrosis and healthy children. Turkish Journal of Pediatrics, 2019, 61, 20.	0.6	6
86	Dollâ€like face: Is it an underestimated clinical presentation of cystic fibrosis?. Pediatric Pulmonology, 2008, 43, 634-637.	2.0	5
87	The value of flexible bronchoscopy in pulmonary infections of immunosuppressed children. Clinical Respiratory Journal, 2020, 14, 78-84.	1.6	5
88	Patients eligible for modulator drugs: Data from cystic fibrosis registry of Turkey. Pediatric Pulmonology, 2020, 55, 2302-2306.	2.0	5
89	Plasma Ceramides and Sphingomyelins of Pediatric Patients Increase in Primary Ciliary Dyskinesia but Decrease in Cystic Fibrosis. Lipids, 2020, 55, 213-223.	1.7	5
90	Impact of mannoseâ€binding lectin 2 gene polymorphisms on disease severity in noncystic fibrosis bronchiectasis in children. Pediatric Pulmonology, 2020, 55, 1190-1198.	2.0	5

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91	Respiratory viruses: What is their role in acute exacerbations in children with cystic fibrosis?. Pediatric Pulmonology, 2020, 55, 1646-1652.	2.0	5
92	The Effect of Self-Efficacy, Social Support and Quality of Life on Readiness for Transition to Adult Care Among Adolescents with Cystic Fibrosis in Turkey. Journal of Pediatric Nursing, 2021, 57, e79-e84.	1.5	5
93	Evaluation of pulmonary complications and affecting factors in children for repaired esophageal atresia and tracheoesophageal fistula. Respiratory Medicine, 2021, 181, 106376.	2.9	5
94	Impact of COVIDâ€19 on pediatric pulmonology healthcare practice. Pediatric Pulmonology, 2021, 56, 2811-2817.	2.0	5
95	Diagnosis of cystic fibrosis with chloride meter (Sherwood M926S chloride analyzerÂ®) and sweat test analysis system (CFÎ" collection systemÂ®) compared to the Gibson Cooke method. Turkish Journal of Pediatrics, 2016, 58, 27-33.	0.6	5
96	Comparison of inhaled mannitol/dornase alfa combination and daily dornase alfa alone in children with cystic fibrosis. Pediatric Pulmonology, 2022, 57, 142-151.	2.0	5
97	The controversy of drug hypersensitivity in patients with cystic fibrosis and review of the literature. Pediatric Allergy and Immunology, 2022, 33, .	2.6	5
98	Impact of Achromobacter spp. isolation on clinical outcomes in children with cystic fibrosis. Pediatric Pulmonology, 2022, 57, 658-666.	2.0	5
99	Chemical pneumonia caused by glutaraldehyde. Pediatrics International, 2001, 43, 701-702.	0.5	4
100	A case of congenital agenesis of the right pulmonary artery presenting with hemoptysis and mimicking pulmonary hemosiderosis. European Journal of Pediatrics, 2009, 168, 217-220.	2.7	4
101	Predominance of hospital-associated MRSA among cystic fibrosis patients in a Turkish reference cystic fibrosis centre. Journal of Chemotherapy, 2012, 24, 195-200.	1.5	4
102	From Diagnosis to Treatment of Pediatric Tuberculosis: Ten Years Experience in a Single Institution. Clinical Pediatrics, 2020, 59, 476-482.	0.8	4
103	Physical fitness and activities of daily living in primary ciliary dyskinesia: A retrospective study. Pediatrics International, 2022, 64, .	0.5	4
104	Differentially expressed genes associated with disease severity in siblings with cystic fibrosis. Pediatric Pulmonology, 2021, 56, 910-920.	2.0	4
105	Long term azithromycin therapy in patients with cystic fibrosis. Turkish Journal of Pediatrics, 2016, 58, 34-40.	0.6	4
106	Aerobic exercise training in Kartagenerâ€™s syndrome: case report. Journal of Exercise Rehabilitation, 2019, 15, 468-471.	1.0	4
107	The success of the Cystic Fibrosis Registry of Turkey for improvement of patient care. Pediatric Pulmonology, 2022, , .	2.0	4
108	The role of serum Pseudomonas aeruginosa antibodies in the diagnosis and follow-up of cystic fibrosis. Turkish Journal of Pediatrics, 2013, 55, 50-7.	0.6	4

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109	Primary Nasal Tuberculosis in a 10-Year-Old Girl. Canadian Journal of Infectious Diseases and Medical Microbiology, 2016, 2016, 1-3.	1.9	3
110	Clinical spectrum of children with interstitial pneumonia with autoimmune features. Respiratory Medicine, 2021, 187, 106566.	2.9	3
111	Sleep disordered breathing in patients with Prader willi syndrome: Impact of underlying genetic mechanism. Respiratory Medicine, 2021, 187, 106567.	2.9	3
112	Swallowing dysfunction as a factor that should be remembered in recurrent pneumonia: videofluoroscopic swallow study. Minerva Pediatrics, 2017, 69, 396-402.	0.4	3
113	Evolution of Primary Ciliary Dyskinesia (PCD) diagnostic testing in Europe. , 2017, , .		3
114	Cystic fibrosis newborn screening: Fiveâ€year experience from a tertiary care center. Pediatric Pulmonology, 2022, 57, 403-410.	2.0	3
115	Pediatric Primary Pulmonary Tuberculosis. Chest, 2002, 121, 1722.	0.8	2
116	Does <scp>Helicobacter pylori</scp> play a role in the pathogenesis of nonâ€cystic fibrosis bronchiectasis?. Pediatrics International, 2016, 58, 894-898.	0.5	2
117	Obstructive sleep apnea in children with hypothalamic obesity: Evaluation of possible related factors. Pediatric Pulmonology, 2020, 55, 3532-3540.	2.0	2
118	Cystic fibrosis in Turkey. Lancet Respiratory Medicine,the, 2020, 8, e17.	10.7	2
119	Mutations of the CFTR gene and novel variants in Turkish patients with cystic fibrosis: 24-years experience. Clinica Chimica Acta, 2020, 510, 252-259.	1.1	2
120	The frequency and related factors of nonâ€tuberculosis mycobacteria infections among patients with cystic fibrosis. Pediatrics International, 2021, 63, 1369-1375.	0.5	2
121	Chronic necrotizing pulmonary aspergillosis in an immunocompetent patient after the surgery of hydatid cyst. Tuberkuloz Ve Toraks, 2017, 65, 157-160.	0.4	2
122	Developmental Functioning Outcomes in Infants With Cystic Fibrosis: A 24- to 36-Month Follow-Up Study. Physical Therapy, 2022, 102, .	2.4	2
123	Clinical radiological and pathological staging of children with hypersensitivity pneumonitis. Pediatric Pulmonology, 2022, 57, 2344-2355.	2.0	2
124	Airway Malacia Disorders in Children. Chest, 2006, 130, 304.	0.8	1
125	Comparison of Bronchoalveolar Lavage and Sputum Microbiology in Patients with Primary Ciliary Dyskinesia. Pediatric, Allergy, Immunology, and Pulmonology, 2017, 30, 14-17.	0.8	1
126	Evaluation of sleepâ€disordered breathing and its relationship with respiratory parameters in children with mucopolysaccharidosis Type IVA and VI. American Journal of Medical Genetics, Part A, 2021, 185, 2306-2314.	1.2	1

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127	Clinical implications of fungal isolation from sputum in adult patients with cystic fibrosis. Turkish Journal of Medical Sciences, 2021, 51, 1191-1200.	0.9	1
128	Plasma ceramides and sphingomyelins increase in primary ciliary dyskinesia but decrease in cystic fibrosis. , 2018, , .		1
129	Neonatal manifestations in Primary Ciliary Dyskinesia: a multinational cohort study. , 2018, , .		1
130	Evaluation of Sleep Disorders In Children with Down Syndrome. , 2019, , .		1
131	Retroperitoneal Abscess in Severe Combined Immunodeficiency Probably Due to BCG Vaccine. Asim, Allerji, Immunoloji, 2020, 18, 98-101.	0.0	1
132	Accidental Overdose and Improper Vaccination with BCG in Childhood: Report of Three Cases. Turkiye Klinikleri Journal of Medical Sciences, 2011, 31, 724-726.	0.1	1
133	Unfinished battle with childhood tuberculosis: is it curable with less drugs and shorter duration?. Tuberkuloz Ve Toraks, 2013, 61, 320-326.	0.4	1
134	An insight of lung cysts with filamin A mutation. , 2015, , .		1
135	Presence of Kartagener syndrome reduces exercise capacity in primary ciliary dyskinesia. , 2016, , .		1
136	Clinical features of pseudo-bartter syndrome in cystic fibrosis. , 2017, , .		1
137	Prevalence and Impact of Lung Resection in Primary Ciliary Dyskinesia: a cohort and nested case-control study. , 2018, , .		1
138	A rare cause of acute abdominal pain in a patient with primary ciliary dyskinesia with situs inversus totalis. Turkish Journal of Pediatrics, 2020, 62, 156.	0.6	1
139	Clinical characteristics of children with cystic fibrosis infected with unusual bacteria. Minerva Pediatrics, 2021, , .	0.4	1
140	Human bocavirus and human metapneumovirus in children with lower respiratory tract infections: Effects on clinical, microbiological features and disease severity. Pediatrics International, 2022, 64, .	0.5	1
141	The effect of Pseudomonas aeruginosa eradication regimens on chronic colonization and clinical outcomes in pediatric patients with cystic fibrosis. Pediatrics International, 2022, 64, .	0.5	1
142	Antimycobacterial prophylaxis regarding Bacillus Calmette-GuÃ©rin -associated complications in children with primary immunodeficiency. Respiratory Medicine, 2022, , 106919.	2.9	1
143	Comparason of extraction methods for PCR detection of Burkholderia cepacia complex (BCC) from cystic fibrosis patients. Open Medicine (Poland), 2008, 3, 157-162.	1.3	0
144	Eosinophilic Lung Diseases: Five Cases In Childhood. , 2010, , .		0

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145	Childhood idiopathic interstitial pneumonia: diagnosis, treatment and follow-up. Turk Pediatri Arsivi, 2013, 48, 281-287.	0.9	0
146	Post-Infectious Bronchiolitis Obliterans in Children: Long-Term Outcome, Prognostic Factors, and Relation with Serum KL-6 Levels. Pediatric, Allergy, Immunology, and Pulmonology, 2015, 28, 152-157.	0.8	0
147	Factors associated with severe lung disease in an adult population with cystic fibrosis: a single-center experience. Turkish Journal of Medical Sciences, 2020, 50, 945-952.	0.9	0
148	Clinical findings of patients with cystic fibrosis according to newborn screening results. Pediatrics International, 2022, 64, .	0.5	0
149	Increased Plasma YKL-40 Level and Chitotriosidase Activity in Cystic Fibrosis Patients. Inflammation, 2022, 45, 627-638.	3.8	0
150	Mekonyum ileusunun 12 ay altÄ±nda tanÄ± alan kistik fibrozlu hastalarda klinik seyire etkisi. Turk Pediatri Arsivi, 2011, 45, 105-110.	0.9	0
151	Neuroendocrine cell hyperplasia in children with idiopathic interstitial pneumonia. , 2015, , .		0
152	Hacettepe University experience: Clinical evaluation of 218 primary ciliary dyskinesia patients. , 2015, , .		0
153	Flexible bronchoscopy findings in children with protracted bacterial bronchitis. , 2015, , .		0
154	A comparison of aerobic and anaerobic capacity in children with primary ciliary dyskinesia and healthy peers. , 2015, , .		0
155	Radiological findings of patients with primary ciliary dyskinesia. , 2015, , .		0
156	Pulmonary involvement in children with rheumatic diseases. , 2016, , .		0
157	A ten year experience of non-cystic fibrosis bronchiectasis from a single center. , 2016, , .		0
158	Comparison of peripheral muscle strength, posture and spinal curvatures for patients with cystic fibrosis and bronchiectasis and healthy subjects. , 2016, , .		0
159	Lung function tests in patients with primary ciliary dyskinesia. , 2016, , .		0
160	Comparison of inspiratory muscle training and home-based rehabilitation approach in patients with primary ciliary dyskinesia. , 2016, , .		0
161	Preliminary Results of Whole Exome Sequencing in Turkish Primary Ciliary Dyskinesia Patients - Hacettepe University Experience: â€œThree candidate genes, five novel and two known mutationsâ€• , 2017, , .		0
162	Postoperative respiratory problems in children with esophageal atresia and tracheoesophageal fistula. , 2017, , .		0

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163	Newborn screening for CF: first tree years experience in Hacettepe University. , 2018, , .		0
164	Assessment of motor repertoire and neurodevelopmental level in infants with cystic fibrosis: a follow-up study. , 2019, , .		0
165	Cystic Fibrosis Patients Eligible for Modulator Drugs: Data from Cystic Fibrosis Registry of Turkey. , 2019, , .		0
166	Vitamin D and LL-37 Levels In Childhood Tuberculosis Disease and Latent Tuberculosis Infection. , 2019, , .		0
167	Severe Langerhans Cell Histiocytosis with Pulmonary Involvement. , 2019, , .		0
168	Lung function, functional capacity, energy cost, and nutritional status in cystic fibrosis with and without inspiratory muscle weakness. , 2019, , .		0
169	The Correlation Between Clinical Characteristics and Molecular Genetic Analysis Results of Primary Ciliary Dyskinesia Patients: Hacettepe University Experience. , 2019, , .		0
170	The Effects Of Respiratory Viruses on The Acute Pulmonary Exacerbations In Cystic Fibrosis Patients. , 2019, , .		0
171	Arm functional capacity and activity of daily living in children with primary ciliary dyskinesia and cystic fibrosis. , 2019, , .		0
172	Risk of the developmental delay and behavior problems in preschool-aged primary ciliary dyskinesia patients. , 2019, , .		0
173	Active video gaming in primary ciliary dyskinesia: a randomized controlled trial. European Journal of Pediatrics, 2022, , .	2.7	0