Hayriye Ugur Ozcelik

List of Publications by Year
in descending order

Version: 2024-02-01

173 papers	2,728 citations	270111 25 h-index	252626 46 g-index
175 all docs	175 docs citations	175 times ranked	3880 citing authors

#	Article	IF	Citations
1	Clinical findings of patients with cystic fibrosis according to newborn screening results. Pediatrics International, 2022, 64, .	0.2	0
2	Physical fitness and activities of daily living in primary ciliary dyskinesia: A retrospective study. Pediatrics International, 2022, 64, .	0.2	4
3	Increased Plasma YKL-40 Level and Chitotriosidase Activity in Cystic Fibrosis Patients. Inflammation, 2022, 45, 627-638.	1.7	0
4	Comparison of inhaled mannitol/dornase alfa combination and daily dornase alfa alone in children with cystic fibrosis. Pediatric Pulmonology, 2022, 57, 142-151.	1.0	5
5	Cystic fibrosis newborn screening: Fiveâ€year experience from a tertiary care center. Pediatric Pulmonology, 2022, 57, 403-410.	1.0	3
6	Human bocavirus and human metapneumovirus in children with lower respiratory tract infections: Effects on clinical, microbiological features and disease severity. Pediatrics International, 2022, 64, .	0.2	1
7	The controversy of drug hypersensitivity in patients with cystic fibrosis and review of the literature. Pediatric Allergy and Immunology, 2022, 33, .	1.1	5
8	The success of the Cystic Fibrosis Registry of Turkey for improvement of patient care. Pediatric Pulmonology, 2022, , .	1.0	4
9	Impact of Achromobacter spp. isolation on clinical outcomes in children with cystic fibrosis. Pediatric Pulmonology, 2022, 57, 658-666.	1.0	5
10	Developmental Functioning Outcomes in Infants With Cystic Fibrosis: A 24- to 36-Month Follow-Up Study. Physical Therapy, 2022, 102, .	1.1	2
11	Active video gaming in primary ciliary dyskinesia: a randomized controlled trial. European Journal of Pediatrics, 2022, , .	1.3	0
12	The effect of Pseudomonas aeruginosa eradication regimens on chronic colonization and clinical outcomes in pediatric patients with cystic fibrosis. Pediatrics International, 2022, 64, .	0.2	1
13	Clinical radiological and pathological staging of children with hypersensitivity pneumonitis. Pediatric Pulmonology, 2022, 57, 2344-2355.	1.0	2
14	Antimycobacterial prophylaxis regarding Bacillus Calmette-GuÃ©rin -associated complications in children with primary immunodeficiency. Respiratory Medicine, 2022, , 106919.	1.3	1
15	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.	0.7	5
16	Nonmyeloablative hematopoietic stem cell transplantation in a patient with hereditary pulmonary alveolar proteinosis. Pediatric Pulmonology, 2021, 56, 341-343.	1.0	7
17	Motor repertoire is age-inadequate in infants with cystic fibrosis. Pediatric Research, 2021, 89, 1291-1296.	1.1	7
18	The frequency and related factors of nonâ€tuberculosis mycobacteria infections among patients with cystic fibrosis. Pediatrics International, 2021, 63, 1369-1375.	0.2	2

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19	Does cystic fibrosis make susceptible to celiac disease?. European Journal of Pediatrics, 2021, 180, 2807-2813.	1.3	8
20	Evaluation of pulmonary complications and affecting factors in children for repaired esophageal atresia and tracheoesophageal fistula. Respiratory Medicine, 2021, 181, 106376.	1.3	5
21	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.	0.7	1
22	Clinical implications of fungal isolation from sputum in adult patients with cystic fibrosis. Turkish Journal of Medical Sciences, 2021, 51, 1191-1200.	0.4	1
23	Impact of COVIDâ€19 on pediatric pulmonology healthcare practice. Pediatric Pulmonology, 2021, 56, 2811-2817.	1.0	5
24	Clinical spectrum of children with interstitial pneumonia with autoimmune features. Respiratory Medicine, 2021, 187, 106566.	1.3	3
25	Sleep disordered breathing in patients with Prader willi syndrome: Impact of underlying genetic mechanism. Respiratory Medicine, 2021, 187, 106567.	1.3	3
26	Differentially expressed genes associated with disease severity in siblings with cystic fibrosis. Pediatric Pulmonology, 2021, 56, 910-920.	1.0	4
27	Study protocol: the earâ€“noseâ€“throat (ENT) prospective international cohort of patients with primary ciliary dyskinesia (EPIC-PCD). BMJ Open, 2021, 11, e051433.	0.8	18
28	Clinical characteristics of children with cystic fibrosis infected with unusual bacteria. Minerva Pediatrics, 2021, , .	0.2	1
29	The value of flexible bronchoscopy in pulmonary infections of immunosuppressed children. Clinical Respiratory Journal, 2020, 14, 78-84.	0.6	5
30	Cystic fibrosis in Turkey: First data from the national registry. Pediatric Pulmonology, 2020, 55, 541-548.	1.0	27
31	Childhood diffuse parenchymal lung diseases: We need a new classification. Clinical Respiratory Journal, 2020, 14, 102-108.	0.6	6
32	Genotype and phenotype evaluation of patients with primary ciliary dyskinesia: First results from Turkey. Pediatric Pulmonology, 2020, 55, 383-393.	1.0	46
33	Obstructive sleep apnea in children with hypothalamic obesity: Evaluation of possible related factors. Pediatric Pulmonology, 2020, 55, 3532-3540.	1.0	2
34	Cystic fibrosis in Turkey. Lancet Respiratory Medicine,the, 2020, 8, e17.	5.2	2
35	Mutations of the CFTR gene and novel variants in Turkish patients with cystic fibrosis: 24-years experience. Clinica Chimica Acta, 2020, 510, 252-259.	0.5	2
36	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.	1.4	9

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37	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.	0.4	0
38	Current Approach in the Diagnosis and Management of Allergic Bronchopulmonary Aspergillosis in Children With Cystic Fibrosis. Frontiers in Pediatrics, 2020, 8, 582964.	0.9	23
39	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.	1.0	39
40	Clinical features and accompanying findings of Pseudoâ€Bartter Syndrome in cystic fibrosis. Pediatric Pulmonology, 2020, 55, 2011-2016.	1.0	12
41	Patients eligible for modulator drugs: Data from cystic fibrosis registry of Turkey. Pediatric Pulmonology, 2020, 55, 2302-2306.	1.0	5
42	Plasma Ceramides and Sphingomyelins of Pediatric Patients Increase in Primary Ciliary Dyskinesia but Decrease in Cystic Fibrosis. Lipids, 2020, 55, 213-223.	0.7	5
43	Impact of mannoseâ€binding lectin 2 gene polymorphisms on disease severity in noncystic fibrosis bronchiectasis in children. Pediatric Pulmonology, 2020, 55, 1190-1198.	1.0	5
44	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.	1.0	8
45	From Diagnosis to Treatment of Pediatric Tuberculosis: Ten Years Experience in a Single Institution. Clinical Pediatrics, 2020, 59, 476-482.	0.4	4
46	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.	1.0	7
47	Respiratory viruses: What is their role in acute exacerbations in children with cystic fibrosis?. Pediatric Pulmonology, 2020, 55, 1646-1652.	1.0	5
48	Retroperitoneal Abscess in Severe Combined Immunodeficiency Probably Due to BCG Vaccine. Asim, Allerji, Immunoloji, 2020, 18, 98-101.	0.2	1
49	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.	0.3	1
50	Prevalence and course of disease after lung resection in primary ciliary dyskinesia: a cohort & nested case-control study. Respiratory Research, 2019, 20, 212.	1.4	23
51	Pulmonary exacerbations in patients with primary ciliary dyskinesia: an expert consensus definition for use in clinical trials. ERJ Open Research, 2019, 5, 00147-2018.	1.1	37
52	Developmental and behavioral problems in preschool-aged primary ciliary dyskinesia patients. European Journal of Pediatrics, 2019, 178, 995-1003.	1.3	12
53	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.	0.4	7
54	Evaluation of Sleep Disorders In Children with Down Syndrome. , 2019, , .		1

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55	Nasal nitric oxide levels in primary ciliary dyskinesia, cystic fibrosis and healthy children. Turkish Journal of Pediatrics, 2019, 61, 20.	0.3	6
56	Pulmonary complications following hematopoietic stem cell transplantation in children. Turkish Journal of Pediatrics, 2019, 61, 59.	0.3	8
57	Aerobic exercise training in Kartagenerâ€™s syndrome: case report. Journal of Exercise Rehabilitation, 2019, 15, 468-471.	0.4	4
58	Assessment of motor repertoire and neurodevelopmental level in infants with cystic fibrosis: a follow-up study. , 2019, , .		0
59	Cystic Fibrosis Patients Eligible for Modulator Drugs: Data from Cystic Fibrosis Registry of Turkey. , 2019, , .		0
60	Vitamin D and LL-37 Levels In Childhood Tuberculosis Disease and Latent Tuberculosis Infection. , 2019, , .		0
61	Severe Langerhans Cell Histiocytosis with Pulmonary Involvement. , 2019, , .		0
62	Lung function, functional capacity, energy cost, and nutritional status in cystic fibrosis with and without inspiratory muscle weakness. , 2019, , .		0
63	The Correlation Between Clinical Characteristics and Molecular Genetic Analysis Results of Primary Ciliary Dyskinesia Patients: Hacettepe University Experience. , 2019, , .		0
64	The Effects Of Respiratory Viruses on The Acute Pulmonary Exacerbations In Cystic Fibrosis Patients. , 2019, , .		0
65	Arm functional capacity and activity of daily living in children with primary ciliary dyskinesia and cystic fibrosis. , 2019, , .		0
66	Risk of the developmental delay and behavior problems in preschool-aged primary ciliary dyskinesia patients. , 2019, , .		0
67	Reduced anaerobic and aerobic performance in children with primary ciliary dyskinesia. European Journal of Pediatrics, 2018, 177, 765-773.	1.3	15
68	A review of the etiology and clinical presentation of non-cystic fibrosis bronchiectasis: A tertiary care experience. Respiratory Medicine, 2018, 137, 35-39.	1.3	20
69	DYNC2H1 mutation causes Jeune syndrome and recurrent lung infections associated with ciliopathy. Clinical Respiratory Journal, 2018, 12, 1017-1020.	0.6	7
70	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.	0.9	22
71	Plasma ceramides and sphingomyelins increase in primary ciliary dyskinesia but decrease in cystic fibrosis. , 2018, , .		1
72	Neonatal manifestations in Primary Ciliary Dyskinesia: a multinational cohort study. , 2018, , .		1

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73	Prevalence and Impact of Lung Resection in Primary Ciliary Dyskinesia: a cohort and nested case-control study. , 2018, , .		1
74	Newborn screening for CF: first tree years experience in Hacettepe University. , 2018, , .		0
75	Comparison of Bronchoalveolar Lavage and Sputum Microbiology in Patients with Primary Ciliary Dyskinesia. Pediatric, Allergy, Immunology, and Pulmonology, 2017, 30, 14-17.	0.3	1
76	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.	0.7	6
77	Chronic necrotizing pulmonary aspergillosis in an immunocompetent patient after the surgery of hydatid cyst. Tuberkuloz Ve Toraks, 2017, 65, 157-160.	0.2	2
78	Swallowing dysfunction as a factor that should be remembered in recurrent pneumonia: videofluoroscopic swallow study. Minerva Pediatrics, 2017, 69, 396-402.	0.2	3
79	Clinical features of pseudo-bartter syndrome in cystic fibrosis. , 2017, , .		1
80	Evolution of Primary Ciliary Dyskinesia (PCD) diagnostic testing in Europe. , 2017, , .		3
81	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
82	Postoperative respiratory problems in children with esophageal atresia and tracheoesophageal fistula. , 2017, , .		0
83	Effects of blood transfusion on cytokine profile and pulmonary function in patients with thalassemia major. Clinical Respiratory Journal, 2016, 10, 153-162.	0.6	14
84	Primary Nasal Tuberculosis in a 10-Year-Old Girl. Canadian Journal of Infectious Diseases and Medical Microbiology, 2016, 2016, 1-3.	0.7	3
85	The success of the different eradication therapy regimens for Pseudomonas aeruginosa in cystic fibrosis. Journal of Clinical Pharmacy and Therapeutics, 2016, 41, 419-423.	0.7	16
86	Does <scp>Helicobacter pylori</scp> play a role in the pathogenesis of nonâ€cystic fibrosis bronchiectasis?. Pediatrics International, 2016, 58, 894-898.	0.2	2
87	Congenital Tuberculosis after in-vitro Fertilization in a Woman Previously Undiagnosed with Tuberculosis Salpingitis. Pediatrics and Neonatology, 2016, 57, 539-540.	0.3	12
88	Diagnosis and treatment of pulmonary alveolar microlithiasis. Pediatrics International, 2016, 58, 805-807.	0.2	10
89	PulmonaryMycobacterium abscessusInfection in a Patient with Triple A Syndrome. Journal of Tropical Pediatrics, 2016, 62, 324-327.	0.7	6
90	Omalizumab Treatment for Allergic Bronchopulmonary Aspergillosis in Cystic Fibrosis. Annals of Pharmacotherapy, 2016, 50, 188-193.	0.9	27

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91	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.	0.3	5
92	Long term azithromycin therapy in patients with cystic fibrosis. Turkish Journal of Pediatrics, 2016, 58, 34-40.	0.3	4
93	Pulmonary involvement in children with rheumatic diseases. , 2016, , .		0
94	A ten year experience of non-cystic fibrosis bronchiectasis from a single center. , 2016, , .		0
95	Comparison of peripheral muscle strength, posture and spinal curvatures for patients with cystic fibrosis and bronchiectasis and healthy subjects. , 2016, , .		0
96	Lung function tests in patients with primary ciliary dyskinesia. , 2016, , .		0
97	Presence of Kartagener syndrome reduces exercise capacity in primary ciliary dyskinesia. , 2016, , .		1
98	Comparison of inspiratory muscle training and home-based rehabilitation approach in patients with primary ciliary dyskinesia. , 2016, , .		0
99	Pulmonary and extrapulmonary features in bronchopulmonary dysplasia: a comparison with healthy children. Journal of Physical Therapy Science, 2015, 27, 1761-1765.	0.2	26
100	Pneumomediastinum, pneumorrhachis and subcutaneous emphysema associated with viral infections: Report of three cases. Pediatrics International, 2015, 57, 1038-1040.	0.2	17
101	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.	1.2	47
102	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.	0.3	0
103	Neuroendocrine cell hyperplasia in children with idiopathic interstitial pneumonia. , 2015, , .		0
104	An insight of lung cysts with filamin A mutation. , 2015, , .		1
105	Hacettepe University experience: Clinical evaluation of 218 primary ciliary dyskinesia patients. , 2015, , .		0
106	Flexible bronchoscopy findings in children with protracted bacterial bronchitis. , 2015, , .		0
107	A comparison of aerobic and anaerobic capacity in children with primary ciliary dyskinesia and healthy peers. , 2015, , .		0
108	Radiological findings of patients with primary ciliary dyskinesia. , 2015, , .		0

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109	Successful treatment of pulmonary hemangioma with propranolol. Pediatric Pulmonology, 2014, 49, 829-833.	1.0	8
110	Childhood idiopathic interstitial pneumonia: diagnosis, treatment and follow-up. Turk Pediatri Arsivi, 2013, 48, 281-287.	0.9	0
111	Unfinished battle with childhood tuberculosis: is it curable with less drugs and shorter duration?. Tuberkuloz Ve Toraks, 2013, 61, 320-326.	0.2	1
112	The role of serum Pseudomonas aeruginosa antibodies in the diagnosis and follow-up of cystic fibrosis. Turkish Journal of Pediatrics, 2013, 55, 50-7.	0.3	4
113	Multicentric analysis of childhood tuberculosis in Turkey. Turkish Journal of Pediatrics, 2013, 55, 121-9.	0.3	12
114	Predominance of hospital-associated MRSA among cystic fibrosis patients in a Turkish reference cystic fibrosis centre. Journal of Chemotherapy, 2012, 24, 195-200.	0.7	4
115	Mycobacterial Disease and Impaired IFN-Î³ Immunity in Humans with Inherited ISG15 Deficiency. Science, 2012, 337, 1684-1688.	6.0	455
116	Different features of lung involvement in Niemann-Pick disease and Gaucher disease. Respiratory Medicine, 2012, 106, 1278-1285.	1.3	45
117	Effects of chest physiotherapy and aerobic exercise training on physical fitness in young children with cystic fibrosis. Italian Journal of Pediatrics, 2012, 38, 2.	1.0	29
118	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
119	Mekonyum ileusunun 12 ay altÄ±nda tanÄ± alan kistik fibrozlu hastalarda klinik seyire etkisi. Turk Pediatri Arsivi, 2011, 45, 105-110.	0.9	0
120	Preventing tuberculosis in children receiving anti-tnf treatment. Clinical Rheumatology, 2010, 29, 389-392.	1.0	20
121	The role of human leucocyte antigens in children with hydatid disease: their association with clinical condition and prognosis. Parasitology Research, 2010, 106, 795-800.	0.6	16
122	Longâ€term results of disodium etidronate treatment in pulmonary alveolar microlithiasis. Pediatric Pulmonology, 2010, 45, 514-517.	1.0	52
123	Eosinophilic Lung Diseases: Five Cases In Childhood. , 2010, , .		0
124	TAP1 and TAP2 gene polymorphisms in childhood cystic echinococcosis. Parasitology International, 2010, 59, 283-285.	0.6	14
125	Does Defective Apoptosis Play A Role in Cystic Fibrosis Lung Disease?. Archives of Medical Research, 2009, 40, 561-564.	1.5	8
126	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.	1.3	4

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127	Gastric organo-axial malrotation coexisting respiratory symptoms. European Journal of Pediatrics, 2009, 168, 491-494.	1.3	9
128	A 4-month-old boy with acrodermatitis enteropathica-like symptoms. European Journal of Pediatrics, 2009, 168, 119-121.	1.3	7
129	Horseshoe lung associated with left-lung hypoplasia, left pulmonary artery sling and bilateral agenesis of upper lobe bronchi. Pediatric Radiology, 2009, 39, 1002-1005.	1.1	16
130	Gorhamâ€“Stout Syndrome with chylothorax: Successful remission by interferon alphaâ€2b. Pediatric Pulmonology, 2009, 44, 613-615.	1.0	47
131	Relation of bone mineral density with clinical and laboratory parameters in preâ€pubertal children with cystic fibrosis. Pediatric Pulmonology, 2009, 44, 706-712.	1.0	16
132	Successful unilateral partial lung lavage in a child with pulmonary alveolar proteinosis. Journal of Clinical Anesthesia, 2009, 21, 127-130.	0.7	15
133	Dollâ€like face: Is it an underestimated clinical presentation of cystic fibrosis?. Pediatric Pulmonology, 2008, 43, 634-637.	1.0	5
134	Comparason of extraction methods for PCR detection of Burkholderia cepacia complex (BCC) from cystic fibrosis patients. Open Medicine (Poland), 2008, 3, 157-162.	0.6	0
135	Pleural Fluid PCR Method for Detection of Staphylococcus aureus, Streptococcus pneumoniae and Haemophilus influenzae in Pediatric Parapneumonic Effusions. Respiration, 2008, 75, 437-442.	1.2	14
136	Environmental Tobacco Smoke Exposure and Respiratory Morbidity in Children. Inhalation Toxicology, 2007, 19, 779-785.	0.8	12
137	The role of TAP1 and TAP2 gene polymorphism in idiopathic bronchiectasis in children. Pediatric Pulmonology, 2007, 42, 237-241.	1.0	18
138	Tuberculin skin test positivity in pediatric allogeneic BMT recipients and donors in Turkey. Pediatric Transplantation, 2007, 11, 414-418.	0.5	11
139	An epidemic of pseudo-Bartter syndrome in cystic fibrosis patients. European Journal of Pediatrics, 2007, 167, 115-116.	1.3	22
140	Mutations in SLC34A2 Cause Pulmonary Alveolar Microlithiasis and Are Possibly Associated with Testicular Microlithiasis. American Journal of Human Genetics, 2006, 79, 650-656.	2.6	226
141	Airway Malacia Disorders in Children. Chest, 2006, 130, 304.	0.4	1
142	Tracheobronchopathia Osteochondroplastica in a 9-year-old Girl. Pediatric Pulmonology, 2006, 41, 95-97.	1.0	32
143	Prenatal period to adolescence: the variable presentations of congenital cystic adenomatoid malformation. Pediatrics International, 2006, 48, 626-630.	0.2	6
144	Childhood Parapneumonic Effusions. Chest, 2005, 128, 1436-1441.	0.4	22

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145	Xanthoma disseminatum: A child with respiratory system involvement and bronchiectasis. Pediatric Pulmonology, 2005, 39, 84-87.	1.0	18
146	Foreign-body aspiration mimicking congenital lobar emphysema in a forty-eight-day-old girl. Pediatric Pulmonology, 2005, 39, 189-191.	1.0	7
147	Childhood diffuse panbronchiolitis: A case report. Pediatric Pulmonology, 2005, 40, 354-357.	1.0	20
148	Clinical features and treatment approaches in cystic fibrosis with pseudo-Bartter syndrome. Annals of Tropical Paediatrics, 2005, 25, 119-124.	1.0	39
149	Tracheomalacia and Bronchomalacia in 34 Children: Clinical and Radiologic Profiles and Associations with Other Diseases. Clinical Pediatrics, 2005, 44, 777-781.	0.4	37
150	Bronchiectasis: the Consequence of Late Diagnosis in Chronic Respiratory Symptoms. Journal of Tropical Pediatrics, 2005, 51, 362-365.	0.7	39
151	Medical treatment of pulmonary hydatid disease: for which child?. Parasitology International, 2005, 54, 135-138.	0.6	57
152	BALF Nitrite as an Indicator of Inflammation in Children with Cystic Fibrosis. Respiration, 2004, 71, 625-629.	1.2	14
153	Hypersensitivity pneumonitis in children: pigeon breeders disease. Annals of Tropical Paediatrics, 2004, 24, 349-355.	1.0	15
154	An unusual pattern of arthritis in a child with Kawasaki syndrome. Clinical Rheumatology, 2004, 23, 73-75.	1.0	25
155	Surgical Management of Infants with Congenital Lobar Emphysema and Concomitant Congenital Heart Disease. Heart Surgery Forum, 2004, 7, E644-E649.	0.2	16
156	Pigeon-breeder's disease in a child with selective IgA deficiency. Pediatrics International, 2003, 45, 216-218.	0.2	8
157	Congenital lobar emphysema: Evaluation and long-term follow-up of thirty cases at a single center. Pediatric Pulmonology, 2003, 35, 384-391.	1.0	101
158	Postinfectious Bronchiolitis obliterans in Children: Clinical and Radiological Profile and Prognostic Factors. Respiration, 2003, 70, 371-375.	1.2	64
159	Cardiac hydatid cyst and pulmonary hydatidosis in a child. Pediatric Infectious Disease Journal, 2002, 21, 1178-1180.	1.1	8
160	Hepatobiliary manifestations of cystic fibrosis in children: correlation of CT and US findings. European Journal of Radiology, 2002, 41, 26-33.	1.2	28
161	Pediatric Primary Pulmonary Tuberculosis. Chest, 2002, 121, 1722.	0.4	2
162	Treatment and follow-up of pulmonary alveolar microlithiasis with disodium editronate: radiological demonstration. Pediatric Radiology, 2002, 32, 380-383.	1.1	20

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163	High resolution CT in children with cystic fibrosis: correlation with pulmonary functions and radiographic scores. European Journal of Radiology, 2001, 37, 54-59.	1.2	68
164	Treatment of Hydatid Disease. Paediatric Drugs, 2001, 3, 123-135.	1.3	68
165	Homozygous and Compound-heterozygous Type I Plasminogen Deficiency Is a Common Cause of Ligneous Conjunctivitis. Thrombosis and Haemostasis, 2001, 85, 1004-1010.	1.8	56
166	Chemical pneumonia caused by glutaraldehyde. Pediatrics International, 2001, 43, 701-702.	0.2	4
167	Pulmonary involvement in a child with ligneous conjunctivitis and homozygous type I plasminogen deficiency. Pediatric Pulmonology, 2001, 32, 179-183.	1.0	19
168	Neutrophil chemotaxis in acutely infected and clinically stable cystic fibrosis patients*. Pediatrics International, 1999, 41, 514-518.	0.2	9
169	Hydatid disease in childhood: A retrospective analysis of 376 cases. , 1998, 26, 190-196.		46
170	Auditory nerve-brainstem responses in cystic fibrosis patients. International Journal of Pediatric Otorhinolaryngology, 1996, 35, 165-169.	0.4	10
171	Clinical status, ocular surface changes and tear ferning in patients with cystic fibrosis. Acta Ophthalmologica, 1996, 74, 563-565.	0.4	8
172	Sodium chloride deficiency in cystic fibrosis patients. European Journal of Pediatrics, 1994, 153, 829-831.	1.3	31
173	Conservative Treatment of Empyema in Children. Respiration, 1993, 60, 182-185.	1.2	50