Deniz Dogru Ersoz

List of Publications by Year
in descending order

136	2,003	346980	406436
papers	citations	22 h-index	35 g-index

141	141	141	2362
all docs	docs citations	times ranked	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	Increased Plasma YKL-40 Level and Chitotriosidase Activity in Cystic Fibrosis Patients. Inflammation, 2022, 45, 627-638.	1.7	0
3	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
4	Cystic fibrosis newborn screening: Fiveâ€year experience from a tertiary care center. Pediatric Pulmonology, 2022, 57, 403-410.	1.0	3
5	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
6	The controversy of drug hypersensitivity in patients with cystic fibrosis and review of the literature. Pediatric Allergy and Immunology, 2022, 33, .	1.1	5
7	The success of the Cystic Fibrosis Registry of Turkey for improvement of patient care. Pediatric Pulmonology, 2022, , .	1.0	4
8	Impact of Achromobacter spp. isolation on clinical outcomes in children with cystic fibrosis. Pediatric Pulmonology, 2022, 57, 658-666.	1.0	5
9	Effectiveness of different eradication treatment protocols for newâ€onset Pseudomonas aeruginosa in children with cystic fibrosis. Pediatric Pulmonology, 2022, , .	1.0	1
10	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
11	Clinical radiological and pathological staging of children with hypersensitivity pneumonitis. Pediatric Pulmonology, 2022, 57, 2344-2355.	1.0	2
12	Risk factors for recurrent pulmonary exacerbation in idiopathic pulmonary hemosiderosis. Pediatric Pulmonology, 2021, 56, 1060-1068.	1.0	10
13	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
14	Nonmyeloablative hematopoietic stem cell transplantation in a patient with hereditary pulmonary alveolar proteinosis. Pediatric Pulmonology, 2021, 56, 341-343.	1.0	7
15	The frequency and related factors of nonâ€tuberculosis mycobacteria infections among patients with cystic fibrosis. Pediatrics International, 2021, 63, 1369-1375.	0.2	2
16	Does cystic fibrosis make susceptible to celiac disease?. European Journal of Pediatrics, 2021, 180, 2807-2813.	1.3	8
17	Evaluation of pulmonary complications and affecting factors in children for repaired esophageal atresia and tracheoesophageal fistula. Respiratory Medicine, 2021, 181, 106376.	1.3	5
18	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

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19	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
20	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
21	Impact of COVIDâ€19 on pediatric pulmonology healthcare practice. Pediatric Pulmonology, 2021, 56, 2811-2817.	1.0	5
22	Respiratory system findings in pediatric patients with primary immunodeficiency. Pediatric Pulmonology, 2021, 56, 4011-4019.	1.0	2
23	Clinical spectrum of children with interstitial pneumonia with autoimmune features. Respiratory Medicine, 2021, 187, 106566.	1.3	3
24	Sleep disordered breathing in patients with Prader willi syndrome: Impact of underlying genetic mechanism. Respiratory Medicine, 2021, 187, 106567.	1.3	3
25	Differentially expressed genes associated with disease severity in siblings with cystic fibrosis. Pediatric Pulmonology, 2021, 56, 910-920.	1.0	4
26	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
27	Clinical characteristics of children with cystic fibrosis infected with unusual bacteria. Minerva Pediatrics, 2021, , .	0.2	1
28	The relationship between oxidative stress markers in exhaled breath condensate and respiratory problems in patients with repaired esophageal atresia. Journal of Pediatric Surgery, 2020, 55, 1516-1521.	0.8	4
29	Cystic fibrosis in Turkey: First data from the national registry. Pediatric Pulmonology, 2020, 55, 541-548.	1.0	27
30	Childhood diffuse parenchymal lung diseases: We need a new classification. Clinical Respiratory Journal, 2020, 14, 102-108.	0.6	6
31	Genotype and phenotype evaluation of patients with primary ciliary dyskinesia: First results from Turkey. Pediatric Pulmonology, 2020, 55, 383-393.	1.0	46
32	Cystic fibrosis in Turkey. Lancet Respiratory Medicine,the, 2020, 8, e17.	5.2	2
33	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
34	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
35	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
36	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

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37	Clinical features and accompanying findings of Pseudoâ€Bartter Syndrome in cystic fibrosis. Pediatric Pulmonology, 2020, 55, 2011-2016.	1.0	12
38	Patients eligible for modulator drugs: Data from cystic fibrosis registry of Turkey. Pediatric Pulmonology, 2020, 55, 2302-2306.	1.0	5
39	Visuomotor reaction time and dynamic balance in children with cystic fibrosis and nonâ€cystic fibrosis bronchiectasis: A caseâ€control study. Pediatric Pulmonology, 2020, 55, 2341-2347.	1.0	9
40	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
41	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
42	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
43	From Diagnosis to Treatment of Pediatric Tuberculosis: Ten Years Experience in a Single Institution. Clinical Pediatrics, 2020, 59, 476-482.	0.4	4
44	Respiratory viruses: What is their role in acute exacerbations in children with cystic fibrosis?. Pediatric Pulmonology, 2020, 55, 1646-1652.	1.0	5
45	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
46	Bilateral Pulmonary Langerhans's Cell Histiocytosis is Surgical Challenge in Children: A Case Report. European Journal of Pediatric Surgery Reports, 2019, 07, e8-e11.	0.1	4
47	Developmental and behavioral problems in preschool-aged primary ciliary dyskinesia patients. European Journal of Pediatrics, 2019, 178, 995-1003.	1.3	12
48	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
49	Nasal nitric oxide levels in primary ciliary dyskinesia, cystic fibrosis and healthy children. Turkish Journal of Pediatrics, 2019, 61, 20.	0.3	6
50	Pulmonary complications following hematopoietic stem cell transplantation in children. Turkish Journal of Pediatrics, 2019, 61, 59.	0.3	8
51	Bk virus associated nephropathy and severe pneumonia in a kidney transplanted adolescent with schimke immuneosseous- dysplasia. Turkish Journal of Pediatrics, 2019, 61, 111.	0.3	1
52	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
53	Psychiatric morbidity and quality of life in children and adolescents with cystic fibrosis. Turkish Journal of Pediatrics, 2018, 60, 32.	0.3	10
54	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

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55	Genotypicâ€phenotypic features and enzyme replacement therapy outcome in patients with mucopolysaccharidosis VI from Turkey. American Journal of Medical Genetics, Part A, 2017, 173, 2954-2967.	0.7	17
56	Chronic necrotizing pulmonary aspergillosis in an immunocompetent patient after the surgery of hydatid cyst. Tuberkuloz Ve Toraks, 2017, 65, 157-160.	0.2	2
57	Flexible Bronchoscopy Findings in Patient with Protracted Bacterial Bronchitis. Turkiye Klinikleri Pediatri, 2017, 26, 39-43.	0.1	0
58	Flow-volume curve in the diagnosis and follow-up of intrathoracic airway obstruction. Turkish Journal of Pediatrics, 2017, 59, 594-597.	0.3	0
59	Swallowing dysfunction as a factor that should be remembered in recurrent pneumonia: videofluoroscopic swallow study. Minerva Pediatrics, 2017, 69, 396-402.	0.2	3
60	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
61	A Global Survey on Whole Lung Lavage in Pulmonary Alveolar Proteinosis. Chest, 2016, 150, 251-253.	0.4	20
62	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
63	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
64	Congenital Tuberculosis after in-vitro Fertilization in a Woman Previously Undiagnosed with Tuberculosis Salpingitis. Pediatrics and Neonatology, 2016, 57, 539-540.	0.3	12
65	Use of serial rigid bronchoscopy in the treatment of plastic bronchitis in children. Journal of Pediatric Surgery, 2016, 51, 1640-1643.	0.8	31
66	Diagnosis and treatment of pulmonary alveolar microlithiasis. Pediatrics International, 2016, 58, 805-807.	0.2	10
67	PulmonaryMycobacterium abscessusInfection in a Patient with Triple A Syndrome. Journal of Tropical Pediatrics, 2016, 62, 324-327.	0.7	6
68	Omalizumab Treatment for Allergic Bronchopulmonary Aspergillosis in Cystic Fibrosis. Annals of Pharmacotherapy, 2016, 50, 188-193.	0.9	27
69	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
70	Long term azithromycin therapy in patients with cystic fibrosis. Turkish Journal of Pediatrics, 2016, 58, 34-40.	0.3	4
71	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
72	Successful treatment of pulmonary hemangioma with propranolol. Pediatric Pulmonology, 2014, 49, 829-833.	1.0	8

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73	Cytomegalovirus infection in immunocompetent wheezy infants: the diagnostic value of CMV PCR in bronchoalveolar lavage fluid. Journal of Clinical Pharmacy and Therapeutics, 2014, 39, 399-403.	0.7	12
74	Reliability and validity of the Cystic Fibrosis Questionnaire-Revised for children and parents in Turkey: cross-sectional study. Quality of Life Research, 2013, 22, 409-414.	1.5	15
75	Prevalence and genetic diversity of Staphylococcus aureus small-colony variants in cystic fibrosis patients. Clinical Microbiology and Infection, 2013, 19, 77-84.	2.8	49
76	Childhood idiopathic interstitial pneumonia: diagnosis, treatment and follow-up. Turk Pediatri Arsivi, 2013, 48, 281-287.	0.9	0
77	Unfinished battle with childhood tuberculosis: is it curable with less drugs and shorter duration?. Tuberkuloz Ve Toraks, 2013, 61, 320-326.	0.2	1
78	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
79	Multicentric analysis of childhood tuberculosis in Turkey. Turkish Journal of Pediatrics, 2013, 55, 121-9.	0.3	12
80	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
81	Different features of lung involvement in Niemann-Pick disease and Gaucher disease. Respiratory Medicine, 2012, 106, 1278-1285.	1.3	45
82	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
83	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
84	Endobronchial, Pulmonary and Liver Leiomyomata in a Child with Primary Immune Deficiency. European Journal of Pediatric Surgery, 2010, 20, 423-425.	0.7	6
85	TAP1 and TAP2 gene polymorphisms in childhood cystic echinococcosis. Parasitology International, 2010, 59, 283-285.	0.6	14
86	Tuberculosis in children with congenital immunodeficiency syndromes. Tuberkuloz Ve Toraks, 2010, 58, 59-63.	0.2	10
87	Does Defective Apoptosis Play A Role in Cystic Fibrosis Lung Disease?. Archives of Medical Research, 2009, 40, 561-564.	1.5	8
88	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
89	Gastric organo-axial malrotation coexisting respiratory symptoms. European Journal of Pediatrics, 2009, 168, 491-494.	1.3	9
90	A 4-month-old boy with acrodermatitis enteropathica-like symptoms. European Journal of Pediatrics, 2009, 168, 119-121.	1.3	7

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91	Gorhamâ€“Stout Syndrome with chylothorax: Successful remission by interferon alphaâ€2b. Pediatric Pulmonology, 2009, 44, 613-615.	1.0	47
92	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
93	Successful unilateral partial lung lavage in a child with pulmonary alveolar proteinosis. Journal of Clinical Anesthesia, 2009, 21, 127-130.	0.7	15
94	Effects of scoliosis on respiratory muscle strength in patients with neuromuscular disorders. Spine Journal, 2009, 9, 981-986.	0.6	37
95	Achalasia-like findings in a case with delayed diagnosis of H-type tracheoesophageal fistula. Pediatric Surgery International, 2008, 24, 965-969.	0.6	8
96	Childhood intrathoracic Hodgkin lymphoma with hypertrophic pulmonary osteoarthropathy: a case report and review of the literature. European Journal of Pediatrics, 2008, 167, 419-423.	1.3	11
97	Dollâ€like face: Is it an underestimated clinical presentation of cystic fibrosis?. Pediatric Pulmonology, 2008, 43, 634-637.	1.0	5
98	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
99	Environmental Tobacco Smoke Exposure and Respiratory Morbidity in Children. Inhalation Toxicology, 2007, 19, 779-785.	0.8	12
100	The role of TAP1 and TAP2 gene polymorphism in idiopathic bronchiectasis in children. Pediatric Pulmonology, 2007, 42, 237-241.	1.0	18
101	An epidemic of pseudo-Bartter syndrome in cystic fibrosis patients. European Journal of Pediatrics, 2007, 167, 115-116.	1.3	22
102	Airway Malacia Disorders in Children. Chest, 2006, 130, 304.	0.4	1
103	Tracheobronchopathia Osteochondroplastica in a 9-year-old Girl. Pediatric Pulmonology, 2006, 41, 95-97.	1.0	32
104	Effects of claritromycin on inflammatory parameters and clinical conditions in children with bronchiectasis1. Journal of Clinical Pharmacy and Therapeutics, 2006, 31, 49-55.	0.7	101
105	Prenatal period to adolescence: the variable presentations of congenital cystic adenomatoid malformation. Pediatrics International, 2006, 48, 626-630.	0.2	6
106	Childhood Parapneumonic Effusions. Chest, 2005, 128, 1436-1441.	0.4	22
107	Xanthoma disseminatum: A child with respiratory system involvement and bronchiectasis. Pediatric Pulmonology, 2005, 39, 84-87.	1.0	18
108	Childhood diffuse panbronchiolitis: A case report. Pediatric Pulmonology, 2005, 40, 354-357.	1.0	20

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109	Clinical features and treatment approaches in cystic fibrosis with pseudo-Bartter syndrome. Annals of Tropical Paediatrics, 2005, 25, 119-124.	1.0	39
110	Tracheomalacia and Bronchomalacia in 34 Children: Clinical and Radiologic Profiles and Associations with Other Diseases. Clinical Pediatrics, 2005, 44, 777-781.	0.4	37
111	Neonatal Hyperparathyroidism Due to Maternal Hypoparathyroidism and Vitamin D Deficiency: A Cause of Multiple Bone Fractures. Clinical Pediatrics, 2005, 44, 267-269.	0.4	22
112	Congenital Hydrocephalus as a Rare Association with Ligneous Conjunctivitis and Type I Plasminogen Deficiency. Neuropediatrics, 2005, 36, 108-111.	0.3	19
113	Bronchiectasis: the Consequence of Late Diagnosis in Chronic Respiratory Symptoms. Journal of Tropical Pediatrics, 2005, 51, 362-365.	0.7	39
114	Medical treatment of pulmonary hydatid disease: for which child?. Parasitology International, 2005, 54, 135-138.	0.6	57
115	Tracheoesophageal Fistula Due to Disc-Battery Ingestion. European Journal of Pediatric Surgery, 2004, 14, 274-278.	0.7	28
116	Yellow Nail Syndrome in an Infant Presenting with Lymphedema of the Eyelids and Pleural Effusion. Clinical Pediatrics, 2004, 43, 569-572.	0.4	13
117	BALF Nitrite as an Indicator of Inflammation in Children with Cystic Fibrosis. Respiration, 2004, 71, 625-629.	1.2	14
118	Hypersensitivity pneumonitis in children: pigeon breeders disease. Annals of Tropical Paediatrics, 2004, 24, 349-355.	1.0	15
119	Spontaneous Neonatal Chylothorax Treated with Octreotide in Turkey: A Case Report. Journal of Perinatology, 2004, 24, 261-262.	0.9	24
120	Pigeon-breeder's disease in a child with selective IgA deficiency. Pediatrics International, 2003, 45, 216-218.	0.2	8
121	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
122	Postinfectious Bronchiolitis obliterans in Children: Clinical and Radiological Profile and Prognostic Factors. Respiration, 2003, 70, 371-375.	1.2	64
123	Use of recombinant human DNase in a premature infant with recurrent atelectasis. Pediatrics International, 2003, 45, 584-586.	0.2	10
124	Cardiac hydatid cyst and pulmonary hydatidosis in a child. Pediatric Infectious Disease Journal, 2002, 21, 1178-1180.	1.1	8
125	Pediatric Primary Pulmonary Tuberculosis. Chest, 2002, 121, 1722.	0.4	2
126	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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127	Hodgkin's disease and ataxia telangiectasia with pulmonary cavities. Pediatric Pulmonology, 2002, 33, 399-403.	1.0	25
128	Antioxidant effect of beta-carotene in cystic fibrosis and bronchiectasis: clinical and laboratory parameters of a pilot study. Acta Paediatrica, International Journal of Paediatrics, 2002, 91, 793-8.	0.7	8
129	Treatment of Hydatid Disease. Paediatric Drugs, 2001, 3, 123-135.	1.3	68
130	Chemical pneumonia caused by glutaraldehyde. Pediatrics International, 2001, 43, 701-702.	0.2	4
131	Pulmonary involvement in a child with ligneous conjunctivitis and homozygous type I plasminogen deficiency. Pediatric Pulmonology, 2001, 32, 179-183.	1.0	19
132	A Case of Tracheobronchomegaly (MounierÂ–Kuhn Syndrome) Diagnosed Via Flexible Bronchoscopy. Journal of Bronchology, 2001, 8, 190-192.	0.2	0
133	Phenylketonuria and cystic fibrosis in the same patient. Pediatrics International, 2000, 42, 92-93.	0.2	8
134	Triple A Syndrome Mimicking Cystic Fibrosis. Journal of Pediatric Endocrinology and Metabolism, 2000, 13, 329-31.	0.4	3
135	Long-term clinical course of patients with idiopathic pulmonary hemosiderosis (1979-1994): Prolonged survival with low-dose corticosteroid therapy. , 1999, 27, 180-184.		107
136	Hydatid disease in childhood: A retrospective analysis of 376 cases. , 1998, 26, 190-196.		46