Nilgun Cakar

List of Publications by Year in descending order

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#	Article	lF	CITATIONS
1	A new set of criteria for the diagnosis of familial Mediterranean fever in childhood. Rheumatology, 2009, 48, 395-398.	0.9	374
2	Spectrum of Steroid-Resistant and Congenital Nephrotic Syndrome in Children. Clinical Journal of the American Society of Nephrology: CJASN, 2015, 10, 592-600.	2.2	225
3	Genotype–phenotype correlation in a large group of Turkish patients with familial Mediterranean fever: evidence for mutation-independent amyloidosis. Rheumatology, 2000, 39, 67-72.	0.9	149
4	Takayasu arteritis in children. Journal of Rheumatology, 2008, 35, 913-9.	1.0	89
5	MEFV mutations in Turkish patients suffering from familial Mediterranean fever. Human Mutation, 2000, 15, 118-119.	1.1	78
6	Reno-vascular hypertension in childhood: a nationwide survey. Pediatric Nephrology, 2007, 22, 1327-1333.	0.9	63
7	<i>MEFV</i> Mutations Modify the Clinical Presentation of Henoch-Schönlein Purpura. Journal of Rheumatology, 2008, 35, 2427-2429.	1.0	62
8	NPHS2 (podicin) mutations in Turkish children with idiopathic nephrotic syndrome. Pediatric Nephrology, 2007, 22, 2031-2040.	0.9	53
9	Familial Mediterranean fever - renal involvement by diseases other than amyloid. Nephrology Dialysis Transplantation, 1999, 14, 475-479.	0.4	46
10	Arthritis in children with familial Mediterranean fever. Rheumatology International, 2002, 21, 213-217.	1.5	44
11	Interleukin-1 targeting treatment in familial Mediterranean fever: an experience of pediatric patients. Modern Rheumatology, 2015, 25, 621-624.	0.9	44
12	Low levels of urinary epidermal growth factorÂpredict chronic kidney disease progressionÂin children. Kidney International, 2019, 96, 214-221.	2.6	43
13	The value of the levels of acute phase reactants for the prediction of familial Mediterranean fever associated amyloidosis: a case control study. Rheumatology International, 2007, 27, 517-522.	1.5	38
14	Serum amyloid A1 and tumor necrosis factor-alpha alleles in Turkish Familial Mediterranean Fever patients with and without amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2003, 10, 12-16.	1.4	36
15	Eculizumab therapy in a child with hemolytic uremic syndrome and CFI mutation. Pediatric Nephrology, 2012, 27, 2327-2331.	0.9	35
16	Markers of Bone Metabolism Are Affected by Renal Function and Growth Hormone Therapy in Children with Chronic Kidney Disease. PLoS ONE, 2015, 10, e0113482.	1.1	33
17	MEFV mutations in multiplex families with familial Mediterranean fever: is a particular genotype necessary for amyloidosis?. Clinical Genetics, 2000, 57, 430-434.	1.0	29
18	Decrease in the rate of secondary amyloidosis in Turkish children with FMF: are we doing better?. European Journal of Pediatrics, 2010, 169, 971-974.	1.3	27

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19	Indoxyl sulfate associates with cardiovascular phenotype in children with chronic kidney disease. Pediatric Nephrology, 2019, 34, 2571-2582.	0.9	27
20	Application of the new pediatric criteria and Tel Hashomer criteria in heterozygous patients with clinical features of FMF. European Journal of Pediatrics, 2011, 170, 1055-1057.	1.3	26
21	Cardiac involvement in childhood polyarteritis nodosa. International Journal of Cardiology, 1997, 60, 257-262.	0.8	25
22	Low-dose erythropoietin is effective and safe in children on continuous ambulatory peritoneal dialysis. Pediatric Nephrology, 1997, 11, 350-352.	0.9	25
23	Keratoconjunctivitis Sicca in Juvenile Rheumatoid Arthritis. Cornea, 2007, 26, 941-944.	0.9	24
24	Global left-ventricular function by tissue Doppler imaging in pediatric dialysis patients. Pediatric Nephrology, 2008, 23, 779-785.	0.9	24
25	The use of low-dose cyclophosphamide followed by AZA/MMF treatment in childhood lupus nephritis. Pediatric Nephrology, 2010, 25, 111-117.	0.9	24
26	The coexistence of familial Mediterranean fever and polyarteritis nodosa; report of a case. Pediatric Nephrology, 1996, 10, 631-633.	0.9	23
27	Familial Mediterranean Fever and Acute Rheumatic Fever: A Pathogenetic Relationship?. Clinical Rheumatology, 1999, 18, 446-449.	1.0	23
28	Primary nephrotic syndrome during childhood in Turkey. Pediatrics International, 2004, 46, 436-438.	0.2	23
29	Analysis of NPHS2 mutations in Turkish steroid-resistant nephrotic syndrome patients. Pediatric Nephrology, 2006, 21, 1093-1096.	0.9	23
30	Effects of nutritional vitamin D supplementation on markers of bone and mineral metabolism in children with chronic kidney disease. Nephrology Dialysis Transplantation, 2018, 33, 2208-2217.	0.4	23
31	Predictors of left ventricular hypertrophy in children on chronic peritoneal dialysis. Pediatric Nephrology, 2010, 25, 1311-1318.	0.9	21
32	Clinical features and disease severity of Turkish FMF children carrying E148Q mutation. Journal of Clinical Laboratory Analysis, 2019, 33, e22852.	0.9	21
33	Sacroiliitis in Children With Familial Mediterranean Fever. Journal of Clinical Rheumatology, 2019, 25, 69-73.	0.5	18
34	Congenital microcephaly and infantile nephrotic syndrome ? a case report. Pediatric Nephrology, 1994, 8, 72-73.	0.9	15
35	Hypertension and Left Ventricular Hypertrophy in Pediatric Peritoneal Dialysis Patients: Ambulatory Blood Pressure Monitoring and Echocardiographic Evaluation. Nephron Clinical Practice, 2006, 104, c101-c106.	2.3	15
36	Assessment of neutrophil to lymphocyte ratio and mean platelet volume in pediatric familial Mediterranean fever patients. Journal of Research in Medical Sciences, 2017, 22, 35.	0.4	14

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37	Renal Involvement in Childhood Vasculitis. Nephron Clinical Practice, 2008, 108, c202-c206.	2.3	13
38	Ocular Findings in Children With Chronic Renal Failure. Cornea, 2009, 28, 5-6.	0.9	13
39	C3 glomerulopathy in NLRP12-related autoinflammatory disorder: case-based review. Rheumatology International, 2018, 38, 1571-1576.	1.5	13
40	The assessment of time-dependent myocardial changes in infants with perinatal hypoxia. Journal of Maternal-Fetal and Neonatal Medicine, 2012, 25, 1564-1568.	0.7	12
41	The effect of colchicine and disease severity on physical growth in children with familial Mediterranean fever. Clinical Rheumatology, 2016, 35, 1603-1607.	1.0	12
42	Hereditary renal tubular disorders in Turkey: demographic, clinical, and laboratory features. Clinical and Experimental Nephrology, 2011, 15, 108-113.	0.7	11
43	Wegener granulomatosis as an uncommon cause of panhypopituitarism in childhood. Journal of Pediatric Endocrinology and Metabolism, 2013, 26, 959-62.	0.4	11
44	Plasma exchange therapy for severe gastrointestinal involvement of Henoch Schonlein purpura in children. Pediatric Rheumatology, 2014, 12, P357.	0.9	11
45	MEFV mutation analysis in Turkish familial Mediterranean fever patients with amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 1999, 6, 301-302.	1.4	10
46	ATP6V1B1mutations in distal renal tubular acidosis and sensorineural hearing loss: clinical and genetic spectrum of five families. Renal Failure, 2013, 35, 1281-1284.	0.8	10
47	<i>HPSE2</i> Mutations in Urofacial Syndrome, Non-Neurogenic Neurogenic Bladder and Lower Urinary Tract Dysfunction. Nephron, 2015, 130, 54-58.	0.9	10
48	The association between obesity, hypertension and left ventricular mass in adolescents. Journal of Pediatric Endocrinology and Metabolism, 2017, 30, 167-174.	0.4	10
49	Atypical Hemolytic Uremic Syndrome in Children Aged <2 Years. Nephron, 2018, 139, 211-218.	0.9	10
50	Right nutcracker syndrome associated with left-sided inferior vena cava, hemiazygos continuation and persistant left superior vena cava: a rare combination. Clinical Imaging, 2014, 38, 340-345.	0.8	9
51	Transplantation within the era of anti-IL-1 therapy: case series of five patients with familial Mediterranean fever-related amyloidosis. Transplant International, 2018, 31, 1181-1184.	0.8	9
52	Familial Mediterranean fever associated with optic neuritis, successfully treated with anti-interleukin 1 agents. Turkish Journal of Pediatrics, 2016, 58, 327-330.	0.3	8
53	The expanded spectrum of arthritis in children with familial Mediterranean fever. Clinical Rheumatology, 2022, 41, 1535-1541.	1.0	8
54	Paediatric membranoproliferative glomerulonephritis is not decreasing in Turkey!. Pediatric Nephrology, 1994, 8, 131-132.	0.9	7

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55	A large intracardiac thrombus in a child with steroid-resistant nephrotic syndrome. Cardiology in the Young, 2013, 23, 440-442.	0.4	7
56	Cytomegalovirus-related hemorrhagic cystitis in an immunocompetent child. Renal Failure, 2014, 36, 1148-1150.	0.8	6
57	Clinico-radiological correlation of nutcracker syndrome: a single centre experience. Archivos Argentinos De Pediatria, 2017, 115, 165-168.	0.3	6
58	Effects of whole blood viscosity and plasma NOx on cardiac function and cerebral blood flow in children with chronic kidney disease. Turkish Journal of Medical Sciences, 2017, 47, 1482-1491.	0.4	6
59	Paracetamol-induced fixed drug eruption in a patient with recurrent fever and rash. Annals of Allergy, Asthma and Immunology, 2016, 117, 210-211.	0.5	5
60	Neonatal onset familial Mediterranean fever. Modern Rheumatology, 2019, 29, 647-650.	0.9	5
61	The effect of genotype on musculoskeletal complaints in patients with familial Mediterranean fever. Postgraduate Medicine, 2020, 132, 220-224.	0.9	5
62	Clinical features and outcomes of childhood polyarteritis nodosa: A single referral center experience. Modern Rheumatology, 2021, 31, 1142-1147.	0.9	5
63	HLAâ€system and the frequency of relapses in childhood minimal change nephrotic syndrome in Turkish children. Pediatrics International, 1995, 37, 419-421.	0.2	4
64	Hemophagocytic lymphohistiocytosis secondary to <scp>V</scp> aricella zoster infection in a child with <scp>H</scp> enoch– <scp>S</scp> chönlein purpura. Pediatrics International, 2015, 57, e37-8.	0.2	4
65	Coexistence of systemic lupus erythematosus and familial Mediterranean fever in a pediatric patient. Lupus, 2016, 25, 1062-1063.	0.8	4
66	Fatigue in pediatric patients with familial Mediterranean fever. Modern Rheumatology, 2018, 28, 1016-1020.	0.9	4
67	Proteinuria in pediatric renal transplant recipients. Pediatric Transplantation, 2018, 22, e13068.	0.5	4
68	Prediction of inactive disease and relapse in oligoarticular juvenile idiopathic arthritis. Modern Rheumatology, 2021, 31, 1025-1030.	0.9	4
69	The Effectiveness of Sirolimus Treatment in Two Rare Disorders with Nonketotic Hypoinsulinemic Hypoglycemia: The Role of mTOR Pathway. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 439-443.	0.4	4
70	Heart rate variability in juvenile systemic lupus erythematosus patients. Turkish Journal of Pediatrics, 2019, 61, 733.	0.3	4
71	Occurrence of Wilms Tumor in a Child with Urofacial (OCHOA) Syndrome. Pediatric Hematology and Oncology, 2011, 28, 616-618.	0.3	3
72	Evaluation of Childhood Health Assessment Questionnaire in Juvenile Idiopathic Arthritis: A Single Center Experience From Turkey. Archives of Rheumatology, 2015, 30, 57-62.	0.3	3

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73	Hemorragia pulmonar, una complicación poco frecuente en una niña con púrpura de Schönlein-Henoch. Archivos Argentinos De Pediatria, 2016, 114, e366-9.	0.3	3
74	Successful methylprednisolone desensitization in a pediatric patient. Pediatric Allergy and Immunology, 2017, 28, 305-306.	1.1	3
75	Nutcracker syndrome: a potentially underdiagnosed cause of proteinuria in children with familial Mediterranean fever. Pediatric Nephrology, 2022, 37, 1615-1621.	0.9	3
76	Ocular inflammatory diseases in children with familial Mediterranean fever: a true association or a coincidence?. International Ophthalmology, 2022, 42, 1249-1257.	0.6	3
77	Juvenile polyarteritis nodosa associated with toxoplasmosis presenting as Kawasaki disease. Pediatrics International, 2014, 56, 262-264.	0.2	2
78	Nephrotic syndrome in a patient with cystinuria: Answers. Pediatric Nephrology, 2020, 35, 979-980.	0.9	2
79	Is the performance of the international severity scoring system for familial mediterranean fever in children better than other scoring systems?. International Journal of Clinical Practice, 2021, 75, e14678.	0.8	2
80	An unexpected diagnostic course of systemic lupus erythematosus. Turkish Journal of Pediatrics, 2016, 58, 223-226.	0.3	2
81	An unusual cause of gross hematuria: Questions. Pediatric Nephrology, 2015, 30, 767-767.	0.9	1
82	Takayasu Arteritis: A Case Presenting With Neurological Symptoms and Proteinuria. Archives of Rheumatology, 2020, 35, 287-291.	0.3	1
83	An unusual cause of gross hematuria: Answers. Pediatric Nephrology, 2015, 30, 769-770.	0.9	Ο
84	Atypical presentation of multicentric Castleman disease in a pediatric patient: pleural and pericardial effusion. European Journal of Pediatrics, 2016, 175, 873-876.	1.3	0
85	Nephrotic syndrome in a patient with cystinuria: Questions. Pediatric Nephrology, 2020, 35, 977-977.	0.9	0
86	A rare cause of postinfectious glomerulonephritis: Questions. Pediatric Nephrology, 2021, 36, 555-556.	0.9	0
87	A rare cause of postinfectious glomerulonephritis: Answers. Pediatric Nephrology, 2021, 36, 557-559.	0.9	ο
88	Thoracic mass lesion in a 14-year-old girl: Questions. Pediatric Nephrology, 2021, 36, 2001-2002.	0.9	0
89	Thoracic mass lesion in a 14-year-old girl: Answers. Pediatric Nephrology, 2021, 36, 2003-2005.	0.9	0
90	Approach to Inherited Cystic Kidney Disease. Guncel Pediatri, 2015, 13, 40-45.	0.1	0

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91	Acute Kidney Injury: Classification and Prognosis. Turkish Journal of Pediatric Disease, 0, , .	0.0	Ο