Zeynep Birsin Ózçakar

List of Publications by Year in descending order

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138 papers 3,339 citations

201385 27 h-index 52 g-index

143 all docs

143
docs citations

times ranked

143

4077 citing authors

#	Article	IF	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	Digestion of Chromatin in Apoptotic Cell Microparticles Prevents Autoimmunity. Cell, 2016, 166, 88-101.	13.5	340
3	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
4	Etiology and outcome of acute kidney injury in children. Pediatric Nephrology, 2010, 25, 1453-1461.	0.9	117
5	<i>DNASE1L3</i> Mutations in Hypocomplementemic Urticarial Vasculitis Syndrome. Arthritis and Rheumatism, 2013, 65, 2183-2189.	6.7	116
6	Antibiotic resistance of urinary tract pathogens and evaluation of empirical treatment in Turkish children with urinary tract infections. International Journal of Antimicrobial Agents, 2006, 28, 413-416.	1.1	88
7	Prevalence of the MEFV Gene Mutations in Childhood Polyarteritis Nodosa. Journal of Pediatrics, 2007, 151, 675-678.	0.9	79
8	Quality of life in children with chronic kidney disease (with child and parent assessments). Pediatric Nephrology, 2010, 25, 1487-1496.	0.9	77
9	Exploring the Clinical and Genetic Spectrum of Steroid Resistant Nephrotic Syndrome: The PodoNet Registry. Frontiers in Pediatrics, 2018, 6, 200.	0.9	77
10	Anti-IL-1 treatment in familial Mediterranean fever and related amyloidosis. Clinical Rheumatology, 2016, 35, 441-446.	1.0	76
11	Prevalence of Hypertension in Children with Early-Stage ADPKD. Clinical Journal of the American Society of Nephrology: CJASN, 2018, 13, 874-883.	2.2	65
12	LRIG2 Mutations Cause Urofacial Syndrome. American Journal of Human Genetics, 2013, 92, 259-264.	2.6	63
13	<i>MEFV</i> Mutations Modify the Clinical Presentation of Henoch-Schönlein Purpura. Journal of Rheumatology, 2008, 35, 2427-2429.	1.0	62
14	Nutcracker Syndrome in Children. Journal of Ultrasound in Medicine, 2007, 26, 573-580.	0.8	57
15	High frequency of kidney and urinary tract anomalies in asymptomatic first-degree relatives of patients with CAKUT. Pediatric Nephrology, 2013, 28, 2143-2147.	0.9	55
16	Isolated nocturnal and isolated daytime hypertension associate with altered cardiovascular morphology and function in children with chronic kidney disease. Journal of Hypertension, 2019, 37, 2247-2255.	0.3	45
17	Prospective evaluation of acute and chronic renal function in children following matched related donor hematopoietic stem cell transplantation. Pediatric Transplantation, 2010, 14, 138-144.	0.5	42
18	Application of the new classification criteria of the Acute Kidney Injury Network: a pilot study in a pediatric population. Pediatric Nephrology, 2009, 24, 1379-1384.	0.9	41

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19	Impact of Everolimus and Low-Dose Cyclosporin on Cytomegalovirus Replication and Disease in Pediatric Renal Transplantation. American Journal of Transplantation, 2016, 16, 921-929.	2.6	41
20	Anemia in pediatric renal transplant recipients. Pediatric Nephrology, 2004, 19, 526-530.	0.9	40
21	Possible effect of subclinical inflammation on daily life in familial Mediterranean fever. Clinical Rheumatology, 2006, 25, 149-152.	1.0	40
22	Hypertension induced reversible posterior leukoencephalopathy syndrome: a report of two cases. European Journal of Pediatrics, 2004, 163, 728-730.	1.3	39
23	Acute kidney injury in a paediatric intensive care unit: comparison of the pRIFLE and AKIN criteria. Acta Paediatrica, International Journal of Paediatrics, 2012, 101, e126-9.	0.7	35
24	Turkish pediatric atypical hemolytic uremic syndrome registry: initial analysis of 146 patients. BMC Nephrology, 2017, 18, 6.	0.8	35
25	Early Effects of Renal Replacement Therapy on Cardiovascular Comorbidity in Children With End-Stage Kidney Disease. Transplantation, 2018, 102, 484-492.	0.5	31
26	Infliximab therapy for familial Mediterranean fever-related amyloidosis: case series with long term follow-up. Clinical Rheumatology, 2012, 31, 1267-1271.	1.0	30
27	Management of antenatal hydronephrosis. Pediatric Nephrology, 2020, 35, 2231-2239.	0.9	30
28	Extra-Renal manifestations of atypical hemolytic uremic syndrome in children. Pediatric Nephrology, 2018, 33, 1395-1403.	0.9	29
29	The "nutcracker phenomenon" with orthostatic proteinuria: case reports. Clinical Nephrology, 2006, 65, 280-283.	0.4	29
30	Familial Mediterranean fever-associated diseases in children. QJM - Monthly Journal of the Association of Physicians, 2017, 110, hcw230.	0.2	27
31	Indoxyl sulfate associates with cardiovascular phenotype in children with chronic kidney disease. Pediatric Nephrology, 2019, 34, 2571-2582.	0.9	27
32	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
33	An infant with severe refractory Crohn's disease and homozygous MEFV mutation who dramatically responded to colchicine. Rheumatology International, 2012, 32, 783-785.	1.5	26
34	Late-onset disease is associated with a mild phenotype in children with familial Mediterranean fever. Clinical Rheumatology, 2016, 35, 1837-1840.	1.0	26
35	Ochoa syndrome: a spectrum of urofacial syndrome. European Journal of Pediatrics, 2010, 169, 431-435.	1.3	25
36	Vascular comorbidities in familial Mediterranean fever. Rheumatology International, 2011, 31, 1275-1281.	1.5	25

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37	Clinical improvement with infliximab in a child with amyloidosis secondary to familial Mediterranean fever. Rheumatology, 2006, 45, 1307-1308.	0.9	24
38	Polyarteritis nodosa: successful diagnostic imaging utilizing pulsed and color Doppler ultrasonography and computed tomography angiography. European Journal of Pediatrics, 2006, 165, 120-123.	1.3	24
39	Global left-ventricular function by tissue Doppler imaging in pediatric dialysis patients. Pediatric Nephrology, 2008, 23, 779-785.	0.9	24
40	Analysis of NPHS2 mutations in Turkish steroid-resistant nephrotic syndrome patients. Pediatric Nephrology, 2006, 21, 1093-1096.	0.9	23
41	Urinary tract infections owing to ESBLâ€producing bacteria: microorganisms change – clinical pattern does not. Acta Paediatrica, International Journal of Paediatrics, 2011, 100, e61-4.	0.7	22
42	Predictors of left ventricular hypertrophy in children on chronic peritoneal dialysis. Pediatric Nephrology, 2010, 25, 1311-1318.	0.9	21
43	Hypocomplementemic urticarial vasculitis syndrome in three siblings. Rheumatology International, 2013, 33, 763-766.	1.5	21
44	Clinical features and disease severity of Turkish FMF children carrying E148Q mutation. Journal of Clinical Laboratory Analysis, 2019, 33, e22852.	0.9	21
45	Can colchicine response be predicted in familial Mediterranean fever patients?. Rheumatology, 2014, 53, 1767-1772.	0.9	20
46	Response to Early Coenzyme Q10 Supplementation Is not Sustained in CoQ10 Deficiency Caused by CoQ2 Mutation. Pediatric Neurology, 2018, 88, 71-74.	1.0	20
47	Discontinuation of RAAS Inhibition in Children with Advanced CKD. Clinical Journal of the American Society of Nephrology: CJASN, 2020, 15, 625-632.	2.2	19
48	Sacroiliitis in Children With Familial Mediterranean Fever. Journal of Clinical Rheumatology, 2019, 25, 69-73.	0.5	18
49	Sex and age as determinants for high blood pressure in pediatric renal transplant recipients: a longitudinal analysis of the CERTAIN Registry. Pediatric Nephrology, 2020, 35, 415-426.	0.9	18
50	Concomitant compression of median and ulnar nerves in a hemophiliac patient: a case report. Joint Bone Spine, 2002, 69, 611-613.	0.8	17
51	Encapsulating peritoneal sclerosis in paediatric peritoneal dialysis patients. Nephrology, 2005, 10, 341-343.	0.7	17
52	Renal replacement therapies in pediatric intensive care patients: Experiences of one center in Turkey. Pediatrics International, 2007, 49, 345-348.	0.2	17
53	Lower urinary tract dysfunction is frequently seen in urinary tract infections in children and is often associated with reduced quality of life. Acta Paediatrica, International Journal of Paediatrics, 2014, 103, e454-e458.	0.7	17
54	The Clinical Characteristics of Pediatric Non-Infectious Uveitis in Two Tertiary Referral Centers in Turkey. Ocular Immunology and Inflammation, 2021, 29, 282-289.	1.0	17

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55	The expanded clinical spectrum of familial Mediterranean fever. Clinical Rheumatology, 2007, 26, 1557-1560.	1.0	16
56	Evaluation and Outcome of Antenatal Hydronephrosis: A Prospective Study. Renal Failure, 2012, 34, 718-721.	0.8	16
57	Vesicoureteral Reflux and Renal Scarring Risk in Children after the First Febrile Urinary Tract Infection. Nephron, 2016, 132, 175-180.	0.9	16
58	Evaluation of Renal Function in Obese Children and Adolescents Using Serum Cystatin C Levels, Estimated Glomerular Filtration Rate Formulae and Proteinuria: Which is most Useful?. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2019, 11, 46-54.	0.4	16
59	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
60	Nutcracker syndrome manifesting with severe proteinuria: a challenging scenario in a single-kidney patient. Pediatric Nephrology, 2011, 26, 987-990.	0.9	15
61	Renal Involvement in Childhood Vasculitis. Nephron Clinical Practice, 2008, 108, c202-c206.	2.3	13
62	Acute kidney injury in congenital cardiac surgery: Pediatric risk–injury–failure–loss–endâ€stage renal disease and Acute Kidney Injury Network. Pediatrics International, 2017, 59, 1252-1260.	0.2	13
63	Familial Mediterranean fever in small children in Turkey. Clinical and Experimental Rheumatology, 2011, 29, S87-90.	0.4	13
64	The effect of colchicine on physical growth in children wıth familial mediterranean fever. European Journal of Pediatrics, 2010, 169, 825-828.	1.3	12
65	Hypertension and improved left ventricular mass index in children after renal transplantation. Pediatric Transplantation, 2017, 21, e13066.	0.5	12
66	Childhood polyarteritis nodosa: diagnosis with non-invasive imaging techniques. Clinical Rheumatology, 2017, 36, 165-171.	1.0	12
67	Oxcarbazepine and valproic acidâ€induced lupus in a 7â€yearâ€old boy. Acta Paediatrica, International Journal of Paediatrics, 2008, 97, 1000-1001.	0.7	11
68	<i>HPSE2</i> Mutations in Urofacial Syndrome, Non-Neurogenic Neurogenic Bladder and Lower Urinary Tract Dysfunction. Nephron, 2015, 130, 54-58.	0.9	10
69	First-Line, Early and Long-Term Eculizumab Therapy in Atypical Hemolytic Uremic Syndrome: A Case Series in Pediatric Patients. Paediatric Drugs, 2016, 18, 413-420.	1.3	10
70	The association between obesity, hypertension and left ventricular mass in adolescents. Journal of Pediatric Endocrinology and Metabolism, 2017, 30, 167-174.	0.4	10
71	Atypical Hemolytic Uremic Syndrome in Children Aged <2 Years. Nephron, 2018, 139, 211-218.	0.9	10
72	Familial Mediterranean fever gene mutation frequencies in a sample Turkish population. Clinical and Experimental Rheumatology, 2016, 34, 97-100.	0.4	10

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73	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
74	<i>Acremonium</i> spp. peritonitis in an infant. Mycoses, 2008, 51, 455-457.	1.8	8
75	Nutcracker syndrome with urolithiasis. Journal of Pediatric Urology, 2010, 6, 519-521.	0.6	8
76	The expanded spectrum of arthritis in children with familial Mediterranean fever. Clinical Rheumatology, 2022, 41, 1535-1541.	1.0	8
77	Is there an association between familial Mediterranean fever and celiac disease?. Clinical Rheumatology, 2008, 27, 1135-1139.	1.0	7
78	A questionnaire survey of radiological diagnosis and management of renal dysplasia in children. Journal of Nephrology, 2018, 31, 95-102.	0.9	7
79	Three siblings with steroid-resistant nephrotic syndrome: New NPHS2 mutations in a Turkish family. American Journal of Kidney Diseases, 2004, 44, e22-e24.	2.1	6
80	The association of cystic nephroma with pulmonary sequestration: is it a coincidence or not?. Pediatric Nephrology, 2006, 21, 1041-1044.	0.9	6
81	The value of procalcitonin measurements in children with familial Mediterranean fever. Rheumatology International, 2012, 32, 3443-3447.	1.5	6
82	Primary Hyperoxaluria Type 1: A Cause for Infantile Renal Failure and Massive Nephrocalcinosis. Klinische Padiatrie, 2015, 227, 293-295.	0.2	6
83	Vitamin B6 deficiency presenting with low alanine aminotransferase in a critically ill child. Pediatrics International, 2009, 51, 597-599.	0.2	5
84	Mayer-Rokitansky-Kýster-Hauser Syndrome Accompanied by Renal Cell Carcinoma. Journal of Pediatric Hematology/Oncology, 2013, 35, e309-e310.	0.3	5
85	Anti-VEGF-related thrombotic microangiopathy in a child presenting with nephrotic syndrome. Pediatric Nephrology, 2016, 31, 1029-1032.	0.9	5
86	Neonatal onset familial Mediterranean fever. Modern Rheumatology, 2019, 29, 647-650.	0.9	5
87	The effect of genotype on musculoskeletal complaints in patients with familial Mediterranean fever. Postgraduate Medicine, 2020, 132, 220-224.	0.9	5
88	The changing resistance patterns of bacterial uropathogens in children. Pediatrics International, 2020, 62, 1058-1063.	0.2	5
89	Physical and mental development of Turkish twins. Pediatrics International, 2003, 45, 712-718.	0.2	4
90	Systemic vascular calcification with retinal calcification in an adolescent treated with long-term peritoneal dialysis. Pediatric Nephrology, 2006, 21, 1915-1916.	0.9	4

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91	De novo amyloidosis in a renal transplant patient. Pediatric Transplantation, 2014, 18, E259-61.	0.5	4
92	Hypertension in children after renal transplantation. Pediatrics International, 2015, 57, 1138-1142.	0.2	4
93	An Analysis of the Levels of the Soluble Form of the Endothelial Protein C Receptor in Children with Henoch–Schönlein Purpura. Pediatric Hematology and Oncology, 2015, 32, 115-122.	0.3	4
94	Fatigue in pediatric patients with familial Mediterranean fever. Modern Rheumatology, 2018, 28, 1016-1020.	0.9	4
95	Proteinuria in pediatric renal transplant recipients. Pediatric Transplantation, 2018, 22, e13068.	0.5	4
96	COL4A3 mutation is an independent risk factor for poor prognosis in children with Alport syndrome. Pediatric Nephrology, 2020, 35, 1941-1952.	0.9	4
97	Renal Biopsy Prognostic Findings in Children With Atypical Hemolytic Uremic Syndrome. Pediatric and Developmental Pathology, 2020, 23, 362-371.	0.5	4
98	Juvenile chronic arthritis in a monozygotic twin couple. Rheumatology International, 2003, 23, 149-150.	1.5	3
99	A favorable outcome of hemolytic uremic syndrome with factor H deficiency. Pediatric Nephrology, 2004, 19, 815-816.	0.9	3
100	Multicystic dysplastic kidney and caliceal diverticulum in a child a coincidence or an association?. International Urology and Nephrology, 2007, 39, 27-29.	0.6	3
101	Pericardial tamponade in a child with nephrotic syndrome. Pediatric Nephrology, 2011, 26, 993-994.	0.9	3
102	Angiodysplasia as a Cause of Severe Hematochezia in a Child with End-Stage Renal Failure. Renal Failure, 2011, 33, 252-254.	0.8	3
103	Pediatric Renal Transplantation: A Single Center Experience. Transplantation Proceedings, 2013, 45, 917-918.	0.3	3
104	Two children with steroid-responsive nephrotic syndrome complicated by cerebral venous sinus thrombosis. Nefrologia, 2015, 35, 497-500.	0.2	3
105	Multiple intraâ€cardiac masses: A lifeâ€threatening complication of Behçet's disease. Echocardiography, 2020, 37, 1077-1079.	0.3	3
106	Nutcracker syndrome: a potentially underdiagnosed cause of proteinuria in children with familial Mediterranean fever. Pediatric Nephrology, 2022, 37, 1615-1621.	0.9	3
107	Ocular inflammatory diseases in children with familial Mediterranean fever: a true association or a coincidence?. International Ophthalmology, 2022, 42, 1249-1257.	0.6	3
108	Eculizumab treatment and discontinuation in pediatric patients with atypical hemolytic uremic syndrome: a multicentric retrospective study. Journal of Nephrology, 2022, , 1.	0.9	3

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109	A girl with microscopic polyangiitis: an unexpected clinical course with long-term follow-up. Pediatric Nephrology, 2005, 20, 694-695.	0.9	2
110	Acute renal failure in a patient with familial Mediterranean fever. Rheumatology International, 2006, 27, 309-310.	1.5	2
111	Training and Deployment as a Basis for Usability Engineering of Mobile Systems. , 2008, , .		2
112	Thrombotic microangiopathy in a pediatric patient: answer. Pediatric Nephrology, 2009, 24, 1139-1141.	0.9	2
113	Transplantation in pediatric aHUS within the era of eculizumab therapy. Pediatric Transplantation, 2021, 25, e13914.	0.5	2
114	Severe hyperphosphatemia after oral laxative administration in a 7-year-old patient. Turkish Journal of Pediatrics, 2016, 58, 116-118.	0.3	2
115	Prevalence and potential relevance of hyperuricemia in pediatric kidney transplant recipients—a CERTAIN registry analysis. Pediatric Transplantation, 2022, 26, e14265.	0.5	2
116	Correspondence. Pediatric Neurology, 2004, 30, 299.	1.0	1
117	A 12-month-old boy with high fever, erythematous lesions and haemorrhagic oedema. European Journal of Pediatrics, 2005, 164, 453-454.	1.3	1
118	An unusual complication of peritoneal dialysis. Pediatric Nephrology, 2006, 21, 129-130.	0.9	1
119	Thrombotic microangiopathy in a pediatric patient: question. Pediatric Nephrology, 2009, 24, 1137-1138.	0.9	1
120	An adolescent girl with hypertension and neuropsychiatric symptoms: Questions. Pediatric Nephrology, 2013, 28, 427-427.	0.9	1
121	Conservative Management of Lumbar Disk Herniations in Adolescents. Southern Medical Journal, 2002, 95, 1454-1455.	0.3	O
122	Soluble CD27 Levels in Children with Acute and Chronic Renal Failure. Journal of Allergy and Clinical Immunology, 2009, 123, S226-S226.	1.5	0
123	Neurogenic bladder in twins: question. Pediatric Nephrology, 2010, 25, 1057-1057.	0.9	O
124	Neurogenic bladder in twins: answer. Pediatric Nephrology, 2010, 25, 1059-1061.	0.9	0
125	Medullary nephrocalcinosis in a pediatric patient: question. Pediatric Nephrology, 2010, 25, 1429-1430.	0.9	O
126	Hypertension and hypokalemia in a 15-year-old boy: question. Pediatric Nephrology, 2010, 25, 2443-2444.	0.9	0

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127	Hypertension and hypokalemia in a 15-year-old boy: answer. Pediatric Nephrology, 2010, 25, 2445-2447.	0.9	O
128	Vascular Calcification in an Adolescent Treated with Long-Term Peritoneal Dialysis. International Journal of Nephrology, 2011, 2011, 1-4.	0.7	O
129	Familial Mediterranean Fever in Siblings. Journal of Rheumatology, 2012, 39, 2170-2174.	1.0	O
130	An adolescent girl with hypertension and neuropsychiatric symptoms: Answers. Pediatric Nephrology, 2013, 28, 429-431.	0.9	0
131	Two children with steroid-responsive nephrotic syndrome complicated by cerebral venous sinus thrombosis. Nefrologia, 2015, 35, 497-500.	0.2	O
132	Proteinuria in a Crohn's disease patient: Questions. Pediatric Nephrology, 2015, 30, 1433-1433.	0.9	0
133	Correction: Primary Hyperoxaluria Type 1: A Cause for Infantile Renal Failure and Massive Nephrocalcinosis. Klinische Padiatrie, 2015, 227, e3-e3.	0.2	O
134	Proteinuria in a Crohn's disease patient: Answers. Pediatric Nephrology, 2015, 30, 1435-1436.	0.9	O
135	Campylobacter jejuni: un agente infrecuente de peritonitis en un niñ0 con diálisis peritoneal. Archivos Argentinos De Pediatria, 2016, 114, e354.	0.3	O
136	Dyslipidemias in the Pediatric Chronic Kidney Disease Patient. , 2014, , 231-253.		O
137	Peritoneal dialysis in acute kidney injury. , 2015, , .		O
138	Membranoproliferative glomerulonephritis in a patient with Wilson's disease. Journal of Nephrology, 2006, 19, 831-3.	0.9	0