Zeynep Birsin Ã“zÃ§akar

List of Publications by Year
in descending order

140 papers	3,339 citations	201674 27 h-index	175258 52 g-index
143 all docs	143 docs citations	143 times ranked	4077 citing authors

#	Article	IF	Citations
1	Nutcracker syndrome: a potentially underdiagnosed cause of proteinuria in children with familial Mediterranean fever. Pediatric Nephrology, 2022, 37, 1615-1621.	1.7	3
2	Ocular inflammatory diseases in children with familial Mediterranean fever: a true association or a coincidence?. International Ophthalmology, 2022, 42, 1249-1257.	1.4	3
3	Eculizumab treatment and discontinuation in pediatric patients with atypical hemolytic uremic syndrome: a multicentric retrospective study. Journal of Nephrology, 2022, , 1.	2.0	3
4	The expanded spectrum of arthritis in children with familial Mediterranean fever. Clinical Rheumatology, 2022, 41, 1535-1541.	2.2	8
5	Prevalence and potential relevance of hyperuricemia in pediatric kidney transplant recipientsâ€”a CERTAIN registry analysis. Pediatric Transplantation, 2022, 26, e14265.	1.0	2
6	Transplantation in pediatric aHUS within the era of eculizumab therapy. Pediatric Transplantation, 2021, 25, e13914.	1.0	2
7	The Clinical Characteristics of Pediatric Non-Infectious Uveitis in Two Tertiary Referral Centers in Turkey. Ocular Immunology and Inflammation, 2021, 29, 282-289.	1.8	17
8	The effect of genotype on musculoskeletal complaints in patients with familial Mediterranean fever. Postgraduate Medicine, 2020, 132, 220-224.	2.0	5
9	Management of antenatal hydronephrosis. Pediatric Nephrology, 2020, 35, 2231-2239.	1.7	30
10	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.	1.7	18
11	COL4A3 mutation is an independent risk factor for poor prognosis in children with Alport syndrome. Pediatric Nephrology, 2020, 35, 1941-1952.	1.7	4
12	Renal Biopsy Prognostic Findings in Children With Atypical Hemolytic Uremic Syndrome. Pediatric and Developmental Pathology, 2020, 23, 362-371.	1.0	4
13	Multiple intraâ€cardiac masses: A lifeâ€threatening complication of BehÃ§etâ€™s disease. Echocardiography, 2020, 37, 1077-1079.	0.9	3
14	The changing resistance patterns of bacterial uropathogens in children. Pediatrics International, 2020, 62, 1058-1063.	0.5	5
15	Discontinuation of RAAS Inhibition in Children with Advanced CKD. Clinical Journal of the American Society of Nephrology: CJASN, 2020, 15, 625-632.	4.5	19
16	CaracterÃsticas clÃnicas y evoluciÃ³n del reflujo vesicoureteral en la infancia. Archivos Argentinos De Pediatria, 2020, 118, .	0.2	0
17	Neonatal onset familial Mediterranean fever. Modern Rheumatology, 2019, 29, 647-650.	1.8	5
18	Indoxyl sulfate associates with cardiovascular phenotype in children with chronic kidney disease. Pediatric Nephrology, 2019, 34, 2571-2582.	1.7	27

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19	Clinical features and disease severity of Turkish FMF children carrying E148Q mutation. Journal of Clinical Laboratory Analysis, 2019, 33, e22852.	2.1	21
20	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.	0.5	45
21	Sacroiliitis in Children With Familial Mediterranean Fever. Journal of Clinical Rheumatology, 2019, 25, 69-73.	0.9	18
22	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.	0.9	16
23	Prevalence of Hypertension in Children with Early-Stage ADPKD. Clinical Journal of the American Society of Nephrology: CJASN, 2018, 13, 874-883.	4.5	65
24	Extra-Renal manifestations of atypical hemolytic uremic syndrome in children. Pediatric Nephrology, 2018, 33, 1395-1403.	1.7	29
25	Fatigue in pediatric patients with familial Mediterranean fever. Modern Rheumatology, 2018, 28, 1016-1020.	1.8	4
26	Atypical Hemolytic Uremic Syndrome in Children Aged <2 Years. Nephron, 2018, 139, 211-218.	1.8	10
27	A questionnaire survey of radiological diagnosis and management of renal dysplasia in children. Journal of Nephrology, 2018, 31, 95-102.	2.0	7
28	Proteinuria in pediatric renal transplant recipients. Pediatric Transplantation, 2018, 22, e13068.	1.0	4
29	Early Effects of Renal Replacement Therapy on Cardiovascular Comorbidity in Children With End-Stage Kidney Disease. Transplantation, 2018, 102, 484-492.	1.0	31
30	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.	1.6	9
31	Response to Early Coenzyme Q10 Supplementation Is not Sustained in CoQ10 Deficiency Caused by CoQ2 Mutation. Pediatric Neurology, 2018, 88, 71-74.	2.1	20
32	Exploring the Clinical and Genetic Spectrum of Steroid Resistant Nephrotic Syndrome: The PodoNet Registry. Frontiers in Pediatrics, 2018, 6, 200.	1.9	77
33	Familial Mediterranean fever-associated diseases in children. QJM - Monthly Journal of the Association of Physicians, 2017, 110, hcw230.	0.5	27
34	Turkish pediatric atypical hemolytic uremic syndrome registry: initial analysis of 146 patients. BMC Nephrology, 2017, 18, 6.	1.8	35
35	The association between obesity, hypertension and left ventricular mass in adolescents. Journal of Pediatric Endocrinology and Metabolism, 2017, 30, 167-174.	0.9	10
36	Hypertension and improved left ventricular mass index in children after renal transplantation. Pediatric Transplantation, 2017, 21, e13066.	1.0	12

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37	Acute kidney injury in congenital cardiac surgery: Pediatric riskâ€“injuryâ€“failureâ€“lossâ€“endâ€stage renal disease and Acute Kidney Injury Network. Pediatrics International, 2017, 59, 1252-1260.	0.5	13
38	Childhood polyarteritis nodosa: diagnosis with non-invasive imaging techniques. Clinical Rheumatology, 2017, 36, 165-171.	2.2	12
39	Anti-IL-1 treatment in familial Mediterranean fever and related amyloidosis. Clinical Rheumatology, 2016, 35, 441-446.	2.2	76
40	Campylobacter jejuni: A rare agent in a child with peritoneal dialysis-related peritonitis. Archivos Argentinos De Pediatria, 2016, 114, .	0.2	0
41	Campylobacter jejuni: un agente infrecuente de peritonitis en un niÃ±o con diÃ¡lisis peritoneal. Archivos Argentinos De Pediatria, 2016, 114, e354.	0.2	0
42	Impact of Everolimus and Low-Dose Cyclosporin on Cytomegalovirus Replication and Disease in Pediatric Renal Transplantation. American Journal of Transplantation, 2016, 16, 921-929.	4.7	41
43	Vesicoureteral Reflux and Renal Scarring Risk in Children after the First Febrile Urinary Tract Infection. Nephron, 2016, 132, 175-180.	1.8	16
44	Anti-VEGF-related thrombotic microangiopathy in a child presenting with nephrotic syndrome. Pediatric Nephrology, 2016, 31, 1029-1032.	1.7	5
45	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.	3.1	10
46	Digestion of Chromatin in Apoptotic Cell Microparticles Prevents Autoimmunity. Cell, 2016, 166, 88-101.	28.9	340
47	Late-onset disease is associated with a mild phenotype in children with familial Mediterranean fever. Clinical Rheumatology, 2016, 35, 1837-1840.	2.2	26
48	Severe hyperphosphatemia after oral laxative administration in a 7-year-old patient. Turkish Journal of Pediatrics, 2016, 58, 116-118.	0.6	2
49	Familial Mediterranean fever gene mutation frequencies in a sample Turkish population. Clinical and Experimental Rheumatology, 2016, 34, 97-100.	0.8	10
50	Hypertension in children after renal transplantation. Pediatrics International, 2015, 57, 1138-1142.	0.5	4
51	Two children with steroid-responsive nephrotic syndrome complicated by cerebral venous sinus thrombosis. Nefrologia, 2015, 35, 497-500.	0.4	0
52	Spectrum of Steroid-Resistant and Congenital Nephrotic Syndrome in Children. Clinical Journal of the American Society of Nephrology: CJASN, 2015, 10, 592-600.	4.5	225
53	Proteinuria in a Crohnâ€™s disease patient: Questions. Pediatric Nephrology, 2015, 30, 1433-1433.	1.7	0
54	Two children with steroid-responsive nephrotic syndrome complicated by cerebral venous sinus thrombosis. Nefrologia, 2015, 35, 497-500.	0.4	3

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55	HPSE2 Mutations in Urofacial Syndrome, Non-Neurogenic Neurogenic Bladder and Lower Urinary Tract Dysfunction. Nephron, 2015, 130, 54-58.	1.8	10
56	An Analysis of the Levels of the Soluble Form of the Endothelial Protein C Receptor in Children with Henochâ€“SchÃ¶nlein Purpura. Pediatric Hematology and Oncology, 2015, 32, 115-122.	0.8	4
57	Primary Hyperoxaluria Type 1: A Cause for Infantile Renal Failure and Massive Nephrocalcinosis. Klinische Padiatrie, 2015, 227, 293-295.	0.6	6
58	Correction: Primary Hyperoxaluria Type 1: A Cause for Infantile Renal Failure and Massive Nephrocalcinosis. Klinische Padiatrie, 2015, 227, e3-e3.	0.6	0
59	Proteinuria in a Crohnâ€™s disease patient: Answers. Pediatric Nephrology, 2015, 30, 1435-1436.	1.7	0
60	Peritoneal dialysis in acute kidney injury. , 2015, , .		0
61	De novo amyloidosis in a renal transplant patient. Pediatric Transplantation, 2014, 18, E259-61.	1.0	4
62	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.	1.5	17
63	Can colchicine response be predicted in familial Mediterranean fever patients?. Rheumatology, 2014, 53, 1767-1772.	1.9	20
64	Dyslipidemias in the Pediatric Chronic Kidney Disease Patient. , 2014, , 231-253.		0
65	High frequency of kidney and urinary tract anomalies in asymptomatic first-degree relatives of patients with CAKUT. Pediatric Nephrology, 2013, 28, 2143-2147.	1.7	55
66	Hypocomplementemic urticarial vasculitis syndrome in three siblings. Rheumatology International, 2013, 33, 763-766.	3.0	21
67	An adolescent girl with hypertension and neuropsychiatric symptoms: Questions. Pediatric Nephrology, 2013, 28, 427-427.	1.7	1
68	An adolescent girl with hypertension and neuropsychiatric symptoms: Answers. Pediatric Nephrology, 2013, 28, 429-431.	1.7	0
69	Pediatric Renal Transplantation: A Single Center Experience. Transplantation Proceedings, 2013, 45, 917-918.	0.6	3
70	LRIG2 Mutations Cause Urofacial Syndrome. American Journal of Human Genetics, 2013, 92, 259-264.	6.2	63
71	DNASE1L3 Mutations in Hypocomplementemic Urticarial Vasculitis Syndrome. Arthritis and Rheumatism, 2013, 65, 2183-2189.	6.7	116
72	Mayer-Rokitansky-KÃ¼ster-Hauser Syndrome Accompanied by Renal Cell Carcinoma. Journal of Pediatric Hematology/Oncology, 2013, 35, e309-e310.	0.6	5

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73	Familial Mediterranean Fever in Siblings. Journal of Rheumatology, 2012, 39, 2170-2174.	2.0	0
74	Infliximab therapy for familial Mediterranean fever-related amyloidosis: case series with long term follow-up. Clinical Rheumatology, 2012, 31, 1267-1271.	2.2	30
75	The value of procalcitonin measurements in children with familial Mediterranean fever. Rheumatology International, 2012, 32, 3443-3447.	3.0	6
76	Evaluation and Outcome of Antenatal Hydronephrosis: A Prospective Study. Renal Failure, 2012, 34, 718-721.	2.1	16
77	An infant with severe refractory Crohnâ€™s disease and homozygous MEFV mutation who dramatically responded to colchicine. Rheumatology International, 2012, 32, 783-785.	3.0	26
78	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.	1.5	35
79	Urinary tract infections owing to ESBLâ€producing bacteria: microorganisms change â€“ clinical pattern does not. Acta Paediatrica, International Journal of Paediatrics, 2011, 100, e61-4.	1.5	22
80	Pericardial tamponade in a child with nephrotic syndrome. Pediatric Nephrology, 2011, 26, 993-994.	1.7	3
81	Nutcracker syndrome manifesting with severe proteinuria: a challenging scenario in a single-kidney patient. Pediatric Nephrology, 2011, 26, 987-990.	1.7	15
82	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.	2.7	26
83	Vascular comorbidities in familial Mediterranean fever. Rheumatology International, 2011, 31, 1275-1281.	3.0	25
84	Angiodysplasia as a Cause of Severe Hematochezia in a Child with End-Stage Renal Failure. Renal Failure, 2011, 33, 252-254.	2.1	3
85	Vascular Calcification in an Adolescent Treated with Long-Term Peritoneal Dialysis. International Journal of Nephrology, 2011, 2011, 1-4.	1.3	0
86	Familial Mediterranean fever in small children in Turkey. Clinical and Experimental Rheumatology, 2011, 29, S87-90.	0.8	13
87	Ochoa syndrome: a spectrum of urofacial syndrome. European Journal of Pediatrics, 2010, 169, 431-435.	2.7	25
88	The effect of colchicine on physical growth in children wÄ±th familial mediterranean fever. European Journal of Pediatrics, 2010, 169, 825-828.	2.7	12
89	Neurogenic bladder in twins: question. Pediatric Nephrology, 2010, 25, 1057-1057.	1.7	0
90	Neurogenic bladder in twins: answer. Pediatric Nephrology, 2010, 25, 1059-1061.	1.7	0

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91	Medullary nephrocalcinosis in a pediatric patient: question. Pediatric Nephrology, 2010, 25, 1429-1430.	1.7	0
92	Hypertension and hypokalemia in a 15-year-old boy: question. Pediatric Nephrology, 2010, 25, 2443-2444.	1.7	0
93	Hypertension and hypokalemia in a 15-year-old boy: answer. Pediatric Nephrology, 2010, 25, 2445-2447.	1.7	0
94	Predictors of left ventricular hypertrophy in children on chronic peritoneal dialysis. Pediatric Nephrology, 2010, 25, 1311-1318.	1.7	21
95	Quality of life in children with chronic kidney disease (with child and parent assessments). Pediatric Nephrology, 2010, 25, 1487-1496.	1.7	77
96	Etiology and outcome of acute kidney injury in children. Pediatric Nephrology, 2010, 25, 1453-1461.	1.7	117
97	Prospective evaluation of acute and chronic renal function in children following matched related donor hematopoietic stem cell transplantation. Pediatric Transplantation, 2010, 14, 138-144.	1.0	42
98	Nutcracker syndrome with urolithiasis. Journal of Pediatric Urology, 2010, 6, 519-521.	1.1	8
99	A new set of criteria for the diagnosis of familial Mediterranean fever in childhood. Rheumatology, 2009, 48, 395-398.	1.9	374
100	Thrombotic microangiopathy in a pediatric patient: answer. Pediatric Nephrology, 2009, 24, 1139-1141.	1.7	2
101	Thrombotic microangiopathy in a pediatric patient: question. Pediatric Nephrology, 2009, 24, 1137-1138.	1.7	1
102	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.	1.7	41
103	Vitamin B6 deficiency presenting with low alanine aminotransferase in a critically ill child. Pediatrics International, 2009, 51, 597-599.	0.5	5
104	Soluble CD27 Levels in Children with Acute and Chronic Renal Failure. Journal of Allergy and Clinical Immunology, 2009, 123, S226-S226.	2.9	0
105	Is there an association between familial Mediterranean fever and celiac disease?. Clinical Rheumatology, 2008, 27, 1135-1139.	2.2	7
106	Global left-ventricular function by tissue Doppler imaging in pediatric dialysis patients. Pediatric Nephrology, 2008, 23, 779-785.	1.7	24
107	Acremonium spp. peritonitis in an infant. Mycoses, 2008, 51, 455-457.	4.0	8
108	Oxcarbazepine and valproic acidâ€induced lupus in a 7â€yearâ€old boy. Acta Paediatrica, International Journal of Paediatrics, 2008, 97, 1000-1001.	1.5	11

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109	Training and Deployment as a Basis for Usability Engineering of Mobile Systems. , 2008, , .		2
110	Renal Involvement in Childhood Vasculitis. Nephron Clinical Practice, 2008, 108, c202-c206.	2.3	13
111	MEFV Mutations Modify the Clinical Presentation of Henoch-SchÃ¶nlein Purpura. Journal of Rheumatology, 2008, 35, 2427-2429.	2.0	62
112	Nutcracker Syndrome in Children. Journal of Ultrasound in Medicine, 2007, 26, 573-580.	1.7	57
113	Renal replacement therapies in pediatric intensive care patients: Experiences of one center in Turkey. Pediatrics International, 2007, 49, 345-348.	0.5	17
114	Prevalence of the MEFV Gene Mutations in Childhood Polyarteritis Nodosa. Journal of Pediatrics, 2007, 151, 675-678.	1.8	79
115	Multicystic dysplastic kidney and caliceal diverticulum in a child a coincidence or an association?. International Urology and Nephrology, 2007, 39, 27-29.	1.4	3
116	The expanded clinical spectrum of familial Mediterranean fever. Clinical Rheumatology, 2007, 26, 1557-1560.	2.2	16
117	Clinical improvement with infliximab in a child with amyloidosis secondary to familial Mediterranean fever. Rheumatology, 2006, 45, 1307-1308.	1.9	24
118	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.	2.5	88
119	An unusual complication of peritoneal dialysis. Pediatric Nephrology, 2006, 21, 129-130.	1.7	1
120	Analysis of NPHS2 mutations in Turkish steroid-resistant nephrotic syndrome patients. Pediatric Nephrology, 2006, 21, 1093-1096.	1.7	23
121	The association of cystic nephroma with pulmonary sequestration: is it a coincidence or not?. Pediatric Nephrology, 2006, 21, 1041-1044.	1.7	6
122	Systemic vascular calcification with retinal calcification in an adolescent treated with long-term peritoneal dialysis. Pediatric Nephrology, 2006, 21, 1915-1916.	1.7	4
123	Possible effect of subclinical inflammation on daily life in familial Mediterranean fever. Clinical Rheumatology, 2006, 25, 149-152.	2.2	40
124	Polyarteritis nodosa: successful diagnostic imaging utilizing pulsed and color Doppler ultrasonography and computed tomography angiography. European Journal of Pediatrics, 2006, 165, 120-123.	2.7	24
125	Acute renal failure in a patient with familial Mediterranean fever. Rheumatology International, 2006, 27, 309-310.	3.0	2
126	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

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127	The Â“nutcracker phenomenonÂ” with orthostatic proteinuria: case reports. Clinical Nephrology, 2006, 65, 280-283.	0.7	29
128	Membranoproliferative glomerulonephritis in a patient with Wilson's disease. Journal of Nephrology, 2006, 19, 831-3.	2.0	0
129	Encapsulating peritoneal sclerosis in paediatric peritoneal dialysis patients. Nephrology, 2005, 10, 341-343.	1.6	17
130	A girl with microscopic polyangiitis: an unexpected clinical course with long-term follow-up. Pediatric Nephrology, 2005, 20, 694-695.	1.7	2
131	A 12-month-old boy with high fever, erythematous lesions and haemorrhagic oedema. European Journal of Pediatrics, 2005, 164, 453-454.	2.7	1
132	Three siblings with steroid-resistant nephrotic syndrome: New NPHS2 mutations in a Turkish family. American Journal of Kidney Diseases, 2004, 44, e22-e24.	1.9	6
133	Anemia in pediatric renal transplant recipients. Pediatric Nephrology, 2004, 19, 526-530.	1.7	40
134	A favorable outcome of hemolytic uremic syndrome with factor H deficiency. Pediatric Nephrology, 2004, 19, 815-816.	1.7	3
135	Hypertension induced reversible posterior leukoencephalopathy syndrome: a report of two cases. European Journal of Pediatrics, 2004, 163, 728-730.	2.7	39
136	Correspondence. Pediatric Neurology, 2004, 30, 299.	2.1	1
137	Juvenile chronic arthritis in a monozygotic twin couple. Rheumatology International, 2003, 23, 149-150.	3.0	3
138	Physical and mental development of Turkish twins. Pediatrics International, 2003, 45, 712-718.	0.5	4
139	Conservative Management of Lumbar Disk Herniations in Adolescents. Southern Medical Journal, 2002, 95, 1454-1455.	0.7	0
140	Concomitant compression of median and ulnar nerves in a hemophiliac patient: a case report. Joint Bone Spine, 2002, 69, 611-613.	1.6	17