Mithat Buyukcelik

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/5657423/publications.pdf

Version: 2024-02-01

24 papers	391 citations	840776 11 h-index	752698 20 g-index
24 all docs	24 docs citations	24 times ranked	611 citing authors

#	Article	IF	Citations
1	Disruption of PTPRO Causes Childhood-Onset Nephrotic Syndrome. American Journal of Human Genetics, 2011, 89, 139-147.	6.2	90
2	Ambulatory blood pressure monitoring and renal functions in children with a solitary kidney. Pediatric Nephrology, 2007, 22, 559-564.	1.7	62
3	Associated anomalies in children with congenital solitary functioning kidney. Pediatric Surgery International, 2005, 21, 456-459.	1.4	39
4	Bartter syndrome and growth hormone deficiency: three cases. Pediatric Nephrology, 2012, 27, 2145-2148.	1.7	24
5	Endothelial nitric oxide synthase gene intron 4 a/b VNTR polymorphism in children with APSGN. Pediatric Nephrology, 2006, 21, 1661-1665.	1.7	16
6	Pediatric postâ€streptococcal glomerulonephritis: Clinical and laboratory data. Pediatrics International, 2018, 60, 645-650.	0.5	16
7	An unusual cause of pleural effusion, urinothorax in a child with urinary stone disease. Pediatric Nephrology, 2005, 20, 1487-1489.	1.7	14
8	ACE Gene Polymorphism in Turkish Children with Nephrotic Syndrome. Renal Failure, 2006, 28, 401-403.	2.1	14
9	May the Best Friend be an Enemy if not Recognized Early. Pediatric Emergency Care, 2005, 21, 445-448.	0.9	12
10	Barttin mutations in antenatal Bartter syndrome with sensorineural deafness. Pediatric Nephrology, 2006, 21, 1056-1057.	1.7	12
11	Changes in Osmolal Gap and Osmolality in Children with Chronic and End-Stage Renal Failure. Nephron Physiology, 2007, 105, p19-p21.	1.2	12
12	Mitral annular calcification and brown tumor of the rib in a child with chronic renal failure. Pediatric Nephrology, 2005, 20, 673-675.	1.7	11
13	Prevalence of hypertension determined by ambulatory blood pressure monitoring (ABPM) and body composition in long-term survivors of childhood cancer. Pediatric Hematology and Oncology, 2018, 35, 1-10.	0.8	10
14	Aggressive angiomyxoma in a child with chronic renal failure. Pediatric Surgery International, 2005, 21, 563-565.	1.4	9
15	Urinary annexin V in children with nephrotic syndrome: a new prognostic marker?. Pediatric Nephrology, 2008, 23, 79-82.	1.7	9
16	Kidney disease profile and encountered problems during follow-up in Syrian refugee children: a multicenter retrospective study. Pediatric Nephrology, 2022, 37, 393-402.	1.7	9
17	Arterial thrombosis associated with factorÂV Leiden and methylenetetrahydrofolate reductase C677T mutation in childhood membranous glomerulonephritis. Pediatric Nephrology, 2008, 23, 491-494.	1.7	8
18	Urotensin-II: More Than a Mediator for Kidney. International Journal of Nephrology, 2012, 2012, 1-7.	1.3	8

#	Article	IF	Citations
19	Association of eNOS gene intron 4 a/b VNTR polymorphisms in children with nephrotic syndrome. Gene, 2013, 522, 192-195.	2.2	6
20	Colchicine Treatment in Children With Familial Mediterranean Fever: Is it a Risk Factor for Neuromyopathy?. Pediatric Neurology, 2013, 49, 417-419.	2.1	4
21	DD Genotype of ACE Gene in Boys: May it be a Risk Factor for Minimal Change Nephrotic Syndrome?. Renal Failure, 2012, 34, 19-23.	2.1	2
22	Autoimmune Polyglandular Syndrome Type 3c with Ectodermal Dysplasia, Immune Deficiency and Hemolytic-Uremic Syndrome. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2014, 6, 47-50.	0.9	2
23	Uteroglobin gene polymorphism (G38A) may be a risk factor in childhood idiopathic nephrotic syndrome. Pediatric Nephrology, 2018, 33, 295-303.	1.7	1
24	Renal biopsy in children with IgA vasculitis. Jornal Brasileiro De Nefrologia: Orgao Oficial De Sociedades Brasileira E Latino-Americana De Nefrologia, 2021, , .	0.9	1