

Mithat Buyukcelik

List of Publications by Year in descending order

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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	Kidney disease profile and encountered problems during follow-up in Syrian refugee children: a multicenter retrospective study. <i>Pediatric Nephrology</i> , 2022, 37, 393-402.	1.7	9
2	Renal biopsy in children with IgA vasculitis. <i>Jornal Brasileiro De Nefrologia: Orgao Oficial De Sociedades Brasileira E Latino-Americana De Nefrologia</i> , 2021, , .	0.9	1
3	Prevalence of hypertension determined by ambulatory blood pressure monitoring (ABPM) and body composition in long-term survivors of childhood cancer. <i>Pediatric Hematology and Oncology</i> , 2018, 35, 1-10.	0.8	10
4	Uteroglobin gene polymorphism (G38A) may be a risk factor in childhood idiopathic nephrotic syndrome. <i>Pediatric Nephrology</i> , 2018, 33, 295-303.	1.7	1
5	Pediatric poststreptococcal glomerulonephritis: Clinical and laboratory data. <i>Pediatrics International</i> , 2018, 60, 645-650.	0.5	16
6	Autoimmune Polyglandular Syndrome Type 3c with Ectodermal Dysplasia, Immune Deficiency and Hemolytic-Uremic Syndrome. <i>JCRPE Journal of Clinical Research in Pediatric Endocrinology</i> , 2014, 6, 47-50.	0.9	2
7	Association of eNOS gene intron 4 a/b VNTR polymorphisms in children with nephrotic syndrome. <i>Gene</i> , 2013, 522, 192-195.	2.2	6
8	Colchicine Treatment in Children With Familial Mediterranean Fever: Is it a Risk Factor for Neuromyopathy?. <i>Pediatric Neurology</i> , 2013, 49, 417-419.	2.1	4
9	Urotensin-II: More Than a Mediator for Kidney. <i>International Journal of Nephrology</i> , 2012, 2012, 1-7.	1.3	8
10	Bartter syndrome and growth hormone deficiency: three cases. <i>Pediatric Nephrology</i> , 2012, 27, 2145-2148.	1.7	24
11	DD Genotype of ACE Gene in Boys: May it be a Risk Factor for Minimal Change Nephrotic Syndrome?. <i>Renal Failure</i> , 2012, 34, 19-23.	2.1	2
12	Disruption of PTPRO Causes Childhood-Onset Nephrotic Syndrome. <i>American Journal of Human Genetics</i> , 2011, 89, 139-147.	6.2	90
13	Urinary annexin V in children with nephrotic syndrome: a new prognostic marker?. <i>Pediatric Nephrology</i> , 2008, 23, 79-82.	1.7	9
14	Arterial thrombosis associated with factor V Leiden and methylenetetrahydrofolate reductase C677T mutation in childhood membranous glomerulonephritis. <i>Pediatric Nephrology</i> , 2008, 23, 491-494.	1.7	8
15	Changes in Osmolal Gap and Osmolality in Children with Chronic and End-Stage Renal Failure. <i>Nephron Physiology</i> , 2007, 105, p19-p21.	1.2	12
16	Ambulatory blood pressure monitoring and renal functions in children with a solitary kidney. <i>Pediatric Nephrology</i> , 2007, 22, 559-564.	1.7	62
17	ACE Gene Polymorphism in Turkish Children with Nephrotic Syndrome. <i>Renal Failure</i> , 2006, 28, 401-403.	2.1	14
18	Barttin mutations in antenatal Bartter syndrome with sensorineural deafness. <i>Pediatric Nephrology</i> , 2006, 21, 1056-1057.	1.7	12

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19	Endothelial nitric oxide synthase gene intron 4 a/b VNTR polymorphism in children with APSGN. <i>Pediatric Nephrology</i> , 2006, 21, 1661-1665.	1.7	16
20	May the Best Friend be an Enemy if not Recognized Early. <i>Pediatric Emergency Care</i> , 2005, 21, 445-448.	0.9	12
21	Aggressive angiomyxoma in a child with chronic renal failure. <i>Pediatric Surgery International</i> , 2005, 21, 563-565.	1.4	9
22	Associated anomalies in children with congenital solitary functioning kidney. <i>Pediatric Surgery International</i> , 2005, 21, 456-459.	1.4	39
23	Mitral annular calcification and brown tumor of the rib in a child with chronic renal failure. <i>Pediatric Nephrology</i> , 2005, 20, 673-675.	1.7	11
24	An unusual cause of pleural effusion, urinotorax in a child with urinary stone disease. <i>Pediatric Nephrology</i> , 2005, 20, 1487-1489.	1.7	14