Deniz Kor

List of Publications by Year
in descending order

23	158	1478505 6	1199594 12
papers	citations	h-index	g-index

27	27	27	238
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	More than tubular dysfunction: cystinosis and kidney outcomes. Journal of Nephrology, 2022, 35, 831-840.	2.0	3
2	Herediter Tirozinemi Tip-1 ve Tip-1 Diabetes Mellitus BirlikteliÄŸinde Diyet YÃ¶netimi: Olgu Sunumu. Journal of Nutrition and Dietetics, 2022, 49, 115-120.	0.2	0
3	Evaluation of bone health in patients with mucopolysaccharidosis. Journal of Bone and Mineral Metabolism, 2022, , 1.	2.7	1
4	Morquio A syndrome and effect of enzyme replacement therapy in different age groups of Turkish patients: a case series. Orphanet Journal of Rare Diseases, 2021, 16, 144.	2.7	5
5	Mucopolysaccharidosis Type-II with Pathognomonic Skin Appearance: A Case with Pebbling Sign. Journal of Pediatric Research, 2021, 8, 206-208.	0.2	0
6	Glycogen storage disease type XII; an ultra rare cause of hemolytic anemia and rhabdomyolysis: one new case report. Journal of Pediatric Endocrinology and Metabolism, 2021, 34, 1335-1339.	0.9	1
7	Mucopolysaccharidosis type VI, 9 sibling pairs and 1 set of three siblings: single center experience from Turkey. Molecular Genetics and Metabolism, 2018, 123, S101.	1.1	0
8	A case with Pallister-Killian syndrome misdiagnosed as mucopolysaccharidosis. Molecular Genetics and Metabolism, 2018, 123, S100.	1.1	0
9	Early onset alpha-mannosidosis: A Turkish case. Molecular Genetics and Metabolism, 2018, 123, S100-S101.	1.1	0
10	Improved metabolic control in tetrahydrobiopterin (BH4), responsive phenylketonuria with sapropterin administered in two divided doses vs. a single daily dose. Journal of Pediatric Endocrinology and Metabolism, 2017, 30, 713-718.	0.9	6
11	P.val452ile mutation of the slc25a13 gene in a turkish patient with citrin deficiency. Turkish Journal of Pediatrics, 2017, 59, 311.	0.6	2
12	Tyrosinemia type 1 and irreversible neurologic crisis after one month discontinuation of nitisone. Metabolic Brain Disease, 2016, 31, 1181-1183.	2.9	12
13	Propionic acidemia: a Turkish case report of a successful pregnancy, labor and lactation. Journal of Pediatric Endocrinology and Metabolism, 2016, 29, 863-6.	0.9	3
14	Brown-Vialetto-Van Laere syndrome: two siblings with a new mutation and dramatic therapeutic effect of high-dose riboflavin. Journal of Pediatric Endocrinology and Metabolism, 2016, 29, 227-31.	0.9	7
15	Homozygous familial hypobetalipoproteinemia: A Turkish case carrying a missense mutation in apolipoprotein B. Clinica Chimica Acta, 2016, 452, 185-190.	1.1	11
16	Genotypic and phenotypic features of the cystinosis patients from the South Eastern part of Turkey. Turkish Journal of Pediatrics, 2016, 58, 362-370.	0.6	3
17	A Case Report of a Very Rare Association of Tyrosinemia type I and Pancreatitis Mimicking Neurologic Crisis of Tyrosinemia Type I. Balkan Medical Journal, 2016, 33, 370-372.	0.8	6
18	Two Novel Missense Mutations in Nonketotic Hyperglycinemia. Journal of Child Neurology, 2015, 30, 789-792.	1.4	5

#	Article	IF	Citations
19	An asymptomatic mother diagnosed with 3-methylcrotonyl-CoA carboxylase deficiency after newborn screening. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 669-71.	0.9	5
20	Primary systemic carnitine deficiency: a Turkish case with a novel homozygous SLC22A5 mutation and 14 years follow-up. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 1179-81.	0.9	5
21	Blue-colored sweating: four infants with apocrine chromhidrosis. Turkish Journal of Pediatrics, 2015, 57, 290-3.	0.6	5
22	Prevalence and correlates of restless legs syndrome in adolescents. Developmental Medicine and Child Neurology, 2011, 53, 40-47.	2.1	78
23	Jansen Syndrome With Unaccountable Severe Hypercalcemia Treated by Biphosphanate. , 2010, 20, 160-161.		0