Halise Neslihan Ã–nenli Mungan

List of Publications by Year
in descending order

42 papers	1,264 citations	840119 11 h-index	360668 35 g-index
46 all docs	46 docs citations	46 times ranked	1563 citing authors

#	Article	IF	Citations
1	TAC3 and TACR3 mutations in familial hypogonadotropic hypogonadism reveal a key role for Neurokinin B in the central control of reproduction. Nature Genetics, 2009, 41, 354-358.	9.4	817
2	Loss-of-Function Mutations inPNPLA6Encoding Neuropathy Target Esterase Underlie Pubertal Failure and Neurological Deficits in Gordon Holmes Syndrome. Journal of Clinical Endocrinology and Metabolism, 2014, 99, E2067-E2075.	1.8	92
3	Wolcott-Rallison syndrome due to the same mutation (W522X) in EIF2AK3 in two unrelated families and review of the literature*. Pediatric Diabetes, 2010, 11, 279-285.	1.2	43
4	Molecular Genetic Analysis of Normosmic Hypogonadotropic Hypogonadism in a Turkish Population: Identification and Detailed Functional Characterization of a Novel Mutation in the Gonadotropin-Releasing Hormone Receptor Gene. Neuroendocrinology, 2006, 84, 301-308.	1.2	41
5	Hashimoto's encephalopathy: four cases and review of literature. International Journal of Neuroscience, 2014, 124, 302-306.	0.8	30
6	Distribution of Gene Mutations Associated with Familial Normosmic Idiopathic Hypogonadotropic Hypogonadism. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2012, 4, 121-126.	0.4	28
7	Seroprevalence of rubella in school girls and pregnant women. European Journal of Epidemiology, 2002, 18, 81-84.	2.5	21
8	Bone calcium changes during diabetic ketoacidosis: A comparison with lactic acidosis due to volume depletion. Bone, 2005, 37, 122-127.	1.4	21
9	A homozygous recurring mutation in WISP3 causing progressive pseudorheumatoid arthropathy. Journal of Pediatric Endocrinology and Metabolism, 2011, 24, 105-8.	0.4	14
10	Three Children with Triple A Syndrome due to a Mutation (R478X) in the AAAS Gene. Hormone Research in Paediatrics, 2004, 61, 3-6.	0.8	13
11	Ambulatory Blood Pressure Monitoring and Serum Nitric Oxide Concentration in Type 1 Diabetic Children. Endocrine Journal, 2009, 56, 477-485.	0.7	12
12	Chromium levels in healthy and newly diagnosed type 1 diabetic children. Pediatrics International, 2012, 54, 780-785.	0.2	12
13	Tyrosinemia type 1 and irreversible neurologic crisis after one month discontinuation of nitisone. Metabolic Brain Disease, 2016, 31, 1181-1183.	1.4	12
14	Homozygous familial hypobetalipoproteinemia: A Turkish case carrying a missense mutation in apolipoprotein B. Clinica Chimica Acta, 2016, 452, 185-190.	0.5	11
15	Multiple sulfatase deficiency: A case series of four children. Annals of Indian Academy of Neurology, 2013, 16, 720.	0.2	10
16	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.	0.4	7
17	Primary Hyperparathyroidism in an Infant with Three Parathyroid Glands and Pulmonary Calcinosis. Journal of Pediatric Endocrinology and Metabolism, 2001, 14, 1173-1176.	0.4	6
18	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.	0.4	6

#	Article	IF	Citations
19	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.	0.3	6
20	The importance of arginine mutation for the evolutionary structure and function of phenylalanine hydroxylase gene. Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis, 2006, 601, 39-45.	0.4	5
21	Two Novel Missense Mutations in Nonketotic Hyperglycinemia. Journal of Child Neurology, 2015, 30, 789-792.	0.7	5
22	An asymptomatic mother diagnosed with 3-methylcrotonyl-CoA carboxylase deficiency after newborn screening. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 669-71.	0.4	5
23	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.	0.4	5
24	Clinical findings and effect of sodium hydrogen carbonate in patients with glutathione synthetase deficiency. Journal of Pediatric Endocrinology and Metabolism, 2016, 29, 481-5.	0.4	5
25	Prospective Turkish Cohort Study to Investigate the Frequency of Niemann-Pick Disease Type C Mutations in Consanguineous Families with at Least One Homozygous Family Member. Molecular Diagnosis and Therapy, 2017, 21, 643-651.	1.6	5
26	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.	1.2	5
27	Congenital erythropoietic porphyria with erythrodontia: A case report. International Journal of Paediatric Dentistry, 2019, 29, 542-548.	1.0	4
28	The outcome of 41 Late-Diagnosed Turkish GA-1 Patients: A Candidate for the Turkish NBS. Neuropediatrics, 2021, 52, 358-369.	0.3	4
29	Bone Mineral Changes in Acute Metabolic Acidosis due to Acute Gastroenteritis. Calcified Tissue International, 2004, 75, 380-383.	1.5	3
30	A 17-Year-Old Girl with Chronic Intermittent Abdominal Pain. Pediatric Annals, 2015, 44, 139-141.	0.3	3
31	Propionic acidemia: a Turkish case report of a successful pregnancy, labor and lactation. Journal of Pediatric Endocrinology and Metabolism, 2016, 29, 863-6.	0.4	3
32	An ignored cause of red urine in children: rhabdomyolysis due to carnitine palmitoyltransferase II (CPT-II) deficiency. Journal of Pediatric Endocrinology and Metabolism, 2017, 30, 237-239.	0.4	3
33	A Desensitization Method to Maintain Enzyme Replacement Therapy in Mucopolysaccharidosis Type VI. Journal of Investigational Allergology and Clinical Immunology, 2016, 26, 130-132.	0.6	3
34	A rare cause of hydrops fetalis in two Gaucher disease type 2 patients with a novel mutation. Metabolic Brain Disease, 2022, , .	1.4	2
35	Combination of two different homozygote mutations in Pompe disease. Pediatrics International, 2016, 58, 241-243.	0.2	1
36	Clinical and molecular findings in 37 Turkish patients with isolated methylmalonic acidemia. Turkish Journal of Medical Sciences, 2021, 51, 1220-1228.	0.4	1

#	Article	IF	Citations
37	Mucopolysaccharidosis type VI, 9 sibling pairs and 1 set of three siblings: single center experience from Turkey. Molecular Genetics and Metabolism, 2018, 123, S101.	0.5	0
38	A case with Pallister-Killian syndrome misdiagnosed as mucopolysaccharidosis. Molecular Genetics and Metabolism, 2018, 123, S100.	0.5	0
39	Early onset alpha-mannosidosis: A Turkish case. Molecular Genetics and Metabolism, 2018, 123, S100-S101.	0.5	0
40	Mucopolysaccharidosis Type-II with Pathognomonic Skin Appearance: A Case with Pebbling Sign. Journal of Pediatric Research, 2021, 8, 206-208.	0.1	0
41	Coexistence of molybdenum cofactor deficiency type A and hypertrophic pyloric stenosis, a new case. Turk Pediatri Arsivi, 2020, 56, 78-80.	0.9	0
42	Herediter Tirozinemi Tip-1 ve Tip-1 Diabetes Mellitus BirlikteliÄŸinde Diyet YÃ¶netimi: Olgu Sunumu. Journal of Nutrition and Dietetics, 2022, 49, 115-120.	0.1	0