Asburce Olgac

List of Publications by Year
in descending order

25	77	1684188 5	1588992 8
papers	citations	h-index	g-index

27	27	27	98
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Congenital Rare Diseases Causing Persistent Diarrhea in the Newborn: A Single Center Experience. Zeitschrift Fur Geburtshilfe Und Neonatologie, 2022, , .	0.4	0
2	Genotypic and phenotypic features in Turkish patients with classic nonketotic hyperglycinemia. Metabolic Brain Disease, 2021, 36, 1213-1222.	2.9	5
3	Retrospective evaluation of patients with X-linked adrenoleukodystrophy with a wide range of clinical presentations: a single center experience. Journal of Pediatric Endocrinology and Metabolism, 2021, 34, 1169-1179.	0.9	1
4	MAN1B1-CDG: novel patients and novel variant. Journal of Pediatric Endocrinology and Metabolism, 2021, 34, 1207-1209.	0.9	3
5	Oxidative Stress in IntoxÄ±cation Type Inborn Errors of Metabolism using Thiol-Disulfide Ratio. Journal of the College of Physicians and Surgeons--Pakistan: JCPSP, 2021, 31, 663-667.	0.4	2
6	The Evaluation of Skeletal Manifestations in Patients with Gaucher Disease. Journal of Pediatric Research, 2021, 8, 257-261.	0.2	0
7	The COVID-19 Pandemic and Enzyme Replacement Therapy in Lysosomal Storage Disorders. Journal of Pediatric Research, 2021, 8, 370-376.	0.2	0
8	A rare case of primary coenzyme Q10 deficiency due to COQ9 mutation. Journal of Pediatric Endocrinology and Metabolism, 2020, 33, 165-170.	0.9	15
9	A rare urea cycle disorder in a neonate: N-acetylglutamate synthetase deficiency. Archivos Argentinos De Pediatria, 2020, 118, e545-e548.	0.2	1
10	Beneficial Effects of Modified Atkins Diet in Glycogen Storage Disease Type IIIa. Annals of Nutrition and Metabolism, 2020, 76, 233-241.	1.9	12
11	Posterior fossa horns; a new calvarial finding of mucopolysaccharidoses with well-known cranial MRI features. Turkish Journal of Medical Sciences, 2020, 50, 1048-1061.	0.9	3
12	Carbonic anhydrase VA deficiency: a very rare case of hyperammonemic encephalopathy. Journal of Pediatric Endocrinology and Metabolism, 2020, 33, 1349-1352.	0.9	6
13	Lysinuric protein intolerance: an overlooked diagnosis. Egyptian Journal of Medical Human Genetics, 2020, 21, .	1.0	3
14	An infant with an extremely rare cobalamin disorder: methionine synthase deficiency and importance of early diagnosis and treatment. Turkish Journal of Pediatrics, 2019, 61, 282.	0.6	2
15	Vitamin D Levels and Bone Mineral Density in Inborn Errors of Metabolism Requiring Specialised Diets. Journal of the College of Physicians and Surgeons--Pakistan: JCPSP, 2019, 29, 1207-1211.	0.4	1
16	Patient With Niemann-Pick Type C Presenting With a Jaw Mass Characterized With Lymph Node Involvement by Niemann-Pick Cells. Journal of Pediatric Hematology/Oncology, 2018, 40, 243-245.	0.6	0
17	Acute Stroke In A Patient With Mucopolysaccharidosis Type 1 With Increased Carotis Intima Media Thickness. Eurasian Journal of Medicine and Oncology, 2018, , .	1.0	2
18	Encephalocraniocutaneous lipomatosis with Wilms' tumor. Pediatrics International, 2017, 59, 835-836.	0.5	1

#	Article	IF	Citations
19	A Newborn with an Alternative Porto-Caval Shunt. Polski Przeglad Radiologii I Medycyny Nuklearnej, 2017, 82, 320-321.	1.0	1
20	Diagnostic Dilemma: Osteopetrosis with superimposed rickets causing Neonatal Hypocalcemia. Journal of Tropical Pediatrics, 2015, 61, 146-150.	1.5	3
21	Recurrence of carbamoyl phosphate synthetase 1 (CPS1) deficiency in Turkish patients: Characterization of a founder mutation by use of recombinant CPS1 from insect cells expression. Molecular Genetics and Metabolism, 2014, 113, 267-273.	1.1	8
22	Successful Treatment of ICE-Rituximab Chemotherapy and Subsequent Bone Marrow Transplantation in a Patient With Early-relapse Burkitt Leukemia and Inverted Duplication of 1q. Journal of Pediatric Hematology/Oncology, 2012, 34, e84-e85.	0.6	1
23	Hyperprolactinemia as a Rare Cause of Hypertension in Chronic Renal Failure. Renal Failure, 2012, 34, 792-794.	2.1	2
24	Acute Disseminated Encephalomyelitis Associated With Acute Rheumatic Fever. Pediatric Neurology, 2011, 44, 233-235.	2.1	5
25	â€œDouble Hitâ€•Homozygous Mutations for Two Different Rare Inborn Errors of Metabolism: A Burden for Countries with High Prevalences of Consangineous Marriages. Journal of Pediatric Research, 0, , 47-50.	0.2	0