

Cigdem Aktuglu-Zeybek

List of Publications by Year in descending order

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Version: 2024-02-01

29
papers

254
citations

1039406

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h-index

1058022

14
g-index

29
all docs

29
docs citations

29
times ranked

375
citing authors

#	ARTICLE	IF	CITATIONS
1	Hereditary tyrosinemia type 1 in Turkey: Twenty year single-center experience. <i>Pediatrics International</i> , 2015, 57, 281-289.	0.2	27
2	The effect of low-carbohydrate diet on left ventricular diastolic function in obese children. <i>Pediatrics International</i> , 2010, 52, 218-223.	0.2	21
3	Tissue Doppler echocardiographic assessment of cardiac function in children with bronchial asthma. <i>Pediatrics International</i> , 2007, 49, 911-917.	0.2	20
4	Right Ventricular Subclinical Diastolic Dysfunction in Obese Children: The Effect of Weight Reduction with a Low-Carbohydrate Diet. <i>Pediatric Cardiology</i> , 2009, 30, 946-953.	0.6	18
5	Early diagnosed cerebrotendinous xanthomatosis patients: clinical, neuroradiological characteristics and therapy results of a single center from Turkey. <i>Acta Neurologica Belgica</i> , 2019, 119, 343-350.	0.5	18
6	Capillary leak syndrome in a 5-month-old infant associated with intractable diarrhoea. <i>Annals of Tropical Paediatrics</i> , 2007, 27, 81-86.	1.0	16
7	Bitterness of Glucose/Galactose. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2014, 58, 57-60.	0.9	14
8	The impact of continuous renal replacement therapy for metabolic disorders in infants. <i>Pediatrics and Neonatology</i> , 2018, 59, 85-90.	0.3	12
9	Determination of NTBC in serum samples from patients with hereditary tyrosinemia type I by capillary electrophoresis†. <i>Talanta</i> , 2010, 80, 1846-1848.	2.9	11
10	Evaluation of dynamic thiol/disulphide homeostasis as a novel indicator of oxidative stress in maple syrup urine disease patients under treatment. <i>Metabolic Brain Disease</i> , 2017, 32, 179-184.	1.4	10
11	Oxidative stress among L-2-hydroxyglutaric aciduria disease patients: evaluation of dynamic thiol/disulfide homeostasis. <i>Metabolic Brain Disease</i> , 2019, 34, 283-288.	1.4	9
12	Challenges of following patients with inherited metabolic diseases during the COVID-19 outbreak. A cross-sectional online survey study. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2021, 34, 103-107.	0.4	9
13	Hereditary Tyrosinemia Type 1 in Turkey. <i>Advances in Experimental Medicine and Biology</i> , 2017, 959, 157-172.	0.8	8
14	Inborn errors of metabolism and coronavirus disease 2019: Evaluation of the metabolic outcome. <i>Pediatrics International</i> , 2022, 64, .	0.2	6
15	Screening of Free Carnitine and Acylcarnitine Status in Children With Familial Mediterranean Fever. <i>Archives of Rheumatology</i> , 2016, 31, 133-138.	0.3	5
16	Coagulation Disturbances in Patients with Argininemia. <i>Acta Haematologica</i> , 2018, 140, 221-225.	0.7	5
17	Continuous Renal Replacement Therapy with High Flow Rate Can Effectively, Safely, and Quickly Reduce Plasma Ammonia and Leucine Levels in Children. <i>Children</i> , 2019, 6, 53.	0.6	5
18	Multimodal imaging including optical coherence tomography angiography in patients with type B Niemann-Pick disease. <i>International Ophthalmology</i> , 2019, 39, 2545-2552.	0.6	5

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19	Impact of sodium phenylbutyrate treatment in acute management of maple syrup urine disease attacks: a single-center experience. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2021, 34, 121-126.	0.4	5
20	Treatment of maple syrup urine disease with high flow hemodialysis in a neonate. <i>Turkish Journal of Pediatrics</i> , 2019, 61, 107.	0.3	5
21	Altered immune response in organic acidemia. <i>Pediatrics International</i> , 2022, 64, .	0.2	5
22	Screening for Fabry Disease in Patients With Juvenile Systemic Lupus Erythematosus. <i>Archives of Rheumatology</i> , 2020, 35, 7-12.	0.3	4
23	Long-term N-carbamylglutamate treatment of hyperammonemia in patients with classic organic acidemias. <i>Molecular Genetics and Metabolism Reports</i> , 2021, 26, 100715.	0.4	4
24	Glutaric acidemia type II patient with thalassemia minor and novel electron transfer flavoprotein-A gene mutations: A case report and review of literature. <i>World Journal of Clinical Cases</i> , 2018, 6, 786-790.	0.3	4
25	Identifying and elucidating the roles of Y198N and Y204F mutations in the PAH enzyme through molecular dynamic simulations. <i>Journal of Biomolecular Structure and Dynamics</i> , 2021, , 1-12.	2.0	3
26	COVID-19 triggered encephalopathic crisis in a patient with glutaric aciduria type 1. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2021, 34, 1611-1614.	0.4	3
27	Evaluation of the effect of chenodeoxycholic acid treatment on skeletal system findings in cerebrotendinous xanthomatosis patients. <i>Turk Pediatri Arsivi</i> , 2019, 54, 113-118.	0.9	2
28	Evaluation of dynamic thiol/disulfide homeostasis in hereditary tyrosinemia type 1 patients. <i>Pediatric Research</i> , 2021, , .	1.1	0
29	Screening Inherited Metabolic Disorders in Patients with Familial Mediterranean Fever. <i>Journal of Pediatric Research</i> , 2015, 2, 201-205.	0.1	0