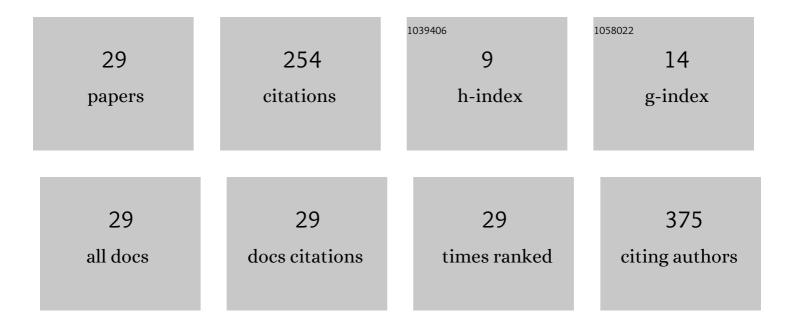
Cigdem Aktuglu-Zeybek

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Inborn errors of metabolism and coronavirus disease 2019: Evaluation of the metabolic outcome. Pediatrics International, 2022, 64, .	0.2	6
2	Altered immune response in organic acidemia. Pediatrics International, 2022, 64, .	0.2	5
3	Long-term N-carbamylglutamate treatment of hyperammonemia in patients with classic organic acidemias. Molecular Genetics and Metabolism Reports, 2021, 26, 100715.	0.4	4
4	Identifying and elucidating the roles of Y198N and Y204F mutations in the PAH enzyme through molecular dynamic simulations. Journal of Biomolecular Structure and Dynamics, 2021, , 1-12.	2.0	3
5	COVID-19 triggered encephalopathic crisis in a patient with glutaric aciduria type 1. Journal of Pediatric Endocrinology and Metabolism, 2021, 34, 1611-1614.	0.4	3
6	Impact of sodium phenylbutyrate treatment in acute management of maple syrup urine disease attacks: a single-center experience. Journal of Pediatric Endocrinology and Metabolism, 2021, 34, 121-126.	0.4	5
7	Challenges of following patients with inherited metabolic diseases during the COVID-19 outbreak. A cross-sectional online survey study. Journal of Pediatric Endocrinology and Metabolism, 2021, 34, 103-107.	0.4	9
8	Evaluation of dynamic thiol/disulfide homeostasis in hereditary tyrosinemia type 1 patients. Pediatric Research, 2021, , .	1.1	0
9	Screening for Fabry Disease in Patients With Juvenile Systemic Lupus Erythematosus. Archives of Rheumatology, 2020, 35, 7-12.	0.3	4
10	Continuous Renal Replacement Therapy with High Flow Rate Can Effectively, Safely, and Quickly Reduce Plasma Ammonia and Leucine Levels in Children. Children, 2019, 6, 53.	0.6	5
11	Multimodal imaging including optical coherence tomography angiography in patients with type B Niemann–Pick disease. International Ophthalmology, 2019, 39, 2545-2552.	0.6	5
12	Oxidative stress among L-2-hydroxyglutaric aciduria disease patients: evaluation of dynamic thiol/disulfide homeostasis. Metabolic Brain Disease, 2019, 34, 283-288.	1.4	9
13	Early diagnosed cerebrotendinous xanthomatosis patients: clinical, neuroradiological characteristics and therapy results of a single center from Turkey. Acta Neurologica Belgica, 2019, 119, 343-350.	0.5	18
14	Treatment of maple syrup urine disease with high flow hemodialysis in a neonate. Turkish Journal of Pediatrics, 2019, 61, 107.	0.3	5
15	Evaluation of the effect of chenodeoxycholic acid treatment on skeletal system findings in cerebrotendinous xanthomatosis patients. Turk Pediatri Arsivi, 2019, 54, 113-118.	0.9	2
16	The impact of continuous renal replacement therapy for metabolic disorders in infants. Pediatrics and Neonatology, 2018, 59, 85-90.	0.3	12
17	Coagulation Disturbances in Patients with Argininemia. Acta Haematologica, 2018, 140, 221-225.	0.7	5
18	Glutaric acidemia type II patient with thalassemia minor and novel electron transfer flavoprotein-A gene mutations: A case report and review of literature. World Journal of Clinical Cases, 2018, 6, 786-790.	0.3	4

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#	Article	IF	CITATIONS
19	Hereditary Tyrosinemia Type 1 in Turkey. Advances in Experimental Medicine and Biology, 2017, 959, 157-172.	0.8	8
20	Evaluation of dynamic thiol/disulphide homeostasis as a novel indicator of oxidative stress in maple syrup urine disease patients under treatment. Metabolic Brain Disease, 2017, 32, 179-184.	1.4	10
21	Screening of Free Carnitine and Acylcarnitine Status in Children With Familial Mediterranean Fever. Archives of Rheumatology, 2016, 31, 133-138.	0.3	5
22	Hereditary tyrosinemia type 1 in <scp>T</scp> urkey: Twenty year singleâ€center experience. Pediatrics International, 2015, 57, 281-289.	0.2	27
23	Screening Inherited Metabolic Disorders in Patients with Familial Mediterranean Fever. Journal of Pediatric Research, 2015, 2, 201-205.	0.1	0
24	Bitterness of Glucose/Galactose. Journal of Pediatric Gastroenterology and Nutrition, 2014, 58, 57-60.	0.9	14
25	The effect of lowâ€carbohydrate diet on left ventricular diastolic function in obese children. Pediatrics International, 2010, 52, 218-223.	0.2	21
26	Determination of NTBC in serum samples from patients with hereditary tyrosinemia type I by capillary electrophoresisâ~†. Talanta, 2010, 80, 1846-1848.	2.9	11
27	Right Ventricular Subclinical Diastolic Dysfunction in Obese Children: The Effect of Weight Reduction with a Low-Carbohydrate Diet. Pediatric Cardiology, 2009, 30, 946-953.	0.6	18
28	Capillary leak syndrome in a 5-month-old infant associated with intractable diarrhoea. Annals of Tropical Paediatrics, 2007, 27, 81-86.	1.0	16
29	Tissue Doppler echocardiographic assessment of cardiac function in children with bronchial asthma. Pediatrics International 2007 49 911-917	0.2	20