

Behzat Ozkan

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/6775873/publications.pdf>

Version: 2024-02-01

28
papers

219
citations

1478280

6
h-index

1058333

14
g-index

28
all docs

28
docs citations

28
times ranked

401
citing authors

#	ARTICLE	IF	CITATIONS
1	Vitamin D intoxication. Turkish Journal of Pediatrics, 2012, 54, 93-8.	0.3	55
2	Novel CYP27B1 Gene Mutations in Patients with Vitamin D-Dependent Rickets Type 1A. PLoS ONE, 2015, 10, e0131376.	1.1	37
3	Molecular diagnosis of maturity-onset diabetes of the young (MODY) in Turkish children by using targeted next-generation sequencing. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 1265-71.	0.4	28
4	Intragastric alendronate therapy in two infants with vitamin D intoxication: A new method. Clinical Toxicology, 2008, 46, 300-302.	0.8	17
5	Melanocortin-4 Receptor Gene Mutations in a Group of Turkish Obese Children and Adolescents. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2017, 9, 216-221.	0.4	11
6	Detection of <i>SHOX</i> Gene Variations in Patients with Skeletal Abnormalities with or without Short Stature. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 358-365.	0.4	11
7	Comparison of Treatment Regimens in Management of Severe Hypercalcemia Due to Vitamin D Intoxication in Children. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2019, 11, 140-148.	0.4	8
8	Ultrasonographic determination of thyroid volume in infants and children from Aegean region of Turkey and comparison with national and international references. Journal of Pediatric Endocrinology and Metabolism, 2021, 34, 457-464.	0.4	7
9	A novel mutation of AMH in three siblings with persistent Mullerian duct syndrome. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 1379-82.	0.4	6
10	Association of Wolfram syndrome with Fallot tetralogy in a girl. Archivos Argentinos De Pediatria, 2016, 114, e163-6.	0.3	6
11	Genetic Diagnosis Using Whole Exome Analysis in Two Cases with Malignant Osteopetrosis of Infancy. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2015, 7, 356-357.	0.4	5
12	Three Siblings with Idiopathic Hypogonadotropic Hypogonadism in a Nonconsanguineous Family: A Novel <i>KISS1R/GPR54</i> Loss-of-Function Mutation. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2019, 11, 444-448.	0.4	5
13	A novel splice site mutation of FGD1 gene in an Aarskog-Scott syndrome patient with a large anterior fontanel. Journal of Pediatric Endocrinology and Metabolism, 2016, 29, 1111-4.	0.4	4
14	Long-term monitoring of Graves' disease in children and adolescents: a single-center experience. Turkish Journal of Medical Sciences, 2019, 49, 464-471.	0.4	4
15	Infantile-onset thiamine responsive megaloblastic anemia syndrome with SLC19A2 mutation: a case report. Archivos Argentinos De Pediatria, 2017, 115, e153-e156.	0.3	4
16	Screening of non-syndromic early-onset child and adolescent obese patients in terms of <i>LEP</i> , <i>LEPR</i> , <i>MC4R</i> and <i>POMC</i> gene variants by next-generation sequencing. Journal of Pediatric Endocrinology and Metabolism, 2022, 35, 1041-1050.	0.4	4
17	A Rare Complication of Insulin Therapy in a Child with Newly Diagnosed Type 1 Diabetes: Insulin Edema. Journal of Pediatric Research, 2021, 8, 506-509.	0.1	2
18	Higher-Than-Conventional Subcutaneous Regular Insulin Doses in Diabetic Ketoacidosis in Children and Adolescents. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2017, 9, 132-137.	0.4	1

#	ARTICLE	IF	CITATIONS
19	Identification of two AMH gene variants in two unrelated patients with persistent Müllerian duct syndrome: one novel variant. <i>Gynecological Endocrinology</i> , 2021, 37, 476-479.	0.7	1
20	Association Between Thyroid Parenchymal Echogenicity and Thyroid Function in Pediatric Population. <i>Iranian Journal of Pediatrics</i> , 2022, 32, .	0.1	1
21	Ultrasonographic measurements of the testicular volume in Turkish boys aged 0-8 years and comparison with international references. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2022, 35, 489-495.	0.4	1
22	Analysis of New Biomarkers for the Diagnosis of Polycystic Ovary Syndrome in Adolescents. <i>Guncel Pediatri</i> , 2021, 19, 311-318.	0.1	1
23	Outcomes of Dyslipidemia Screening Program in School-aged Children. <i>Journal of Pediatric Research</i> , 2021, 8, 155-160.	0.1	0
24	Investigating the Efficiency of Vitamin D Administration with Buccal Spray in the Treatment of Vitamin D Deficiency in Children and Adolescents. <i>JCRPE Journal of Clinical Research in Pediatric Endocrinology</i> , 2021, 13, 426-432.	0.4	0
25	Spectrum of Clinical Manifestations in Turkish Patients with Williams-Beuren Syndrome: A Monocentric Study. <i>Journal of Pediatric Research</i> , 2021, 8, 297-302.	0.1	0
26	A Case of Late-onset Hyperinsulinemic Hypoglycemia: HNF4A Mutation. <i>Journal of Pediatric Research</i> , 2020, 7, 168-171.	0.1	0
27	Detection of Gene Variations in Patients with Skeletal Abnormalities with or without Short Stature. <i>JCRPE Journal of Clinical Research in Pediatric Endocrinology</i> , 2020, 12, 358-365.	0.4	0
28	Sonographic evaluation of normal thyroid volume and thyroid isthmus depth among infants in the west coast of Turkey. <i>Endokrynologia Polska</i> , 2022, , .	0.3	0