

Doga Turkkahraman

List of Publications by Year in descending order

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Version: 2024-02-01

14
papers

177
citations

1307594

7
h-index

1125743

13
g-index

14
all docs

14
docs citations

14
times ranked

264
citing authors

#	ARTICLE	IF	CITATIONS
1	Serum alpha-melanocyte-stimulating hormone (a-MSH), brain-derived neurotrophic factor (BDNF), and agouti-related protein (AGRP) levels in children with Prader-Willi or Bardet-Biedl syndromes. <i>Journal of Endocrinological Investigation</i> , 2022, , 1.	3.3	4
2	A novel <i>EZH2</i> gene variant in a case of Weaver syndrome with postaxial polydactyly. <i>American Journal of Medical Genetics, Part A</i> , 2021, 185, 2234-2237.	1.2	3
3	A Case of Congenital Central Hypothyroidism Caused by a Novel Variant (Gln1255Ter) in <i>IGSF1</i> . <i>JCRPE Journal of Clinical Research in Pediatric Endocrinology</i> , 2021, 13, 353-357.	0.9	3
4	Novel Mutations in Obesity-related Genes in Turkish Children with Non-syndromic Early Onset Severe Obesity: A Multicentre Study. <i>JCRPE Journal of Clinical Research in Pediatric Endocrinology</i> , 2019, 11, 341-349.	0.9	20
5	A Large <i>PROP1</i> Gene Deletion in a Turkish Pedigree. <i>Case Reports in Endocrinology</i> , 2018, 2018, 1-5.	0.4	3
6	Basal Serum Neurokinin B Levels in Differentiating Idiopathic Central Precocious Puberty from Premature Thelarche. <i>JCRPE Journal of Clinical Research in Pediatric Endocrinology</i> , 2017, 9, 101-105.	0.9	5
7	<i>AVP-NP11</i> gene mutations and clinical characteristics of the patients with autosomal dominant familial central diabetes insipidus. <i>Pituitary</i> , 2015, 18, 898-904.	2.9	21
8	High-Dose Hook Effect in 17-Hydroxyprogesterone Assay in a Patient with 21-Hydroxylase Deficiency. <i>JCRPE Journal of Clinical Research in Pediatric Endocrinology</i> , 2015, 7, 329-332.	0.9	5
9	Letter to the Editor regarding "Testotoxicosis: Report of Two Cases, One with a Novel Mutation in Luteinizing Hormone/Choriogonadotropin Receptor Gene". <i>JCRPE Journal of Clinical Research in Pediatric Endocrinology</i> , 2015, 7, 355-355.	0.9	0
10	A novel <i>DAX-1</i> mutation presented with precocious puberty and hypogonadotropic hypogonadism in different members of a large pedigree. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2013, 26, 551-5.	0.9	13
11	P300 auditory event-related potentials in children with obesity: is childhood obesity related to impairment in cognitive functions?. <i>Pediatric Diabetes</i> , 2011, 12, 589-595.	2.9	34
12	Analysis of <i>TPO</i> gene in Turkish children with iodide organification defect: identification of a novel mutation. <i>Endocrine</i> , 2010, 37, 124-128.	2.3	10
13	Final Diagnosis in Children with Subclinical Hypothyroidism and Mutation Analysis of the Thyroid Peroxidase Gene (<i>TPO</i>). <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2009, 22, 845-51.	0.9	10
14	Permanent Neonatal Diabetes Mellitus Caused by a Novel Homozygous (T168A) Glucokinase (<i>GCK</i>) Mutation: Initial Response to Oral Sulphonylurea Therapy. <i>Journal of Pediatrics</i> , 2008, 153, 122-126.	1.8	46