Doga Turkkahraman

List of Publications by Year
in descending order

14	177	1307594 7	1125743 13
papers	citations	h-index	g-index

14	14	14	264
all docs	docs citations	times ranked	citing authors

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