Eda Mengen

List of Publications by Year
in descending order

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		1163117	940533
21	276	8	16
papers	citations	h-index	g-index

21	21	21	418
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	<i>DLG2</i> Mutations in the Etiology of Pubertal Delay and Idiopathic Hypogonadotropic Hypogonadism. Hormone Research in Paediatrics, 2021, 94, 364-368.	1.8	2
2	The Significance of Thiol/Disulfide Homeostasis and Ischemia-modified Albumin Levels in Assessing Oxidative Stress in Obese Children and Adolescents. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 45-54.	0.9	7
3	A Rare Etiology of 46,XY Disorder of Sex Development and Adrenal Insufficiency: A Case of MIRAGE Syndrome Caused by Mutations in the <i>SAMD9</i> Gene. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 206-211.	0.9	9
4	A Duplication Upstream of SOX9 Associated with <i>SRY</i> Negative 46,XX Ovotesticular Disorder of Sex Development: A Case Report. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 308-314.	0.9	4
5	Evaluation of the relationship between the one-hour plasma glucose concentration and beta-cell functions and cardiometabolic parameters during oral glucose tolerance test in obese children and adolescents. Journal of Pediatric Endocrinology and Metabolism, 2020, 33, 767-775.	0.9	1
6	Catecholamine-induced Myocarditis in a Child with Pheochromocytoma. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 202-205.	0.9	3
7	Mutations Within the Transcription Factor <i>PROP1</i> in a Cohort of Turkish Patients with Combined Pituitary Hormone Deficiency. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 261-268.	0.9	4
8	The Effects of Risk Behaviors and Orthorexic Behavior on Glycemic Control in Adolescents with Type 1 Diabetes. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 233-240.	0.9	1
9	The Effects of Risk Behaviors and Orthorexic Behavior on Glycemic Control in Adolescents with Type 1 Diabetes. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 233-240.	0.9	3
10	Clinical and Laboratory Characteristics of Hyperprolactinemia in Children and Adolescents: National Survey. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2019, 11, 149-156.	0.9	13
11	A Rare Etiology of 46, XY Disorder of Sex Development and Adrenal Insufficiency: A case of MIRAGE syndrome caused by mutations in SAMD9 gene. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2019, .	0.9	1
12	Subclinical Myocardial Dysfunction Demonstrated by Speckle Tracking Echocardiography in Children with Euthyroid Hashimotoâ€™s Thyroiditis. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2019, 11, 410-418.	0.9	5
13	Dynamic Thiol/Disulphide Homeostasis in Children and Adolescents with Non-Autoimmune Subclinical Hypothyroidism. Medical Principles and Practice, 2018, 27, 44-48.	2.4	2
14	Evaluation of the Children with Hypoparathyroidism and Pseudohypoparathyroidism Presenting to an Endocrinology Outpatient Clinic. Journal of Ankara University Faculty of Medicine, 2018, 71, 212-216.	0.1	0
15	CCDC141 Mutations in Idiopathic Hypogonadotropic Hypogonadism. Journal of Clinical Endocrinology and Metabolism, 2017, 102, 1816-1825.	3.6	33
16	A novel genetic mutation in a Turkish family with GCK-MODY. International Journal of Diabetes in Developing Countries, 2017, 37, 323-326.	0.8	1
17	Complete Idiopathic Hypogonadotropic Hypogonadism due to Homozygous <i>GNRH1</i> Mutations in the Mutational Hot Spots in the Region Encoding the Decapeptide. Hormone Research in Paediatrics, 2016, 85, 107-111.	1.8	18
18	CCDC141 Mutation Identified in Anosmic Hypogonadotropic Hypogonadism (Kallmann Syndrome) Alters GnRH Neuronal Migration. Endocrinology, 2016, 157, 1956-1966.	2.8	47

#	Article	IF	Citations
19	Effects of methylphenidate on appetite and growth in children diagnosed with attention deficit and hyperactivity disorder. Journal of Pediatric Endocrinology and Metabolism, 2016, 29, 85-92.	0.9	25
20	Mutations in FEZF1 Cause Kallmann Syndrome. American Journal of Human Genetics, 2014, 95, 326-331.	6.2	69
21	Distribution of Gene Mutations Associated with Familial Normosmic Idiopathic Hypogonadotropic Hypogonadism. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2012, 4, 121-126.	0.9	28