Nursel Muratoglu Sahin

List of Publications by Year
in descending order

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Version: 2024-02-01

15	114	1683354 5	1372195 10
papers	citations	h-index	g-index

15	15	15	266
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Investigation of MKRN3 Mutation in Patients with Familial Central Precocious Puberty. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2018, 10, 223-229.	0.4	22
2	Clinical and genetic characterisation of a series of patients with triple A syndrome. European Journal of Pediatrics, 2018, 177, 363-369.	1.3	20
3	A Rare Cause of Short Stature: 3M Syndrome in a Patient with Novel Mutation in OBSL1 Gene. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2017, 9, 91-94.	0.4	14
4	AMH levels at central precocious puberty and premature thelarche: is it a parameter?. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 1351-6.	0.4	13
5	How does thiol/disulfide homeostasis change in children with type 1 diabetes mellitus?. Diabetes Research and Clinical Practice, 2019, 149, 64-68.	1.1	8
6	Cushing syndrome related to leukemic infiltration of the central nervous system: a case report and a possible role of LIF. Journal of Pediatric Endocrinology and Metabolism, 2013, 26, 967-70.	0.4	6
7	Subnormal Growth Velocity and Related Factors During GnRH Analog Therapy for Idiopathic Central Precocious Puberty. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2018, 10, 239-246.	0.4	6
8	Congenital Hypothyroidism and Bone Remodeling Cycle. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2017, 9, 106-110.	0.4	6
9	OGTT results in obese adolescents with normal HOMA-IR values. Journal of Pediatric Endocrinology and Metabolism, 2013, 26, 285-91.	0.4	5
10	Î²-3AR W64R Polymorphism and 30-Minute Post-Challenge Plasma Glucose Levels in Obese Children. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2015, 7, 7-12.	0.4	5
11	Congenital hyperinsulinism in a newborn with a novel homozygous mutation (p.Q392H) in the ABCC8 gene. Journal of Pediatric Endocrinology and Metabolism, 2014, 27, 1253-5.	0.4	3
12	Vaginal bleeding and a giant ovarian cyst in an infant with 21-hydroxylase deficiency. Journal of Pediatric Endocrinology and Metabolism, 2018, 31, 229-233.	0.4	3
13	AntimÃ¼llerian Hormone Levels of Infants with Premature Thelarche. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2019, 11, 287-292.	0.4	2
14	Experience of intravenous calcium treatment and long-term responses to treatment in a patient with hereditary vitamin D-resistant rickets resulting from a novel mutation. Journal of Pediatric Endocrinology and Metabolism, 2019, 32, 647-651.	0.4	1
15	Type 1 rhizomelic chondrodysplasia punctata with a homozygous PEX7 mutation. Journal of Pediatric Endocrinology and Metabolism, 2017, 30, 889-892.	0.4	0