Ayfer AlikaÅifoÄlu

List of Publications by Year
in descending order

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30 papers	416 citations	933447 10 h-index	794594 19 g-index
30 all docs	30 docs citations	30 times ranked	663 citing authors

#	Article	IF	Citations
1	Feminizing Adrenocortical Tumors as a Rare Etiology of Isosexual/Contrasexual Pseudopuberty. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2022, 14, 17-28.	0.9	2
2	Management of prolactinomas in children and adolescents; which factors define the response to treatment?. Pituitary, 2022, 25, 167-179.	2.9	8
3	Alphaâ€Melanocyteâ€“Stimulating Hormone is Elevated in Hypothalamic Obesity Associated with Childhood Craniopharyngioma. Obesity, 2021, 29, 402-408.	3.0	5
4	Adrenocortical tumours in children: a review of surgical management at a tertiary care centre. ANZ Journal of Surgery, 2021, 91, 992-999.	0.7	5
5	Which parameters predict the beneficial effect of GnRHa treatment on height in girls with central precocious puberty?. Clinical Endocrinology, 2021, 94, 804-810.	2.4	10
6	Approach to pheochromocytoma and paraganglioma in children and adolescents: A retrospective clinical study from a tertiary care center. Journal of Pediatric Urology, 2021, 17, 400.e1-400.e7.	1.1	1
7	Basal Serum Thyroxine Level should Guide Initial Thyroxine Replacement Dose in Neonates with Congenital Hypothyroidism. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, 13, 269-275.	0.9	0
8	Long-term effect of conventional phosphate and calcitriol treatment on metabolic recovery and catch-up growth in children with PHEX mutation. Journal of Pediatric Endocrinology and Metabolism, 2021, 34, 1573-1584.	0.9	3
9	Long-term effects of GnRH agonist treatment on body mass index in girls with idiopathic central precocious puberty. Journal of Pediatric Endocrinology and Metabolism, 2020, 33, 99-105.	0.9	20
10	Obstructive sleep apnea in children with hypothalamic obesity: Evaluation of possible related factors. Pediatric Pulmonology, 2020, 55, 3532-3540.	2.0	2
11	Hyperinsulinemic Hypoglycemia in a Patient with Costello Syndrome: An Etiology to Consider in Hypoglycemia. Molecular Syndromology, 2020, 11, 207-216.	0.8	5
12	Novel insights into diabetes mellitus due to <scp>DNAJC3</scp> â€• defect: Evolution of neurological and endocrine phenotype in the pediatric age group. Pediatric Diabetes, 2020, 21, 1176-1182.	2.9	9
13	Poikiloderma with Neutropenia, Clericuzio-Type Accompanied by Loss of Digits Due to Severe Osteomyelitis. Journal of Clinical Immunology, 2020, 40, 934-939.	3.8	1
14	Clinical and Molecular Analysis in 2 Families With Novel Compound Heterozygous SBP2 (SECISBP2) Mutations. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e6-e11.	3.6	9
15	Treatment response to long term antiresorptive therapy in osteogenesis imperfecta type VI: does genotype matter?. Journal of Pediatric Endocrinology and Metabolism, 2020, 33, 1617-1624.	0.9	1
16	Treatment with Depot Leuprolide Acetate in Girls with Idiopathic Precocious Puberty: What Parameter should be Used in Deciding on the Initial Dose?. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 37-44.	0.9	5
17	Central nervous system imaging in girls with central precocious puberty: when is necessary?. Archives of Endocrinology and Metabolism, 2020, 64, 591-596.	0.6	2
18	Can having a sibling with type 1 diabetes cause disordered eating behaviors?. Journal of Pediatric Endocrinology and Metabolism, 2018, 31, 711-716.	0.9	5

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19	Novel and prevalent CYP11B1 gene mutations in Turkish patients with 11-Î² hydroxylase deficiency. Journal of Steroid Biochemistry and Molecular Biology, 2017, 165, 57-63.	2.5	26
20	Clinical, genetic, and structural basis of congenital adrenal hyperplasia due to 11Î²-hydroxylase deficiency. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, E1933-E1940.	7.1	106
21	Clinical and laboratory parameters predicting a requirement for the reevaluation of growth hormone status during growth hormone treatment. Growth Hormone and IGF Research, 2017, 34, 31-37.	1.1	13
22	Changing Etiological Trends in Male Precocious Puberty: Evaluation of 100 Cases with Central Precocious Puberty over the Last Decade. Hormone Research in Paediatrics, 2015, 83, 340-344.	1.8	28
23	Severe Undervirilisation in a 46,XY Case Due to a Novel Mutation in HSD17B3 Gene. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2015, 7, 249-252.	0.9	6
24	Need for Comprehensive Hormonal Workup in the Management of Adrenocortical Tumors in Children. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2014, 6, 68-73.	0.9	9
25	17Î²-Hydroxysteroid dehydrogenase type 3 deficiency as a result of a homozygous 7 base pair deletion in 17Î²HSD3 gene. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, 561-3.	0.9	13
26	The Relationship Between Serum Adiponectin, Tumor Necrosis Factor-Alpha, Leptin Levels and Insulin Sensitivity in Childhood and Adolescent Obesity: Adiponectin is a Marker of Metabolic Syndrome. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2009, 1, 233-239.	0.9	46
27	Bone mineral density and serum bone turnover markers in survivors of childhood acute lymphoblastic leukemia: Comparison of megadose methylprednisolone and conventional-dose prednisolone treatments. American Journal of Hematology, 2005, 80, 113-118.	4.1	25
28	Neonatal Hyperparathyroidism Due to Maternal Hypoparathyroidism and Vitamin D Deficiency: A Cause of Multiple Bone Fractures. Clinical Pediatrics, 2005, 44, 267-269.	0.8	22
29	Feminizing Sertoli Cell Tumor Associated with Peutz-Jeghers Syndrome. Journal of Pediatric Endocrinology and Metabolism, 2002, 15, 449-52.	0.9	20
30	Serum insulin-like growth factor-I (IGF-I) and IGF-binding protein-3 levels in severe iodine deficiency. Turkish Journal of Pediatrics, 2002, 44, 215-8.	0.6	9