Zehra YavaÅ AbalÄ±

List of Publications by Year
in descending order

27 papers	365 citations	1163065 8 h-index	839512 18 g-index
27 all docs	27 docs citations	27 times ranked	678 citing authors

#	Article	IF	Citations
1	Comparison of National Growth Standards for Turkish Infants and Children with World Health Organization Growth Standards. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2022, , .	0.9	3
2	Mutations in AR or SRD5A2 Genes: Clinical Findings, Endocrine Pitfalls, and Genetic Features of Children With 46,XY DSD. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2022, , .	0.9	1
3	Evaluation of growth, puberty, osteoporosis, and the response to longâ€term bisphosphonate therapy in four patients with osteoporosisâ€pseudoglioma syndrome. American Journal of Medical Genetics, Part A, 2022, , .	1.2	2
4	LRBA deficiency: a rare cause of type 1 diabetes, colitis, and severe immunodeficiency. Hormones, 2021, 20, 389-394.	1.9	5
5	International practice of corticosteroid replacement therapy in congenital adrenal hyperplasia: data from the I-CAH registry. European Journal of Endocrinology, 2021, 184, 553-563.	3.7	21
6	Clinical and Hormonal Profiles Correlate With Molecular Characteristics in Patients With 11Î²-Hydroxylase Deficiency. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e3714-e3724.	3.6	20
7	Rapid molecular diagnosis of ALB gene variants prevents unnecessary interventions in familial dysalbuminemic hyperthyroxinemia. Journal of Pediatric Endocrinology and Metabolism, 2021, 34, 1201-1205.	0.9	2
8	Efficacy of the Novel Degludec/Aspart Insulin Co-formulation in Children and Adolescents with Type 1 Diabetes: A Real-life Experience with 1-year IDeg/Asp Therapy in Poorly Controlled and Non-compliant Patients. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, .	0.9	1
9	Broad-spectrum XX and XY gonadal dysgenesis in patients with a homozygous L193S variant in PPP2R3C. European Journal of Endocrinology, 2021, 186, 65-72.	3.7	1
10	Familial Hypomagnesemia with Hypercalciuria and Nephrocalcinosis Due to CLDN16 Gene Mutations: Novel Findings in Two Cases with Diverse Clinical Features. Calcified Tissue International, 2021, , 1.	3.1	1
11	Rare cause of severe hypertension in an adolescent boy presenting with short stature: Questions. Pediatric Nephrology, 2020, 35, 403-404.	1.7	1
12	Rare cause of severe hypertension in an adolescent boy presenting with short stature: Answers. Pediatric Nephrology, 2020, 35, 405-407.	1.7	4
13	A rare cause of hypertension in childhood: Questions. Pediatric Nephrology, 2020, 35, 77-78.	1.7	1
14	A rare cause of hypertension in childhood: Answers. Pediatric Nephrology, 2020, 35, 79-82.	1.7	5
15	Clinical Characteristics, Molecular Features, and Long-Term Follow-Up of 15 Patients with Neonatal Diabetes: A Single-Centre Experience. Hormone Research in Paediatrics, 2020, 93, 423-432.	1.8	2
16	Genotypic Sex and Severity of the Disease Determine the Time of Clinical Presentation in Steroid 17Î±-Hydroxylase/17,20-Lyase Deficiency. Hormone Research in Paediatrics, 2020, 93, 558-566.	1.8	11
17	Low DHEAS Concentration in a Girl Presenting with Short Stature and Premature Pubarche: A Novel PAPSS2 Gene Mutation. Hormone Research in Paediatrics, 2019, 92, 262-268.	1.8	6
18	Exome Sequencing of a Primary Ovarian Insufficiency Cohort Reveals Common Molecular Etiologies for a Spectrum of Disease. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 3049-3067.	3.6	53

#	ARTICLE	IF	CITATIONS
19	Comparison of the Clinical and Anthropometric Features of Treated and Untreated Girls with Borderline Early Puberty. Journal of Pediatric and Adolescent Gynecology, 2019, 32, 264-270.	0.7	4
20	Persistent MÃ¼llerian Duct Syndrome: A Rare But Important Etiology of Inguinal Hernia and Cryptorchidism. Sexual Development, 2019, 13, 264-270.	2.0	6
21	Evaluation of the Efficacy and Safety of 3 Different Management Protocols in Pediatric Diabetic Ketoacidosis. Pediatric Emergency Care, 2019, Publish Ahead of Print, e707-e712.	0.9	4
22	Prevalence, clinical characteristics and long-term outcomes of classical 11 Î²-hydroxylase deficiency (11BOHD) in Turkish population and novel mutations in CYP11B1 gene. Journal of Steroid Biochemistry and Molecular Biology, 2018, 181, 88-97.	2.5	23
23	Precocious or early puberty in patients with combined pituitary hormone deficiency due to POU1F1 gene mutation: case report and review of possible mechanisms. Hormones, 2018, 17, 581-588.	1.9	13
24	Body mass index at the presentation of premature adrenarche is associated with components of metabolic syndrome at puberty. European Journal of Pediatrics, 2018, 177, 1593-1601.	2.7	20
25	Incidence of Type 1 Diabetes in Children Aged Below 18 Years During 2013-2015 in Northwest Turkey. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2018, 10, 336-342.	0.9	5
26	Two novel mutations in XYLT2 cause spondyloocular syndrome. American Journal of Medical Genetics, Part A, 2017, 173, 3195-3200.	1.2	22
27	Rare Causes of Primary Adrenal Insufficiency: Genetic and Clinical Characterization of a Large Nationwide Cohort. Journal of Clinical Endocrinology and Metabolism, 2016, 101, 284-292.	3.6	128