Ayla Guven

List of Publications by Year
in descending order

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67 papers	1,083 citations	516710 16 h-index	477307 29 g-index
67 all docs	67 docs citations	67 times ranked	1926 citing authors

#	Article	IF	Citations
1	Real-World Estimates of Adrenal Insufficiencyâ€“Related Adverse Events in Children With Congenital Adrenal Hyperplasia. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e192-e203.	3.6	20
2	Hypogonadism: Is It Always Hypogonadotropic in an Adolescent With a Cleft Palate? A Surprising Case of Klinefelter Syndrome. Cleft Palate-Craniofacial Journal, 2021, 58, 787-790.	0.9	0
3	Surgical Practice in Girls with Congenital Adrenal Hyperplasia: An International Registry Study. Sexual Development, 2021, 15, 229-235.	2.0	4
4	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
5	Gonadectomy in conditions affecting sex development: a registry-based cohort study. European Journal of Endocrinology, 2021, 184, 791-801.	3.7	9
6	Molecular Analysis of CYP27B1 Mutations in Vitamin D-Dependent Rickets Type 1A: c.590G > A (p.G197D) Missense Mutation Causes a RNA Splicing Error. Frontiers in Genetics, 2020, 11, 607517.	2.3	5
7	Revisiting Classical 3Î²-hydroxysteroid Dehydrogenase 2 Deficiency: Lessons from 31 Pediatric Cases. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e1718-e1728.	3.6	20
8	Idiopathic infantile hypercalcemia: mutations in SLC34A1 and CYP24A1 in two siblings and fathers. Journal of Pediatric Endocrinology and Metabolism, 2020, 33, 1353-1358.	0.9	3
9	Different Potent Glucocorticoids, Different Routes of Exposure but the Same Result: Iatrogenic Cushingâ€™s Syndrome and Adrenal Insufficiency. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 383-392.	0.9	6
10	Abatacept as a Long-Term Targeted Therapy for LRBA Deficiency. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 2790-2800.e15.	3.8	112
11	Lipid accumulation product is a predictor of nonalcoholic fatty liver disease in childhood obesity. Korean Journal of Pediatrics, 2019, 62, 450-455.	1.9	12
12	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
13	Associated clinical abnormalities among patients with Turner syndrome. Ä°stanbul Kuzey Klinikleri, 2019, 7, 226-230.	0.3	5
14	A Case with Chickenpox and Epididymoorchitis: More Than a Coincidence?. Cocuk Enfeksiyon Dergisi, 2019, 13, 88-91.	0.1	0
15	EpididimoorÅŸitle Seyreden Bir SuÃ§iÃ§eÄŸi Olgusu, Bir TesadÃ¼ften FazlasÄ± mÄ±?. Cocuk Enfeksiyon Dergisi, 2019, 13, 105-108.	0.1	0
16	Five novel ALMS1 gene mutations in six patients with AlstrÃ¶m syndrome. Journal of Pediatric Endocrinology and Metabolism, 2018, 31, 681-687.	0.9	8
17	A novel <scp>NEUROG3</scp> mutation in neonatal diabetes associated with a neuroâ€intestinal syndrome. Pediatric Diabetes, 2018, 19, 381-387.	2.9	17
18	Cardiac examination in children with Laron syndrome undergoing mecasermin therapy. Journal of Pediatric Endocrinology and Metabolism, 2018, 31, 675-679.	0.9	2

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19	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
20	Seven cases with Williams-Beuren syndrome: endocrine evaluation and long-term follow-up. Journal of Pediatric Endocrinology and Metabolism, 2017, 30, 159-165.	0.9	8
21	Mutational analysis of PHEX, FGF23 and CLCN5 in patients with hypophosphataemic rickets. Clinical Endocrinology, 2017, 87, 103-112.	2.4	21
22	Testicular Adrenal Rest Tumor in Two Brothers with a Novel Mutation in the 3-Beta-Hydroxysteroid Dehydrogenase-2 Gene. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2017, 9, 85-90.	0.9	13
23	LRBA EksikliÄŸi TanÄ±sÄ±nda Akan HÃ¼cre Ã–lÃ§erde Protein Ä°fadesi. Asim, Allerji, Immunoloji, 2017, , .	0.0	0
24	Clinical and Genetic Characteristics, Management and Long-Term Follow-Up of Turkish Patients with Congenital Hyperinsulinism. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2016, 8, 197-204.	0.9	4
25	Spondyloenchondrodysplasia Due to Mutations in ACP5: A Comprehensive Survey. Journal of Clinical Immunology, 2016, 36, 220-234.	3.8	71
26	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
27	Assessment of Anti-MÃ¼llerian Hormone Level in Management of Adolescents with Polycystic Ovary Syndrome. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2016, 8, 55-60.	0.9	6
28	The Growth Characteristics of Patients with Noonan Syndrome: Results of Three Years of Growth Hormone Treatment: A Nationwide Multicenter Study. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2016, 8, 305-312.	0.9	18
29	Current Practice in Diagnosis and Treatment of Growth Hormone Deficiency in Childhood: A Survey from Turkey. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2015, 7, 37-44.	0.9	13
30	Gonadotropin releasing hormone analog treatment in children with congenital adrenal hyperplasia complicated by central precocious puberty. Hormones, 2014, 14, 265-71.	1.9	15
31	Symptomatic cerebral infarction in a child with severe diabetic ketoacidosis. Journal of Pediatric Endocrinology and Metabolism, 2014, 27, 1001-4.	0.9	6
32	Detection of vascular risk markers in children and adolescents with type 1 diabetes. Journal of Pediatric Endocrinology and Metabolism, 2014, 27, 87-92.	0.9	2
33	Obesity and Increasing Rate of Infantile Blount Disease. Clinical Pediatrics, 2014, 53, 539-543.	0.8	15
34	Bilateral galactocele in a male infant with Down syndrome and congenital hypothyroidism. Pediatrics International, 2013, 55, e116-8.	0.5	5
35	Next-Generation Sequencing Reveals Deep Intronic Cryptic ABCC8 and HADH Splicing Founder Mutations Causing Hyperinsulinism by Pseudoexon Activation. American Journal of Human Genetics, 2013, 92, 131-136.	6.2	76
36	Remission with Cabergoline in Adolescent Boys with Cushingâ€™s Disease. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2013, 5, 194-198.	0.9	4

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37	Profile of Hypothyroidism in Downâ€™s Syndrome. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2013, 5, 116-120.	0.9	38
38	Complete androgen insensitivity syndrome and discordant MÃ¼llerian remnants: two cases with novel mutation in the androgen receptor. Journal of Pediatric Endocrinology and Metabolism, 2013, 26, 909-14.	0.9	7
39	Hoffmannâ€™s syndrome and pituitary hyperplasia in an adolescent secondary to Hashimoto thyroiditis. Journal of Pediatric Endocrinology and Metabolism, 2013, 26, 747-51.	0.9	6
40	Delirium in Diabetic Ketoacidosis: A Case Report. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2012, 4, 39-41.	0.9	7
41	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
42	Developmental Defects of the Thyroid Gland: Relationship with Advanced Maternal Age. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2012, 4, 72-75.	0.9	17
43	Clinical features and management of diabetic ketoacidosis in different age groups of children: children less than 5 years of age are at higher risk of metabolic decompensation. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, 917-25.	0.9	4
44	Warburg Micro syndrome. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, 379-82.	0.9	12
45	Kocher-DebrÃ©-SÃ©mÃ©laigne syndrome with pericardial effusion. Journal of Pediatric Endocrinology and Metabolism, 2011, 24, 1099-101.	0.9	3
46	Multiple pterygium syndrome: mimicking the findings of Turner syndrome. Journal of Pediatric Endocrinology and Metabolism, 2011, 24, .	0.9	3
47	3M Syndrome: A Report of Four Cases in Two Families. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2011, 3, 154-159.	0.9	12
48	Multiple pterygium syndrome: mimicking the findings of Turner syndrome. Journal of Pediatric Endocrinology and Metabolism, 2011, 24, 1089-93.	0.9	2
49	Interaction of leptin and nitric oxide pathway on penicillin-induced epileptiform activity in rats. Brain Research, 2010, 1321, 117-124.	2.2	14
50	Association Between the Corrected QT Interval and Carotid Artery Intima-Media Thickness in Obese Children. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2010, 2, 21-27.	0.9	17
51	Adiponectin and resistin concentrations after glucose load in adolescents with polycystic ovary syndrome. Gynecological Endocrinology, 2010, 26, 30-38.	1.7	8
52	Characteristics of Polycystic Ovarian Syndrome and Relationship with Ghrelin in Adolescents. Journal of Pediatric and Adolescent Gynecology, 2010, 23, 285-289.	0.7	16
53	The role of nitric oxide in the inhibitory effect of ghrelin against penicillin-induced epileptiform activity in rat. Neuropeptides, 2009, 43, 295-302.	2.2	32
54	Precocious puberty in a girl with Down syndrome due to primary hypothyroidism. Turkish Journal of Pediatrics, 2009, 51, 381-3.	0.6	4

#	Article	IF	Citations
55	The Mayerâ€“Rokitanskyâ€“Kusterâ€“Hauser and gonadal dysgenesis anomaly in a girl with 45,X/46,X,del(X)(p11.21). American Journal of Medical Genetics, Part A, 2008, 146A, 128-131.	1.2	10
56	Effects of individual factors on adolescent obesity: Study in Turkey. Pediatrics International, 2008, 50, 356-362.	0.5	11
57	Pubertal Progression and Serum Lipid Profile in Obese Children. Journal of Pediatric Endocrinology and Metabolism, 2008, 21, 135-46.	0.9	4
58	Coexistence of Craniovertebral Junction Stenosis with Pancake Kidney in an Adolescent with Acromesomelic Dysplasia. Journal of Pediatric Endocrinology and Metabolism, 2008, 21, 941-2.	0.9	1
59	Complete vaginal agenesis in a girl with Mayer-Rokitansky-KÃ¼ster-Hauser syndrome type II. Journal of Pediatric Endocrinology and Metabolism, 2008, 21, 409.	0.9	0
60	Acromegaly Symptoms Without Pituitary Adenoma in an Adolescent. Journal of Pediatric Endocrinology and Metabolism, 2007, 20, 945-9.	0.9	1
61	Cushing's Syndrome and Adrenocortical Insufficiency Caused by Topical Steroids: Misuse or Abuse?. Journal of Pediatric Endocrinology and Metabolism, 2007, 20, 1173-82.	0.9	35
62	Is premature adrenarche a risk factor for atherogenesis?. Pediatrics International, 2005, 47, 20-25.	0.5	37
63	Leptin and Soluble Leptin Receptor Levels in Obese Children in Fasting and Satiety States. Journal of Pediatric Endocrinology and Metabolism, 2005, 18, 303-7.	0.9	17
64	Are Growth Factors and Leptin Involved in the Pathogenesis of Premature Adrenarche in Girls?. Journal of Pediatric Endocrinology and Metabolism, 2005, 18, 785-91.	0.9	14
65	Severe Hypothyroidism Caused by Hepatic Hemangioendothelioma in an Infant of a Diabetic Mother. Hormone Research in Paediatrics, 2005, 63, 86-89.	1.8	21
66	Fatal Disseminated Cytomegalovirus Infection in an Infant with Cushingâ€™s Syndrome Caused by Topical Steroid. Hormone Research in Paediatrics, 2005, 64, 35-38.	1.8	22
67	LYMPHOMATOID GRANULOMATOSIS IN A BOY WITH LONG-TERM FOLLOW-UP. Pediatric Hematology and Oncology, 2001, 18, 377-382.	0.8	6