Zehra Aycan

List of Publications by Year
in descending order

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177	2,241	346980	406436
papers	citations	22 h-index	35 g-index

178	178	178	3444
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Investigation of early puberty prevalence and time of addition thelarche to pubarche in girls with premature pubarche: two-year follow-up results. Clinical Pediatric Endocrinology, 2022, 31, 25-32.	0.4	2
2	Evaluation of permanent and transient congenital hypothyroidism in cases referred from National Neonatal Screening Program. Journal of Paediatrics and Child Health, 2022, 58, 1431-1438.	0.4	1
3	An evaluation of the knowledge and attitudes of school staff related to diabetes care at school: The 10th year of the â€œdiabetes program at schoolâ€•in Turkey. Pediatric Diabetes, 2021, 22, 233-240.	1.2	6
4	Evaluation of the pathophysiological role of Fetuin A levels in adolescents with polycystic ovary syndrome. Journal of Pediatric Endocrinology and Metabolism, 2021, 34, 911-916.	0.4	6
5	The Diagnostic Value of Free Androgen Index in Obese Adolescent Females with Idiopathic Hirsutism and Polycystic Ovary Syndrome. Journal of Academic Research in Medicine, 2021, 11, 81-85.	0.1	2
6	Care and Support of Children with Type 1 Diabetes at School: The Turkish Experience. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, 13, 370-374.	0.4	5
7	Evaluation of Growth Hormone Results in Different Diagnosis and Trend Over 10 Year of Follow-up: A Single Center Experience. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, 13, 332-341.	0.4	1
8	Long-term Clinical Follow-up of Patients with Familial Hypomagnesemia with Secondary Hypocalcemia. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, 13, 300-307.	0.4	6
9	Recommendations for Clinical Decision-making in Children with Type 1 Diabetes and Celiac Disease: Type 1 Diabetes and Celiac Disease Joint Working Group Report. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, .	0.4	0
10	Clinical Characteristics and Growth Hormone Treatment in Patients with Prader-Willi Syndrome. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, 13, 308-319.	0.4	9
11	Relationship Between the Levels of 25-hydroxyvitamin D at Presentation and the Clinical, Laboratory and Follow-up Data of Children and Adolescents with Type-1 Diabetes Mellitus. Journal of Academic Research in Medicine, 2021, 11, 143-148.	0.1	0
12	Evaluation of Children and Adolescents with Thyroid Nodules: A Single Center Experience. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, 13, 276-284.	0.4	5
13	The effect of GnRH stimulation on AMH regulation in central precocious puberty and isolated premature thelarche. Journal of Pediatric Endocrinology and Metabolism, 2021, 34, 1385-1391.	0.4	1
14	Clinical Characteristics of 46,XX Males with Congenital Adrenal Hyperplasia. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, 13, 180-186.	0.4	0
15	Is Bioavailable Vitamin D Better Than Total Vitamin D to Evaluate Vitamin D Status in Obese Children?. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, 13, 391-399.	0.4	1
16	Alterations of Tear Film and Ocular Surface in Children with Type 1 Diabetes Mellitus. Ocular Immunology and Inflammation, 2020, 28, 362-369.	1.0	8
17	Local Ocular Surface Alterations in Children with Hashimotoâ€™s Thyroiditis. Ocular Immunology and Inflammation, 2020, 28, 791-797.	1.0	0
18	Early visual field changes in patients with type 1 diabetes mellitus. European Journal of Ophthalmology, 2020, 30, 1467-1472.	0.7	0

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19	The alteration of IGF-1 levels and relationship between IGF-1 levels and growth velocity during GnRH analogue therapy. Journal of Pediatric Endocrinology and Metabolism, 2020, 33, 113-120.	0.4	2
20	Serum Fetuin-A and Insulin Levels in Classic Congenital Adrenal Hyperplasia. Hormone and Metabolic Research, 2020, 52, 654-659.	0.7	4
21	Is There Any Association Between Hirsutism and Serum Zinc Levels in Adolescents?. Biological Trace Element Research, 2020, 198, 403-409.	1.9	4
22	Effects of 5-Hydroxymethylfurfural on Pubertal Development of Female Wistar Rats. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 79-85.	0.4	7
23	A case of prohormone convertase deficiency diagnosed with type 2 diabetes. Turk Pediatri Arsivi, 2020, 56, 81-84.	0.9	2
24	Infants with failure to thrive, only symptom of a rare disorder: pseudohypoaldosteronism, case series. Turk Pediatri Arsivi, 2020, 56, 75-77.	0.9	1
25	Recombinant GH treatment in a case of Costello syndrome with a 5-year follow-up. Clinical Pediatric Endocrinology, 2020, 29, 195-199.	0.4	1
26	Intramuscular Short-term ACTH Test for the Determination of Adrenal Function in Children: Safe, Effective and Reliable. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 241-247.	0.4	0
27	Intramuscular Short-term ACTH Test for the Determination of Adrenal Function in Children: Safe, Effective and Reliable. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 241-247.	0.4	1
28	Functional assessment of variants associated with Wolfram syndrome. Human Molecular Genetics, 2019, 28, 3815-3824.	1.4	10
29	Detection of the SRY gene in patients with Turner Syndrome. Journal of Gynecology Obstetrics and Human Reproduction, 2019, 48, 265-267.	0.6	3
30	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
31	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.	1.8	53
32	Contribution of functionally assessed GHRHR mutations to idiopathic isolated growth hormone deficiency in patients without GH1 mutations. Human Mutation, 2019, 40, 2033-2043.	1.1	9
33	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
34	Changes in Retinal Microcirculation Precede the Clinical Onset of Diabetic Retinopathy in Children With Type 1 Diabetes Mellitus. American Journal of Ophthalmology, 2019, 207, 37-44.	1.7	60
35	Evaluation of long-term follow-up and methimazole therapy outcomes of pediatric Gravesâ€™ disease: a single-center experience. Journal of Pediatric Endocrinology and Metabolism, 2019, 32, 341-346.	0.4	12
36	Vitamin D deficiency in adolescent pregnancy and obstetric outcomes. Taiwanese Journal of Obstetrics and Gynecology, 2019, 58, 778-783.	0.5	8

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37	Rare Cause of Infantile Hypercalcemia: A Novel Mutation in the SLC34A1 Gene. Hormone Research in Paediatrics, 2019, 91, 278-284.	0.8	7
38	The effect of 2000 Ä±u/day vitamin d supplementation on insulin resistance and cardiovascular risk parameters in vitamin d deficient obese adolescents. Turkish Journal of Pediatrics, 2019, 61, 723.	0.3	4
39	Comparison of Treatment Regimens in Management of Severe Hypercalcemia Due to Vitamin D Intoxication in Children. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2019, 11, 140-148.	0.4	8
40	Management of Thyrotoxicosis in Children and Adolescents: A Turkish Multi-center Experience. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2019, 11, 164-172.	0.4	8
41	AntimÃ¼llerian Hormone Levels of Infants with Premature Thelarche. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2019, 11, 287-292.	0.4	2
42	Congenital long-qt syndrome in type 1 diabetes: a unique association. Turkish Journal of Pediatrics, 2019, 61, 791.	0.3	2
43	Cigarette smoking in adolescents with type 1 diabetes mellitus and congenital adrenal hyperplasia. Turkish Journal of Pediatrics, 2019, 61, 236.	0.3	0
44	A Myasthenia Gravis Case Diagnosed Simultaneously with Diabetic Ketoacidosis. Journal of Pediatric Research, 2019, 6, 73-76.	0.1	1
45	Cases Referred from the Turkish National Screening Program: Frequency of Congenital Hypothyroidism and Etiological Distribution. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2019, 11, 240-246.	0.4	5
46	Quantitative evaluation of early retinal changes in children with type 1 diabetes mellitus without retinopathy. Australasian journal of optometry, The, 2018, 101, 680-685.	0.6	25
47	Reply. Echocardiography, 2018, 35, 580-580.	0.3	0
48	Clinical follow-up data and the rate of development of precocious and rapidly progressive puberty in patients with premature thelarche. Journal of Pediatric Endocrinology and Metabolism, 2018, 31, 305-312.	0.4	8
49	Response to growth hormone treatment in very young patients with growth hormone deficiencies and mini-puberty. Journal of Pediatric Endocrinology and Metabolism, 2018, 31, 175-184.	0.4	12
50	Evaluation of anterior segment parameters in patients with Turner syndrome using Scheimpflug imaging. Journal of AAPOS, 2018, 22, 56-60.	0.2	1
51	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
52	Clinical and genetic characterisation of a series of patients with triple A syndrome. European Journal of Pediatrics, 2018, 177, 363-369.	1.3	20
53	Subclinical left ventricular systolic and diastolic dysfunction in type 1 diabetic children and adolescents with good metabolic control. Echocardiography, 2018, 35, 227-233.	0.3	30
54	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

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55	Abnormal Uterine Bleeding In Adolescents. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2018, 10, 191-197.	0.4	36
56	Follow-up in children with non-obese and non-autoimmune subclinical hypothyroidism. Journal of Pediatric Endocrinology and Metabolism, 2018, 31, 1133-1138.	0.4	4
57	SHOX gene deletion screening by FISH in children with short stature and Madelung deformity and their characteristics. Journal of Pediatric Endocrinology and Metabolism, 2018, 31, 1273-1278.	0.4	0
58	Near final height in patients with idiopathic growth hormone deficiency: A singleâ€centre experience. Journal of Paediatrics and Child Health, 2018, 54, 1221-1226.	0.4	5
59	Low hemoglobin a1c levels in a patient with diabetic ketoacidosis: fulminant type 1 diabetes mellitus. Turkish Journal of Pediatrics, 2018, 60, 201.	0.3	1
60	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
61	Evaluation of the Ovarian Reserve in Adolescents with Hashimoto's Thyroiditis Using Serum anti-MÃ¼llerian Hormone Levels. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2018, 10, 331-335.	0.4	13
62	A Patient with Proopiomelanocortin Deficiency: An Increasingly Important Diagnosis to Make. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2018, 10, 68-73.	0.4	17
63	AMH levels in girls with various pubertal problems. Journal of Pediatric Endocrinology and Metabolism, 2017, 30, 333-335.	0.4	11
64	The variable clinical phenotype of three patients with hepatic glycogen synthase deficiency. Journal of Pediatric Endocrinology and Metabolism, 2017, 30, 459-462.	0.4	23
65	Objective Evaluation of Corneal and Lens Clarity in Children With Type 1 Diabetes Mellitus. American Journal of Ophthalmology, 2017, 179, 190-197.	1.7	20
66	Digenic DUOX1 and DUOX2 Mutations in Cases With Congenital Hypothyroidism. Journal of Clinical Endocrinology and Metabolism, 2017, 102, 3085-3090.	1.8	53
67	Effects of 1-year growth hormone replacement therapy on thyroid volume and function of the children and adolescents with idiopathic growth hormone deficiency. Journal of Pediatric Endocrinology and Metabolism, 2017, 30, 1187-1190.	0.4	12
68	Novel mutations in the LRP5 gene in patients with Osteoporosisâ€pseudoglioma syndrome. American Journal of Medical Genetics, Part A, 2017, 173, 3132-3135.	0.7	13
69	Clinical, biochemical and genetic features with nonclassical 21-hydroxylase deficiency and final height. Journal of Pediatric Endocrinology and Metabolism, 2017, 30, 759-766.	0.4	16
70	Optic Nerve Parameters in Obese Children as Measured by Spectral Domain Optical Coherence Tomography. Seminars in Ophthalmology, 2017, 32, 743-747.	0.8	4
71	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
72	Cardiac MRI and 3D contrast-enhanced MR angiography in pediatric and young adult patients with Turner syndrome. Turkish Journal of Medical Sciences, 2017, 47, 127-133.	0.4	5

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73	Conventional insulin pump therapy in two neonatal diabetes patients harboring the homozygous PTF1A enhancer mutation: Need for a novel approach for the management of neonatal diabetes. Turkish Journal of Pediatrics, 2017, 59, 458-462.	0.3	5
74	Type 1 rhizomelic chondrodysplasia punctata with a homozygous PEX7 mutation. Journal of Pediatric Endocrinology and Metabolism, 2017, 30, 889-892.	0.4	0
75	Evaluation of Iodine Deficiency in Children with Attention Deficit/Hyperactivity Disorder. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2016, 8, 61-66.	0.4	10
76	Bone mineral density and growth in children with coeliac disease on a gluten free-diet. Turkish Journal of Medical Sciences, 2016, 46, 1816-1821.	0.4	10
77	Efficacy and safety of pamidronate in children with vitamin D intoxication. Pediatrics International, 2016, 58, 562-568.	0.2	10
78	Comparison of Anterior Segment Parameters in Juvenile Diabetes Mellitus and Healthy Eyes. European Journal of Ophthalmology, 2016, 26, 618-622.	0.7	8
79	Treatment experience and long-term follow-up data in two severe neonatal hyperparathyroidism cases. Journal of Pediatric Endocrinology and Metabolism, 2016, 29, 1103-10.	0.4	20
80	Comprehensive Screening of Eight Known Causative Genes in Congenital Hypothyroidism With Gland-in-Situ. Journal of Clinical Endocrinology and Metabolism, 2016, 101, 4521-4531.	1.8	82
81	Do the Anti-MÃ¼llerian Hormone Levels of Adolescents with Polycystic Ovary Syndrome, Those Who Are at Risk for Developing Polycystic Ovary Syndrome, and Those Who Exhibit Isolated Oligomenorrhea Differ from Those of Adolescents with Normal Menstrual Cycles?. Hormone Research in Paediatrics, 2016, 85, 406-411.	0.8	12
82	The Etiology and Clinical Features of Non-CAH Gonadotropin-Independent Precocious Puberty: A Multicenter Study. Journal of Clinical Endocrinology and Metabolism, 2016, 101, 1980-1988.	1.8	20
83	Co-Existence of Thyroid Nodule and Thyroid Cancer in Children and Adolescents with Hashimoto Thyroiditis: A Single-Center Study. Hormone Research in Paediatrics, 2016, 85, 181-187.	0.8	25
84	Maturity onset diabetes of youth (MODY) in Turkish children: sequence analysis of 11 causative genes by next generation sequencing. Journal of Pediatric Endocrinology and Metabolism, 2016, 29, 487-96.	0.4	28
85	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.	1.8	128
86	The first childhood case with coexisting Hashimoto thyroiditis, vitiligo and autoimmune hepatitis. Turkish Journal of Pediatrics, 2016, 58, 432-435.	0.3	2
87	A case with atrophic autoimmune thyroiditis-related hypothyroidism causing multisystem involvement in early childhood. Turkish Journal of Pediatrics, 2016, 58, 446-451.	0.3	4
88	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.	0.4	18
89	Gonadoblastoma in a patient with 46, XY complete gonadal dysgenesis. Turkish Journal of Pediatrics, 2016, 58, 538-540.	0.3	3
90	Is Chromosomal Study Necessary for Girls with Inguinal Hernia?. Journal of Clinical and Diagnostic Research JCDR, 2016, 10, SL01.	0.8	0

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91	Ophthalmic Manifestations in Children With Congenital Hypothyroidism. Journal of Pediatric Ophthalmology and Strabismus, 2016, 53, 29-34.	0.3	3
92	Complete androgen insensitivity syndrome associated with bilateral sertoli cell adenomas and unilateral paratesticular leiomyoma: a case report. Turkish Journal of Pediatrics, 2016, 58, 654.	0.3	3
93	Evaluating the Efficacy of Treatment with a GnRH Analogue in Patients with Central Precocious Puberty. International Journal of Endocrinology, 2015, 2015, 1-7.	0.6	11
94	Effects of GnRH analogue treatment on anterior pituitary hormones in children with central precocious puberty. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 1145-51.	0.4	6
95	Urinary C-Peptide/Creatinine Ratio Can Distinguish Maturity-Onset Diabetes of the Young from Type 1 Diabetes in Children and Adolescents: A Single-Center Experience. Hormone Research in Paediatrics, 2015, 84, 54-61.	0.8	18
96	Molecular analysis of PROP1, POU1F1, LHX3, and HESX1 in Turkish patients with combined pituitary hormone deficiency: a multicenter study. Endocrine, 2015, 49, 479-491.	1.1	27
97	Rett syndrome and precocious puberty association. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 1197.	0.4	2
98	17Î²HSD-3 enzyme deficiency due to novel mutations in the HSD17B3 gene diagnosed in a neonate. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 957-9.	0.4	6
99	The use of pamidronate for acute vitamin D intoxication, clinical experience with three cases. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 709-12.	0.4	10
100	Investigation of androgen receptor gene mutations in a series of 21 patients with 46,XY disorders of sex development. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 1257-63.	0.4	11
101	Investigation of autoimmune diseases accompanying Hashimotoâ€™s thyroiditis in children and adolescents and evaluation of cardiac signs. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 767-71.	0.4	4
102	17Î±-Hydroylase/17,20-lyase deficiency related to P.Y27*(c.81C>A) mutation in CYP17A1 gene. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 919-21.	0.4	10
103	Risk factors affecting the development of nephrocalcinosis, the most common complication of hypophosphatemic rickets. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 1333-7.	0.4	15
104	Clinical Review of 95 Patients with 46,XX Disorders of Sex Development Based on the New Chicago Classification. Journal of Pediatric and Adolescent Gynecology, 2015, 28, 6-11.	0.3	14
105	The evaluation of transient hypothyroidism in patients diagnosedwith congenital hypothyroidism. Turkish Journal of Medical Sciences, 2015, 45, 745-750.	0.4	8
106	Effect of Obesity on Left Ventricular Longitudinal Myocardial Strain by Speckle Tracking Echocardiography in Children and Adolescents. Balkan Medical Journal, 2015, 32, 56-63.	0.3	15
107	The evaluation of transient hypothyroidism in patients diagnosed with congenital hypothyroidism. Turkish Journal of Medical Sciences, 2015, 45, 745-50.	0.4	5
108	Decreased Retinal Nerve Fiber Layer Thickness in Patients with Congenital Isolated Growth Hormone Deficiency. European Journal of Ophthalmology, 2014, 24, 873-878.	0.7	7

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109	Adherence to Growth Hormone Therapy: Results of a Multicenter Study. Endocrine Practice, 2014, 20, 46-51.	1.1	67
110	Increased Central Corneal Thickness in Patients with Turner Syndrome. European Journal of Ophthalmology, 2014, 24, 309-313.	0.7	7
111	A nonsense thyrotropin receptor gene mutation (R609X) is associated with congenital hypothyroidism and heart defects. Journal of Pediatric Endocrinology and Metabolism, 2014, 27, 1101-5.	0.4	10
112	A common thyroid peroxidase gene mutation (G319R) in Turkish patients with congenital hypothyroidism could be due to a founder effect. Journal of Pediatric Endocrinology and Metabolism, 2014, 27, 383-7.	0.4	12
113	A truncating DUOX2 mutation (R434X) causes severe congenital hypothyroidism. Journal of Pediatric Endocrinology and Metabolism, 2014, 27, 323-7.	0.4	17
114	Diseases accompanying congenital hypothyroidism. Journal of Pediatric Endocrinology and Metabolism, 2014, 27, 485-9.	0.4	14
115	Evaluation of Asymmetric Dimethylarginine (ADMA) Levels in Children with Growth Hormone Deficiency. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2014, 6, 22-27.	0.4	6
116	Prader-Willi Syndrome and Growth Hormone Deficiency. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2014, 6, 62-67.	0.4	48
117	An Uncommon Cause of Hypoglycemia: Insulin Autoimmune Syndrome. Hormone Research in Paediatrics, 2014, 82, 278-282.	0.8	17
118	An evaluation of heart rate variability and its modifying factors in children with type 1 diabetes. Cardiology in the Young, 2014, 24, 872-879.	0.4	8
119	Corneal biomechanical characteristics in children with diabetes mellitus. International Ophthalmology, 2014, 34, 881-886.	0.6	6
120	Tip-1 Diyabetes Mellitusun Ã‡ocuklarda Merkezi Kornea KalÄ±nlÄ±ÄŸÄ±na Etkisi. TÃ¼rk Oftalmoloji Dergisi, 2014, 44, 445-448.	0.4	0
121	What has national screening program changed in cases with congenital hypothyroidism?. Iranian Journal of Pediatrics, 2014, 24, 255-60.	0.1	4
122	Thyroid dyshormonogenesis is mainly caused by <scp>TPO</scp> mutations in consanguineous community. Clinical Endocrinology, 2013, 79, 275-281.	1.2	47
123	Early Subclinical Left-Ventricular Dysfunction in Obese Nonhypertensive Children: A Tissue Doppler Imaging Study. Pediatric Cardiology, 2013, 34, 1482-1490.	0.6	22
124	The Relationship Between Pediatric Nonalcoholic Fatty Liver Disease and Cardiovascular Risk Factors and Increased Risk of Atherosclerosis in Obese Children. Pediatric Cardiology, 2013, 34, 308-315.	0.6	40
125	Peroxisome proliferator activated receptor (PPAR)-gamma concentrations in childhood obesity. Scandinavian Journal of Clinical and Laboratory Investigation, 2013, 73, 355-360.	0.6	16
126	Evaluation of bone mineral density in children with type 1 diabetes mellitus. Journal of Pediatric Endocrinology and Metabolism, 2013, 26, 1077-81.	0.4	12

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127	Prevalence of hyperthyrotropinemia in obese children before and after weight loss. Eating and Weight Disorders, 2013, 18, 87-90.	1.2	11
128	Prevalence and longâ€term followâ€up outcomes of testicular adrenal rest tumours in children and adolescent males with congenital adrenal hyperplasia. Clinical Endocrinology, 2013, 78, 667-672.	1.2	67
129	Value of pelvic sonography in the diagnosis of various forms of precocious puberty in girls. Journal of Clinical Ultrasound, 2013, 41, 84-93.	0.4	26
130	Diabetes Care, Glycemic Control, Complications, and Concomitant Autoimmune Diseases in Children with Type 1 Diabetes in Turkey: A Multicenter Study. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2013, 5, 20-26.	0.4	32
131	Long-Term Follow-Up of Cushingâ€™s Disease: A Case Report. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2013, 5, 202-205.	0.4	0
132	Hyperinsulinemic Hypoglycemia: Experience in A Series of 17 Cases. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2013, 5, 150-155.	0.4	8
133	Diabetes mellitus with Laron syndrome: case report. Journal of Pediatric Endocrinology and Metabolism, 2013, 26, 955-8.	0.4	12
134	Report of the first case of precocious puberty in Rett syndrome. Journal of Pediatric Endocrinology and Metabolism, 2013, 26, 937-9.	0.4	10
135	Assessment of the Knowledge of Diabetes Mellitus Among School Teachers within the Scope of the Managing Diabetes at School Program. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2012, 4, 199-203.	0.4	25
136	A Rare Combination: Congenital Adrenal Hyperplasia Due To 21 Hydroxylase Deficiency and Turner Syndrome. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2012, 4, 213-215.	0.4	4
137	Mild and severe congenital primary hypothyroidism in two patients by thyrotropin receptor (TSHR) gene mutation. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, 1153-6.	0.4	12
138	A pediatric Conn syndrome case. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, 203-6.	0.4	5
139	Seizure due to somatostatin analog discontinuation in a case diagnosed as congenital hyperinsulinism novel mutation. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, .	0.4	4
140	Audiologic evaluation in pediatric patients with type 1 diabetes mellitus. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, 503-8.	0.4	8
141	TSHR is the main causative locus in autosomal recessively inherited thyroid dysgenesis. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, 419-26.	0.4	27
142	Assessment of the 21-hydroxylase deficiency and the adrenal functions in young females with Turner syndrome. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, 681-5.	0.4	2
143	Thyroid nodules in children and adolescents: a single institutionâ€™s experience. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, 633-8.	0.4	8
144	The Exon 3-Deleted/Full-Length Growth Hormone Receptor Polymorphism and Response to Growth Hormone Therapy in Growth Hormone Deficiency and Turner Syndrome: A Multicenter Study. Hormone Research in Paediatrics, 2012, 77, 85-93.	0.8	14

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145	Iatrogenic Cushing syndrome due to nasal steroid drops. European Journal of Pediatrics, 2012, 171, 735-736.	1.3	14
146	Thiamine-Responsive Megaloblastic Anemia Syndrome With Atrial Standstill. Journal of Pediatric Hematology/Oncology, 2011, 33, 144-147.	0.3	21
147	Osteogenesis imperfecta associated with partial trisomy 20p. Current Opinion in Biotechnology, 2011, 22, S104-S105.	3.3	0
148	Vitamin D status and insulin requirements in children and adolescent with type 1 diabetes. Journal of Pediatric Endocrinology and Metabolism, 2011, 24, .	0.4	16
149	A new variant of a known mutation in two siblings with permanent neonatal diabetes mellitus. Journal of Pediatric Endocrinology and Metabolism, 2011, 24, 373-5.	0.4	0
150	A case of Langerhans cell histiocytosis with thyroid involvement. Journal of Pediatric Endocrinology and Metabolism, 2011, 24, .	0.4	1
151	Eight-Year Follow-up of a Girl with McCune-Albright Syndrome. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2011, 3, 40-42.	0.4	4
152	Does Pseudohypoaldosteronism Mask the Diagnosis of Congenital Adrenal Hyperplasia?. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2011, 3, 219-221.	0.4	9
153	A novel mutation of 5Î±-steroid reductase 2 deficiency (CD 65 ALA-PRO) with severe virilization defect in a Turkish family and difficulty in gender assignment. European Journal of Pediatrics, 2010, 169, 991-995.	1.3	10
154	Novel TSHR mutations in consanguineous families with congenital nongoitrous hypothyroidism. Clinical Endocrinology, 2010, 73, 671-677.	1.2	28
155	A Case Report of Neonatal Diabetes Due to Neonatal Hemochromatosis. Journal of Pediatric Endocrinology and Metabolism, 2010, 23, 521-4.	0.4	1
156	The Evaluation of the Adrenal and Thyroid Axes and Glucose Metabolism after Burn Injury in Children. Journal of Pediatric Endocrinology and Metabolism, 2010, 23, 481-9.	0.4	8
157	Celiac Disease and Autoimmune Thyroid Disease in Children with Type 1 Diabetes Mellitus: Clinical and HLA-Genotyping Results-Original Article. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2010, 2, 151-154.	0.4	29
158	Refractive Errors and Strabismus in Children With Laurenceâ€“Moonâ€“Biedl Syndrome. Journal of Pediatric Ophthalmology and Strabismus, 2010, 47, 26-28.	0.3	4
159	Testicular adrenal rest tumor in a patient with 11beta-hydroxylase deficient congenital adrenal hyperplasia. Journal of Pediatric Endocrinology and Metabolism, 2010, 23, 729-32.	0.4	5
160	Donohue Syndrome in a Neonate with Homozygous Deletion of Exon 3 of the Insulin Receptor Gene. Journal of Pediatric Endocrinology and Metabolism, 2009, 22, 669-74.	0.4	9
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