Semra Ãetinkaya

List of Publications by Year
in descending order

157	1,370	394421	552781
papers	citations	19 h-index	26 g-index

160	160	160	2092
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Urinary levels of phthalate esters and heavy metals in adolescents with thyroid colloid cysts. International Journal of Environmental Health Research, 2022, 32, 1359-1372.	2.7	3
2	Diagnostic Value of Bilateral Petrosal Sinus Sampling in Children with Cushing Disease: A Multi-center Study. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2022, 14, 29-36.	0.9	4
3	A major health problem facing immigrant children: nutritional rickets. Journal of Pediatric Endocrinology and Metabolism, 2022, 35, 223-229.	0.9	1
4	Sleep disorder and behavior problems in children with type 1 diabetes mellitus. Journal of Pediatric Endocrinology and Metabolism, 2022, 35, 29-38.	0.9	1
5	Ergenlerden OluÅŸan bir Klinik Ã–rneklemde DÃ¼rtÃ¼sellik ile Ä°nternet BaÄŸÄ±mlÄ±lÄ±ÄŸÄ± AÅŸermesi ArasÄ±ndaki Ä°liÅŸki. Journal of Dependence, 2022, 23, 1-1.	0.8	0
6	Unfavorbale Effects of Low-Carbonhydrate Diet In a Pediatric Patient With Type 1 Diabetes Mellitus. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2022, .	0.9	1
7	PLXNB1 mutations in the etiology of idiopathic hypogonadotropic hypogonadism. Journal of Neuroendocrinology, 2022, 34, e13103.	2.6	5
8	Insights From Long-term Follow-up of a Girl With Adrenal Insufficiency and Sphingosine-1-Phosphate Lyase Deficiency. Journal of the Endocrine Society, 2022, 6, bvac020.	0.2	6
9	DELETE. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2022, , .	0.9	1
10	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.	0.8	1
11	Hypergonadotrophic hypogonadism in a patient with transaldolase deficiency: novel mutation in the pentose phosphate pathway. Hormones, 2021, 20, 581-585.	1.9	5
12	A Newborn Admitted with Hyponatremia and Hyperkalemia Clinic and Diagnosed with Primary Hypoaldosteronism. Trends in Pediatrics, 2021, 2, 94-98.	0.1	0
13	Langerhans Cell Histiocytosis with Isolated Central Diabetes Insipidus, Low Grade Fever and Sellar Erosion. Trends in Pediatrics, 2021, 2, 104-108.	0.1	0
14	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.	0.9	6
15	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
16	Retrospective evaluation of patients with X-linked adrenoleukodystrophy with a wide range of clinical presentations: a single center experience. Journal of Pediatric Endocrinology and Metabolism, 2021, 34, 1169-1179.	0.9	1
17	Perinatal outcomes of high-dose vitamin D administration in the last trimester. TâŠÂºrk Jinekoloji Ve Obstetrik Dernei Dergisi, 2021, 18, 159-162.	0.8	4
18	Mild phenotype in two siblings with a missense GHR variant. Journal of Pediatric Endocrinology and Metabolism, 2021, 34, 1349-1354.	0.9	0

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19	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.	0.9	1
20	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.	0.9	6
21	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
22	Evaluation of Children and Adolescents with Thyroid Nodules: A Single Center Experience. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, 13, 276-284.	0.9	5
23	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.	0.9	1
24	Clinical Characteristics of 46,XX Males with Congenital Adrenal Hyperplasia. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, 13, 180-186.	0.9	0
25	Effects of subclinical emotional and behavioral problems on metabolic control in adolescents with type 1 diabetes: Role of maternal and adolescent personality traits. Archives De Pediatrie, 2021, 28, 626-631.	1.0	1
26	Clinical Management in Systemic Type Pseudohypoaldosteronism Due to <i>SCNN1B</i> Variant and Literature Review. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, 13, 446-451.	0.9	4
27	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.	0.9	2
28	Serum Fetuin-A and Insulin Levels in Classic Congenital Adrenal Hyperplasia. Hormone and Metabolic Research, 2020, 52, 654-659.	1.5	4
29	Effects of 5-Hydroxymethylfurfural on Pubertal Development of Female Wistar Rats. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 79-85.	0.9	7
30	Nationwide Turkish Cohort Study of Hypophosphatemic Rickets. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 150-159.	0.9	8
31	A case of prohormone convertase deficiency diagnosed with type 2 diabetes. Turk Pediatri Arsivi, 2020, 56, 81-84.	0.9	2
32	Infants with failure to thrive, only symptom of a rare disorder: pseudohypoaldosteronism, case series. Turk Pediatri Arsivi, 2020, 56, 75-77.	0.9	1
33	Clitoromegaly caused by ovarian stimulation in a preterm newborn: ovarian hyperstimulation syndrome of preterm babies. Turkish Journal of Pediatrics, 2020, 62, 1088.	0.6	1
34	Effect of gonadotropin releasing hormone analog treatment on final height in girls aged 6-10 years with central precocious and early puberty. Turk Pediatri Arsivi, 2020, 55, 361-369.	0.9	1
35	Evaluation of Thiol/Disulfide Homeostasis in Pediatric Patients with Diabetic Ketoacidosis. Combinatorial Chemistry and High Throughput Screening, 2020, 23, 185-190.	1.1	0
36	Functional assessment of variants associated with Wolfram syndrome. Human Molecular Genetics, 2019, 28, 3815-3824.	2.9	10

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37	Detection of the SRY gene in patients with Turner Syndrome. Journal of Gynecology Obstetrics and Human Reproduction, 2019, 48, 265-267.	1.3	3
38	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.	0.9	1
39	Rare Cause of Infantile Hypercalcemia: A Novel Mutation in the SLC34A1 Gene. Hormone Research in Paediatrics, 2019, 91, 278-284.	1.8	7
40	Level of Certain Oxidants and Antioxidants in Patients with Uterine Fibroids. Gynecology Obstetrics & Reproductive Medicine (gorm), 2019, 25, 158-162.	0.3	2
41	Role of Ischemia Modified Albumin Serum Levels as an Oxidative Stress Marker in Children with Diabetic Ketoacidosis. Combinatorial Chemistry and High Throughput Screening, 2019, 22, 577-581.	1.1	8
42	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.	0.6	4
43	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.	0.9	8
44	AntimÃ¼llerian Hormone Levels of Infants with Premature Thelarche. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2019, 11, 287-292.	0.9	2
45	Congenital long-qt syndrome in type 1 diabetes: a unique association. Turkish Journal of Pediatrics, 2019, 61, 791.	0.6	2
46	Bone Loss in Pediatric Survivors of Acute Lymphoblastic Leukemia. Eurasian Journal of Medicine, 2019, 51, 38-41.	0.6	5
47	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.	0.9	5
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.	0.9	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.	0.9	12
50	Vaginal bleeding and a giant ovarian cyst in an infant with 21-hydroxylase deficiency. Journal of Pediatric Endocrinology and Metabolism, 2018, 31, 229-233.	0.9	3
51	Clinical and genetic characterisation of a series of patients with triple A syndrome. European Journal of Pediatrics, 2018, 177, 363-369.	2.7	20
52	Renal complications of lipodystrophy: A closer look at the natural history of kidney disease. Clinical Endocrinology, 2018, 89, 65-75.	2.4	16
53	Investigation of MKRN3 Mutation in Patients with Familial Central Precocious Puberty. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2018, 10, 223-229.	0.9	22
54	Follow-up in children with non-obese and non-autoimmune subclinical hypothyroidism. Journal of Pediatric Endocrinology and Metabolism, 2018, 31, 1133-1138.	0.9	4

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55	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.	0.9	0
56	Near final height in patients with idiopathic growth hormone deficiency: A singleâ€centre experience. Journal of Paediatrics and Child Health, 2018, 54, 1221-1226.	0.8	5
57	Low hemoglobin a1c levels in a patient with diabetic ketoacidosis: fulminant type 1 diabetes mellitus. Turkish Journal of Pediatrics, 2018, 60, 201.	0.6	1
58	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.	0.9	6
59	A Patient with Proopiomelanocortin Deficiency: An Increasingly Important Diagnosis to Make. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2018, 10, 68-73.	0.9	17
60	Sublingual ectopic thyroid gland diagnosed with subclinical hypothyroidism in the pubertal period. Endocrinology&Metabolism International Journal, 2018, 6, .	0.1	0
61	AMH levels in girls with various pubertal problems. Journal of Pediatric Endocrinology and Metabolism, 2017, 30, 333-335.	0.9	11
62	A Newborn with Transient Diabetes Mellitus Accompanied by Ketoacidosis Attributable to a ZFP57 Mutation. Journal of Tropical Pediatrics, 2017, 63, 399-401.	1.5	2
63	Clinical, biochemical and genetic features with nonclassical 21-hydroxylase deficiency and final height. Journal of Pediatric Endocrinology and Metabolism, 2017, 30, 759-766.	0.9	16
64	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.	0.9	14
65	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.	0.6	5
66	Contents Vol. 85, 2016. Hormone Research in Paediatrics, 2016, 85, I-VI.	1.8	0
67	Efficacy and safety of pamidronate in children with vitamin D intoxication. Pediatrics International, 2016, 58, 562-568.	0.5	10
68	Treatment experience and long-term follow-up data in two severe neonatal hyperparathyroidism cases. Journal of Pediatric Endocrinology and Metabolism, 2016, 29, 1103-10.	0.9	20
69	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.	1.8	12
70	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.	3.6	20
71	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.	0.9	28
72	The first childhood case with coexisting Hashimoto thyroiditis, vitiligo and autoimmune hepatitis. Turkish Journal of Pediatrics, 2016, 58, 432-435.	0.6	2

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73	A case with atrophic autoimmune thyroiditis-related hypothyroidism causing multisystem involvement in early childhood. Turkish Journal of Pediatrics, 2016, 58, 446-451.	0.6	4
74	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
75	Gonadoblastoma in a patient with 46, XY complete gonadal dysgenesis. Turkish Journal of Pediatrics, 2016, 58, 538-540.	0.6	3
76	Complete androgen insensitivity syndrome associated with bilateral sertoli cell adenomas and unilateral paratesticular leiomyoma: a case report. Turkish Journal of Pediatrics, 2016, 58, 654.	0.6	3
77	Evaluating the Efficacy of Treatment with a GnRH Analogue in Patients with Central Precocious Puberty. International Journal of Endocrinology, 2015, 2015, 1-7.	1.5	11
78	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.	0.9	6
79	Molecular analysis of PROP1, POU1F1, LHX3, and HESX1 in Turkish patients with combined pituitary hormone deficiency: a multicenter study. Endocrine, 2015, 49, 479-491.	2.3	27
80	Selenium, zinc, and copper levels and their relation with HbA1c status in children with type 1 diabetes mellitus. International Journal of Diabetes in Developing Countries, 2015, 35, 514-518.	0.8	5
81	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.	0.9	6
82	The use of pamidronate for acute vitamin D intoxication, clinical experience with three cases. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 709-12.	0.9	10
83	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.	0.9	10
84	Risk factors affecting the development of nephrocalcinosis, the most common complication of hypophosphatemic rickets. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 1333-7.	0.9	15
85	The evaluation of transient hypothyroidism in patients diagnosedwith congenital hypothyroidism. Turkish Journal of Medical Sciences, 2015, 45, 745-750.	0.9	8
86	The evaluation of transient hypothyroidism in patients diagnosed with congenital hypothyroidism. Turkish Journal of Medical Sciences, 2015, 45, 745-50.	0.9	5
87	Adherence to Growth Hormone Therapy: Results of a Multicenter Study. Endocrine Practice, 2014, 20, 46-51.	2.1	67
88	Increased Central Corneal Thickness in Patients with Turner Syndrome. European Journal of Ophthalmology, 2014, 24, 309-313.	1.3	7
89	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.	0.9	12
90	Diseases accompanying congenital hypothyroidism. Journal of Pediatric Endocrinology and Metabolism, 2014, 27, 485-9.	0.9	14

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91	Kidney growth and renal functions under the growth hormone replacement therapy in children. Renal Failure, 2014, 36, 508-513.	2.1	10
92	Evaluation of Asymmetric Dimethylarginine (ADMA) Levels in Children with Growth Hormone Deficiency. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2014, 6, 22-27.	0.9	6
93	An Uncommon Cause of Hypoglycemia: Insulin Autoimmune Syndrome. Hormone Research in Paediatrics, 2014, 82, 278-282.	1.8	17
94	Corneal biomechanical characteristics in children with diabetes mellitus. International Ophthalmology, 2014, 34, 881-886.	1.4	6
95	Corneal properties in children with congenital isolated growth hormone deficiency. International Journal of Ophthalmology, 2014, 7, 317-20.	1.1	3
96	What has national screening program changed in cases with congenital hypothyroidism?. Iranian Journal of Pediatrics, 2014, 24, 255-60.	0.3	4
97	Thyroid dyshormonogenesis is mainly caused by <scp>TPO</scp> mutations in consanguineous community. Clinical Endocrinology, 2013, 79, 275-281.	2.4	47
98	Peroxisome proliferator activated receptor (PPAR)-gamma concentrations in childhood obesity. Scandinavian Journal of Clinical and Laboratory Investigation, 2013, 73, 355-360.	1.2	16
99	Evaluation of bone mineral density in children with type 1 diabetes mellitus. Journal of Pediatric Endocrinology and Metabolism, 2013, 26, 1077-81.	0.9	12
100	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.	2.4	67
101	Value of pelvic sonography in the diagnosis of various forms of precocious puberty in girls. Journal of Clinical Ultrasound, 2013, 41, 84-93.	0.8	26
102	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.	0.9	32
103	Long-Term Follow-Up of Cushingâ€™s Disease: A Case Report. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2013, 5, 202-205.	0.9	0
104	Hyperinsulinemic Hypoglycemia: Experience in A Series of 17 Cases. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2013, 5, 150-155.	0.9	8
105	Diabetes mellitus with Laron syndrome: case report. Journal of Pediatric Endocrinology and Metabolism, 2013, 26, 955-8.	0.9	12
106	Report of the first case of precocious puberty in Rett syndrome. Journal of Pediatric Endocrinology and Metabolism, 2013, 26, 937-9.	0.9	10
107	A girl with steroid cell ovarian tumor misdiagnosed as non-classical congenital adrenal hyperplasia. Turkish Journal of Pediatrics, 2013, 55, 443-6.	0.6	13
108	The first case of Bruck syndrome associated with gastroschisis. Turkish Journal of Pediatrics, 2013, 55, 651-4.	0.6	1

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109	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.	0.9	25
110	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.	0.9	4
111	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.	0.9	12
112	A pediatric Conn syndrome case. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, 203-6.	0.9	5
113	Seizure due to somatostatin analog discontinuation in a case diagnosed as congenital hyperinsulinism novel mutation. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, .	0.9	4
114	TSHR is the main causative locus in autosomal recessively inherited thyroid dysgenesis. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, 419-26.	0.9	27
115	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.	0.9	2
116	Thyroid nodules in children and adolescents: a single institutionâ€™s experience. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, 633-8.	0.9	8
117	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.	1.8	14
118	Iatrogenic Cushing syndrome due to nasal steroid drops. European Journal of Pediatrics, 2012, 171, 735-736.	2.7	14
119	Problematic Eating Behaviour in Turkish Children Aged 12-72 Months: Characteristics of Mothers and Children. Central European Journal of Public Health, 2012, 20, 257-261.	1.1	25
120	Seizure due to somatostatin analog discontinuation in a case diagnosed as congenital hyperinsulinism novel mutation. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, 553-5.	0.9	4
121	Evaluation of adrenocortical function in 3-7 aged asthmatic children treated with moderate doses of fluticasone propionate: reliability of dehydroepiandrosterone sulphate (dhea-s) as a screening test. Allergologia Et Immunopathologia, 2011, 39, 154-158.	1.7	8
122	Rhabdomyolysis Without Detectable Myoglobulinuria Due to Severe Hypophosphatemia in Diabetic Ketoacidosis. Pediatric Emergency Care, 2011, 27, 537-538.	0.9	22
123	Thiamine-Responsive Megaloblastic Anemia Syndrome With Atrial Standstill. Journal of Pediatric Hematology/Oncology, 2011, 33, 144-147.	0.6	21
124	Alendronate for the treatment of hypercalcaemia due to neonatal subcutaneous fat necrosis. European Journal of Pediatrics, 2011, 170, 1085-1086.	2.7	6
125	The effect of glucocorticoid replacement therapy on bone mineral density in children with congenital adrenal hyperplasia. Journal of Pediatric Endocrinology and Metabolism, 2011, 24, 265-9.	0.9	9
126	Vitamin D status and insulin requirements in children and adolescent with type 1 diabetes. Journal of Pediatric Endocrinology and Metabolism, 2011, 24, .	0.9	16

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127	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.	0.9	0
128	A case of Langerhans cell histiocytosis with thyroid involvement. Journal of Pediatric Endocrinology and Metabolism, 2011, 24, .	0.9	1
129	Eight-Year Follow-up of a Girl with McCune-Albright Syndrome. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2011, 3, 40-42.	0.9	4
130	Successful treatment of pulmonary arteriovenous malformation and infantile hepatic hemangioendothelioma with alpha-interferon. Anatolian Journal of Cardiology, 2011, 11, 181-3.	0.4	1
131	Novel TSHR mutations in consanguineous families with congenital nongoitrous hypothyroidism. Clinical Endocrinology, 2010, 73, 671-677.	2.4	28
132	Cord blood thyroidâ€stimulating hormone and free T₄ levels in Turkish neonates: Is iodine deficiency still a continuing problem?. Pediatrics International, 2010, 52, 762-768.	0.5	11
133	A Case Report of Neonatal Diabetes Due to Neonatal Hemochromatosis. Journal of Pediatric Endocrinology and Metabolism, 2010, 23, 521-4.	0.9	1
134	Insulin Oedema in Newly Diagnosed Type 1 Diabetes Mellitus. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2010, 2, 46-48.	0.9	27
135	Hypothyroidism Due to Hepatic Hemangioendothelioma: A Case Report. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2010, 2, 126-130.	0.9	13
136	Sporadic Nonautoimmune Neonatal Hyperthyroidism Due to A623V Germline Mutation in the Thyrotropin Receptor Gene-Case Report. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2010, 2, 168-172.	0.9	9
137	Testicular adrenal rest tumor in a patient with 11beta-hydroxylase deficient congenital adrenal hyperplasia. Journal of Pediatric Endocrinology and Metabolism, 2010, 23, 729-32.	0.9	5
138	Hyperparathyroidism secondary to maternal hypoparathyroidism and vitamin D deficiency: an uncommon cause of neonatal respiratory distress. Annals of Tropical Paediatrics, 2009, 29, 149-154.	1.0	25
139	Kidney growth in children with congenital hypothyroidism. Pediatric Nephrology, 2009, 24, 333-340.	1.7	8
140	The effects of metabolic control on oxidized low-density lipoprotein antibodies in children and adolescents with type 1 diabetes mellitus. Pediatric Diabetes, 2008, 9, 17-22.	2.9	10
141	Inappropriate Use of Potent Topical Glucocorticoids in Infants. Journal of Pediatric Endocrinology and Metabolism, 2007, 20, 219-25.	0.9	28
142	Diagnostic Value of Salivary Cortisol in Children with Abnormal Adrenal Cortex Functions. Hormone Research in Paediatrics, 2007, 67, 301-306.	1.8	20
143	Central diabetes insipidus associated with Caroli syndrome. Indian Journal of Pediatrics, 2007, 74, 419-420.	0.8	1
144	Effect of total attending period on infection episode rate in a child-care center. Pediatrics International, 2004, 46, 555-560.	0.5	6

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145	Antiseizure activity of insulin: insulin inhibits pentylenetetrazole, penicillin and kainic acid-induced seizures in rats. Epilepsy Research, 1996, 25, 185-190.	1.6	24
146	2015-2019 yÄ±llarÄ± arasÄ±nda Ã§ocuk endokrinoloji servisine yatÄ±rÄ±lmÄ±ÅŸ diyabetes mellitus tanÄ±lÄ± olgularÄ±n yatÄ±ÅŸ durumlarÄ±nÄ±n deÄŸerlendirilmesi. Turkish Journal of Clinics and Laboratory, 0, , .	0.4	0
147	Ã–TÄ°ROÄ°D HASHÄ°MOTO TÄ°ROÄ°DÄ°TLÄ° Ã‡OCUK VE ADÃ–LESANLARDA BÄ°R Ä°NFLAMASYON BELÄ°RTECÄ° OLARAK NÃ–TROFÄ°L/LENFOSÄ°T VE PLATELET/LENFOSÄ°T ORANLARI. KÄ±rÄ±kkale Ãœniversitesi TÄ±p FakÃ¼ltesi Dergisi, 0, , 293-299.	0.3	1
148	Multisystemic Severe Form Pseudohypoaldosteronism: Can Gastrostomy be Useful in the Management?. Turkish Journal of Pediatric Disease, 0, , .	0.0	0
149	Vitamin D Deficiency/Insufficiency in Children and Adolescents with Chronic Disease. Turkish Journal of Pediatric Disease, 0, , .	0.0	0
150	Evaluation of Patients Diagnosed with Nutritional Rickets: A Single Center Study. Turkish Journal of Pediatric Disease, 0, , .	0.0	1
151	Evaluation of Premature Pubarche Cases: A Single Center Experience. Turkish Journal of Pediatric Disease, 0, , .	0.0	0
152	Severâ€™s Disease in a Patient Receiving Growth Hormone with no Causative Relation. Journal of Clinical and Diagnostic Research JCDR, 0, , .	0.8	0
153	Detection of SRY gene in patients with turner syndrome. Endocrine Abstracts, 0, , .	0.0	0
154	Comparison of Treatment Regimens for the Management of Severe Hypercalcemia due to Vitamin D Intoxication in Children. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 0, , .	0.9	1
155	The Evaluation of Etiological Distribution and the Rate of Congenital Hypothyroidism among the Cases Referred from National Screening Program. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 0, , .	0.9	0
156	Metformin ile tedavi edilen non-diyabetik obez Ã§ocuk ve adolesanlarÄ±n uzun sureli izlemi. Turkish Journal of Clinics and Laboratory, 0, , .	0.4	0
157	Puberte prekokslu kÄ±zlarda anti-MÃ¼llerian hormon(AMH) dÃ¼zeyinin folikÃ¼l sayÄ±sÄ± ve bÃ¼yÃ¼klÃ¼ÄŸÃ¼ ile iliÅŸkisi. Turkish Journal of Clinics and Laboratory, 0, , .	0.4	0