Ayhan Abaci

List of Publications by Year
in descending order

150	2,095	257450	330143
papers	citations	24 h-index	37 g-index

150	150	150	3468
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	The immunostimulant OM-85 BV prevents wheezing attacks in preschool children. Journal of Allergy and Clinical Immunology, 2010, 126, 763-769.	2.9	149
2	Maturity-onset diabetes of the young (MODY): an update. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 251-63.	0.9	137
3	Vaspin and its correlation with insulin sensitivity indices in obese children. Diabetes Research and Clinical Practice, 2009, 84, 325-328.	2.8	63
4	Trace Elements in Obese Turkish Children. Biological Trace Element Research, 2011, 143, 188-195.	3.5	61
5	The relation of serum nesfatin-1 level with metabolic and clinical parameters in obese and healthy children. Pediatric Diabetes, 2013, 14, n/a-n/a.	2.9	54
6	Disease characteristics of MCT8 deficiency: an international, retrospective, multicentre cohort study. Lancet Diabetes and Endocrinology,the, 2020, 8, 594-605.	11.4	50
7	Epidemiology, Classification and Management of Undescended Testes: Does Medication Have Value in its Treatment?. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2013, 5, 65-72.	0.9	45
8	Hepatic glycogenosis: a rare cause of hepatomegaly in Type 1 diabetes mellitus. Journal of Diabetes and Its Complications, 2008, 22, 325-328.	2.3	43
9	Gonadal malignancy risk and prophylactic gonadectomy in disorders of sexual development. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 1019-27.	0.9	43
10	Turner Syndrome and Associated Problems in Turkish Children: A Multicenter Study. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2015, 7, 27-36.	0.9	42
11	The Role of Initial Clinical and Laboratory Findings in Infants With Hyperthyrotropinemia to Predict Transient or Permanent Hypothyroidism. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2013, 5, 170-173.	0.9	40
12	The Relationship between Serum Zonulin Level and Clinical and Laboratory Parameters of Childhood Obesity. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2017, 9, 31-38.	0.9	36
13	P300 auditory event-related potentials in children with obesity: is childhood obesity related to impairment in cognitive functions?. Pediatric Diabetes, 2011, 12, 589-595.	2.9	34
14	Echocardiographic Measurement of Epicardial Adipose Tissue in Obese Children. Pediatric Cardiology, 2010, 31, 853-860.	1.3	33
15	Low Omentin-1 Levels Are Related with Clinical and Metabolic Parameters in Obese Children. Experimental and Clinical Endocrinology and Diabetes, 2013, 121, 595-600.	1.2	33
16	Metabolic Alterations During Valproic Acid Treatment: A Prospective Study. Pediatric Neurology, 2009, 41, 435-439.	2.1	32
17	A Case of Rapid-Onset Obesity with Hypothalamic Dysfunction, Hypoventilation, Autonomic Dysregulation, and Neural Crest Tumor: Rohhadnet Syndrome. Endocrine Practice, 2013, 19, e12-e16.	2.1	32
18	46,XX Male Disorder of Sexual Development: A Case Report. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2013, 5, 258-260.	0.9	31

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19	Subclinical hypothyroidism in childhood and adolescense. Journal of Pediatric Endocrinology and Metabolism, 2014, 27, 1049-57.	0.9	30
20	Decreased Plasma Apelin Levels in Pubertal Obese Children. Journal of Pediatric Endocrinology and Metabolism, 2010, 23, 1039-46.	0.9	29
21	Homozygous Loss-of-function Mutations inSOHLH1in Patients With Nonsyndromic Hypergonadotropic Hypogonadism. Journal of Clinical Endocrinology and Metabolism, 2015, 100, E808-E814.	3.6	29
22	Incidence of maternal vitamin D deficiency in a region of Ankara, Turkey: a preliminary study. Turkish Journal of Medical Sciences, 2014, 44, 616-623.	0.9	28
23	The relation of leptin and soluble leptin receptor levels with metabolic and clinical parameters in obese and healthy children. Peptides, 2014, 56, 72-76.	2.4	28
24	Molecular diagnosis of maturity-onset diabetes of the young (MODY) in Turkish children by using targeted next-generation sequencing. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 1265-71.	0.9	28
25	Relation of serum irisin level with metabolic and antropometric parameters in obese children. Journal of Diabetes and Its Complications, 2016, 30, 1560-1565.	2.3	27
26	Threshold value of subepicardial adipose tissue to detect insulin resistance in obese children. International Journal of Obesity, 2009, 33, 440-446.	3.4	25
27	Circulating betatrophin concentration is negatively correlated with insulin resistance in obese children and adolescents. Diabetes Research and Clinical Practice, 2016, 114, 37-42.	2.8	25
28	Importance of Plasma N-Terminal Pro B-Type Natriuretic Peptide, Epicardial Adipose Tissue, and Carotid Intima-Media Thicknesses in Asymptomatic Obese Children. Pediatric Cardiology, 2010, 31, 792-799.	1.3	24
29	Targeted next generation sequencing in patients with maturity-onset diabetes of the young (MODY). Journal of Pediatric Endocrinology and Metabolism, 2018, 31, 1295-1304.	0.9	23
30	Acute Vitamin D Intoxication Possibly Due to Faulty Production of a Multivitamin Preparation. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2013, 5, 136-139.	0.9	21
31	Significance of serum neurokinin B and kisspeptin levels in the differential diagnosis of premature thelarche and idiopathic central precocious puberty. Peptides, 2015, 64, 29-33.	2.4	21
32	Sensorineural hearing loss in pediatric celiac patients. International Journal of Pediatric Otorhinolaryngology, 2011, 75, 65-68.	1.0	20
33	Hyperprolactinemia in children: clinical features and long-term results. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, 1123-8.	0.9	20
34	A novel GATA6 mutation leading to congenital heart defects and permanent neonatal diabetes: A case report. Diabetes and Metabolism, 2013, 39, 370-374.	2.9	20
35	Levothyroxine replacement in primary congenital hypothyroidism: the higher the initial dose the higher the rate of overtreatment. Journal of Pediatric Endocrinology and Metabolism, 2016, 29, 133-8.	0.9	20
36	Clinical and genetic characteristics of 15 families with hereditary hypophosphatemia: Novel Mutations in PHEX and SLC34A3. PLoS ONE, 2018, 13, e0193388.	2.5	20

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37	Clinical and laboratory findings in the differential diagnosis of central precocious puberty and premature thelarche. Turk Pediatri Arsivi, 2015, 50, 20-26.	0.9	20
38	Psychomotor Retardation Caused by a Defective Thyroid Hormone Transporter: Report of Two Families with Different <i>MCT8</i> Mutations. Hormone Research in Paediatrics, 2014, 82, 261-271.	1.8	19
39	The association of serum lipocalin-2 levels with metabolic and clinical parameters in obese children: a pilot study. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, 525-8.	0.9	18
40	Genotypeâ€“phenotype correlation, gonadal malignancy risk, gender preference, and testosterone/dihydrotestosterone ratio in steroid 5-alpha-reductase type 2 deficiency: a multicenter study from Turkey. Journal of Endocrinological Investigation, 2019, 42, 453-470.	3.3	18
41	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
42	Treatment and control of hypertension in Turkish population: a survey on high blood pressure in primary care (the TURKSAHA study). Journal of Human Hypertension, 2006, 20, 355-361.	2.2	17
43	The Effect of L-Thyroxine Treatment on Hypothyroid Symptom Scores and Lipid Profile in Children with Subclinical Hypothyroidism. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2014, 6, 238-244.	0.9	16
44	The effect of L-thyroxine treatment on left ventricular functions in children with subclinical hypothyroidism. Archives of Disease in Childhood, 2015, 100, 130-137.	1.9	16
45	Gynecomastia: review. Pediatric Endocrinology Reviews, 2007, 5, 489-99.	1.2	16
46	Evaluation of neutrophil gelatinase-associated lipocalin in normoalbuminuric normotensive type 1 diabetic adolescents. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, .	0.9	15
47	Clinical profile and etiologies of children with central diabetes insipidus: a single-center experience from Turkey. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, 499-502.	0.9	15
48	Effect of gonadotropin-releasing hormone agonist therapy on body mass index and growth in girls with idiopathic central precocious puberty. Indian Journal of Endocrinology and Metabolism, 2015, 19, 267.	0.4	15
49	An unusual complication of dorsal penile nerve block for circumcision. Paediatric Anaesthesia, 2006, 16, 1094-1095.	1.1	14
50	Neonatal Diabetes Mellitus Accompanied by Diabetic Ketoacidosis and Mimicking Neonatal Sepsis: A Case Report. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2010, 2, 131-13.	0.9	14
51	Relationship between hair cadmium levels, indoor ETS exposure and wheezing frequency in children. Allergologia Et Immunopathologia, 2012, 40, 51-59.	1.7	14
52	Neuron-specific enolase and S100B protein in children with carbon monoxide poisoning: children are not just small adults. American Journal of Emergency Medicine, 2013, 31, 524-528.	1.6	14
53	Fasting and postprandial levels of a novel anorexigenic peptide nesfatin in childhood obesity. Journal of Pediatric Endocrinology and Metabolism, 2014, 27, 623-8.	0.9	13
54	Cross-reactivity of adrenal steroids with aldosterone may prevent the accurate diagnosis of congenital adrenal hyperplasia. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 701-4.	0.9	13

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55	Relationship between oxidative stress and blood glucose fluctuations evaluated with daily glucose monitoring in children with type 1 diabetes mellitus. Journal of Pediatric Endocrinology and Metabolism, 2016, 29, 435-9.	0.9	13
56	Serum galectin-1 levels are positively correlated with body fat and negatively with fasting glucose in obese children. Peptides, 2017, 95, 51-56.	2.4	13
57	Altered regional grey matter volume and appetite-related hormone levels in adolescent obesity with or without binge-eating disorder. Eating and Weight Disorders, 2021, 26, 2555-2562.	2.5	13
58	Clinical and Laboratory Characteristics of Hyperprolactinemia in Children and Adolescents: National Survey. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2019, 11, 149-156.	0.9	13
59	Audiological Findings in Celiac Disease. Orl, 2011, 73, 82-87.	1.1	12
60	Assessment of cardiac functions using tissue Doppler imaging in children with familial Mediterranean fever. Cardiology in the Young, 2012, 22, 188-193.	0.8	12
61	Isolated hyperthyrotropinemia in childhood obesity and its relation with metabolic parameters. Journal of Endocrinological Investigation, 2014, 37, 799-804.	3.3	12
62	Serum nesfatin-1 and leptin levels in non-obese girls with premature thelarche. Journal of Endocrinological Investigation, 2015, 38, 909-913.	3.3	12
63	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
64	Celiac disease in children and adolescents with Hashimoto Thyroiditis. Turk Pediatri Arsivi, 2016, 51, 100-105.	0.9	12
65	Oxytocin receptor gene polymorphism and low serum oxytocin level are associated with hyperphagia and obesity in adolescents. International Journal of Obesity, 2021, 45, 2064-2073.	3.4	11
66	Two Cases Presenting With Pubertal Delay and Diagnosed as Celiac Disease. Clinical Pediatrics, 2008, 47, 607-609.	0.8	10
67	Clinical and diagnostic characteristics of hyperprolactinemia in childhood and adolescence. Journal of Pediatric Endocrinology and Metabolism, 2013, 26, 1-11.	0.9	10
68	Decreased vitamin <scp>D</scp> levels in children with familial <scp>M</scp>editerranean fever. International Journal of Rheumatic Diseases, 2014, 17, 321-326.	1.9	10
69	A Comparison of Multiple Daily Insulin Therapy with Continuous Subcutaneous Insulin Infusion Therapy in Adolescents with Type 1 Diabetes Mellitus: A Single-Center Experience From Turkey. Journal of Pediatric Endocrinology and Metabolism, 2009, 22, 539-45.	0.9	9
70	A novel missense mutation in HSD17B3 gene in a 46, XY adolescent presenting with primary amenorrhea and virilization at puberty. Clinica Chimica Acta, 2015, 438, 154-156.	1.1	9
71	Social cognition and emotion regulation may be impaired in adolescents with obesity independent of the presence of binge eating disorder: a two-center study. Journal of Theoretical Social Psychology, 2019, 29, 887-894.	1.9	8
72	Nationwide Turkish Cohort Study of Hypophosphatemic Rickets. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 150-159.	0.9	8

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73	Serum chitotriosidase activity: is it a new inflammatory marker in obese children?. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, 63-7.	0.9	7
74	Low serum nesfatin-1 levels may be a contributing factor for monogenic obesity due to prohormone convertase 1 deficiency. Medical Hypotheses, 2013, 81, 172-174.	1.5	7
75	Anthropometric findings from birth to adulthood and their relation with karyotpye distribution in Turkish girls with Turner syndrome. American Journal of Medical Genetics, Part A, 2016, 170, 942-948.	1.2	7
76	A novel mutation in steroidogenic factor (SF1/NR5A1) gene in a patient with 46 XY DSD without adrenal insufficiency. Andrologia, 2017, 49, e12589.	2.1	7
77	A Synopsis of Current Practice in the Diagnosis and Management of Patients with Turner Syndrome in Turkey: A Survey of 18 Pediatric Endocrinology Centers. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2018, 10, 230-238.	0.9	7
78	Positive correlation of galanin with insulin resistance and triglyceride levels in obese children. Turkish Journal of Medical Sciences, 2018, 48, 560-568.	0.9	7
79	Increased concentrations of serum nesfatin-1 levels in childhood with idiopathic chronic malnutrition. Turkish Journal of Medical Sciences, 2018, 48, 378-385.	0.9	7
80	Cushing syndrome related to leukemic infiltration of the central nervous system: a case report and a possible role of LIF. Journal of Pediatric Endocrinology and Metabolism, 2013, 26, 967-70.	0.9	6
81	Association Between Vitamin D Receptor Polymorphism and Familial Mediterranean Fever Disease in Turkish Children. Biochemical Genetics, 2016, 54, 169-176.	1.7	6
82	Intratubular large cell hyalinizing Sertoli cell tumor of the testis presenting with prepubertal gynecomastia: a case report. Acta Clinica Belgica, 2017, 72, 254-258.	1.2	6
83	Gravesâ€™ disease following allogenic hematopoietic stem cell transplantation for severe aplastic anemia: case report and literature review. Journal of Pediatric Endocrinology and Metabolism, 2018, 31, 589-593.	0.9	6
84	A Mutation in INSR in a Child Presenting with Severe Acanthosis Nigricans. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2017, 9, 371-374.	0.9	6
85	Low Complement C1q/TNF-related Protein-13 Levels are Associated with Childhood Obesity But not Binge Eating Disorder. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2022, 14, 179-187.	0.9	6
86	Cardiac Functions in Children With Growth Hormone Deficiency Before and During Growth Hormoneâ€“Replacement Therapy. Pediatric Cardiology, 2011, 32, 766-771.	1.3	5
87	A novel activating ABCC8 mutation underlying neonatal diabetes mellitus in an infant presenting with cerebral sinovenous thrombosis. Journal of Pediatric Endocrinology and Metabolism, 2014, 27, 533-7.	0.9	5
88	Neonatal diabetes mellitus due to a novel mutation in the GATA6 gene accompanying renal dysfunction: A case report. American Journal of Medical Genetics, Part A, 2015, 167, 925-927.	1.2	5
89	Horseshoe kidney with growth retardation: Don't forget Turner syndrome. Turkish Journal of Pediatrics, 2016, 58, 227-229.	0.6	5
90	A nonsense variant in FGFR1: a rare cause of combined pituitary hormone deficiency. Journal of Pediatric Endocrinology and Metabolism, 2020, 33, 1613-1615.	0.9	5

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91	PLXNB1 mutations in the etiology of idiopathic hypogonadotropic hypogonadism. Journal of Neuroendocrinology, 2022, 34, e13103.	2.6	5
92	Endocrine cancer syndromes: an update. Minerva Pediatrica, 2014, 66, 533-47.	2.7	5
93	Relation of subepicardial adipose tissue thickness and clinical and metabolic parameters in obese prepubertal children. Pediatric Diabetes, 2010, 11, 556-562.	2.9	4
94	Factors associated with left atrial size in obese children: an observational study. Anatolian Journal of Cardiology, 2011, 11, 633-7.	0.4	4
95	Permanent neonatal diabetes caused by a novel mutation in the INS gene. Diabetes Research and Clinical Practice, 2013, 99, e5-e8.	2.8	4
96	Olfactory dysfunction in children with Kallmann syndrome: relation of smell tests with brain magnetic resonance imaging. Hormones, 2014, 14, 293-9.	1.9	4
97	Serum Level of Biotin Rather than the Daily Dosage Is the Main Determinant of Interference on Thyroid Function Assays. Hormone Research in Paediatrics, 2019, 92, 92-98.	1.8	4
98	Molecular Diagnosis of Monogenic Diabetes and Their Clinical/Laboratory Features in Turkish Children. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, 13, 0-0.	0.9	4
99	Growth curves for Turkish Girls with Turner Syndrome: Results of the Turkish Turner Syndrome Study Group. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2015, 7, 183-191.	0.9	4
100	Neonatal adrenal insufficiency: Turkish Neonatal and Pediatric Endocrinology and Diabetes Societies consensus report. Turk Pediatri Arsivi, 2019, 53, 239-243.	0.9	4
101	A Non-Endocrine Cause of Testicular Enlargement Mimicking Precocious Puberty: Testicular Microlithiasis. Journal of Pediatric Endocrinology and Metabolism, 2007, 20, 1237-40.	0.9	3
102	Osteopetrosis and Congenital Hypothyroidism Complicated by Slipped Capital Femoral Epiphysis. Endocrine Practice, 2010, 16, 646-649.	2.1	3
103	Acceleration of Puberty During Growth Hormone Therapy in a Child with Septo-Optic Dysplasia. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2014, 6, 116-118.	0.9	3
104	Presentation of central precocious puberty in two patients with Tay-Sachs disease. Hormones, 2018, 17, 415-418.	1.9	3
105	Comparison of the effectiveness of simple carbohydrates on hypoglycemic episodes in children and adolescents with type 1 diabetes mellitus: A randomized study in a diabetes camp. Pediatric Diabetes, 2020, 21, 1249-1255.	2.9	3
106	Clinical, genetic characteristics and treatment outcomes of children and adolescents with osteogenesis imperfecta: a two-center experience. Connective Tissue Research, 2022, 63, 349-358.	2.3	3
107	A 2-Year-Old Boy with a Testicular Mass. Pediatric Annals, 2010, 39, 471-474.	0.8	3
108	The Missense Alteration A5T of the Thyroid Peroxidase Gene is Pathogenic and Associated with Mild Congenital Hypothyroidism. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2015, 7, 238-241.	0.9	3

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109	Evaluation of Thyroid Function Tests in Children with Chronic Liver Diseases. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 143-149.	0.9	3
110	Anemia and Its Effect on Cardiovascular Findings in Obese Adolescents. Turkish Journal of Haematology, 2018, 35, 192-196.	0.5	3
111	Subepicardial adipose tissue thickness and its relation with anthropometric and clinical parameters in pubertal obese children. Journal of Endocrinological Investigation, 2010, 33, 715-719.	3.3	2
112	Recovery of Central Fever after GH Therapy in a Patient with GH Deficiency Secondary to Posttraumatic Brain Injury. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2015, 7, 77-79.	0.9	2
113	Relation of fetuin A levels with cardiac, subcutaneous lipid accumulation and insulin resistance parameters in Turkish obese children. Journal of Pediatric Endocrinology and Metabolism, 2016, 29, 669-73.	0.9	2
114	Identification of an AR mutation in Klinefelter syndrome during evaluation for penoscrotal hypospadias. Hormones, 2017, 16, 313-317.	1.9	2
115	Impaired systolic and diastolic left ventricular function in children and adolescents with congenital adrenal hyperplasia receiving corticosteroid therapy. Cardiology in the Young, 2019, 29, 319-324.	0.8	2
116	Comparison of the Effectiveness of Adult Height Prediction Methods in Children with Growth Hormone Deficiency. Endocrine Research, 2021, 46, 140-147.	1.2	2
117	Prevalence of anticardiolipin antibodies in type 1 diabetes and autoimmune thyroiditis. Polish Archives of Internal Medicine, 2010, 120, 71-75.	0.4	2
118	A Nove L Mutation in the AVPR2 Gene (222delA) Associated with X-Linked Nephrogenic Diabetes Insipidus In A Boy with Growth Failure. Endocrine Practice, 2010, 16, 231-236.	2.1	2
119	Prevalence of anticardiolipin antibodies in type 1 diabetes and autoimmune thyroiditis. , 2010, 120, 71-5.		2
120	Initial neutrophil/lymphocyte and lymphocyte/monocyte ratios can predict future insulin need in newly diagnosed type 1 diabetes mellitus. Journal of Pediatric Endocrinology and Metabolism, 2022, .	0.9	2
121	Obez ve saÄŸlÄ±klÄ± Ã§ocuklarda leptin ve leptin reseptÃ¶r gen polimorfizmleri. Cukurova Medical Journal, 2022, 47, 71-78.	0.2	2
122	Atypical comorbidities in a child considered to have type 1 diabetes led to the diagnosis of SLC29A3 spectrum disorder. Hormones, 2022, 21, 501-506.	1.9	2
123	Complex Urogenital Malformation Associated with Female Pseudohermaphroditism: Caudal Dysgenesis Syndrome. Journal of Pediatric Endocrinology and Metabolism, 2006, 19, 1171-4.	0.9	1
124	A 6-Year-Old Boy with a Nodule on His Hand. Pediatric Annals, 2012, 41, 360-361.	0.8	1
125	Brain injury markers: S100 calcium-binding protein B, neuron-specific enolase and glial fibrillary acidic protein in children with diabetic ketoacidosis. Pediatric Diabetes, 2018, 19, 1000-1006.	2.9	1
126	The Role of Thyrotropin-Releasing Hormone Stimulation Test in Management of Hyperthyrotropinemia in Infants. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2015, 7, 211-216.	0.9	1

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127	Child with RET proto-oncogene codon 634 mutation. Turkish Journal of Pediatrics, 2017, 59, 590-593.	0.6	1
128	Changes in the Frequency of Diabetic Ketoacidosis in Type I Diabetes Mellitus Cases at Diagnosis: A Fifteen-Year Single Center Experience. Journal of Pediatric Research, 2017, 4, 143-148.	0.2	1
129	A Novel De Novo Missense Mutation in HNF4A Resulting in Sulfonylurea-Responsive Maturity-onset Diabetes of the Young. Journal of Pediatric Research, 2018, 5, 156-160.	0.2	1
130	Case report of two siblings with familial ovarian dysgenesis. Minerva Pediatrica, 2007, 59, 57-9.	2.7	1
131	A 4-hour Profile of 17-hydroxyprogesterone in Salt-wasting Congenital Adrenal Hyperplasia: Is the Serial Monitoring Strategy Worth the Effort?. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, .	0.9	1
132	Does estimated glomerular filtration rate affect left ventricular function after ST elevation myocardial infarction?. European Heart Journal, 2013, 34, P1313-P1313.	2.2	0
133	A case of infantile-onset Graves. Turk Pediatri Arsivi, 2013, 48, 332-335.	0.9	0
134	A Case of Central Precocious Puberty Due to Concomitant Hypothalamic Hamartoma and Juvenile Pilocytic Astrocytoma. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2014, 6, 183-186.	0.9	0
135	A Novel Androgen Receptor Gene Mutation in a 46,XY Patient: Complete Androgen Insensitivity Syndrome. AACE Clinical Case Reports, 2016, 2, e202-e205.	1.1	0
136	Colchicine resistance and low vitamin D in familial Mediterranean fever. Renal Failure, 2016, 38, 340-341.	2.1	0
137	Comparison of the effects of the l-dopa and insulin tolerance tests on cortisol secretion. Journal of Endocrinological Investigation, 2018, 41, 901-907.	3.3	0
138	P6581Decrease in prevalence of hypertension in 15 years: a success story of a population through salt reduction initiatives. European Heart Journal, 2018, 39, .	2.2	0
139	Personality and subjective psychiatric symptoms of parents of obese youth: a controlled study. Journal of Theoretical Social Psychology, 2019, 29, 618-623.	1.9	0
140	The relationship of carotid intima-media thickness with anthropometric and metabolic parameters in patients with classic congenital adrenal hyperplasia. Turkish Journal of Medical Sciences, 2021, 51, 1738-1746.	0.9	0
141	Does fludrocortisone treatment cause hypomagnesemia in children with primary adrenal insufficiency?. Turkish Journal of Medical Sciences, 2021, 51, 231-237.	0.9	0
142	Vitamin D Dependent Rickets Type I: Two Cases Report. Turkiye Klinikleri Journal of Medical Sciences, 2012, 32, 1786-1790.	0.1	0
143	A Rare Karyotype of Turner Syndrome: 45.X/47.XXX. Guncel Pediatri, 2014, 12, 43-47.	0.1	0
144	Boy KÄ±salÄ±ÄŸÄ± ile BaÅŸvuran Kleidokranial Displazi TanÄ±sÄ± Alan Bir Olgu. Journal of Pediatric Research, 2014, 1, 92-94.	0.2	0

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145	A Rare Cause of Hypercalcemia in Childhood; Parathyroid Adenoma: Case Report and Review of the Literature. Guncel Pediatri, 2014, 12, 107-111.	0.1	0
146	A Rare Cause of a 46,XY Disorders of Sex Development: Persistent Mullerian Duct Syndrome. Journal of Dr Behcet Uz Children S Hospital, 2017, , .	0.1	0
147	Re: HLA genes as a predictive screening tool for celiac disease. Turk Pediatri Arsivi, 2017, 52, 184-184.	0.9	0
148	Early-Onset Isolated Bilateral Pheochromocytoma As a Major Clinical Manifestation of von-Hippel Lindau Syndrome Type 2C. Journal of Pediatric Research, 0, , 48-51.	0.2	0
149	Pediatric Bilateral Pheochromocytoma and Experience of Laparoscopic Cortical Sparing Adrenalectomy. Journal of Pediatric Research, 2018, 5, 218-220.	0.2	0
150	Clinical, Genetic Features and Treatment Results in Patients with Congenital Hyperinsulinemic Hypoglycemia: A Single Center Experience. Guncel Pediatri, 2020, 18, 317-335.	0.1	0