Serap Turan

List of Publications by Year in descending order

153 papers	3,762 citations	136740 32 h-index	168136 53 g-index
186 all docs	186 docs citations	186 times ranked	4867 citing authors

#	Article	IF	Citations
1	Catch-up Growth and Discontinuation of Fludrocortisone Treatment in Aldosterone Synthase Deficiency. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e106-e117.	1.8	5
2	Long-Term Efficacy of T3 Analogue Triac in Children and Adults With MCT8 Deficiency: A Real-Life Retrospective Cohort Study. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e1136-e1147.	1.8	15
3	Lack of GNAS Remethylation During Oogenesis May Be a Cause of Sporadic Pseudohypoparathyroidism Type Ib. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e1610-e1619.	1.8	5
4	A novel deletion involving the first GNAS exon encoding GsÎ± causes PHP1A without methylation changes at exon A/B. Bone, 2022, 157, 116344.	1.4	0
5	Steroid Hormone Profiles and Molecular Diagnostic Tools in Pediatric Patients With non-CAH Primary Adrenal Insufficiency. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e1924-e1931.	1.8	5
6	Dysosteosclerosis: Clinical and Radiological Evolution Reflecting Genetic Heterogeneity. JBMR Plus, 2022, 6, .	1.3	2
7	Does Genotypeâ€“Phenotype Correlation Exist in Vitamin D-Dependent Rickets Type IA: Report of 13 New Cases and Review of the Literature. Calcified Tissue International, 2021, 108, 576-586.	1.5	17
8	Endocrine disrupting chemicals and bone. Best Practice and Research in Clinical Endocrinology and Metabolism, 2021, 35, 101495.	2.2	20
9	Use of Insulin Degludec/Insulin Aspart in the Management of Diabetes Mellitus: Expert Panel Recommendations on Appropriate Practice Patterns. Frontiers in Endocrinology, 2021, 12, 616514.	1.5	10
10	Cranial MRI Abnormalities and Long-term Follow-up of the Lesions in 770 Girls With Central Precocious Puberty. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e2557-e2566.	1.8	9
11	Clinical and Hormonal Profiles Correlate With Molecular Characteristics in Patients With 11Î²-Hydroxylase Deficiency. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e3714-e3724.	1.8	20
12	Dysgenesis and Dysfunction of the Pancreas and Pituitary Due toFOXA2Gene Defects. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e4142-e4154.	1.8	6
13	Efficacy of the Novel Degludec/Aspart Insulin Co-formulation in Children and Adolescents with Type 1 Diabetes: A Real-life Experience with 1-year IDeg/Asp Therapy in Poorly Controlled and Non-compliant Patients. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, .	0.4	1
14	Adrenal steroids reference ranges in infancy determined by LC-MS/MS. Pediatric Research, 2021, , .	1.1	3
15	Broad-spectrum XX and XY gonadal dysgenesis in patients with a homozygous L193S variant in PPP2R3C. European Journal of Endocrinology, 2021, 186, 65-72.	1.9	1
16	Familial Hypomagnesemia with Hypercalciuria and Nephrocalcinosis Due to CLDN16 Gene Mutations: Novel Findings in Two Cases with Diverse Clinical Features. Calcified Tissue International, 2021, , 1.	1.5	1
17	Non-hormonal Clitoromegaly due to Clitoral Priapism Caused by Appendicitis/Appendectomy. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, .	0.4	1
18	Rare cause of severe hypertension in an adolescent boy presenting with short stature: Questions. Pediatric Nephrology, 2020, 35, 403-404.	0.9	1

#	Article	IF	Citations
19	Rare cause of severe hypertension in an adolescent boy presenting with short stature: Answers. Pediatric Nephrology, 2020, 35, 405-407.	0.9	4
20	A rare cause of hypertension in childhood: Questions. Pediatric Nephrology, 2020, 35, 77-78.	0.9	1
21	A rare cause of hypertension in childhood: Answers. Pediatric Nephrology, 2020, 35, 79-82.	0.9	5
22	Recommendations for improving clinical trial design to facilitate the study of youth-onset type 2 diabetes. Clinical Trials, 2020, 17, 87-98.	0.7	2
23	Adrenocortical carcinoma in atypical Beckwithâ€Wiedemann syndrome due to loss of methylation at imprinting control region 2. Pediatric Blood and Cancer, 2020, 67, e28042.	0.8	8
24	Rapid progression of type 2 diabetes and related complications in children and young peopleâ€”A literature review. Pediatric Diabetes, 2020, 21, 158-172.	1.2	34
25	Disease characteristics of MCT8 deficiency: an international, retrospective, multicentre cohort study. Lancet Diabetes and Endocrinology,the, 2020, 8, 594-605.	5.5	50
26	Recommendations for Diagnosis and Treatment of Pseudohypoparathyroidism and Related Disorders: An Updated Practical Tool for Physicians and Patients. Hormone Research in Paediatrics, 2020, 93, 182-196.	0.8	42
27	Cinacalcet as a First-Line Treatment in Neonatal Severe Hyperparathyroidism Secondary to Calcium Sensing Receptor (CaSR) Mutation. Hormone Research in Paediatrics, 2020, 93, 313-321.	0.8	14
28	Clinical Significance of Hypophosphatasemia in Children. Calcified Tissue International, 2020, 106, 608-615.	1.5	3
29	Hereditary vitamin D-resistant rickets: a report of four cases with two novel variants in the VDR gene and successful use of intermittent intravenous calcium via a peripheral route. Journal of Pediatric Endocrinology and Metabolism, 2020, 33, 557-562.	0.4	5
30	Revisiting Classical 3Î²-hydroxysteroid Dehydrogenase 2 Deficiency: Lessons from 31 Pediatric Cases. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e1718-e1728.	1.8	20
31	Management of Systemic Hypersensitivity Reactions to Gonadotropin-Releasing Hormone Analogues during Treatment of Central Precocious Puberty. Hormone Research in Paediatrics, 2020, 93, 66-72.	0.8	7
32	Once-Weekly Somapacitan vs Daily GH in Children With GH Deficiency: Results From a Randomized Phase 2 Trial. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e1847-e1861.	1.8	37
33	A Rare Cause of Hypophosphatemia: Raine Syndrome Changing Clinical Features with Age. Calcified Tissue International, 2020, 107, 96-103.	1.5	10
34	Nationwide Turkish Cohort Study of Hypophosphatemic Rickets. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 150-159.	0.4	8
35	Genotypic Sex and Severity of the Disease Determine the Time of Clinical Presentation in Steroid 17Î±-Hydroxylase/17,20-Lyase Deficiency. Hormone Research in Paediatrics, 2020, 93, 558-566.	0.8	11
36	Restoration of Height after 11 Years of Letrozole Treatment in 11Î²-Hydroxylase Deficiency. Hormone Research in Paediatrics, 2019, 92, 203-208.	0.8	5

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37	Low DHEAS Concentration in a Girl Presenting with Short Stature and Premature Pubarche: A Novel PAPSS2 Gene Mutation. Hormone Research in Paediatrics, 2019, 92, 262-268.	0.8	6
38	Letter to the Editor: Dysosteosclerosis related to the unique mutation in SLC29A3. Bone, 2019, 128, 115057.	1.4	0
39	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
40	Characteristics of Turkish children with Type 2 diabetes at onset: a multicentre, crossâ€sectional study. Diabetic Medicine, 2019, 36, 1243-1250.	1.2	6
41	Liraglutide in Children and Adolescents with Type 2 Diabetes. New England Journal of Medicine, 2019, 381, 637-646.	13.9	209
42	Ptosis as a unique hallmark for autosomal recessive WNT1â€associated osteogenesis imperfecta. American Journal of Medical Genetics, Part A, 2019, 179, 908-914.	0.7	16
43	Evaluation of growth and puberty in a child with a novel TBX19 gene mutation and review of the literature. Hormones, 2019, 18, 229-236.	0.9	6
44	Persistent MÃ¼llerian Duct Syndrome: A Rare But Important Etiology of Inguinal Hernia and Cryptorchidism. Sexual Development, 2019, 13, 264-270.	1.1	6
45	Fibroblast Growth Factor-23 and Matrix Extracellular Phosphoglycoprotein Levels in Healthy Children and, Pregnant and Puerperal Women. Hormone Research in Paediatrics, 2019, 92, 302-310.	0.8	2
46	PPP2R3C gene variants cause syndromic 46,XY gonadal dysgenesis and impaired spermatogenesis in humans. European Journal of Endocrinology, 2019, 180, 291-309.	1.9	18
47	Clinical and Laboratory Characteristics of Hyperprolactinemia in Children and Adolescents: National Survey. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2019, 11, 149-156.	0.4	13
48	MON-LB061 Cushing Syndrome Due to an Adrenocortical Carcinoma in a Baby with Atypical Beckwith-Wiedemann Syndrome. Journal of the Endocrine Society, 2019, 3, .	0.1	0
49	Acquired modification of sphingosine-1-phosphate lyase activity is not related to adrenal insufficiency. BMC Neurology, 2018, 18, 48.	0.8	3
50	Mutations in the mitochondrial ribosomal protein MRPS22 lead to primary ovarian insufficiency. Human Molecular Genetics, 2018, 27, 1913-1926.	1.4	39
51	Biallelic and monoallelic ESR2 variants associated with 46,XY disorders of sex development. Genetics in Medicine, 2018, 20, 717-727.	1.1	28
52	Diagnosis and management of pseudohypoparathyroidism and related disorders: first international Consensus Statement. Nature Reviews Endocrinology, 2018, 14, 476-500.	4.3	224
53	The Distribution of Different Types of Diabetes in Childhood: A Single Center Experience. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2018, 10, 125-130.	0.4	12
54	GNAS Complex Locus. , 2018, , 2173-2185.		1

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55	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.	0.4	5
56	Relation of serum IGF-1 and IGFBP3 levels with acute exacerbation in cystic fibrosis. , 2018, , .		0
57	Evaluation and Treatment Results of Ovarian Cysts in Childhood and Adolescence: A Multicenter, Retrospective Study of 100 Patients. Journal of Pediatric and Adolescent Gynecology, 2017, 30, 449-455.	0.3	11
58	Heterotrimeric G proteins in the control of parathyroid hormone actions. Journal of Molecular Endocrinology, 2017, 58, R203-R224.	1.1	28
59	Pycnodysostosis at otorhinolaryngology. International Journal of Pediatric Otorhinolaryngology, 2017, 95, 91-96.	0.4	7
60	The diagnostic value of soluble urokinase plasminogen activator receptor (suPAR) compared to C-reactive protein (CRP) and procalcitonin (PCT) in children with systemic inflammatory response syndrome (SIRS). Journal of Infection and Chemotherapy, 2017, 23, 17-22.	0.8	16
61	Hypoglycemia is common in children with cystic fibrosis and seen predominantly in females. Pediatric Diabetes, 2017, 18, 607-613.	1.2	20
62	Persistent hyperglycemia in a neonate: is it a complication of therapeutic hypothermia?. Turkish Journal of Pediatrics, 2017, 59, 193.	0.3	1
63	Current Nomenclature of Pseudohypoparathyroidism: Inactivating Parathyroid Hormone/Parathyroid Hormone-Related Protein Signaling Disorder. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2017, 9, 58-68.	0.4	12
64	<scp>GCK</scp> gene mutations are a common cause of childhoodâ€onset <scp>MODY</scp> (maturityâ€onset diabetes of the young) in Turkey. Clinical Endocrinology, 2016, 85, 393-399.	1.2	21
65	Cathepsin K osteoporosis trials, pycnodysostosis and mouse deficiency models: Commonalities and differences. Expert Opinion on Drug Discovery, 2016, 11, 457-472.	2.5	51
66	The diagnostic value of soluble urokinase plasminogen activator receptor compared with C-reactive protein and procalcitonin in children with febrile neutropenia. Pediatric Hematology and Oncology, 2016, 33, 200-208.	0.3	0
67	From pseudohypoparathyroidism to inactivating PTH/PTHrP signalling disorder (iPPSD), a novel classification proposed by the EuroPHP network. European Journal of Endocrinology, 2016, 175, P1-P17.	1.9	117
68	Risk factors for mortality caused by hypothalamic obesity in children with hypothalamic tumours. Pediatric Obesity, 2016, 11, 383-388.	1.4	11
69	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.	0.7	7
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.	1.8	20
71	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
72	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

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73	GNAS Complex Locus. , 2016, , 1-13.		0
74	Higher insulin detemir doses are required for the similar glycemic control: comparison of insulin detemir and glargine in children with type 1 diabetes mellitus. Pediatric Diabetes, 2015, 16, 361-366.	1.2	16
75	H Syndrome: A Multifaceted Histiocytic Disorder with Hyperpigmentation and Hypertrichosis. Acta Dermato-Venereologica, 2015, 95, 1021-1023.	0.6	21
76	Premature Pubarche, Hyperinsulinemia and Hypothyroxinemia: Novel Manifestations of Congenital Portosystemic Shunts (Abernethy Malformation) in Children. Hormone Research in Paediatrics, 2015, 83, 282-287.	0.8	30
77	Effects of leukemia inhibitory receptor gene mutations on human hypothalamoâ€“pituitaryâ€“adrenal function. Pituitary, 2015, 18, 456-460.	1.6	7
78	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.	0.4	13
79	Turner Syndrome and Associated Problems in Turkish Children: A Multicenter Study. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2015, 7, 27-36.	0.4	42
80	Normative Data of Thyroid Volume-Ultrasonographic Evaluation of 422 Subjects Aged 0-55 Years. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2015, 7, 98-101.	0.4	32
81	Homozygous Loss-of-function Mutations inSOHLH1in Patients With Nonsyndromic Hypergonadotropic Hypogonadism. Journal of Clinical Endocrinology and Metabolism, 2015, 100, E808-E814.	1.8	29
82	GNAS Spectrum of Disorders. Current Osteoporosis Reports, 2015, 13, 146-158.	1.5	147
83	European guidance for the molecular diagnosis of pseudohypoparathyroidism not caused by point genetic variants at GNAS: an EQA study. European Journal of Human Genetics, 2015, 23, 438-444.	1.4	27
84	Evidence of hormone resistance in a pseudo-pseudohypoparathyroidism patient with a novel paternal mutation in GNAS. Bone, 2015, 71, 53-57.	1.4	29
85	CLINICAL AND CEPHALOMETRIC ANALYSIS OF THREE CASES WITH PYCNODYSOSTOSIS: CASE REPORTS. Journal of Istanbul University Faculty of Dentistry, 2015, 49, 51.	0.2	1
86	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.	0.4	4
87	Current research on pycnodysostosis. Intractable and Rare Diseases Research, 2014, 3, 91-93.	0.3	39
88	Does common channel length affect surgical choice in female congenital adrenal hyperplasia patients?. Journal of Pediatric Urology, 2014, 10, 948-954.	0.6	10
89	<scp>AR</scp> and <scp>SRD</scp>5A2 gene mutations in a series of 51 Turkish 46,<scp>XY DSD</scp> children with a clinical diagnosis of androgen insensitivity. Andrology, 2014, 2, 572-578.	1.9	43
90	Identification of PENDRIN (SLC26A4) Mutations in Patients With Congenital Hypothyroidism and â€œApparentâ€•Thyroid Dysgenesis. Journal of Clinical Endocrinology and Metabolism, 2014, 99, E169-E176.	1.8	44

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91	Postnatal Establishment of Allelic GÎ±s Silencing as a Plausible Explanation for Delayed Onset of Parathyroid Hormone Resistance Owing to Heterozygous GÎ±s Disruption. Journal of Bone and Mineral Research, 2014, 29, 749-760.	3.1	64
92	The Frequency and the Effects of 21-Hydroxylase Gene Defects in Congenital Adrenal Hyperplasia Patients. Annals of Human Genetics, 2014, 78, 399-409.	0.3	11
93	Cathepsin K analysis in a pycnodysostosis cohort: demographic, genotypic and phenotypic features. Orphanet Journal of Rare Diseases, 2014, 9, 60.	1.2	35
94	Novel homozygous inactivating mutation of the calcium-sensing receptor gene (CASR) in neonatal severe hyperparathyroidismâ€"lack of effect of cinacalcet. Bone, 2014, 64, 102-107.	1.4	31
95	Prevalence of acne in primary school children and the relationship of acne with pubertal maturation. Turkderm, 2014, 48, 182-186.	0.0	0
96	Infantile loss of teeth: odontohypophosphatasia or childhood hypophosphatasia. European Journal of Pediatrics, 2013, 172, 851-853.	1.3	8
97	A novel homozygous TMEM70 mutation results in congenital cataract and neonatal mitochondrial encephalo-cardiomyopathy. Gene, 2013, 515, 197-199.	1.0	15
98	Maternal Thyroid Dysfunction and Neonatal Thyroid Problems. International Journal of Endocrinology, 2013, 2013, 1-6.	0.6	33
99	The <i>GNAS</i> Complex Locus and Human Diseases Associated with Loss-of-Function Mutations or Epimutations within This Imprinted Gene. Hormone Research in Paediatrics, 2013, 80, 229-241.	0.8	60
100	Pitfalls in the diagnosis of thyroid dysgenesis by thyroid ultrasonography and scintigraphy. European Journal of Endocrinology, 2012, 166, 43-48.	1.9	34
101	Clinical and molecular characterization of Turkish patients with familial hypomagnesaemia: novel mutations in TRPM6 and CLDN16 genes. Nephrology Dialysis Transplantation, 2012, 27, 667-673.	0.4	38
102	De NovoSTX16 Deletions: An Infrequent Cause of Pseudohypoparathyroidism Type Ib that Should Be Excluded in Sporadic Cases. Journal of Clinical Endocrinology and Metabolism, 2012, 97, E2314-E2319.	1.8	32
103	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
104	Loss of XLÎ±s (extra-large Î±s) imprinting results in early postnatal hypoglycemia and lethality in a mouse model of pseudohypoparathyroidism Ib. Proceedings of the National Academy of Sciences of the United States of America, 2012, 109, 6638-6643.	3.3	19
105	An Atypical Case of Familial Glucocorticoid Deficiency without Pigmentation Caused by Coexistent Homozygous Mutations inMC2R(T152K) andMC1R(R160W). Journal of Clinical Endocrinology and Metabolism, 2012, 97, E771-E774.	1.8	26
106	The prevalence and risk factors of premature thelarche and pubarche in 4â€•to 8â€yearâ€old girls. Acta Paediatrica, International Journal of Paediatrics, 2012, 101, e71-5.	0.7	46
107	Cognitive and psychosocial development in children with familial hypomagnesaemia. Magnesium Research, 2011, 24, 7-12.	0.4	4
108	Prevalence of type 1 diabetes mellitus in 6-18-yr-old school children living in Istanbul, Turkey. Pediatric Diabetes, 2011, 12, no-no.	1.2	16

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109	Exclusion of the GNAS locus in PHP-Ib patients with broad GNAS methylation changes: Evidence for an autosomal recessive form of PHP-Ib?. Journal of Bone and Mineral Research, 2011, 26, 1854-1863.	3.1	34
110	Radiologic and hormonal evaluation of pituitary abnormalities in patients with Bardetâ€“Biedl syndrome. Clinical Dysmorphology, 2011, 20, 26-31.	0.1	15
111	Extra-long GÎ±s Variant XLÎ±s Protein Escapes Activation-induced Subcellular Redistribution and Is Able to Provide Sustained Signaling. Journal of Biological Chemistry, 2011, 286, 38558-38569.	1.6	26
112	Puberty and Influencing Factors in Schoolgirls Living in Istanbul: End of the Secular Trend?. Pediatrics, 2011, 128, e40-e45.	1.0	54
113	Serum Alkaline Phosphatase Levels in Healthy Children and Evaluation of Alkaline Phosphatase z-scores in Different Types of Rickets. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2011, 3, 7-11.	0.4	118
114	Constitutional Growth Delay Pattern of Growth in Velo-Cardio-Facial Syndrome: Longitudinal follow up and final height of two cases. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2011, 1, 43-48.	0.4	4
115	Long-term clinical outcome and carrier phenotype in autosomal recessive hypophosphatemia caused by a novel DMP1 mutation. Journal of Bone and Mineral Research, 2010, 25, 2165-2174.	3.1	53
116	Content analysis of food advertising in Turkish television. Journal of Paediatrics and Child Health, 2010, 46, 427-430.	0.4	23
117	Recessive versus imprinted disorder: consanguinity can impede establishing the diagnosis of autosomal dominant pseudohypoparathyroidism type Ib. European Journal of Endocrinology, 2010, 163, 489-493.	1.9	5
118	T4 plus T3 Treatment in Children with Hypothyroidism and Inappropriately Elevated Thyroid-Stimulating Hormone despite Euthyroidism on T4 Treatment. Hormone Research in Paediatrics, 2010, 73, 108-114.	0.8	9
119	Circulating Insulin-like Growth Factor Binding Protein-4. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2010, 2, 17-20.	0.4	22
120	Deletion of the NoncodingGNASAntisense Transcript Causes Pseudohypoparathyroidism Type Ib and Biparental Defects ofGNASMethylationin cis. Journal of Clinical Endocrinology and Metabolism, 2010, 95, 3993-4002.	1.8	113
121	Deletion of the NoncodingGNASAntisense Transcript Causes Pseudohypoparathyroidism Type Ib and Biparental Defects ofGNASMethylationin cis. Molecular Endocrinology, 2010, 24, 1305-1306.	3.7	0
122	Identification of a novel dentin matrix protein-1 (DMP-1) mutation and dental anomalies in a kindred with autosomal recessive hypophosphatemia. Bone, 2010, 46, 402-409.	1.4	55
123	Deletion of the Noncoding GNAS Antisense Transcript Causes Pseudohypoparathyroidism Type Ib and Biparental Defects of GNAS Methylation in cis. Endocrine Reviews, 2010, 31, 400-400.	8.9	1
124	Cushing's Syndrome Due to a Non-Adrenal Ectopic Adrenocorticotropin-Secreting Ewing's Sarcoma in a Child. Journal of Pediatric Endocrinology and Metabolism, 2009, 22, 363-8.	0.4	7
125	9 Years Follow-up of a Patient with Pituitary Form of Resistance to Thyroid Hormones Comparison of Two Treatment Periods of D-Thyroxine and Triiodothyroacetic Acid. Journal of Pediatric Endocrinology and Metabolism, 2009, 22, 971-8.	0.4	20
126	Alopecia: Association with Resistance to Thyroid Hormones. Journal of Pediatric Endocrinology and Metabolism, 2009, 22, 1075-81.	0.4	10

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127	Hypogonadotropic Hypogonadism due to a Novel Missense Mutation in the First Extracellular Loop of the Neurokinin B Receptor. Journal of Clinical Endocrinology and Metabolism, 2009, 94, 3633-3639.	1.8	122
128	Puberty in a case with novel 17-hydroxylase mutation and the putative role of estrogen in development of pubic hair. European Journal of Endocrinology, 2009, 160, 325-330.	1.9	17
129	The role of leptin, soluble leptin receptor, resistin, and insulin secretory dynamics in the pathogenesis of hypothalamic obesity in children. European Journal of Pediatrics, 2009, 168, 1043-1048.	1.3	30
130	Comparison of capillary blood ketone measurement by electrochemical method and urinary ketone in treatment of diabetic ketosis and ketoacidosis in children. Acta Diabetologica, 2008, 45, 83-85.	1.2	30
131	Significance of acanthosis nigricans in childhood obesity. Journal of Paediatrics and Child Health, 2008, 44, 338-341.	0.4	51
132	Bone Mineral Density in Children with Non-Cystic Fibrosis Bronchiectasis. Respiration, 2008, 75, 432-436.	1.2	10
133	Screening of Parents and Siblings of Patients with Thyroid Dysgenesis by Thyroid Function Tests and Ultrasound. Hormone Research in Paediatrics, 2008, 70, 329-339.	0.8	5
134	Evaluation of Diagnosis and Treatment Results in Children with Graves' Disease with Emphasis on the Pubertal Status of Patients. Journal of Pediatric Endocrinology and Metabolism, 2008, 21, 745-51.	0.4	16
135	Alendronate treatment in children with osteogenesis imperfecta. Indian Pediatrics, 2008, 45, 105-9.	0.2	14
136	Adult height in Turkish patients with Turner syndrome without growth hormone treatment. Turkish Journal of Pediatrics, 2008, 50, 415-7.	0.3	4
137	Detection of Y Chromosomal Material in Patients with a 45,X Karyotype by PCR Method. Tohoku Journal of Experimental Medicine, 2007, 211, 243-249.	0.5	21
138	The effect of the mode of delivery on neonatal thyroid function. Journal of Maternal-Fetal and Neonatal Medicine, 2007, 20, 473-476.	0.7	10
139	The effect of economic status on height, insulin-like growth factor (IGF)-I and IGF binding protein-3 concentrations in healthy Turkish children. European Journal of Clinical Nutrition, 2007, 61, 752-758.	1.3	20
140	Reference data for bone speed of sound measurement by quantitative ultrasound in healthy children. Archives of Osteoporosis, 2007, 1, 37-41.	1.0	12
141	Severe diabetic ketoacidosis: hyperventilation or relative hypoventilation. Pediatric Critical Care Medicine, 2006, 7, 291.	0.2	0
142	Serum IGF-I and IGFBP-3 Levels of Turkish Children during Childhood and Adolescence: Establishment of Reference Ranges with Emphasis on Puberty. Hormone Research in Paediatrics, 2006, 65, 96-105.	0.8	47
143	Effect of Zinc Supplementation on Growth Hormone Secretion, IGF-I, IGFBP-3, Somatomedin Generation, Alkaline Phosphatase, Osteocalcin and Growth in Prepubertal Children with Idiopathic Short Stature. Journal of Pediatric Endocrinology and Metabolism, 2005, 18, 69-74.	0.4	52
144	Height, Weight, IGF-I, IGFBP-3 and Thyroid Functions in Prepubertal Children with Attention Deficit Hyperactivity Disorder: Effect of Methylphenidate Treatment. Hormone Research in Paediatrics, 2005, 63, 159-164.	0.8	40

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145	A Patient with Hypopituitarism and Isochromosome 18q Mosaicism. Hormone Research in Paediatrics, 2005, 64, 261-265.	0.8	3
146	Upper segment/lower segment ratio and armspanâ€“height difference in healthy Turkish children. Acta Paediatrica, International Journal of Paediatrics, 2005, 94, 407-413.	0.7	34
147	Addition of orlistat to conventional treatment in adolescents with severe obesity. European Journal of Pediatrics, 2004, 163, 738-741.	1.3	101
148	Two Patients with Kabuki Syndrome Presenting with Endocrine Problems. Journal of Pediatric Endocrinology and Metabolism, 2001, 14, 215-20.	0.4	27
149	Dysosteosclerosis from a unique mutation in SLC29A3. Bone Abstracts, 0, , .	0.0	2
150	From pseudohypoparathyroidism to inactivating PTH/PTHrP signaling disorder (iPPSD), a novel classification proposed by the European EuroPHP-network. Endocrine Abstracts, 0, , .	0.0	0
151	Diagnosis and management of pseudohypoparathyroidism and related disorders: first international consensus statement. Endocrine Abstracts, 0, , .	0.0	1
152	Clinical and Laboratory Characteristics of Hyperprolactinemia in Children and Adolescents: National Survey. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 0, , .	0.4	1
153	The Spectrum From Classic to Non-Classic 11Î²-Hydroxylase Deficiency. SSRN Electronic Journal, 0, , .	0.4	0