Samim Ozen

List of Publications by Year in descending order

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86 1,351 21 33
papers citations h-index g-index

88 88 1970
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#	Article	IF	CITATIONS
1	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
2	Effects of Environmental Endocrine Disruptors on Pubertal Development. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2011, 3, 1-6.	0.4	92
3	Natural History of Congenital Generalized Lipodystrophy: A Nationwide Study From Turkey. Journal of Clinical Endocrinology and Metabolism, 2016, 101, 2759-2767.	1.8	67
4	New Features for Child Metrics: Further Growth References and Blood Pressure Calculations. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 125-129.	0.4	61
5	Clinical presentations, metabolic abnormalities and end-organ complications in patients with familial partial lipodystrophy. Metabolism: Clinical and Experimental, 2017, 72, 109-119.	1.5	54
6	A Comprehensive Online Calculator for Pediatric Endocrinologists: ÇEDD Çözüm/TPEDS Metrics. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2017, 9, 182-184.	0.4	53
7	The relation of vitamin D deficiency with puberty and insulin resistance in obese children and adolescents. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, 83-7.	0.4	45
8	Clinical Course of Hashimoto's Thyroiditis and Effects of Levothyroxine Therapy on the Clinical Course of the Disease in Children and Adolescents. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2011, 3, 192-197.	0.4	44
9	The effects of Pilates on metabolic control and physical performance in adolescents with type 1 diabetes mellitus. Journal of Diabetes and Its Complications, 2012, 26, 348-351.	1.2	44
10	Turner Syndrome and Associated Problems in Turkish Children: A Multicenter Study. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2015, 7, 27-36.	0.4	42
11	Reliability and Validity of the Diabetes Eating Problem Survey in Turkish Children and Adolescents with Type 1 Diabetes Mellitus. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2017, 9, 323-328.	0.4	41
12	Frequency and Risk Factors of Endocrine Complications in Turkish Children and Adolescents with Sickle Cell Anemia. Turkish Journal of Haematology, 2013, 30, 25-31.	0.2	33
13	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
14	Effects of pesticides used in agriculture on the development of precocious puberty. Environmental Monitoring and Assessment, 2012, 184, 4223-4232.	1.3	30
15	Rapid Molecular Genetic Diagnosis with Next-Generation Sequencing in 46,XY Disorders of Sex Development Cases: Efficiency and Cost Assessment. Hormone Research in Paediatrics, 2017, 87, 81-87.	0.8	30
16	Idiopathic Hypogonadotropic Hypogonadism Caused by Inactivating Mutations in SRA1. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2016, 8, 125-134.	0.4	30
17	Distribution of Gene Mutations Associated with Familial Normosmic Idiopathic Hypogonadotropic Hypogonadism. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2012, 4, 121-126.	0.4	28
18	Aromatase Deficiency, a Rare Syndrome: Case Report. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2013, 5, 129-132.	0.4	28

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19	Neurodevelopment evaluation in children with congenital hypothyroidism by Bayley-III. Brain and Development, 2013, 35, 392-397.	0.6	27
20	Acquired partial lipodystrophy is associated with increased risk for developing metabolic abnormalities. Metabolism: Clinical and Experimental, 2015, 64, 1086-1095.	1.5	25
21	Gonadotropin-Dependent Precocious Puberty in a Patient with X-Linked Adrenal Hypoplasia Congenita Caused by a Novel DAX-1 Mutation. Hormone Research in Paediatrics, 2011, 75, 153-156.	0.8	22
22	A New Cause of Obesity Syndrome Associated with a Mutation in the Carboxypeptidase Gene Detected in Three Siblings with Obesity, Intellectual Disability and Hypogonadotropic Hypogonadism. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, 13, 52-60.	0.4	21
23	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
24	Serum Adiponectin and hsCRP Levels and Non-Invasive Radiological Methods in the Early Diagnosis of Cardiovascular System Complications in Children and Adolescents with Type 1 Diabetes Mellitus. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2013, 5, 174-181.	0.4	18
25	Does COVID-19 predispose patients to type 1 diabetes mellitus?. Clinical Pediatric Endocrinology, 2022, 31, 33-37.	0.4	17
26	Unexpected clinical features in a female patient with proopiomelanocortin (POMC) deficiency. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 691-4.	0.4	16
27	Renal complications of lipodystrophy: A closer look at the natural history of kidney disease. Clinical Endocrinology, 2018, 89, 65-75.	1.2	16
28	Current Diagnosis, Treatment and Clinical Challenges in the Management of Lipodystrophy Syndromes in Children and Young People. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 17-28.	0.4	16
29	Effectiveness of Continuous Subcutaneous Insulin Infusion Pump Therapy During Five Years of Treatment on Metabolic Control in Children and Adolescents with Type 1 Diabetes Mellitus. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2018, 10, 147-152.	0.4	15
30	Melanocortin 4 receptor (MC4R) gene variants in children and adolescents having familial early-onset obesity: genetic and clinical characteristics. European Journal of Pediatrics, 2020, 179, 1445-1452.	1.3	15
31	The Impact of Psycho-Educational Training on the Psychosocial Adjustment of Caregivers of Osteogenesis Imperfecta Patients. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2014, 6, 84-92.	0.4	14
32	Functional analyses of a novel missense and other mutations of the vitamin D receptor in association with alopecia. Scientific Reports, 2017, 7, 5102.	1.6	14
33	A large cohort of disorders of sex development and their genetic characteristics: 6 novel mutations in known genes. European Journal of Medical Genetics, 2021, 64, 104154.	0.7	14
34	Management of Childhood Thyroid Nodules: Surgical and Endocrinological Findings in a Large Group of Cases. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2017, 9, 222-228.	0.4	14
35	Psychiatric Approaches for Disorders of Sex Development: Experience of a Multidisciplinary Team. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2013, 5, 229-235.	0.4	13
36	Analysis of the GCK gene in 79 MODY type 2 patients: A multicenter Turkish study, mutation profile and description of twenty novel mutations. Gene, 2018, 641, 186-189.	1.0	12

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37	Exploring emotional dysregulation characteristics and comorbid psychiatric disorders in type 1 diabetic children with disordered eating behavior risk. Journal of Psychosomatic Research, 2020, 131, 109960.	1.2	11
38	Agricultural Pesticides and Precocious Puberty. Vitamins and Hormones, 2014, 94, 27-40.	0.7	10
39	A novel thyroid hormone receptor alpha gene mutation, clinic characteristics, and follow-up findings in a patient with thyroid hormone resistance. Hormones, 2019, 18, 223-227.	0.9	9
40	Wiedemann–Rautenstrauch syndrome: Report of a variant case. American Journal of Medical Genetics, Part A, 2012, 158A, 1434-1436.	0.7	8
41	The spectrum of HNF1A gene mutations in patients with MODY 3 phenotype and identification of three novel germline mutations in Turkish Population. Diabetes and Metabolic Syndrome: Clinical Research and Reviews, 2017, 11, S491-S496.	1.8	8
42	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
43	A rare cause of syndromic short stature: <scp>3M</scp> syndrome in three families. American Journal of Medical Genetics, Part A, 2021, 185, 461-468.	0.7	7
44	Parental Perception of Terminology of Disorders of Sex Development in Western Turkey. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2018, 10, 216-222.	0.4	7
45	The Value of Telemedicine for the Follow-up of Patients with New Onset Type 1 Diabetes Mellitus During COVID-19 Pandemic in Turkey: A Report of Eight Cases. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, 13, 468-472.	0.4	7
46	Aromatase Deficiency in Two Siblings with 46,XX Karyotype Raised as Different Genders: A Novel Mutation (p.R115X) in the <i>CYP19A1</i> Gene. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 109-112.	0.4	6
47	Local complications of insulin administration sites and effect on diabetes management. Journal of Clinical Nursing, 2022, 31, 2530-2538.	1.4	6
48	Loss of thymidine phosphorylase activity disrupts adipocyte differentiation and induces insulin-resistant lipoatrophic diabetes. BMC Medicine, 2022, 20, 95.	2.3	6
49	Rare Types of Turner Syndrome: Clinical Presentation and Cytogenetics in Five Cases. Laboratory Medicine, 2012, 43, 197-204.	0.8	5
50	Intraoperative Parathyroid Hormone Monitoring Corroborates the Success of Parathyroidectomy in Children. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2014, 6, 158-162.	0.4	5
51	The Gonadotropin-Releasing Hormone Analogue Therapy May Not Impact Final Height in Precocious Puberty of Girls With Onset of Puberty Aged 6 - 8 Years. Journal of Clinical Medicine Research, 2019, 11, 133-136.	0.6	5
52	Effect of Education on Impaired Hypoglycemia Awareness and Glycemic Variability in Children and Adolescents with Type 1 Diabetes Mellitus. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2019, 11, 189-195.	0.4	5
53	An unusual manifestation: Papillary thyroid carcinoma in a patient with ataxia-telengiectasia. Turkish Journal of Pediatrics, 2016, 58, 442-445.	0.3	5
54	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.	0.4	4

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55	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
56	Diagnostic and therapeutic approach in newborns with ambiguous genitale with disorder of sex development: consensus report of Turkish Neonatal and Pediatric Endocrinology and Diabetes Societies. Turk Pediatri Arsivi, 2019, 53, 198-208.	0.9	4
57	Initial Basal and Bolus Rates and Basal Rate Variability During Pump Treatment in Children and Adolescents. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, 13, 198-203.	0.4	4
58	A Neurofibromatosis Noonan Syndrome Patient Presenting with Abnormal External Genitalia. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 113-116.	0.4	4
59	Many admissions to the emergency departments with recurrent syncope attacks and seizures in an adolescent boy. European Journal of Pediatrics, 2009, 168, 761-763.	1.3	3
60	Primary Hyperparathyroidism in a Young Adult Presenting with Severe Hypertension. Journal of Pediatric Endocrinology and Metabolism, 2010, 23, 193-6.	0.4	3
61	Persistent Mýllerian Duct Syndrome with Transverse Testicular Ectopia: A Novel Anti-Müllerian Hormone Receptor Mutation. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2017, 9, 179-181.	0.4	3
62	The utility of reverse phenotyping: a case of lysinuric protein intolerance presented with childhood osteoporosis. Journal of Pediatric Endocrinology and Metabolism, 2021, 34, 957-960.	0.4	2
63	The Role of Cardiac Magnetic Resonance Imaging in the Determination of Cardiovascular Anomalies in Children and Young Adults with Turner Syndrome. Journal of Pediatric Research, 2019, 6, 203-207.	0.1	2
64	Glycated hemoglobin variability and microvascular complications in patients with typeÂ1 diabetes mellitus. Journal of Pediatric Endocrinology and Metabolism, 2020, 33, 1533-1537.	0.4	2
65	Renal Anomalies Associated with Ectopic Neurohypophysis - Original Article. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2011, 3, 56-59.	0.4	1
66	Multiple cutaneous hemangiomas in a patient with combined pituitary hormone deficiency. Journal of Pediatric Endocrinology and Metabolism, 2014, 27, 171-4.	0.4	1
67	Quality of Life and Psychological Well-being in Children and Adolescents with Disorders of Sex Development. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, 13, 23-33.	0.4	1
68	Is anti-Mullerian hormone an indicator of potential polycystic ovary syndrome in prepubertal girls with simple obesity?. Turkish Journal of Pediatrics, 2016, 58, 406-412.	0.3	1
69	Çocukluk Çağı Hashimoto Tiroiditi Tanılı Olguların Klinik Özellikleri ve İzlem Bulguları-Retrospektif Merkez Deneyimi. Konuralp Tip Dergisi, 2019, 11, 89-94.	Tek	1
70	Treatment and long-term follow-up of patients diagnosed with type 1 diabetes mellitus before age 5. Journal of Pediatric Endocrinology and Metabolism, 2021, 34, 201-207.	0.4	1
71	The Impact of the CEDD-NET on the Evaluation of Rare Disorders: A Multicenter Scientific Research Platform in the Field of Pediatric Endocrinology. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2022, 14, 216-220.	0.4	1
72	Prediction of Transient or Permanent Congenital Hypothyroidism. Journal of Pediatric Research, 2022, 9, 38-45.	0.1	1

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73	A Remarkable Coexistence of Systemic Capillary Leak Syndrome and Diabetes in an 11-Year-Old Boy: A Case Report and Review of the Literature. Case Reports in Immunology, 2020, 2020, 1-5.	0.2	o
74	Predictive low-glucose suspend system and glycemic variability. International Journal of Diabetes in Developing Countries, 0 , 1 .	0.3	0
75	The Effect of Diabetes Camp on Glycemic Variability in Children and Adolescents with Type 1 Diabetes Mellitus. Journal of Pediatric Research, 2021, 8, 303-308.	0.1	O
76	Insulin Pump Therapy: Its Effect on Metabolic Control in Different Age Groups in Children with Type 1 Diabetes. Journal of Pediatric Research, 2014, 1, 147-151.	0.1	0
77	Acquired partial lipodystrophy is associated with increased risk for metabolic complications. Endocrine Abstracts, 0, , .	0.0	O
78	Familial partial lipodystrophy linked to a novel peroxisome proliferator activator receptor-[gamma] mutation, H449L. Endocrine Abstracts, 0, , .	0.0	0
79	A case of Swyer syndrome with gonadoblastoma and dysgerminoma. Endocrine Abstracts, 0, , .	0.0	O
80	Fibrous dysplasia in McCune Albright syndrome; treatment and follow up. Bone Abstracts, 0, , .	0.0	0
81	Treatment and Follow-up in a Case with Diazoxide Treatment-Resistant Hyperinsulinemic Hypoglycaemia. Journal of Pediatric Research, 0, , 245-248.	0.1	0
82	Bannayan-Riley-Ruvalcaba Syndrome in a Case Evaluated Due to Multinodular Goiter. Journal of Pediatric Research, 2018, 5, 214-217.	0.1	0
83	Pediatric Bilateral Pheochromocytoma and Experience of Laparoscopic Cortical Sparing Adrenalectomy. Journal of Pediatric Research, 2018, 5, 218-220.	0.1	0
84	HİPOKALSEMİ İLE BAŞVURAN BİR OTOİMMUN POLİGLANDÜLER SENDROM TİP 1 OLGUSU. Kır Fakültesi Dergisi, 0, , 109-114.	ıkkale Ã	.œnjyersitesi T.
85	Psychiatric view for disorders of sex development: a 12-year experience of a multidisciplinary team in a university hospital. Journal of Pediatric Endocrinology and Metabolism, 2020, 33, 605-611.	0.4	0
86	The Risk Factors of Puberty Precocious in Girls: Is the Condition Related with Polychlorobiphenyls?. Journal of Pediatric Research, 2021, 8, 408-413.	0.1	0