Mustafa Kendirci

List of Publications by Year in descending order

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49 papers	1,844 citations	566801 15 h-index	264894 42 g-index
49 all docs	49 docs citations	49 times ranked	2855 citing authors

#	Article	IF	Citations
1	Evaluation of micronutrient levels in children and adolescents with obesity and their correlation with the components of metabolic syndrome. Turkish Journal of Pediatrics, 2021, 63, 48.	0.3	10
2	Endocrine Disruptors and Polycystic Ovary Syndrome: Phthalates. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 393-400.	0.4	18
3	The effect of thiaminepyrophosphate levels on mortality and morbidity Ä±n patients with stress hyperglycemia. , 2020, , 25-29.		0
4	Neutropenia and Increased Mean Corpuscular Volume (MCV) With Abnormal Neurologic Findings: A Case of Cobalamin D Deficiency. Journal of Pediatric Hematology/Oncology, 2019, 41, e54-e56.	0.3	5
5	The Incidence of Cystic Fibrosis in the Central Region of Anatolia in Turkey Between 2015 and 2016. Balkan Medical Journal, 2019, 36, 179-183.	0.3	15
6	Comparison of Treatment Regimens in Management of Severe Hypercalcemia Due to Vitamin D Intoxication in Children. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2019, 11, 140-148.	0.4	8
7	Screening Inherited Metabolic Disorder in Children with Intellectual Disability and Epilepsy. Turk Noroloji Dergisi = Turkish Journal of Neurology, 2019, 25, 135-139.	0.1	0
8	Menstrual Characteristics and Related Problems in 9- to 18-Year-Old Turkish School Girls. Journal of Pediatric and Adolescent Gynecology, 2018, 31, 350-355.	0.3	21
9	Arthropathy-like findings and a carpal tunnel syndrome as the presenting features of Scheie syndrome: Three cases from the same family. Turkish Journal of Pediatrics, 2018, 60, 344-347.	0.3	4
10	A Rare Cause of Hypothalamic Obesity, Rohhad Syndrome: 2 Cases. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2018, 10, 382-386.	0.4	6
11	Urinary levels of pyridinoline and deoxypyridinoline and bone mineral density in children with type 1 diabetes mellitus. Endocrine Research, 2017, 42, 281-286.	0.6	4
12	Evaluation of vitamin D prophylaxis in 3â€“36-month-old infants and children. Journal of Pediatric Endocrinology and Metabolism, 2017, 30, 543-549.	0.4	3
13	Tyrosinemia type <scp>II</scp>: Novel mutations in <scp>TAT</scp> in a boy with unusual presentation. Pediatrics International, 2016, 58, 1069-1072.	0.2	7
14	Long-term efficacy of lipoprotein apheresis in the management of familial hypercholesterolaemia: Application of two different apheresis techniques in childhood. Transfusion and Apheresis Science, 2016, 54, 282-288.	0.5	4
15	The outcome of seven patients with hereditary tyrosinemia type 1. Journal of Pediatric Endocrinology and Metabolism, 2016, 29, 1151-1157.	0.4	8
16	Increased Serum Phthalates (MEHP, DEHP) and Bisphenol A Concentrations in Children With Autism Spectrum Disorder. Journal of Child Neurology, 2016, 31, 629-635.	0.7	75
17	Pseudotumour Cerebri Presentation in a Child Under the Gonadotropin-Releasing Hormone Agonist Treatment. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2016, 8, 365-367.	0.4	14
18	Complex Glycerol Kinase Deficiency and Adrenocortical Insufficiency in Two Neonates. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2016, 8, 468-471.	0.4	8

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19	Plasma glutamine and cystine are decreased and negatively correlated with endomysial antibody in children with celiac disease. Asia Pacific Journal of Clinical Nutrition, 2016, 25, 452-6.	0.3	2
20	The Effects of Ketogenic Diet on Seizures, Cognitive Functions, and Other Neurological Disorders in Classical Phenotype of Glucose Transporter 1 Deficiency Syndrome. Neuropediatrics, 2015, 46, 313-320.	0.3	21
21	The endocrine disruptor bisphenol A may play a role in the aetiopathogenesis of polycystic ovary syndrome in adolescent girls. Acta Paediatrica, International Journal of Paediatrics, 2015, 104, e171-7.	0.7	63
22	AMINO ACID LEVELS IN CHILDREN WITH CELIAC DISEASE. Nutricion Hospitalaria, 2015, 32, 139-43.	0.2	6
23	An Unusual Presentation of Parathyroid Adenoma in an Adolescent: Calcific Achilles Tendinitis. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2015, 7, 333-335.	0.4	6
24	Premature thelarche related to fennel tea consumption?. Journal of Pediatric Endocrinology and Metabolism, 2014, 27, 175-9.	0.4	5
25	The absence of insulin resistance in metabolic syndrome definition leads to underdiagnosing of metabolic risk in obese patients. European Journal of Pediatrics, 2012, 171, 1331-1337.	1.3	29
26	Hemophagocytic syndrome in a 4â€•monthâ€•old infant with biotinidase deficiency. Pediatric Blood and Cancer, 2012, 59, 191-193.	0.8	23
27	Pituitary duplication: a rare cause of precocious puberty. Child's Nervous System, 2011, 27, 1157-1160.	0.6	11
28	The Association of Serum Sialic Acid with Carotid Intima-Media Thickness and Anthropometric and Metabolic Parameters in Obese Children and Adolescents. Annals of Nutrition and Metabolism, 2011, 59, 139-144.	1.0	4
29	Lupus Nephritis in a Child with Type I Diabetes Mellitus. Journal of Tropical Pediatrics, 2011, 57, 396-398.	0.7	3
30	Does Early Treatment Prevent Deafness in Thiamine-Responsive Megaloblastic Anaemia Syndrome?. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2011, 3, 36-39.	0.4	22
31	Primary adrenal failure due to viral infection in an infant. European Journal of Pediatrics, 2010, 169, 887-889.	1.3	12
32	Neck circumference: an additional tool of screening overweight and obesity in childhood. European Journal of Pediatrics, 2010, 169, 733-739.	1.3	97
33	Neurofibromatosis Type 1 with Overlap Turner Syndrome and Klinefelter Syndrome. Journal of Tropical Pediatrics, 2010, 56, 69-72.	0.7	8
34	Hook Effect: A Pitfall Leading to Misdiagnosis of Hypoaldosteronism in an Infant with Pseudohypoaldosteronism. Hormone Research in Paediatrics, 2010, 74, 72-75.	0.8	15
35	Familial Glucocorticoid Deficiency Type 2: A Case Report - Case Report. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2010, 2, 122-125.	0.4	11
36	Vitamin D Deficiency Rickets Mimicking Pseudohypoparathyroidism-Case Report. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2010, 2, 173-175.	0.4	35

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37	Insulin Resistance in Obese Children and Adolescents: HOMA-IR Cut-Off Levels in the Prepubertal and Pubertal Periods - Original Article. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2010, 2, 100-106.	0.4	230
38	An autosomal recessive adducted thumb-club foot syndrome observed in Turkish cousins. Clinical Genetics, 2008, 51, 61-64.	1.0	42
39	Thiamine Withdrawal Can Lead to Diabetic Ketoacidosis in Thiamine Responsive Megaloblastic Anemia: Report of Two Siblings. Journal of Pediatric Endocrinology and Metabolism, 2008, 21, 393-7.	0.4	18
40	An occasional side effect in the treatment of congenital hypothyroidism: hair loss. European Journal of Pediatrics, 2006, 165, 500-501.	1.3	2
41	Protein oxidation in obesity and insulin resistance. European Journal of Pediatrics, 2006, 165, 753-756.	1.3	64
42	Homeostasis Model Assessment Is More Reliable Than the Fasting Glucose/Insulin Ratio and Quantitative Insulin Sensitivity Check Index for Assessing Insulin Resistance Among Obese Children and Adolescents. Pediatrics, 2005, 115, e500-e503.	1.0	851
43	Vulvovaginal Candidiasis in Children and Adolescents with Type I Diabetes Mellitus. Journal of Pediatric Endocrinology and Metabolism, 2004, 17, 1545-9.	0.4	14
44	Insulin-like Growth Factor-I and Insulin-like Growth Factor Binding Protein-3 Levels of Children Living in an Iodine- and Selenium-Deficient Endemic Goiter Area. Biological Trace Element Research, 2002, 90, 25-30.	1.9	15
45	Iodine and selenium deficiency in school-children in an endemic goiter area in Turkey. Journal of Pediatric Endocrinology and Metabolism, 2002, 15, 1027-31.	0.4	10
46	Gliclazide-Induced Hepatitis, Hemiplegia and Dysphasia in a Suicide Attempt. Journal of Pediatric Endocrinology and Metabolism, 2001, 14, 1157-9.	0.4	11
47	A case of malignant histiocytosis associated with skin involvement mimicking kwashiorkor. Pediatric Dermatology, 2001, 18, 545-546.	0.5	4
48	Congenital alacrima in a patient with G (Opitz Frias) syndrome. Human Genetics, 1996, 97, 540-542.	1.8	0
49	Late Diagnosed Argininemia. The Journal of Pediatric Academy, 0, , 39-42.	0.1	0