

Prof. ERDAL EREN

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Education Information

Post Doctorate of Medicine, Bursa Uludağ University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, Turkey 2007 - 2011

Expertise In Medicine, Suleyman Demirel University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, Turkey 2001 - 2006

Undergraduate, Akdeniz University, Faculty Of Medicine, Tıp Pr., Turkey 1992 - 1998

Research Areas

Medicine, Health Sciences, Internal Medicine Sciences, Child Health and Diseases, Pediatric Endocrinology and Metabolism

Academic Titles / Tasks

Associate Professor, Bursa Uludağ University, Tıp Fakültesi, Dahili Tıp Bilimleri, 2015 - Continues

Assistant Professor, Bursa Uludağ University, Tıp Fakültesi, Dahili Tıp Bilimleri, 2013 - 2015

Research Assistant, Bursa Uludağ University, Tıp Fakültesi, Dahili Tıp Bilimleri, 2007 - 2011

Research Assistant, Suleyman Demirel University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2000 - 2005

Academic and Administrative Experience

Bursa Uludağ University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2007 - 2011

Courses

Hipofiz yetmezliği, Postgraduate, 2016 - 2017, 2017 - 2018

Ananmez alma, Postgraduate, 2016 - 2017, 2017 - 2018

Büyüme geriliği, Postgraduate, 2016 - 2017, 2017 - 2018

Advising Theses

EREN E., Çocuk endokrinoloji kliniğinde takip edilen hipofiz yetmezlikli olguların geriye dönük değerlendirilmesi, Expertise In Medicine, K.DİNÇ(Student), 2018

Articles Published in Journals That Entered SCI, SSCI and AHCI Indexes

1. The attitudes, experiences, and self-competencies of pediatric endocrinology fellows and attending

physicians regarding diabetes technology: the Turkey experience

Mutlu G. Y. , EREN E., Eviz E., Gokce T., Sakarya S., Hatun S.

JOURNAL OF PEDIATRIC ENDOCRINOLOGY & METABOLISM, vol.35, pp.611-616, 2022 (Journal Indexed in SCI)

- II. **Novel homozygous missense mutation in NARS1 gene: A new neurodevelopmental disorder with microcephaly**
TEMEL Ş. G. , ÖZEMRİ SAĞ Ş., EREN E., Deniz E.
EUROPEAN JOURNAL OF HUMAN GENETICS, vol.30, no.SUPPL 1, pp.352, 2022 (Journal Indexed in SCI)
- III. **Maltodextrin May Be a Promising Treatment Modality After Near-total Pancreatectomy in Infants Younger Than Six Months with Persistent Hyperinsulinism: A Case Report.**
Denkboy Ongen Y., Eren E., Sağlam H.
Journal of clinical research in pediatric endocrinology, 2021 (Journal Indexed in SCI Expanded)
- IV. **Compound Heterozygous Variants in <i>FAM111A</i> Cause Autosomal Recessive Kenny-Caffey Syndrome Type 2.**
Eren E., Tezcan Unlu H., Ceylaner S., Tarim Ö. F.
Journal of clinical research in pediatric endocrinology, 2021 (Journal Indexed in SCI Expanded)
- V. **Telemedicine experiences at a pediatric endocrinology clinic during the COVID-19 pandemic**
DENKBOY ÖNGEN Y., EREN E., ŞAHİN K. C. , BUHUR PİRİMOĞLU M., SAĞLAM H., TARIM Ö. F.
IRISH JOURNAL OF MEDICAL SCIENCE, 2021 (Journal Indexed in SCI)
- VI. **Molecular Diagnosis of Monogenic Diabetes and Their Clinical/Laboratory Features in Turkish Children**
Gökşen D., Yeşilkaya E., Özen S., Kor Y., Eren E., Korkmaz Ö., Berberoğlu M., Karagüzel G., Er E., Abacı A., et al.
Journal of clinical research in pediatric endocrinology, vol.13, pp.433-438, 2021 (Journal Indexed in SCI)
- VII. **Genotype and Phenotype Heterogeneity in Neonatal Diabetes: A Single Centre Experience in Turkey.**
Denkboy Ongen Y., Eren E., Demirbas O., Sobu E., Ellard S., Tarim Ö. F. , De Franco E.
Journal of clinical research in pediatric endocrinology, vol.13, pp.80-87, 2021 (Journal Indexed in SCI)
- VIII. **Magnetic resonance spectroscopy to assess hepatic steatosis in patients with lipodystrophy**
ALTAY C., SEÇİL M., Adiyaman S. C. , Saydam B. O. , Demir T., Akinci G., YILDIRIM ŞİMŞİR İ., EREN E., Keskin E. T. , Demir L., et al.
TURKISH JOURNAL OF GASTROENTEROLOGY, vol.31, no.8, pp.588-595, 2020 (Journal Indexed in SCI)
- IX. **Nationwide Turkish Cohort Study of Hypophosphatemic Rickets**
ŞIKLAR Z., DEMİRCİOĞLU S., BEREKET A., Bas F., GÜRAN T., Akberzade A., ABACI A., DEMİR K., BÖBER E., Ozbek M. N. , et al.
Journal of clinical research in pediatric endocrinology, vol.12, no.2, pp.150-159, 2020 (Journal Indexed in SCI Expanded)
- X. **Three different faces of TAC1 mutations.**
ÇEKİÇ Ş., Cicek F., Karali Y., Gorukmez O., EREN E., Kilic S. Ş.
Scandinavian journal of immunology, vol.91, no.6, 2020 (Journal Indexed in SCI Expanded)
- XI. **Novel Mutations in Obesity-related Genes in Turkish Children with Non-syndromic Early Onset Severe Obesity: A Multicentre Study**
AKINCI A., Turkkahraman D., TEKEDERELİ İ., Ozer L., EVREN B., ŞAHİN İ., Kalkan T., Curek Y., ÇAMTOSUN E., DÖĞER E., et al.
JOURNAL OF CLINICAL RESEARCH IN PEDIATRIC ENDOCRINOLOGY, vol.11, no.4, pp.341-349, 2019 (Journal Indexed in SCI)
- XII. **Evaluation of AGP reports in patients with type 1 diabetes using intermittently viewed continuous glucose measurement system (iCGM)**
EREN E., DENKBOY ÖNGEN Y., DEMİRBAŞ Ö., Tarim Ö. F.
HORMONE RESEARCH IN PAEDIATRICS, vol.91, pp.293, 2019 (Journal Indexed in SCI)
- XIII. **Clinical and Laboratory Characteristics of Hyperprolactinemia in Children and Adolescents: National Survey**
EREN E., Ergur A. T. , İŞGÜVEN Ş. P. , Bitkin E. C. , BERBEROĞLU M., ŞIKLAR Z., Bas F., Yel S., Bas S., Sobu E., et al.
JOURNAL OF CLINICAL RESEARCH IN PEDIATRIC ENDOCRINOLOGY, vol.11, no.2, pp.149-156, 2019 (Journal

Indexed in SCI)

- XIV. **Comparison of Treatment Regimens in Management of Severe Hypercalcemia Due to Vitamin D Intoxication in Children**
DEMİR K., DÖNERAY H., Kara C., Atay Z., Cetinkaya S., Cayir A., ANIK A., EREN E., Ucakurk A., Yilmaz G. C. , et al.
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- XV. **Liver Involvement in Congenital Hypopituitarism.**
ALTAY D., EREN E., ÖZKAN T. M. , ÖZGÜR T., Tarim Ö. F.
Indian journal of pediatrics, vol.86, no.5, pp.412-416, 2019 (Journal Indexed in SCI Expanded)
- XVI. **Genotype-phenotype correlation, gonadal malignancy risk, gender preference, and testosterone/dihydrotestosterone ratio in steroid 5-alpha-reductase type 2 deficiency: a multicenter study from Turkey**
Abaci A., Catli G., Kirbiyik O., Sahin N. M. , Abali Z. Y. , Unal E., Siklar Z., Mengen E., Ozen S., Guran T., et al.
JOURNAL OF ENDOCRINOLOGICAL INVESTIGATION, vol.42, no.4, pp.453-470, 2019 (Journal Indexed in SCI)
- XVII. **Perceived Expressed Emotion, Emotional and Behavioral Problems and Self-Esteem in Obese Adolescents: A Case-Control Study**
Colpan M., ERAY Ş., EREN E., VURAL A. P.
JOURNAL OF CLINICAL RESEARCH IN PEDIATRIC ENDOCRINOLOGY, vol.10, no.4, pp.357-363, 2018 (Journal Indexed in SCI)
- XVIII. **Incidence of Type 1 Diabetes in Children Aged Below 18 Years during 2013-2015 in Northwest Turkey**
Poyrazoglu S., Bundak R., Abali Z. Y. , Onal H., Sarikaya S., Akgun A., Bas S., Abali S., BEREKET A., EREN E., et al.
Journal of clinical research in pediatric endocrinology, vol.10, no.4, pp.336-342, 2018 (Journal Indexed in SCI Expanded)
- XIX. **Homozygous loss-of-function mutations in SLC26A7 cause goitrous congenital hypothyroidism.**
CANGÜL H., Liao X., Schoenmakers E., Kero J., Barone S., Srichomkwun P., Iwayama H., Serra E. G. , SAĞLAM H., EREN E., et al.
JCI insight, vol.3, no.20, 2018 (Journal Indexed in SCI Expanded)
- XX. **Renal complications of lipodystrophy: A closer look at the natural history of kidney disease**
AKINCI B., ÜNLÜ Ş. M. , ÇELİK A., YILDIRIM ŞİMŞİR I., ŞEN S., NUR B., Keskin F. E. , Saydam B. O. , Ozdemir N. K. , ŞARER YÜREKLİ B. P. , et al.
CLINICAL ENDOCRINOLOGY, vol.89, no.1, pp.65-75, 2018 (Journal Indexed in SCI)
- XXI. **Pituitary Stalk Interruption Syndrome (PSIS) is not a Rare Cause of the Congenital Hypopituitarism**
EREN E., YAZICI Z., DEMİRBAŞ Ö., Gulleroglu N. B. , Tarim Ö. F.
HORMONE RESEARCH IN PAEDIATRICS, vol.90, pp.514-515, 2018 (Journal Indexed in SCI)
- XXII. **A Neurological Disease Mimicking Central Hypothyroidism: MCT8 Deficiency**
DEMİRBAŞ Ö., EREN E., Tarim O.
HORMONE RESEARCH IN PAEDIATRICS, vol.90, pp.610, 2018 (Journal Indexed in SCI)
- XXIII. **Nationwide Hypophosphatemic Rickets Study**
ŞIKLAR Z., DEMİRCİOĞLU S., BEREKET A., ABACI A., Bas F., DEMİR K., GÜRAN T., Akberzade A., BÖBER E., Ozbek M. N. , et al.
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- XXIV. **Analysis of the GCK gene in 79 MODY type 2 patients: A multicenter Turkish study, mutation profile and description of twenty novel mutations**
AYKUT A., KARACA E., ONAY H., Goksen D., ÇETİNKALP Ş., EREN E., ERSOY B., Cakir E. P. , Buyukinan M., Kara C., et al.
GENE, vol.641, pp.186-189, 2018 (Journal Indexed in SCI)
- XXV. **Determining residual adipose tissue characteristics with MRI in patients with various subtypes of lipodystrophy**
ALTAY C., SEÇİL M., Demir T., Atik T., Akinci G., Kutbay N. O. , Temeloglu E. K. , YILDIRIM ŞİMŞİR I., Ozisik S., Demir L., et al.

- DIAGNOSTIC AND INTERVENTIONAL RADIOLOGY, vol.23, no.6, pp.428-434, 2017 (Journal Indexed in SCI)
- XXVI. **The Relationship Between Perceived Family Climate and Glycemic Control in Type 1 Diabetes Mellitus Adolescent Patients**
Eray Ş., Ucar H. N. , Cetinkaya F., EREN E., Vural P.
JOURNAL OF CLINICAL RESEARCH IN PEDIATRIC ENDOCRINOLOGY, vol.9, no.3, pp.253-259, 2017 (Journal Indexed in SCI)
- XXVII. **Effects of long-term consumption of high fructose corn syrup containing peach nectar on body weight gain in sprague dawley rats**
ÖZCAN SİNİR G., SUNA S., Inan S., Bagdas D., TAMER C. E. , ÇOPUR Ö. U. , SIĞIRLI D., SARANDÖL E., SÖNMEZ G., ERCAN İ., et al.
FOOD SCIENCE AND TECHNOLOGY, vol.37, no.2, pp.337-343, 2017 (Journal Indexed in SCI)
- XXVIII. **PHENOTYPIC, HORMONAL AND MOLECULAR GENETIC CHARACTERISTICS OF 5-ALPHA REDUCTASE TYPE 2 DEFICIENCY PATIENTS: A MULTICENTER STUDY FROM TURKEY**
ABACI A., ÇATLI G., Kirbiyik O., Sahin N. M. , Abali Z. Y. , Unal E., ŞIKLAR Z., Ucakturk E. M. , ÖZEN S., GÜRAN T., et al.
HORMONE RESEARCH IN PAEDIATRICALS, vol.88, pp.551-552, 2017 (Journal Indexed in SCI)
- XXIX. **CLINICAL AND LABORATORY CHARACTERISTICS OF HYPERPROLACTINEMIC CHILDREN AND ADOLESCENTS: NATIONAL SURVEY**
EREN E., Ergur A. T. , İŞGÜVEN Ş. P. , Bitkin E. C. , BERBEROĞLU M., ŞIKLAR Z., Genens M., Doğan M., Yel S., Bas S., et al.
HORMONE RESEARCH IN PAEDIATRICALS, vol.88, pp.313-314, 2017 (Journal Indexed in SCI)
- XXX. **Homozygous Ala65Pro Mutation with V89L Polymorphism in SRD5A2 Deficiency**
EREN E., EDGÜNLÜ T., Asut E., Celik S. K.
JOURNAL OF CLINICAL RESEARCH IN PEDIATRIC ENDOCRINOLOGY, vol.8, no.2, pp.218-223, 2016 (Journal Indexed in SCI)
- XXXI. **The Etiology and Clinical Features of Non-CAH Gonadotropin-Independent Precocious Puberty: A Multicenter Study**
Atay Z., Yesilkaya E., Erdeve S. S. , DEMİRCİOĞLU S., AKIN L., EREN E., DÖĞER E., Aycan Z., Abali Z. Y. , AKINCI A., et al.
JOURNAL OF CLINICAL ENDOCRINOLOGY & METABOLISM, vol.101, no.5, pp.1980-1988, 2016 (Journal Indexed in SCI)
- XXXII. **Anthropometric findings from birth to adulthood and their relation with karyotype distribution in Turkish girls with Turner syndrome.**
Sari E., BEREKET A., Yesilkaya E., Bas F., Bundak R., Aydin B. K. , DARCAN Ş., Dundar B., Buyukinan M., Kara C., et al.
American journal of medical genetics. Part A, no.4, pp.942-8, 2016 (Journal Indexed in SCI Expanded)
- XXXIII. **Mucormycosis and Type 1 Diabetes: A Case Report**
EREN E., ÇELEBİ S., Sali E., Topac T., Tarim Ö. F.
HORMONE RESEARCH IN PAEDIATRICALS, vol.86, pp.234, 2016 (Journal Indexed in SCI)
- XXXIV. **Retrospective evaluation of pubertal development and linear growth of girls with Turner Syndrome treated with oral and transdermal estrogen**
Cakir E. D. P. , SAĞLAM H., EREN E., Ozgur T., TARIM Ö. F.
JOURNAL OF PEDIATRIC ENDOCRINOLOGY & METABOLISM, vol.28, pp.1219-1226, 2015 (Journal Indexed in SCI)
- XXXV. **Pediatric patients with pheochromocytoma: Experience of a tertiary health center.**
EREN E., SAĞLAM H., Caliskan Y., KIRIŞTIOĞLU İ., Tarim Ö. F.
Pediatrics international : official journal of the Japan Pediatric Society, vol.57, no.5, pp.875-9, 2015 (Journal Indexed in SCI Expanded)
- XXXVI. **Growth curves for Turkish Girls with Turner Syndrome: Results of the Turkish Turner Syndrome Study Group.**
Darendeliler F., Yesilkaya E., BEREKET A., Bas F., Bundak R., Sari E., Aydin B. K. , DARCAN Ş., Dundar B., Buyukinan M., et al.
Journal of clinical research in pediatric endocrinology, vol.7, no.3, pp.183-91, 2015 (Journal Indexed in SCI Expanded)

- XXXVII. **Turner syndrome and associated problems in Turkish children: a multicenter study.**
Yesilkaya E., BEREKET A., Darendeliler F., Bas F., Poyrazoglu S., Aydin B. K. , DARCAN Ş., Dundar B., Buyukinan M., Kara C., et al.
Journal of clinical research in pediatric endocrinology, vol.7, no.1, pp.27-36, 2015 (Journal Indexed in SCI Expanded)
- XXXVIII. **GPR30 Gene Polymorphisms Are Associated with Gynecomastia Risk in Adolescents**
Korkmaz H. A. , Edgunlu T., EREN E., Demir K., Cakir E. D. P. , Celik S. K. , Ozkan B.
HORMONE RESEARCH IN PAEDIATRICS, vol.83, no.3, pp.177-182, 2015 (Journal Indexed in SCI)
- XXXIX. **Transnasal Transsphenoidal Surgical Method in Pediatric Pituitary Adenomas.**
TAŞKAPILIOĞLU M. Ö. , YILMAZLAR S., EREN E., Tarim Ö. F. , Guler T. M.
Pediatric neurosurgery, vol.50, no.3, pp.128-32, 2015 (Journal Indexed in SCI Expanded)
- XL. **Assessment of Growth and Development in Children With Hepatitis B Positivity.**
Sari T., Eren E., Koruk S. T.
Gastroenterology research, vol.7, pp.131-136, 2014 (Journal Indexed in SCI Expanded)
- XLI. **Loss-of-Function Mutations in PNPLA6 Encoding Neuropathy Target Esterase Underlie Pubertal Failure and Neurological Deficits in Gordon Holmes Syndrome**
TOPALOĞLU A. K. , Lomniczi A., Kretschmar D., Dissen G. A. , KOTAN L. D. , McArdle C. A. , KOÇ A. F. , Hamel B. C. , Guclu M., Papatya E. D. , et al.
JOURNAL OF CLINICAL ENDOCRINOLOGY & METABOLISM, vol.99, no.10, 2014 (Journal Indexed in SCI)
- XLII. **An essential splice site mutation (c.317+1G>A) in the TSHR gene leads to severe thyroid dysgenesis.**
Cangul H., SAĞLAM H., Saglam Y., EREN E., Dogan D., Kendall M., Tarim Ö. F. , Maher E. R. , Barrett T. G.
Journal of pediatric endocrinology & metabolism : JPEM, vol.27, pp.1021-5, 2014 (Journal Indexed in SCI Expanded)
- XLIII. **Etiological evaluation of adolescents with primary amenorrhea.**
EREN E., SAĞLAM H., Cakir E. D. P. , Tarim Ö. F.
Indian journal of pediatrics, vol.81, no.9, pp.861-5, 2014 (Journal Indexed in SCI Expanded)
- XLIV. **Serum Paraoxonase/Arylesterase Activity and Oxidative Stress Status in Children with Metabolic Syndrome**
EREN E., Abuhandan M., Solmaz A., Taskin A.
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- XLV. **Epicardial adiposity in children with obesity and metabolic syndrome.**
Eren E., Koca B., Ture M., Guzel B.
Iranian journal of pediatrics, vol.24, pp.411-7, 2014 (Journal Indexed in SCI)
- XLVI. **A deletion including exon 2 of the TSHR gene is associated with thyroid dysgenesis and severe congenital hypothyroidism.**
Cangul H., Schoenmakers N. A. , SAĞLAM H., Doganlar D., Saglam Y., EREN E., Kendall M., Tarim Ö. F. , Barrett T. G. , Chatterjee K., et al.
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- XLVII. **Loss of Function Mutations in PNPLA6 Encoding Neuropathy Target Esterase Cause Pubertal Failure and Cerebellar Ataxia (Gordon Holmes Syndrome)**
KOTAN L. D. , Lomniczi A., Kretschmar D., Dissen G. A. , McArdle C. A. , Koc F., Hamel B. C. , Guclu M., Papatya E. D. , EREN E., et al.
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- XLVIII. **Genetic variants of estrogen beta and leptin receptors may cause gynecomastia in adolescent**
EREN E., EDGÜNLÜ T., Korkmaz H. A. , Cakir E. D. P. , Demir K., ÇETİN E. S. , Celik S. K.
GENE, vol.541, no.2, pp.101-106, 2014 (Journal Indexed in SCI)
- XLIX. **Novel Fructose-1,6-bisphosphatase Gene Mutation in Two Siblings**
EREN E., Edgunlu T., Abuhandan M., Yetkin I.
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- L. **Evaluation of the endocrine functions in pediatric patients with cyanotic congenital heart disease**
EREN E., Cakir E. D. P., Bostan O., SAĞLAM H., Tarim Ö. F.
BIOMEDICAL RESEARCH-INDIA, vol.24, no.2, pp.211-215, 2013 (Journal Indexed in SCI)
- LI. **Metformin intoxication in an adolescent**
EREN E., Ture M., Almaz V., Bilgic T., Yildirim I., Gider T.
TURK PEDIATRI ARSIVI-TURKISH ARCHIVES OF PEDIATRICS, vol.47, no.4, pp.313-314, 2012 (Journal Indexed in SCI)
- LII. **Association of infantile osteopetrosis and rickets: A case report İnfantil Osteopetrozis ve Rikets Birlikteliği: Olgu Sunumu**
Sancak Y., Çakir D., Deniz Aygün F., EREN E., TARIM Ö. F., ÇELEBİ S.
Guncel Pediatri, vol.10, no.1, pp.40-43, 2012 (Journal Indexed in SCI Expanded)
- LIII. **Testicular adrenal rest tumors in patients with congenital adrenal hyperplasia.**
Cakir E. D. P., Mutlu F. S., EREN E., Pasa A. O., SAĞLAM H., Tarim Ö. F.
Journal of clinical research in pediatric endocrinology, vol.4, no.2, pp.94-100, 2012 (Journal Indexed in SCI Expanded)
- LIV. **TSHR is the main causative locus in autosomal recessively inherited thyroid dysgenesis.**
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- LV. **Resolution of Autoimmune Oophoritis after Thymectomy in a Myasthenia Gravis Patient**
Cakir E. D. P., Ozdemir O., EREN E., SAĞLAM H., Okan M., TARIM Ö. F.
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- LVI. **Evaluation of the patients with congenital hypothyroidism: Effect of the national screening program Konjenital Hipotiroidili Olguların Değerlendirilmesi: Ulusal Tarama Programının Etkisi**
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- LVII. **Clinical course of hyperprolactinemia in children and adolescents: a review of 21 cases.**
EREN E., Yapici S., Cakir E. D. P., Ceylan L. A., SAĞLAM H., Tarim Ö. F.
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- LVIII. **Efficacy of biphosphonates in patients with osteogenesis imperfecta Osteogenesis imperfektalı hastalarda bifosfonatların etkinliği**
EREN E., Sincar Ş., Cakir E. D. P., SAĞLAM H., Tarim Ö.
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- LIX. **A retrospective evaluation of patients with rickets Riketsli olguların retrospektif değerlendirilmesi**
Ocak M., EREN E., Istanbul K., SAĞLAM H.
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- LX. **The efficacy of alendronate in children with secondary osteoporosis Sekonder osteoporozlu çocuklarda alendronat tedavisinin etkinliği**
SAĞLAM H., EREN E., Çakir E. D. P., Özboyacı E., Özboyacı A., ÖZGÜR T., Tarim Ö.
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- LXI. **Novel TSHR mutations in consanguineous families with congenital nongoitrous hypothyroidism.**
Cangul H., Morgan N. V., Forman J. R., SAĞLAM H., Ayca Z., Yakut T., Gulden T., Tarim Ö. F., BÖBER E., Cesur Y., et al.
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- LXII. **Evaluation of congenital heart diseases and thyroid abnormalities in children with Down syndrome**
MIHÇI E., Akcurin G., EREN E., Kardelen F., Akcurin S., KESER İ., Ertug H.
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- LXIII. **Amniotic band syndrome associated with sacral meningocele and anal atresia**

- EREN E., Buyukyavuz I., Ayata A.
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- LXIV. **Robinow syndrome Robinow sendromu**
GÖKALP G., EREN E., YAZICI Z., SAĞLAM H.
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- LXV. **Factors influencing glyceic control in children with type 1 diabetes Tip 1 diyabetli çocuklarda glisemik kontrolü etkileyen faktörler**
Çakir S., SAĞLAM H., ÖZGÜR T., EREN E., Tarım Ö.
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- LXVI. **Acquired generalized lipodystrophy associated with autoimmune hepatitis and low serum C4 level.**
Eren E., Özkan T. B. , Çakır E. D. P. , Sağlam H., Tarım Ö. F.
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- LXVII. **A Rare Cause of Precocious Puberty Hepatoblastoma**
Eren E., Demirkaya M., Esra Deniz Ç., Sevinir B. B. , Sağlam H., Tarım Ö. F.
JOURNAL OF CLINICAL RESEARCH IN PEDIATRIC ENDOCRINOLOGY, vol.1, no.6, pp.281-283, 2009 (Journal Indexed in SCI Expanded)
- LXVIII. **Changes in hematologic parameters after second and third doses of measles immunization in children İkinci ve üçüncü kez kızamık aşisi yapılan çocuklarda hematolojik parametrelerin değişimleri**
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