

Prof. SERAP DEMİRCİOĞLU

Personal Information

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Biography

Marmara Üniversitesi Çocuk Sağlığı ve Hastalıkları Bilim Dalında pediatri uzmanlık eğitimimi tamamladıktan sonra, aynı üniversitede Çocuk Endokrinoloji yan dal eğitimi aldım. Yan dal eğitimi sırasında uluslararası pek çok eğitim faaliyetine bilimsel toplantı ve bilim okulları olarak katılarak sunumlar yaptım. Yan dal eğitimimi tamamladıktan sonra klinik eğitimimi temel bilimler ile birleştirmek istedim ve de TÜBİTAK ve Fulbright'tan aldığım burslar ile Amerika Birleşik Devletleri Massachusetts General Hospital-Harvard Medical School'da hipofosfatemik rikets, psödohipoparatiroidi ve metilasyon bozuklukları alanında moleküler genetik çalışmalar yaptım. Türkiye'ye döndükten sonra bu konuda çalışmalarım ve farklı merkezler ile işbirliğim devam etti. 2010 yılında European Society for Paediatric Endocrinology (ESPE) 'Sabbatical Leave Programme' desteği ile 1 yıl süre yine aynı merkezde 'visiting scientist' olarak psödohipoparatiroidi (PHP) etiyo-patogenezi konusunda GNAS knock-out fare modellerinde çalışmalar yaptım. Halen, ESPE bünyesinde kurulan PHP grubunun aktif bir üyesi olarak Avrupa'da yapılan çalışmalara ve konsensus toplantılarına katılmaktayım. Avrupa Birliği Projesi olan COST Action BM10208- European Network for Human Congenital Imprinting Disorders çalışma grubunun aksiyon ortağı olarak Türkiye'yi temsil etmekteyim.

European Society for Paediatric Endocrinology (ESPE) tarafından Annual ESPE Meetingleri'nin program organizasyon komitesinde 4 yıl boyunca görev yapmak üzere 2010 yılında seçildim ve bu görevi 2014 yılında tamamladım. ESPE'nin bilim okulu olan Winter School'da 'Teaching Faculty' olarak 2015 yılında göreve başladım. European Society for Paediatric Endocrinology 1962 yılında Avrupa Pediatrik Endokrinoloji camiası tarafından kurulmuş olup, görev aldığım her iki pozisyon için de görev verilen ilk Türk bilim insanı olmam önem arz etmektedir.

İlk tip 1 diyabet prevalans çalışmasının ülkemizde okul çalışması kapsamında yapılmasında, sorumlu araştırmacı olarak görev aldım ve bu çalışmanın sonuçlarına göre okulda diyabet programının oluşturulmasında çalıştım. Diyabet 2020 projesi ve Sağlık Bakanlığı Diyabet programında raportör olarak görev aldım.

Journal of Pediatric Endocrinology and Metabolism ve Journal of Clinical Research in Pediatric Endocrinology dergilerinde yardımcı editörlük yapmanın yanı sıra pek çok dergi için yayın değerlendirmesi yapmaktayım. 2008'den bu yana Faculty of 1000 Prime'de Associate Faculty Member olarak yayın değerlendirmesi yapıyorum.

Ayrıca, Avrupa Birliği Çerçeve Programı 'imprinting Disorders' EURCOST çerçevesinde proje yürüttüm.

Education Information

Undergraduate Minor, Marmara University, School of Medicine, Internal Medical Sciences, Turkey 2001 - 2004

Under Graduate, Marmara University, School of Medicine, Internal Medical Sciences, Turkey 1995 - 2000

Associate Degree, Hacettepe University, Tıp Fakültesi (İngilizce), Turkey 1988 - 1995

Foreign Languages

English, A1 Beginner

Research Areas

Health Sciences

Academic Titles / Tasks

Professor, Marmara University, School of Medicine, Internal Medical Sciences, 2015 - Continues

Professor, Marmara University, School of Medicine, Internal Medical Sciences, 2015 - Continues

Professor, Marmara University, School of Medicine, 2014 - Continues

Associate Professor, Marmara University, School of Medicine, Internal Medical Sciences, 2009 - Continues

Advising Theses

Demircioğlu Turan S., Obez çocuklarda bazal metabolik hız ve subklinik hipotiroidinin bazal metabolik hıza etkisi, Expertise In Medicine, N.MUZAFFAROVA(Student), 2016

Demircioğlu Turan S., Biyokimya laboratuvarında alkalen fosfataz düşüklüğü tespit edilen hastaların hipofosfatazya açısından incelenmesi, Expertise In Medicine, R.BAYRAMLI(Student), 2016

Demircioğlu Turan S., Boy kısalığı olan hastalarda özellikler ve etiyolojik dağılım, Expertise In Medicine, S.ÖZCAN(Student), 2015

Demircioğlu S., Fibroblast büyüme faktörü-23 (FGF-23) ve matriks ekstraselüler fosfoglikoprotein (MEPE) sağlıklı bireylerde yaşa ve cinse özgü referans değerlerinin belirlenerek fosfat regülasyonundaki rollerinin araştırılması, Expertise In Medicine, A.ÖZŞEN(Student), 2012

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

- I. **Persistent Mullerian Duct Syndrome: A Rare But Important Etiology of Inguinal Hernia and Cryptorchidism**
Bugrul F., Abali Z. Y., KIRKGÖZ T., KARADENİZ CERİT K., CANMEMİŞ A., DEMİRCİOĞLU S., Tugtepe H., Picard J., BEREKET A., GÜRAN T.
SEXUAL DEVELOPMENT, vol.13, pp.264-270, 2020 (Journal Indexed in SCI)
- II. **Recommendations for Diagnosis and Treatment of Pseudohypoparathyroidism and Related Disorders: An Updated Practical Tool for Physicians and Patients**
Mantovani G., Bastepe M., Monk D., De Sanctis L., Thiele S., Ahmed S. F., Bufo R., Choplin T., De Filippo G., Devernois G., et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.93, pp.182-196, 2020 (Journal Indexed in SCI)
- III. **Management of Systemic Hypersensitivity Reactions to Gonadotropin-Releasing Hormone Analogues during Treatment of Central Precocious Puberty**
Kirkgoz T., Karakoc-Aydiner E., Bugrul F., Yavas Abali Z., Helvacioğlu D., Kiykim A., Bilgic Eltan S., Aruci Kasap N., Baris S., Ozen A. O., et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.93, pp.66-72, 2020 (Journal Indexed in SCI)
- IV. **A Rare Cause of Hypophosphatemia: Raine Syndrome Changing Clinical Features with Age**
ELTAN M., ALAVANDA C., Yavas Abali Z., Ergenekon P., Yalindag Ozturk N., SAKAR M., DAĞÇINAR A., KIRKGÖZ T., KAYGUSUZ S. B., GÖKDEMİR Y., et al.
CALCIFIED TISSUE INTERNATIONAL, vol.107, pp.96-103, 2020 (Journal Indexed in SCI)
- V. **Disease characteristics of MCT8 deficiency: an international, retrospective, multicentre cohort study**
Groeneweg S., van Geest F. S., ABACI A., Alcantud A., Ambegaonkar G. P., Armour C. M., Bakhtiani P., Barca D., Bertini E. S., van Beynum I. M., et al.
LANCET DIABETES & ENDOCRINOLOGY, vol.8, pp.594-605, 2020 (Journal Indexed in SCI)

- VI. **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, pp.150-159, 2020 (Journal Indexed in SCI)
- VII. **Clinical Significance of Hypophosphatasemia in Children**
Bayramli R., Cevlik T., GÜRAN T., Atay Z., Bas S., HAKLAR G., BEREKET A., DEMİRCİOĞLU S.
CALCIFIED TISSUE INTERNATIONAL, vol.106, pp.608-615, 2020 (Journal Indexed in SCI)
- VIII. **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**
Abali S., Tamura M., DEMİRCİOĞLU S., Atay Z., Isguven P., GÜRAN T., Haliloglu B., Bas S., Isojima T., Kitanaka S., et al.
JOURNAL OF PEDIATRIC ENDOCRINOLOGY & METABOLISM, vol.33, pp.557-562, 2020 (Journal Indexed in SCI)
- IX. **Revisiting Classical 3 beta-hydroxysteroid Dehydrogenase 2 Deficiency: Lessons from 31 Pediatric Cases**
GÜRAN T., Kara C., Yildiz M., Bitkin E. C. , HAKLAR G., Lin J., KESKİN M., Barnard L., ANIK A., ÇATLI G., et al.
JOURNAL OF CLINICAL ENDOCRINOLOGY & METABOLISM, vol.105, 2020 (Journal Indexed in SCI)
- X. **Fibroblast Growth Factor-23 and Matrix Extracellular Phosphoglycoprotein Levels in Healthy Children and, Pregnant and Puerperal Women**
Ozsen A., FURMAN A., GÜRAN T., BEREKET A., DEMİRCİOĞLU S.
HORMONE RESEARCH IN PAEDIATRICS, vol.92, pp.302-310, 2020 (Journal Indexed in SCI)
- XI. **Rare cause of severe hypertension in an adolescent boy presenting with short stature: Questions**
Yavas A., Yesil G., Kirkgoz T., Cicek N., Alpay H., Turan S., Bereket A., Guran T.
PEDIATRIC NEPHROLOGY, vol.35, pp.403-404, 2020 (Journal Indexed in SCI)
- XII. **Rapid progression of type 2 diabetes and related complications in children and young people-A literature review**
Barrett T., Jalaludin M. Y. , DEMİRCİOĞLU S., Hafez M., Shehadeh N.
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- XIII. **Rare cause of severe hypertension in an adolescent boy presenting with short stature: Answers**
Yavas A., Yesil G., Kirkgoz T., Cicek N., Alpay H., Turan S., Bereket A., Guran T.
PEDIATRIC NEPHROLOGY, vol.35, pp.405-407, 2020 (Journal Indexed in SCI)
- XIV. **Low DHEAS Concentration in a Girl Presenting with Short Stature and Premature Pubarche: A Novel PAPSS2 Gene Mutation**
ELTAN M., Yavas Abali Z., Arslan Ates E., KIRKGÖZ T., KAYGUSUZ S. B. , Türkyllmaz A., BEREKET A., Turan S., GÜRAN T.
Hormone Research in Paediatrics, vol.92, pp.262-268, 2020 (Journal Indexed in SCI)
- XV. **Restoration of Height after 11 Years of Letrozole Treatment in 11 β -Hydroxylase Deficiency**
Atay Z., Turan S., Buğdaycı O., GÜRAN T., BEREKET A.
Hormone Research in Paediatrics, vol.92, pp.203-208, 2020 (Journal Indexed in SCI)
- XVI. **Recommendations for improving clinical trial design to facilitate the study of youth-onset type 2 diabetes**
Jalaludin M. Y. , Barrientos-Pérez M., Hafez M., Lynch J., Shehadeh N., Turan S., Weghuber D.
Clinical Trials, vol.17, pp.87-98, 2020 (Journal Indexed in SCI)
- XVII. **A rare cause of hypertension in childhood: Answers**
Kucuk N., Yavas Abali Z., ABALI S., Canpolat N., YEŞİL G., Turan S., BEREKET A., GÜRAN T.
Pediatric Nephrology, vol.35, pp.79-82, 2020 (Journal Indexed in SCI)
- XVIII. **A rare cause of hypertension in childhood: Questions**
Kucuk N., Yavas A., Abali S., Canpolat N., Yesil G., Turan S., Bereket A., Guran T.
PEDIATRIC NEPHROLOGY, vol.35, pp.77-78, 2020 (Journal Indexed in SCI)
- XIX. **Adrenocortical carcinoma in atypical Beckwith-Wiedemann syndrome due to loss of methylation at imprinting control region 2**
Eltan M., Arslan A., Cerit K., Menevse T., Kaygusuz S., Eker N., Bagci P., Ergelen R., Turan S., Bereket A., et al.

PEDIATRIC BLOOD & CANCER, vol.67, 2020 (Journal Indexed in SCI)

- XX. **Cinacalcet as a First-Line Treatment in Neonatal Severe Hyperparathyroidism Secondary to Calcium Sensing Receptor (CaSR) Mutation**
Gulcan-Kersin S, Kirkgoz T, ELTAN M, Rzayev T, ATA P, Bilgen H, ÖZEK E, BEREKET A, Turan S.
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- XXI. **Letter to the Editor: Dysosteosclerosis related to the unique mutation in SLC29A3**
Turan S.
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- XXII. **Characteristics of Turkish children with Type 2 diabetes at onset: a multicentre, cross-sectional study.**
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- XXIII. **Simplifying the interpretation of steroid metabolome data by a machine-learning approach**
KIRKGÖZ T, Kilic S, Abali Z. Y. , Yaman A, KAYGUSUZ S. B. , ELTAN M, Turan S, HAKLAR G, Sagiroglu M. S. , BEREKET A, et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.91, pp.128, 2019 (Journal Indexed in SCI)
- XXIV. **A rare cause of hypophosphatemia: Raine Syndrome**
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HORMONE RESEARCH IN PAEDIATRICS, vol.91, pp.384, 2019 (Journal Indexed in SCI)
- XXV. **Characteristics of puberty, pubertal height gain and final height in children with classical 21 hydroxylase deficiency**
Abali Z. Y. , Yildiz M, Bas F, Onal H, ABALI S, Cilsaat G, Uyguner Z. O. , Turan S, Darendeliler F, BEREKET A, et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.91, pp.205-206, 2019 (Journal Indexed in SCI)
- XXVI. **Hypergonadotropic hypogonadism in 46, XX adolescents without gonadotoxic therapy: Clinical features and molecular etiologies**
Abali Z. Y. , Jolly A, GÜRAN T, Bayram Y, ABALI S, Bas S, Akdemir Z. C. , Posey J. E. , Helvacioğlu D, KIRKGÖZ T, et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.91, pp.265, 2019 (Journal Indexed in SCI)
- XXVII. **Evaluation of brain MRI lesions in 381 girls with central precocious puberty**
Helvacioğlu D, GÜRAN T, KIRKGÖZ T, Atay Z, Abali Z. Y. , ELTAN M, KAYGUSUZ S. B. , Seven T, Gurpinar B, Turan S, et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.91, pp.329-330, 2019 (Journal Indexed in SCI)
- XXVIII. **Age of obesity onset could be the first indicator of future metabolic complications - preliminary data of prospective multicenter study**
Gawlik A, Wasniewska M, BEREKET A, Antosz A, Aversa T, Corica D, KIRKGÖZ T, Turan S, GÜRAN T, Shmoish M, et al.
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- XXIX. **Cushing Syndrome due to an adrenocortical carcinoma in a baby with atypical Beckwith-Wiedemann Syndrome**
ELTAN M, Cerit K, KAYGUSUZ S. B. , Ates E, EKER N, Bagci P, ERGELEN R, Turan S, BEREKET A, GÜRAN T.
HORMONE RESEARCH IN PAEDIATRICS, vol.91, pp.371, 2019 (Journal Indexed in SCI)
- XXX. **Genotype and Phenotype Characterization of Turkish Patients with Vitamin D Dependent Rickets Type IA**
KAYGUSUZ S. B. , ATA P, KIRKGÖZ T, Abali Z. Y. , ELTAN M, Tosun B. G. , Menevse T. S. , Helvacioğlu D., GÜRAN T, ARMAN A, et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.91, pp.215, 2019 (Journal Indexed in SCI)
- XXXI. **How to Approach Systemic Hypersensitivity reactions to Gonadotropin Releasing Hormone Analogues during treatment of Central Precocious Puberty**
KIRKGÖZ T, Karakoc E. A. , Kiykim A, Bugrul F, Helvacioğlu D, Eltan S. B. , Kasap N, Ozen A. O. , BARIŞ S, GÜRAN T,

et al.

HORMONE RESEARCH IN PAEDIATRICS, vol.91, pp.185, 2019 (Journal Indexed in SCI)

- XXXII. **Evaluation of molecular characteristics and steroid metabolomics in a large cohort of children with 3 beta-hydroxysteroid dehydrogenase 2 deficiency**
GÜRAN T., Kara C., Yildiz M., Bitkin E. C., HAKLAR G., Lin J., Gilligan L. C., Barnard L., Keskin M., Anik A., et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.91, pp.207, 2019 (Journal Indexed in SCI)
- XXXIII. **Rare Causes of Osteogenesis Imperfecta are Common in Consanguineous Pedigrees**
KAYGUSUZ S. B., ARMAN A., ABALI S., ATA P., KIRKGÖZ T., ELTAN M., Abali Z. Y., Helvacioğlu D., Tosun B. G., Menevse T. S., et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.91, pp.214, 2019 (Journal Indexed in SCI)
- XXXIV. **A Case Of Syndromic Hypopituitarism**
KAYGUSUZ S. B., Ates E. A., KIRKGÖZ T., ELTAN M., Abali Z. Y., Helvacioğlu D., Menevse T. S., Tosun B. G., Tutar E., Volkan B., et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.91, pp.589, 2019 (Journal Indexed in SCI)
- XXXV. **A Real-Life Experience with A New Insulin Co-Formulation Degludec/Aspart For One Year In Poorly Controlled Children And Adolescents With Type 1 Diabetes**
KIRKGÖZ T., ELTAN M., KAYGUSUZ S. B., Abali Z. Y., GÜRAN T., BEREKET A., Turan S.
HORMONE RESEARCH IN PAEDIATRICS, vol.91, pp.534-535, 2019 (Journal Indexed in SCI)
- XXXVI. **Liraglutide in children and adolescents with type 2 diabetes**
Tamborlane W. V., Barrientos-Pérez M., Fainberg U., Frimer-Larsen H., Hafez M., Hale P. M., Jalaludin M. Y., Kovarenko M., Libman I., Lynch J. L., et al.
New England Journal of Medicine, vol.381, pp.637-646, 2019 (Journal Indexed in SCI)
- XXXVII. **Exome Sequencing of a Primary Ovarian Insufficiency Cohort Reveals Common Molecular Etiologies for a Spectrum of Disease**
Jolly A., Bayram Y., Turan S., Aycan Z., Tos T., Abali Z. Y., Hacıhamdioğlu B., Akdemir Z. H. C., Hijazi H., Bas S., et al.
Journal of Clinical Endocrinology and Metabolism, vol.104, pp.3049-3067, 2019 (Journal Indexed in SCI)
- XXXVIII. **Ptois as a unique hallmark for autosomal recessive WNT1-associated osteogenesis imperfecta**
Nampoothiri S., Guillemin B., Elcioglu N., Jagadeesh S., Yesodharan D., Suresh B., Turan S., Symoens S., Malfait F.
American Journal of Medical Genetics, Part A, vol.179, pp.908-914, 2019 (Journal Indexed in SCI)
- XXXIX. **Evaluation of growth and puberty in a child with a novel TBX19 gene mutation and review of the literature**
Abali Z. Y., YEŞİL G., KIRKGÖZ T., KAYGUSUZ S. B., ELTAN M., Turan S., BEREKET A., GÜRAN T.
HORMONES-INTERNATIONAL JOURNAL OF ENDOCRINOLOGY AND METABOLISM, vol.18, pp.229-236, 2019 (Journal Indexed in SCI)
- XL. **Clinical and Laboratory Characteristics of Hyperprolactinemia in Children and Adolescents: National Survey**
Eren E., Törel Ergür A., Pınar İşgüven Ş., Çelebi Bitkin E., Berberoğlu M., Şiklar Z., Baş F., Yel S., Baş S., Söbü E., et al.
Journal of clinical research in pediatric endocrinology, vol.11, pp.149-156, 2019 (Journal Indexed in SCI Expanded)
- XLI. **PPP2R3C gene variants cause syndromic 46,XY gonadal dysgenesis and impaired spermatogenesis in humans**
GÜRAN T., Yesil G., Turan S., Atay Z., BOZKURTLAR E., Aghayev A., Gul S., TİNAY İ., Aru B., Arslan S., et al.
EUROPEAN JOURNAL OF ENDOCRINOLOGY, vol.180, pp.291-309, 2019 (Journal Indexed in SCI)
- XLII. **Incidence of Type 1 Diabetes in Children Aged Below 18 Years during 2013-2015 in Northwest Turkey**
Poyrazoğlu S., Bundak R., Abali Z. Y., Onal H., Sarıkaya S., Akgun A., Bas S., Abali S., BEREKET A., EREN E., et al.
JOURNAL OF CLINICAL RESEARCH IN PEDIATRIC ENDOCRINOLOGY, vol.10, pp.336-342, 2018 (Journal Indexed in SCI)
- XLIII. **THE CO-EXISTENCE OF TWO RARE DISEASES: A CASE REPORT**
Saki M., KIRKGÖZ T., GÖKCE İ., Cicek N., ATA P., Turan S., YILDIZ N., ALPAY H.
PEDIATRIC NEPHROLOGY, vol.33, pp.1881, 2018 (Journal Indexed in SCI)

- XLIV. Diagnosis and management of pseudohypoparathyroidism and related disorders: First international Consensus Statement**
Mantovani G., Bastepe M., Monk D., De Sanctis L., Thiele S., Usardi A., Ahmed S. F. , Bufo R., Choplin T., De Filippo G., et al.
Nature Reviews Endocrinology, vol.14, pp.476-500, 2018 (Journal Indexed in SCI)
- XLV. Biallelic and monoallelic ESR2 variants associated with 46,XY disorders of sex development**
Baetens D., Güran T., Mendonca B. B. , Gomes N. L. , De Cauwer L., Peelman F., Verdin H., Vuylsteke M., Van Der Linden M., Atay Z., et al.
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- XLVI. Mutations in the mitochondrial ribosomal protein MRPS22 lead to primary ovarian insufficiency**
Chen A., Tiosano D., Guran T., Baris H. N. , Bayram Y., Mory A., Shapiro-Kulnane L., Hodges C. A. , Akdemir Z. C. , Turan S., et al.
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- XLVII. The distribution of different types of diabetes in childhood: A single center experience**
Haliloğlu B., ABALI S., Buğrul F., Çelik E., Baş S., Atay Z., GÜRAN T., Turan S., BEREKET A.
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- XLVIII. Acquired modification of sphingosine-1-phosphate lyase activity is not related to adrenal insufficiency**
SÜNTER G., Enver E. O. , Akbarzade A., Turan S., Vatansever P., GÜNAL D., HAKLAR G., BEREKET A., AĞAN YILDIRIM K., GÜRAN T.
BMC NEUROLOGY, vol.18, 2018 (Journal Indexed in SCI)
- XLIX. An 18 Month Old Boy with Hypoglycemic Convulsion and Obesity Due to POMC Deficiency**
KAYGUSUZ S. B. , Yesil G., KIRKGÖZ T., Turan S., BEREKET A., GÜRAN T.
HORMONE RESEARCH IN PAEDIATRICS, vol.90, pp.514, 2018 (Journal Indexed in SCI)
- L. An Unusual Cause of Short Stature**
KAYGUSUZ S. B. , Atay Z., KIRKGÖZ T., GÜRAN T., BEREKET A., Turan S.
HORMONE RESEARCH IN PAEDIATRICS, vol.90, pp.184-185, 2018 (Journal Indexed in SCI)
- LI. Nationwide Hypophosphatemic Rickets Study**
ŞIKLAR Z., Turan S., BEREKET A., ABACI A., Bas F., DEMİR K., GÜRAN T., Akberzade A., BÖBER E., Ozbek M. N. , et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.90, pp.109-110, 2018 (Journal Indexed in SCI)
- LII. Isolated Congenital Central Hypothyroidism due to a Novel Mutation in TSH Beta Subunit Gene**
KIRKGÖZ T., ÖZHAN B., Cetin O., KAYGUSUZ S. B. , Turan S., BEREKET A., GÜRAN T.
HORMONE RESEARCH IN PAEDIATRICS, vol.90, pp.597, 2018 (Journal Indexed in SCI)
- LIII. Recurrent Hypoglycemia in a Preschooler Girl with Overgrowth: Isolated ACTH-Deficiency with a Novel TPIT Mutation**
Abali Z. Y. , Yesil G., KIRKGÖZ T., KAYGUSUZ S. B. , Turan S., BEREKET A., GÜRAN T.
HORMONE RESEARCH IN PAEDIATRICS, vol.90, pp.132, 2018 (Journal Indexed in SCI)
- LIV. Persistent Mullerian duct syndrome: Rare But Important Aetiology of an Inguinal Hernia and Cryptorchidism in Boys**
BEREKET A., Bugrul F., KIRKGÖZ T., KARADENİZ CERİT K., CANMEMİŞ A., Turan S., Picard J., Tugtepe H., GÜRAN T.
HORMONE RESEARCH IN PAEDIATRICS, vol.90, pp.568, 2018 (Journal Indexed in SCI)
- LV. Diagnosis and Management of Pseudohypoparathyroidism and Related Disorders: First International Consensus Statement**
Mantovani G., Bastepe M., Monk D., de Sanctis L., Thiele S., Usardi A., Ahmed F., Bufo R., Choplin T., DeFillipo G., et al.
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- LVI. Delayed Diagnosis of a Patient with Antley-Bixler Syndrome**
KIRKGÖZ T., Bas S., Abali Z. Y. , Turan S., BEREKET A., GÜRAN T.
HORMONE RESEARCH IN PAEDIATRICS, vol.90, pp.166-167, 2018 (Journal Indexed in SCI)
- LVII. Current Nomenclature of Pseudohypoparathyroidism: Inactivating Parathyroid Hormone/Parathyroid Hormone-Related Protein Signaling Disorder**
Turan S.

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- XVI. **Çocuklukta Osteogenesis İmperfekta Klinik Yaklaşım ve Tedavide İlkeler :**
DEMİRCİOĞLU S.

3.Ege Endokrin Hastalıklar ve Genetik Sempozyumu, Turkey, 7 - 09 March 2019

- XVII. Disorders of adrenocortical excess and adrenal medulla**
DEMİRCİOĞLU S.
ESPE Winter School, 22 - 28 February 2019
- XVIII. Congenital Adrenal Hyperplasia**
DEMİRCİOĞLU S.
ESPE Winter School, 22 - 28 February 2019
- XIX. Disorders of sexual differentiation**
DEMİRCİOĞLU S.
Disorders of sexual differentiation, 22 - 28 February 2019
- XX. Introduction to molecular endocrinology**
DEMİRCİOĞLU S.
ESPE Winter School, 22 - 28 February 2019
- XXI. Metabolic bone diseases in children, diagnosis related to oral and dental manifestations**
DEMİRCİOĞLU S.
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- XXII. Psödohipoparatiroidi tanı ve tedavisi**
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- XXIII. Düşük renin düzeyi: endokrin hipertansiyon**
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- XXVI. Nationwide Hypophosphatemic Rickets Study**
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cryptorchidism in boys**
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GURAN T.
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DEMİRCİOĞLU S.
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- XXXVIII. **Düşük Alkali Fosfotaz Düzeyinin Önemi**
DEMİRCİOĞLU S.
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- XLI. **CONGENITAL ADRENAL HYPERPLASIA**
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TURAN S.
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- XLIX. **Disorders of adrenocortical excess and adrenal medulla**
TURAN S.
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- L. **Congenital Adrenal Hyperplasia**
TURAN S.
European Society for Paediatric Endocrinology-Winter School, 10 - 16 February 2017
- LI. **Introduction to molecular endocrinology**
TURAN S.
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- LII. **b hCG from an Occult Source Causing Peripheral Precocious Puberty Identification of the Tumour 6 Years After Presentation**
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- LIII. **Neonatal Hypocalcemia**
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- LIV. **From Pseudohypoparathyroidism to Inactivating PTH PTHrP Signaling Disorder iPPSD a Novel Classification Proposed by the European EuroPHP Network**
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- LVI. **Frequency of Recessive Osteogenesis Imperfecta in a Turkish Cohort and Genetic Causes**
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- LVII. **A European Survey to Identify New Roads for Care Training and Research Around Rare Metabolic Bone Diseases**
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- LX. Reconsideration of Mid Parental Height Calculation**
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- LXI. The Effect of Subclinical Hypothyroidism SH and Treatment of SH with L T4 on Basal Metabolic Rate in Obese Children A Prospective Study**
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- LXII. beta hCG from an Occult Source Causing Peripheral Precocious Puberty Identification of the Tumour 6 Years After Presentation**
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- LXIII. Premature Puberche Hyperinsülinemia Hypothyroxenemia and Hyperintensities in Basal Ganglia All Caused by a Single Congenital Defect**
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- LXIV. Current Perspective on Pseudohypoparathyroidism New Classification**
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- LXV. Late Breaking news**
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- LXVII. Disoerder of Adrenocortical Excess and Adrenal Medulla**
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- LXVIII. CAH Congenital Adrenal Hyperplasia**
TURAN S.
ESPE Winter School, 18 - 24 March 2016
- LXIX. Disorders of SExual Differentiation**
TURAN S.
ESPE Winter School, 18 - 24 March 2016
- LXX. Introduction to Molecular Endocrinology**
TURAN S.
ESPE Winter School, 18 - 24 March 2016
- LXXI. HPP Genel Değerlendirme**
TURAN S.
HİPOFOSFATAZYA DANIŞMA KURULU TOPLANTISI, Turkey, 16 December 2015
- LXXII. HİPOTİROİDİ VE TİROTOKSİKOZUN YENİDOĞAN ÜZERİNE ETKİSİ**
TURAN S.
GEBELİK VE TİROİD SEMPOZYUMU, Turkey, 12 December 2015
- LXXIII. Diagnostic and prognostic value of serum urokinase plasminogen activation receptor supar**

- procalcitonin pct and C reactive protein CRP in children with SIRS sepsis and febrile neutropenia**
SİRİNOĞLU M., SOYSAL A., CİNEL İ. H. , KEPENEKLİ KADAYIFÇI E., KARAASLAN A., YALINDAĞ N., KOÇ A., TOKUÇ A. G. , YAMAN A., HAKLAR G., et al.
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- LXXIV. **Diagnostic and prognostic value of serum urokinase plasminogen activation receptor SUPAR procalcitonin PCT and C reactive protein CRP in children with SIRS sepsis and febrile neutropenia**
Şirinoğlu M., SOYSAL A., CİNEL İ. H. , KEPENEKLİ E., KARAASLAN A., ÖZTÜRK M. N. , KOÇ A., TOKUÇ A. G. , YAMAN A., HAKLAR G., et al.
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- LXXVI. **Diyabet ile yaşam**
TURAN S.
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- LXXVII. **Growth Hormone Deficiency and Treatment**
TURAN S.
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- LXXVIII. **Piknodisostozis Otorinolaringolojik bulgular**
BAĞLAM T., BİNNETOĞLU A., TOPUZ M. F. , TURAN S., SARI M.
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- LXXIX. **Nonklasik Konjenital Adrenal Hiperplazi Hastalarının Genotip Ve Fenotip Özellikleri**
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19. PEDIATRİK ENDOKRİNOLOJİ VE DİYABET KONGRESİ, Turkey, 22 - 24 October 2015
- LXXX. **Ulusal Düzeyde Noonan Sendromlu Olguların Klinik Özellikleri ve Büyüme İzlemlerinin Değerlendirilmesi**
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- LXXXI. **Periferik Puberte Prekokslu 129 Çocukta Etiyolojik Dağılım Ve Klinik Özellikler**
ATAY Z., YEŞİLKAYA E., SAVAŞ Ş., TURAN S., AKIN L., EREN E., DÖĞER E., AYCAN Z., YAVAŞ ABALI Z., AKINCI A., et al.
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- LXXXII. **Temel genetik kavramlar ve kalıtım şekilleri**
TURAN S.
1. Pediatrik Endokrinolojiye Giriş Kursu, Turkey, 21 - 22 October 2015
- LXXXIII. **Boy Kısaldığı Olan Hastalarda Özellikler ve Etiyolojik Dağılım Bir Çocuk Endokrinoloji Kliniği Verileri**
ÖZCAN S., ABALI S., ATAY Z., HALİLOĞLU B., BAŞ S., ÖZTÜRK G., AKÇAY T., GÜRAN T., TURAN S., BERKET A.
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- LXXXIV. **Merkezi Yenidogan Tarama Programi ile Tani Almis Konjenital Hipotiroidili Vakalarimizin İzlemi**
BAŞ S., ABALI S., ATAY Z., GURBANOV Z., HALİLOĞLU B., GÜRAN T., TURAN S., BERKET A.
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- LXXXV. **Serbest T4 Düşük TSH Hafif Yüksek Olgularda Santral Hipotiroidi Primer Hipotiroidi Ayrımını Nasıl Yapabiliriz**
TURAN S., Gurbanov Z., BAŞ S., ABALI S., ATAY Z.

19. Ulusal Pediatrik Endokrinoloji ve Diyabet Kongresi, Turkey, 22 - 24 October 2015
- LXXXVI. **Otozomal Resesif Osteogenezis İmpperfekta Populasyonumuzdaki Sıklığı Ve Genetik Nedenleri**
ABALI S., ARMAN A., ATAY Z., BAŞ S., GÜRAN T., GÖRMEZ Z., DEMİRCİ H., BERKET A., TURAN S.
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- LXXXVII. **YILLIK**
TURAN S.
19. Ulusal Pediatrik Endokrinoloji ve Diyabet Kongresi, Turkey, 22 - 24 October 2015
- LXXXVIII. **Aetiological Spectrum and Clinical Characteristics of 129 Children with Gonadotropin Independent Precocious Puberty A Nationwide Cohort Study**
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- LXXXIX. **Hereditary Vitamin D Resistant Rickets Report of Four Cases with Successful Use of Intermittent Intravenous Calcium Via Peripheral Route**
ABALI S., TAMURA M., ATAY Z., İŞGÜVEN Ş. P., GÜRAN T., HALİLOĞLU T., BAŞ S., ISOJIMA T., TURAN S., KITANAKA S., et al.
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- XC. **Friedreich s Ataxia Presenting with Diabetes Mellitus in an Adolescent**
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- XCI. **Factors Effecting Response to Growth Hormone Treatment in Children with Turner Syndrome**
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- XCII. **Primary Adrenal Insufficiency in Children without Congenital Adrenal Hyperplasia Molecular and Clinical Characterisation of a Nationwide Cohort**
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- XCIII. **Central or Primary Hypothyroidism How to Differentiate in Patients with Low T4 but Mildly Elevated TSH Levels**
TURAN S., GURBANOV Z., BAŞ S., ABALI S., ATAY Z., BERKET A.
54th Annual Meeting of the European Society for Paediatric Endocrinology (ESPE), 1 - 03 October 2015
- XCIV. **Evaluating First Year Response and Final Height to Growth Hormone Treatment in Growth Hormone Deficiency Based on Peak GH Levels on Testing**
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- XCV. **The Diagnostic Treatment and Follow Up Features of Childhood Thyroid Malignancies A Preliminary Report**
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Activities in Scientific Journals

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Congenital Adrenal Hyperplasia, Seminar, European Society for Paediatric Endocrinology Winter School, Azerbaijan, February 2019
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Disorders of sexual differentiation, Seminar, European Society for Paediatric Endocrinology Winter School, Azerbaijan, February 2019
Disorder of Adrenal excess and Adrenal Medulla, Seminar, European Society for Paediatric Endocrinology Winter School, Azerbaijan, February 2019