

## Prof. Dr. ABDULLAH BEREKET

### Kişisel Bilgiler

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### Uluslararası Araştırmacı ID'leri

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Publons / Web Of Science ResearcherID: V-3793-2018

ScopusID: 7004903391

Yoksis Araştırmacı ID: 9240

### Biyografi

Ankara Üniversitesi Tıp Fakültesi 1986 mezunudur. Çocuk Sağlığı ve Hastalıkları uzmanlığını 1991 yılında almıştır. 1991-1996 yılları arasında ABD New York State Üniversitesi (SUNY at Stony Brook) Medical School da Pediatrik Endokrinoloji üst ihtisası ve takiben research fellow olarak çalıştıktan sonra Türkiye'ye dönüp Hacettepe Üniversitesinde iki yıl öğretim üyeliği yapmıştır. 1997 de Doçent olan Dr Bereket, 1999 yılında Marmara Üniversitesine geçerek Pediatrik Endokrinoloji Bilim dal'ını kurmuş ve halen bilim dalı başkanlığını yürütmektedir. Türkiye Bilimler Akademisi tarafından 2001 yılında "Seçkin Genç Bilimci" ödülü ile ödüllendirilmiştir. 2003 yılında Profesör unvanı alan Dr Bereket 2004-2005 yıllarında Türkiye Pediatrik Endokrinoloji Derneğinin başkanlığını yapmıştır. ESPE (Avrupa Pediatrik Endokrinoloji Cemiyeti), Amerikan Diabetes Association ve, Endocrin Society üyesidir. 2010-2014 yılları arası ESPE Summer School Steering Committee üyesi olarak görev yapmıştır. Uluslararası üç dergide yardımcı editör bir dergide de board üyesi olarak görev yapmaktadır. Uluslararası dergilerde yayınlanmış 150 nin üzerinde makalesi bulunmaktadır.

### Eğitim Bilgileri

Tıpta Yandal Uzmanlık, State University of New York at Stony Brook, Pediatrik Endokrinoloji, Amerika Birleşik Devletleri  
1992 - 1996

Tıpta Uzmanlık, Diğer (Kurumlar,hastaneler Vb.), Türkiye 1987 - 1991

Lisans, Ankara Üniversitesi, Ankara Tıp Fakültesi, Türkiye 1980 - 1986

### Yabancı Diller

İngilizce, C2 Ustalık

### Yaptığı Tezler

Tıpta Yandal Uzmanlık, Tip-1 diyabetli çocuklarda IGF-I ve bağlayıcı proteinleri, State University of New York at Stony Brook, 1996

Tıpta Uzmanlık, Tekrarlayan idrar yolu enfeksiyonlu çocuklarda üriner ultrasonografinin yeri, Diğer

(Kurumlar,hastaneler Vb.), 1991

## Araştırma Alanları

Pediyatrik Endokrinoloji ve Metabolizma

## Akademik Unvanlar / Görevler

Prof. Dr., Marmara Üniversitesi, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2003 - Devam Ediyor

Doç. Dr., Marmara Üniversitesi, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 1999 - 2003

Dr. Öğr. Üyesi, Hacettepe Üniversitesi, Tıp Fakültesi (İngilizce), Dahili Tıp Bilimleri Bölümü, 1996 - 1999

Araştırma Görevlisi Dr., Academic medical center at State University of New York at Stony Brook, Health Sciences Center, Pediatric Endocrinology, 1992 - 1996

## Yönetilen Tezler

Bereket A., Santral puberte prekoks kız çocukların klinik ve laboratuvar özellikleri ve aylık ve üç aylık GnRH analog tedavisi alan hastaların enjeksiyon sonrası gonadotropin düzeylerinin karşılaştırılması, Tıpta Uzmanlık, N.TÜRKMEN(Öğrenci), 2023

BEREKET A., Türk toplumunda anne-baba boyunun çocuğun final boyuna etkisi ve mid-parental boy formülünün validasyonu, Tıpta Uzmanlık, İ.SİNAN(Öğrenci), 2016

BEREKET A., Puberte prekoks nedeniyle polikliniğimize başvuran kız çocukların klinik ve laboratuvar özellikleri, Tıpta Uzmanlık, Ü.ACAR(Öğrenci), 2016

Bereket A., Kistik fibroz hastalarında hiperglisemi ve diyabetin CGMS ile araştırılması ve optimal tedavi rejiminin saptanması , Tıpta Yandal Uzmanlık, B.Haliloğlu(Öğrenci), 2013

BEREKET A., Çocukluk çağındaki obezitenin demir eksikliği anemisine neden olmasının araştırılması, Tıpta Uzmanlık, D.HELVACIOĞLU(Öğrenci), 2012

Bereket A., İstanbul'da yaşayan 6-18 yaş arası kız çocuklarında ergenlik gelişim basamaklarının değerlendirilmesi, Tıpta Yandal Uzmanlık, Z.ATAY(Öğrenci), 2011

BEREKET A., İstanbul'da 6-18 yaş arası okul çocuklarında tip 1 diyabet prevalansı çalışması, Tıpta Uzmanlık, E.Akesen(Öğrenci), 2010

BEREKET A., İstanbulda yaşayan 6-18 yaş arası okul çocuklarında Tip 1 Diyabet Prevalansı, Tıpta Uzmanlık, E.AKESEN(Öğrenci), 2010

Bereket A., Hipotalamik obezite patogenezinde, leptin, solubl leptin reseptörü, rezistin ve insülin salgım dinamiklerinin rolü., Tıpta Yandal Uzmanlık, T.Güran(Öğrenci), 2009

Bereket A., Androjen duyarsızlığı ön tanılı 46,XY cinsel farklılaşma bozukluğu olan (Erkek Psödohermafroditizmi) olgularımızın klinik ve hormonal özellikleri ile androjen reseptör geni ve 5-alfa redüktaz geni mutasyon analizleri, Tıpta Yandal Uzmanlık, T.AKÇAY(Öğrenci), 2009

BEREKET A., Puberte prekoksda leuprolide asetat tedavisinin final boya etkisi, Tıpta Uzmanlık, E.BALANLI(Öğrenci), 2009

BEREKET A., Tiroid disgenezili çocukların aile bireylerinde hormonal ve ultrasonografik olarak tiroid disgenezisi taraması, Tıpta Uzmanlık, E.KARAKOÇ(Öğrenci), 2006

Bereket A., Sağlıklı Türk çocuklarında yaş ve puberteye göre insülin benzeri büyüme faktörü-I ve insülin benzeri büyüme faktör bağlayıcı protein-3 düzeyleri: sosyo-ekonomik durumun büyüme ve büyüme faktörleri üzerine etkisi, Tıpta Yandal Uzmanlık, S.Turan(Öğrenci), 2004

BEREKET A., Sağlıklı Türk çocuklarında yaş ve puberteye göre insülin benzeri büyüme faktörü-1 ve insülin benzeri büyüme faktör bağlayıcı protein-3 düzeyleri: sosyo-ekonomik durumun büyüme ve büyüme faktörleri üzerine etkisi, Tıpta Uzmanlık, S.TURAN(Öğrenci), 2004

BEREKET A., Ultrasonografik kemik ses hızı iletimi (SOS) ile kemik sağlığının değerlendirilmesi: sağlıklı Türk çocukları için referans değerler ve osteopenik çocuklarda dual enerji X-ışın karşılaştırma, Tıpta Uzmanlık, A.Omar(Öğrenci), 2004

BEREKET A., Ultrasonografik kemik ses hızı iletimi(SOS)ile kemik sağlığının değerlendirilmesi:Sağlıklı Türk çocukları için referans değerler ve osteopenik çocuklarda dual enerji x-ışın absorpsiyometri (DEXA)yöntemi ile karşılaştırma, Tıpta Uzmanlık, A.OMAR(Öğrenci), 2004

BEREKET A., Boy kısıklığı olan prepubertal çocuklarda çinko suplemantasyonunun büyüme hormonu salınımı, IGF-I, IGFBP-3, somatomedin jenerasyon testi ve kemik formasyon belirteçleri üzerine etkisi, Tıpta Uzmanlık, S.İMAMOĞLU(Öğrenci), 2001

## Jüri Üyelikleri

Doçentlik Sınavı, Doçentlik Sınavı, Marmara Üniversitesi, Nisan, 2020

## SCI, SSCI ve AHCI İndekslerine Giren Dergilerde Yayımlanan Makaleler

- I. **Evaluating breast ultrasonography as a complementary diagnostic method in girls with central precocious puberty**  
BIYIKLI E., Helvacioğlu D., BUĞDAYCI O., Tosun B. G., DEMİRCİOĞLU S., GÜRAN T., BEREKET A.  
Pediatric Radiology, cilt.54, sa.7, ss.1156-1167, 2024 (SCI-Expanded)
- II. **Development of external genitalia during mini-puberty: is it related to somatic growth or reproductive hormones?**  
Gacemer H. A., Tosun B. G., Helvacioğlu D., Yaman A., Abali Z. Y., HALİLOĞLU B., DEMİRCİOĞLU S., HAKLAR G., BEREKET A., GÜRAN T.  
European Journal of Pediatrics, cilt.183, sa.3, ss.1325-1332, 2024 (SCI-Expanded)
- III. **Venous Thrombosis in a Pseudohypoparathyroidism Patient with a Novel GNAS Frameshift Mutation and Complete Resolution of Vascular Calcifications with Acetazolamide Treatment.**  
Menevse T. S., Iwasaki Y., Abali Z. Y., Tosun B. G., Helvacioğlu D., DOĞRU Ö., BUĞDAYCI O., Cyr S. M., GÜRAN T., BEREKET A., et al.  
Hormone research in paediatrics, cilt.97, sa.4, ss.404-415, 2024 (SCI-Expanded)
- IV. **Challenges in the management of a 7 years old child with thyrotropin-secreting pituitary adenoma and the review of the literature**  
KIRKGÖZ T., Abali S., Seker A., GÜRPINAR TOSUN B., ELTAN M., Helvacioğlu D., HALİLOĞLU B., KAYGUSUZ S. B., Yavas Abali Z., SEVEN MENEVŞE T., et al.  
HORMONE RESEARCH IN PAEDIATRICS, cilt.96, sa.5, ss.527-537, 2023 (SCI-Expanded)
- V. **Associations between sleep characteristics and glycemic variability in youth with type 1 diabetes**  
İpar N., BORAN P., BARIŞ H. E., Us M. C., Aygün B., HALİLOĞLU B., Gökçe T., Can E., Eviz E., İnan N. G., et al.  
Sleep Medicine, cilt.109, ss.132-142, 2023 (SCI-Expanded)
- VI. **Decline in the Age of Menarche in Istanbul Schoolgirls Over the Last 12 Years**  
GÜRAN T., HELVACIOĞLU D., TOSUN B. G., ABALI Z. Y., Alır F., Arslan Y. t., Molla G., Şahin B., Sayar M. E., Atay Z., et al.  
Journal of Clinical Research in Pediatric Endocrinology, cilt.15, sa.2, ss.154-159, 2023 (SCI-Expanded)
- VII. **Molecular analysis of MKRN3 gene in Turkish girls with sporadic and familial idiopathic central**  
KIRKGÖZ T., KAYGUSUZ S. B., ALAVANDA C., Helvacioğlu D., Abali Z. Y., GÜRPINAR TOSUN B., ELTAN M., SEVEN MENEVŞE T., GÜRAN T., ARMAN A., et al.  
JOURNAL OF PEDIATRIC ENDOCRINOLOGY & METABOLISM, cilt.36, sa.4, ss.401-408, 2023 (SCI-Expanded)
- VIII. **The views of teenagers with obesity, their caregivers, and doctors: a plain language summary of the ACTION Teens global survey**  
Mooney V., Baur L. A., BEREKET A., Bin-Abbas B., Chen W., Fernandez-Aranda F., Nieto N. G., Sigüero J. P. L., Maffei C., Osorio C. K., et al.  
JOURNAL OF COMPARATIVE EFFECTIVENESS RESEARCH, cilt.12, sa.1, 2023 (SCI-Expanded)

- IX. **Misalignment among adolescents living with obesity, caregivers, and healthcare professionals: ACTION Teens global survey study**  
Halford J. C. G., BEREKET A., Bin-Abbas B., Chen W., Fernandez-Aranda F., Garibay Nieto N., Lopez Siguero J. P., Maffei C., Mooney V., Osorto C. K., et al.  
PEDIATRIC OBESITY, cilt.17, sa.11, 2022 (SCI-Expanded)
- X. **Etiological analysis of hypophosphatemia: A single-center experience**  
Eltan M., Alavanda C., Abali Z. Y., Bayramoglu E., Kaygusuz S. B., Helvacioğlu D., Tosun B. G., Menevse T. S., Ata P., Guran T., et al.  
HORMONE RESEARCH IN PAEDIATRICS, cilt.95, sa.SUPPL 2, ss.141-142, 2022 (SCI-Expanded)
- XI. **Differences due to the variant type in the inheritance pattern of BMP15 gene-related primary ovarian insufficiency: a girl with a homozygous null BMP15 gene variant**  
Abali Z. Y., Ates E. A., ELTAN M., GÜRPINAR TOSUN B., BEREKET A., GÜRAN T., DEMİRCİOĞLU S.  
HORMONE RESEARCH IN PAEDIATRICS, cilt.95, sa.SUPPL 2, ss.377-378, 2022 (SCI-Expanded)
- XII. **Basal cortisol measurements in the prediction of low-dose ACTH stimulation test outcomes**  
Gacemer H., Gürpınar Tosun B., Abali Z. Y., Helvacioğlu D., Haliloğlu B., Demircioğlu S., Bereket A., Güran T.  
HORMONE RESEARCH IN PAEDIATRICS, cilt.95, sa.SUPPL 2, ss.139, 2022 (SCI-Expanded)
- XIII. **Genotype, phenotype characteristics and long-term follow-up of patients with Vitamin D Dependent Rickets Type IA (VDDR1a): A nationwide multicentre retrospective cross-sectional study**  
Cayir A., DEMİRBİLEK H., TÜRKİYILMAZ A., DEMİRCİOĞLU S., BEREKET A., Darendeliler F., Ozbek M. N., Unal E., Okdemir D., Esen I., et al.  
HORMONE RESEARCH IN PAEDIATRICS, cilt.95, sa.SUPPL 2, ss.100, 2022 (SCI-Expanded)
- XIV. **Effect of unawareness of obesity on perceptions of obesity and food/diet among adolescents living with obesity and their caregivers: subanalysis from the ACTION Teens global study**  
BEREKET A., Baur L. A., Bin-Abbas B., Chen W., Fernandez-Aranda F., Garibay Nieto N., Halford J. C. G., Lopez Siguero J. P., Maffei C., Mooney V., et al.  
HORMONE RESEARCH IN PAEDIATRICS, cilt.95, sa.SUPPL 2, ss.211, 2022 (SCI-Expanded)
- XV. **Diagnostic Features and Risk Factors for Childhood Thyroid Cancers**  
ŞAHİN P., GÜRPINAR TOSUN B., YUMUŞAKHUYLU A. C., GÜRAN T., Helvacioğlu D., Abali Z. Y., HALİLOĞLU B., OYSU Ç., BEREKET A., DEMİRCİOĞLU S.  
HORMONE RESEARCH IN PAEDIATRICS, cilt.95, sa.SUPPL 2, ss.407, 2022 (SCI-Expanded)
- XVI. **Effect of gender and age on the mental well-being and self-esteem of adolescents living with obesity: subanalysis from the ACTION Teens global study**  
Lopez Siguero J. P., Baur L. A., BEREKET A., Bin-Abbas B., Chen W., Fernandez-Aranda F., Garibay Nieto N., Halford J. C. G., Maffei C., Mooney V., et al.  
HORMONE RESEARCH IN PAEDIATRICS, cilt.95, sa.SUPPL 2, ss.210, 2022 (SCI-Expanded)
- XVII. **Change of menarcheal age in schoolgirls living in Istanbul over the last 12 years**  
Güran T., Alir F., Arslan Y. T., Molla G., Sahin B., Sayar M. E., Atay Z., Helvacioğlu D., Gürpınar Tosun B., Haliloğlu B., et al.  
HORMONE RESEARCH IN PAEDIATRICS, cilt.95, sa.SUPPL 2, ss.364, 2022 (SCI-Expanded)
- XVIII. **Single Nucleotide Polymorphisms (SNPs) Profile as Predictive Markers of Lifestyle Modification Outcomes in Pediatric Obesity Treatment**  
Gawlik A., Sobalska-Kwapis M., Antosz A., Strapagiel D., Seweryn M., Shmoish M., BEREKET A., Wasniewska M., KIRKGÖZ T., DEMİRCİOĞLU S., et al.  
HORMONE RESEARCH IN PAEDIATRICS, cilt.95, sa.SUPPL 2, ss.231-232, 2022 (SCI-Expanded)
- XIX. **Circulating mRNA and miRNA Signatures as Predictive Markers of Lifestyle Modification Outcomes in Pediatric Obesity Treatment**  
Gawlik A., Shmoish M., BEREKET A., Wasniewska M., Antosz A., KIRKGÖZ T., DEMİRCİOĞLU S., GÜRAN T., Aversa T., Corica D., et al.  
HORMONE RESEARCH IN PAEDIATRICS, cilt.95, sa.SUPPL 2, ss.213, 2022 (SCI-Expanded)
- XX. **Low-dose ACTH Stimulation Test: Comparison of Cortisol Response at 30, 40, and 60 Minutes**  
Gürpınar Tosun B., Arıkan H., Demircioğlu S., Bereket A., Güran T.

- HORMONE RESEARCH IN PAEDIATRICS, cilt.95, sa.SUPPL 2, ss.117-118, 2022 (SCI-Expanded)
- XXI. **Breast ultrasonography: How useful in the diagnosis of precocious puberty?**  
Helvacioğlu D., BIYIKLI E., BUĞDAYCI O., DEMİRCİOĞLU S., GÜRAN T., BEREKET A.  
HORMONE RESEARCH IN PAEDIATRICS, cilt.95, sa.SUPPL 2, ss.46-47, 2022 (SCI-Expanded)
- XXII. **Wide phenotypical spectrum with the same karyotype: Mixed gonadal dysgenesis**  
Seven Menevşe T., Gürpınar Tosun B., Helvacioğlu D., Abali Z. Y., Kirmizibekmez H., Dursun F., Demircioğlu S., Bereket A., Güran T.  
HORMONE RESEARCH IN PAEDIATRICS, cilt.95, sa.SUPPL 2, ss.566, 2022 (SCI-Expanded)
- XXIII. **Glucagon response to hypoglycemia during extended oral glucose tolerance test in children with cystic fibrosis and comparing with healthy peers**  
HALİLOĞLU B., SEVEN MENEVŞE T., GÜRPINAR TOSUN B., GÜRAN T., DEMİRCİOĞLU S., İspir T., GÖKDEMİR Y., ERDEM ERALP E., BEREKET A.  
HORMONE RESEARCH IN PAEDIATRICS, cilt.95, sa.SUPPL 2, ss.199-200, 2022 (SCI-Expanded)
- XXIV. **A rare cause of monogenic obesity: Schaaf-Yang syndrome due to a novel MAGEL2 gene variant**  
Abali Z. Y., Ates E. A., GÜRAN T., BEREKET A., DEMİRCİOĞLU S.  
HORMONE RESEARCH IN PAEDIATRICS, cilt.95, sa.SUPPL 2, ss.230, 2022 (SCI-Expanded)
- XXV. **Adrenal steroids reference ranges in infancy determined by LC-MS/MS**  
Enver E. O., Vatanserver P., Guran O., Bilgin L., Boran P., Demircioğlu S., Haklar G., Bereket A., Güran T.  
PEDIATRIC RESEARCH, cilt.92, sa.1, ss.265-274, 2022 (SCI-Expanded)
- XXVI. **Homozygosity for a novel INHA mutation in two male siblings with hypospadias, primary hypogonadism, and high normal testicular volume**  
Guran T., Ates E. A., Eltan M., Sahin B., Tosun B. G., Seven Menevşe T., Geckinli B. B., Greenfield A., Turan S., Bereket A.  
SEXUAL DEVELOPMENT, cilt.16, sa.SUPPL 1, ss.61-62, 2022 (SCI-Expanded)
- XXVII. **Steroid hormone profiles and molecular diagnostic tools in pediatric patients with non-CAH primary adrenal insufficiency.**  
Seven Menevşe T., Kendir Demirkol Y., Gurpınar Tosun B., Bayramoğlu E., Yıldız M., Acar S., Erisen Karaca S., Orbak Z., Onder A., Sobu E., et al.  
The Journal of clinical endocrinology and metabolism, cilt.107, 2022 (SCI-Expanded)
- XXVIII. **Familial Hypomagnesemia with Hypercalciuria and Nephrocalcinosis Due to CLDN16 Gene Mutations: Novel Findings in Two Cases with Diverse Clinical Features**  
Eltan M., Abali Z. Y., Turkyılmaz A., Gökçe İ., Abali S., Alavanda C., Arman A., Kırkgöz T., Güran T., Hatun S., et al.  
CALCIFIED TISSUE INTERNATIONAL, cilt.110, sa.4, ss.441-450, 2022 (SCI-Expanded)
- XXIX. **A novel deletion involving the first GNAS exon encoding G $\alpha$  causes PHP1A without methylation changes at exon A/B**  
Campbell D., Reyes M., Kaygusuz S. B., Abali S., Güran T., Bereket A., Kagami M., Turan S., Jüppner H.  
Bone, cilt.157, 2022 (SCI-Expanded)
- XXX. **Homozygosity for a novel INHA mutation in two male siblings with hypospadias, primary hypogonadism, and high-normal testicular volume.**  
Arslan Ateş E., Eltan M., Sahin B., Gurpınar Tosun B., Seven Menevşe T., Geckinli B. B., Greenfield A., Turan S., Bereket A., Güran T.  
European journal of endocrinology, cilt.186, sa.5, 2022 (SCI-Expanded)
- XXXI. **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.**  
Kirkgoz T., Eltan M., Kaygusuz S. B., Yavas Abali Z., Helvacioğlu D., Seven Menevşe T., Gurpınar Tosun B., Guran T., Bereket A., Turan S.  
Journal of clinical research in pediatric endocrinology, cilt.14, ss.10-16, 2022 (SCI-Expanded)
- XXXII. **Broad-spectrum XX and XY gonadal dysgenesis in patients with a homozygous L193S variant in PPP2R3C.**  
Cicek D., Warr N., Yesil G., Kocak Eker H., Bas F., Poyrazoglu S., Darendeliler F., Direk G., Hatipoglu N., Eltan M., et al.

- European journal of endocrinology, cilt.186, ss.65-72, 2022 (SCI-Expanded)
- XXXIII. **Catch-up growth and discontinuation of fludrocortisone treatment in aldosterone synthase deficiency.**  
Gurpinar Tosun B., Kendir Demirkol Y., Seven Menevse T., Kaygusuz S. B., Ozbek M. N., Altincik S. A., Mammadova J., Cayir A., Doger E., Bayramoglu E., et al.  
The Journal of clinical endocrinology and metabolism, cilt.107, 2022 (SCI-Expanded)
- XXXIV. **Non-hormonal Clitoromegaly due to Clitoral Priapism Caused by Appendicitis/Appendectomy.**  
Gurpinar Tosun B., Karagozlu Akgul A., Almus E., Abidoglu S., Turan S., Bereket A., Guran T.  
Journal of clinical research in pediatric endocrinology, sa.4, 2021 (SCI-Expanded)
- XXXV. **Personalized approach to childhood obesity: Lessons from gut microbiota and omics studies. Narrative review and insights from the 29th European childhood obesity congress**  
Gawlik A., Salonen A., Jian C., Yanover C., Antosz A., Shmoish M., Wasniewska M., Bereket A., Wudy S. A., Hartmann M. F., et al.  
PEDIATRIC OBESITY, cilt.16, 2021 (SCI-Expanded)
- XXXVI. **Dysgenesis and Dysfunction of the Pancreas and Pituitary Due to FOXA2 Gene Defects.**  
Kaygusuz S. B., Arslan Ates E., Vignola M. L., Volkan B., Geckinli B. B., Turan S., Bereket A., Gaston-Massuet C., Guran T.  
The Journal of clinical endocrinology and metabolism, cilt.106, sa.10, 2021 (SCI-Expanded)
- XXXVII. **A rare cause of hypercalcemia: Congenital Lactase Deficiency**  
Eltan M., Alavanda C., Abali S., Abali Z. Y., Kaygusuz S. B., Gürpınar Tosun B., Seven Menevşe T., Helvacioğlu D., Güran T., Ata P., et al.  
HORMONE RESEARCH IN PAEDIATRICS, cilt.94, sa.SUPPL 1, ss.75, 2021 (SCI-Expanded)
- XXXVIII. **Hyperinsulinemic hypoglycemia due to biallelic mutations in the DNAJC3 gene**  
Gürpınar Tosun B., Seven Menevşe T., Esen N., Demircioğlu S., Yesilyurt A., Güran T., Bereket A.  
HORMONE RESEARCH IN PAEDIATRICS, cilt.94, sa.SUPPL 1, ss.107, 2021 (SCI-Expanded)
- XXXIX. **Global survey study of awareness, care and treatment of adolescents living with obesity, their caregivers and healthcare professionals: ACTION Teens**  
Halford J. C. G., Baur L., BEREKET A., Bin-Abbas B., Chen W., Fernandez-Aranda F., Garibay-Nieto N., Lopez Siguero J. P., Maffei C., Mooney V., et al.  
HORMONE RESEARCH IN PAEDIATRICS, cilt.94, sa.SUPPL 1, ss.291, 2021 (SCI-Expanded)
- XL. **Rahman Syndrome: Hypopituitarism might be responsible for changing height pattern**  
Kaygusuz S. B., Demircioğlu S., Esen N., Bereket A., Yesilyurt A., Güran T.  
HORMONE RESEARCH IN PAEDIATRICS, cilt.94, sa.SUPPL 1, ss.359, 2021 (SCI-Expanded)
- XLI. **46,XY DSD due to biallelic DHX37 gene mutations**  
Eltan M., Helvacioğlu D., Ates E. A., Abali Z. Y., Demircioğlu S., Bereket A., Güran T.  
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- XLII. **Acetazolamide treatment in a patient with pseudohypoparathyroidism with venous calcification**  
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- XLIII. **Is quail egg a potential endocrine disruptor?**  
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- XLIV. **Biallelic PPP2R3C mutations are associated with partial and complete gonadal dysgenesis in 46,XY and 46,XX individuals**  
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- XLV. **Machine Learning Quest for Predictive Markers of Lifestyle Modification Outcomes in Pediatric Obesity Treatment**  
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- XLVI. **Clinical and hormonal evolution of aldosterone synthase deficiency: Is complete remission possible?**  
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- XLVII. **Genotypic Sex and Severity of the Disease Determine the Time of Clinical Presentation in Steroid 17 alpha-Hydroxylase/17,20-Lyase Deficiency**  
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- XLVIII. **Correlation of 11-oxygenated C19 androgens with the clinical and biochemical characteristics in premature adrenarche**  
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- XLIX. **Exploring Urinary Bile Acids as Potential Markers of Metabolism: Reference Values in Children by Targeted LC-MS/MS**  
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- LI. **Clinical and Hormonal Profiles Correlate With Molecular Characteristics in Patients With 11 beta-Hydroxylase Deficiency**  
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- LIX. Management of Systemic Hypersensitivity Reactions to Gonadotropin-Releasing Hormone Analogues during Treatment of Central Precocious Puberty**  
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- LX. A Rare Cause of Hypophosphatemia: Raine Syndrome Changing Clinical Features with Age**  
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- LXIII. Fibroblast Growth Factor-23 and Matrix Extracellular Phosphoglycoprotein Levels in Healthy Children and, Pregnant and Puerperal Women**  
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- LXX. Restoration of Height after 11 Years of Letrozole Treatment in 11 beta-Hydroxylase Deficiency**  
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- LXXIII. **Postoperative and Long-Term Endocrinologic Complications of Craniopharyngioma.**  
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- LXXXIII. **A Case Of Syndromic Hypopituitarism**  
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- LXXXIV. **A rare cause of hypophosphatemia: Raine Syndrome**  
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- LXXXVI. **Genotype and Phenotype Characterization of Turkish Patients with Vitamin D Dependent Rickets Type IA**  
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- LXXXVII. **Characteristics of puberty, pubertal height gain and final height in children with classical 21 hydroxylase deficiency**  
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- XC. **CHD7 mutations in patients with anosmic or normosmic idiopathic hypogonadotropic hypogonadism**  
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- XCII. **Evaluation of growth and puberty in a child with a novel TBX19 gene mutation and review of the literature**  
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- XCIII. **Comparison of Treatment Regimens in Management of Severe Hypercalcemia Due to Vitamin D Intoxication in Children**  
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- XCV. **PPP2R3C gene variants cause syndromic 46,XY gonadal dysgenesis and impaired spermatogenesis in humans**  
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- XCVI. **Editorial: Neonatal Screening for Congenital Adrenal Hyperplasia in Turkey**  
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- CVII. Recurrent Hypoglycemia in a Preschooler Girl with Overgrowth: Isolated ACTH-Deficiency with a Novel TPIT Mutation**  
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- XLIII. **Presentation of coeliac disease diagnosed by screening in children with type 1 diabetes: Is it different?**  
Akkelle B. S., Sengul O., Volkan B., TUTAR E., ÇELİKEL Ç., BEREKET A., ERTEM ŞAHİNOĞLU D.  
52th ESPGHAN, Birleşik Krallık, 05 Haziran 2019 - 08 Haziran 2019, cilt.68, sa.1
- XLIV. **Segmental Aşırı büyüme kliniği olan olguda somatik PIK3CA mutasyonu**  
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3. Ege Endokrin Hastalıklar ve Genetik Sempozyumu, İzmir, Türkiye, 7 - 09 Mart 2019
- XLV. **Tip 1 diyabet ve çölyak hastalığı birlikteliği gösteren hastalarda prezentasyon klasik çölyak hastalarından farklı mı?**  
ŞAHİN AKKELLE B., KALAYCIK ŞENGÜL Ö., VOLKAN B., TUTAR E., ÇELİKEL Ç., BEREKET A., ERTEM ŞAHİNOĞLU D.  
6. Marmara Pediatri kongresi, 21-23 Şubat 2019, İstanbul, 21 Şubat 2019
- XLVI. **Düşük renin düzeyi: endokrin hipertansiyon**  
Abalı Z., ABALI S., KÜÇÜK N., CANPOLAT N., ÇİÇEK N., ALPAY H., BEREKET A., turan S., yeşil g., GÜRAN T.  
Çocuk Endokrinolojisi Olgu Sunumları-9, İstanbul, Türkiye, 19 - 20 Ekim 2018
- XLVII. **Nationwide Hypophosphatemic Rickets Study**  
ŞIKLAR Z., TURAN S., BEREKET A., ABACI A., BAŞ F., DEMİR K., GÜRAN T., AKBERZADE A., BÖBER E., ÖZBEK M. N., et al.

57.th Annual Meeting of the European Society for Paediatric Endocrinology(ESPE), ATİNA, Yunanistan, 27 - 29 Eylül 2018

- XLVIII. Persistan Mullerian Duct Syndrome: Rare but important aetiology of an inguinal hernia and cryptorchidism in boys**  
Bereket A., Buğrul F., Kirkgöz T., Karadeniz Cerit K., Canmemiş A., Turan S., Picard J., Tuğtepe H., Guran T.  
57th Annual European Society for Paediatric Endocrinology(ESPE), Athens, Yunanistan, 27 - 29 Eylül 2018
- XLIX. Relation of serum IGF-1 and IGFBP3 levels with acute exacerbation in cystic fibrosis**  
Eser F., Ergenekon P., Atag E., İkizoglu N. B., ERDEM ERALP E., GÖKDEMİR Y., Turan S., BEREKET A., Karadağ B.  
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- L. Isolated Congenital Central Hypothyroidism due to a Novel Mutation in TSH Beta Subunit Gene**  
Kirkgöz T., Özhan B., Çetin G. O., Kaygusuz S. B., Demircioğlu S., Bereket A., Guran T.  
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- LI. An 18 Month Old Boy with Hypoglycemic Convulsion and Obesity Due to POMC Deficiency**  
KAYGUSUZ S. B., YEŞİL G., KIRKGÖZ T., DEMİRCİOĞLU S., BEREKET A.  
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- LII. 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., CANMEMİS A., DEMİRCİOĞLU S., PICARD J. Y., TUĞTEPE H., GÜRAN T.  
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- LIII. Recurrent Hypoglycemia in a Preschooler Girl with Overgrowth: Isolated ACTH-Deficiency with a Novel TPIT Mutation**  
Yavaş Abalı Z., Yeşil G., Kirkgöz T., Kaygusuz S. B., Demircioğlu S., Bereket A., Guran T.  
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- LIV. Evaluation of molecular characteristics and steroid metabolomics in a large cohort of children with 3β-Hydroxysteroid Dehydrogenase 2 deficiency.**  
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- LV. Çok Merkezli Olarak Hipofosfatemik Riketsli Olguların Değerlendirilmesi**  
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22. PEDIATRİK ENDOKRİNOLOJİ VE DİYABET KONGRESİ, Antalya, Türkiye, 18 - 22 Nisan 2018
- LVI. Relation of serum IGF-1 and IGFBP3 levels with acute exacerbation in cystic fibrosis**  
Ergenekon P., Eser F., Atağ E., Baş İkizoglu N., Gökdemir Y., Erdem Eralp E., Demircioğlu S., Bereket A., Karadağ B. T.  
2nd Annual Middle East Cystic Fibrosis Congress, İzmir, Türkiye, 22 - 23 Mart 2018
- LVII. Neuroendocrin Regulation of Energy Homeostasis and Hypothalamic Obesity in Children**  
BEREKET A.  
3rd International Congress of the Turkish-Neuroendocrinology-Society, Malatya, Türkiye, 29 Haziran - 01 Temmuz 2018, cilt.107, ss.2
- LVIII. 17OH-PREGNENOLONE SEEMS A MAJOR DRIVE OF ANDROGEN EXCESS IN PATIENTS WITH 11β-HYDROXYLASE DEFICIENCY**  
Yildiz M., Turan S., Akcay T., Atay Z., Onal H., Baris T., Haklar G., Bereket A., Guran T.  
. 10th Individual Abstracts for International Meeting of Pediatric Endocrinology, Washington, Amerika Birleşik Devletleri, 14 - 17 Eylül 2017
- LIX. SIMULTANEOUS PROFILING OF 17 STEROID HORMONES USING LIQUID CHROMATOGRAPHY TANDEM MASS SPECTROMETRY IN NEWBORN AND EARLY INFANCY.**  
Enver E. Ö., Vatanserver P., Guran O., Boran P., Turan S., Haklar G., Bereket A., Guran T.  
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- LX. **17OH-pregnenolone seems a major drive of androgen excess in patient with 11 beta hydroxylase deficiency**  
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- LXI. **CLINICAL AND LABORATORY CHARACTERISTICS OF HYPERPROLACTINEMIC CHILDREN AND ADOLESCENTS: NATIONAL SURVEY.**  
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- LXII. **Evaluation of Clinical, Genetical, and Steroid Profile Features of Cases with 3Beta-Hydroxysteroid Dehydrogenase Type 2 Deficiency**  
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- LXIII. **Çocukluk çağındaki over kisti vakalarının incelenmesi: çok merkezli çalışma**  
Küçükemre Aydın B., Saka N., BAŞ F., POYRAZOĞLU Ş., BUNDAK R., YEŞİLTEPE MUTLU R. G., ÇİZMECİOĞLU F. M., HATUN Ş., HALİLOĞLU B., DEMİRCİOĞLU S., et al.  
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- LXIV. **b hCG from an Occult Source Causing Peripheral Precocious Puberty Identification of the Tumour 6 Years After Presentation**  
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- LXV. **Frequency of Recessive Osteogenesis Imperfecta in a Turkish Cohort and Genetic Causes**  
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- LXVI. **Reconsideration of Mid Parental Height Calculation**  
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- LXIX. **Nonsense Mutation in SPARC Gene Causing Autosomal Recessive Osteogenesis Imperfecta**  
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- LXX. **Thiamine Responsive Megaloblastic Anemia Due to SLCA19A2 Gene Mutation Another Cause of Neonatal Diabetes with Succesfull Switch from Insulin to Thiamine**  
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- LXXI. **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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- LXXII. **Premature Puberche Hyperinsülinemia Hypothyroxenemia and Hyperintensities in Basal Ganglia All Caused by a Single Congenital Defect**  
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- LXXIV. **Boy Kısaldığı Olan Hastalarda Özellikler ve Etiyolojik Dağılım Bir Çocuk Endokrinoloji Kliniği Verileri**  
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- LXXV. **Nonklasik Konjenital Adrenal Hiperplazi Hastalarının Genotip Ve Fenotip Özellikleri**  
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- LXXVI. **Otozomal Resesif Osteogenezis İmperfekta Populasyonumuzdaki Sıklığı Ve Genetik Nedenleri**  
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19. PEDIATRİK ENDOKRİNOLOJİ VE DİYABET KONGRESİ, Türkiye, 22 - 24 Ekim 2015
- LXXVII. **Periferal Puberte Prekokslu 129 Çocukta Etiyolojik Dağılım Ve Klinik Özellikler**  
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- LXXVIII. **Merkezi Yenidogan Tarama Programi ile Tani Almis Konjenital Hipotiroidili Vakalarimizin İzlemi**  
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- LXXIX. **Friedreich s Ataxia Presenting with Diabetes Mellitus in an Adolescent**  
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- LXXX. **Aetiological Spectrum and Clinical Characteristics of 129 Children with Gonadotropin Independent Precocious Puberty A Nationwide Cohort Study**  
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- LXXXI. **Primary Adrenal Insufficiency in Children without Congenital Adrenal Hyperplasia Molecular and Clinical Characterisation of a Nationwide Cohort**  
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- LXXXII. **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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- LXXXIII. **Central or Primary Hypothyroidism How to Differentiate in Patients with Low T4 but Mildly Elevated TSH Levels**  
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- LXXXIV. **Factors Effecting Response to Growth Hormone Treatment in Children with Turner Syndrome**  
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- LXXXV. **Hereditary Vitamin D Resistant Rickets Report of Four Cases with Successful Use of Intermittent Intravenous Calcium Via Peripheral Route**  
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- LXXXVI. ANTLEY BİXLER SENDROMLU BİR OLGUMUZ**  
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- LXXXVII. Hipoglisemi: Kistik Fibroze bağlı diyabet tanısında CGMS (sürekli glukoz izleme sistemi) ile OGTT nin karşılaştırılması**  
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- LXXXVIII. Maternal Thyroid Dysfunction and Neonatal Problems**  
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- LXXXIX. Comparison of insulin detemir and NPH insulin in children and adolescents with type 1 diabetes mellitus aged 2-16 years: a 52-week randomised clinical trial**  
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## **Desteklenen Projeler**

Boran P., Ipar N., Bereket A., Hatun Ş., TÜBİTAK Projesi, Tip 1 Diyabetli Çocuk ve Ergenlerde Uyku Bozukluğu ve Kronotipinin Belirlenmesi; Glisemik Kontrol Üzerindeki Etkisi ve Uygulanan Tedavinin Farkının Gözlemlenmesi, 2020 - 2021

Bereket A., Gawlik A., Wasnievska M., Hochberg Z., Wudy S. A., Diğer Uluslararası Fon Programları, ÇOCUKLUK ÇAĞI ŞİŞMANLIĞINDA ÇOKLU BİYOLOJİK VERİ ANALİZİ İLE HASTALIK ÖZELLİKLERİ SAPTANARAK KİŞİSELLEŞTİRİLMİŞ YAKLAŞIM , 2018 - 2021



## Bilimsel Dergilerdeki Faaliyetler

Journal Of Pediatric Endocrinology & Metabolism, Editör, 2016 - Devam Ediyor

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## Bilimsel Kuruluşlardaki Üyelikler / Görevler

Obezite Araştırma Derneği, Bilim Kurulu Üyesi, 2015 - Devam Ediyor , Türkiye

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Pediyatrik Endokrinoloji Derneği, Yönetim Kurulu Üyesi, 2001 - 2003, Türkiye

## Bilimsel Hakemlikler

JCRPE JOURNAL OF CLINICAL RESEARCH IN PEDIATRIC ENDOCRINOLOGY, SCI Kapsamındaki Dergi, Ekim 2021

Hormone Research In Paediatrics, SCI Kapsamındaki Dergi, Haziran 2021

## Bilimsel Danışmanlıklar

ACURARE (Acıbadem Üniversitesi, Nadir Hastalıklar Araştırma Merkezi), Kurum veya Organizasyonlar İçin Yapılan Danışmanlık, Marmara Üniversitesi, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, Türkiye, 2019 - Devam Ediyor

## Etkinlik Organizasyonlarındaki Görevler

Bereket A., Haliloğlu B., Güran T., Demircioğlu S., 27. Ulusal Pediyatrik Endokrinoloji ve Diyabet Kongresi, Bilimsel Kongre / Sempozyum Organizasyonu, Antalya, Türkiye, Mayıs 2023

Bereket A., IX. Pediyatrik Endokrinolojiye Giriş Kursu, Çalıştay Organizasyonu, Antalya, Türkiye, Mayıs 2023

Bereket A., Puberte Sempozyumu, Bilimsel Kongre / Sempozyum Organizasyonu, İstanbul, Türkiye, Aralık 2022

Bereket A., 11. Pediyatrik Endokrinoloji Olgu Sunumları sempozyumu, Bilimsel Kongre / Sempozyum Organizasyonu, İstanbul, Türkiye, Mayıs 2022

Bereket A., 1. Çocuk Endokrinolojide Araştırma Günleri, Bilimsel Kongre / Sempozyum Organizasyonu, Erzurum, Türkiye, Nisan 2022

Bereket A., Koç Üniversitesi hastanesi Pediyatrik Diyabet Günleri, Bilimsel Kongre / Sempozyum Organizasyonu, İstanbul, Türkiye, Kasım 2021

Bereket A., 9. Pediyatrik Endokrinoloji olgu sunumları , Bilimsel Kongre / Sempozyum Organizasyonu, İstanbul, Türkiye, Ekim 2018

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Bereket A., 18. European Congress of Obesity , Bilimsel Kongre / Sempozyum Organizasyonu, İstanbul, Türkiye, Mayıs 2011

Bereket A., Hypothalamic Obesity Symposium, Bilimsel Kongre / Sempozyum Organizasyonu, İstanbul, Türkiye, Mayıs

## Metrikler

Yayın: 389

Atıf (WoS): 3397

Atıf (Scopus): 3958

H-İndeks (WoS): 32

H-İndeks (Scopus): 35

## Kongre ve Sempozyum Katılımı Faaliyetleri

25. Ulusal Pediatrik Endokrinoloji Kongresi, Davetli Konuşmacı, Antalya, Türkiye, 2021

59. Annual European Society for Pediatric Endocrinology Meeting, İzleyici / Dinleyici, Kobenhavn, Danimarka, 2021

58. Annual European Society for Pediatric Endocrinology Meeting, Moderatör, Vienna, Avusturya, 2019

57. Annual European Society for Pediatric Endocrinology Meeting, Moderatör, Athens, Yunanistan, 2018

55. Annual ESPE meeting, Moderatör, Paris, Fransa, 2016

4th International multidisciplinary Postgraduate Course on Childhood Craniopharyngioma, Davetli Konuşmacı, Oldenburg, Almanya, 2016

## Davetli Konuşmalar

Action Teens: background and Introduction. ESPE Satellite symposium: "An ACTION to take to improve the lives of adolescents living with obesity", Konferans, European Society for Pediatric Endocrinology (ESPE), İtalya, Eylül 2022  
The ACTION Teens study. Rationale and Methodology., Konferans, International Congress on Obesity and Metabolic Syndrome (ICOMES)., Güney Kore, Eylül 2022

The ACTION Teens global survey: objectives and methods.: Treatment. V, Konferans, International Meeting on Childhood and Adolescent Obesity and Diabetes, İtalya, Haziran 2022

Growth outcome in children with Turner syndrome, SGA and Noonan syndrome on GH treatment, Seminer, MEDİCO ACADEME virtual Conference, Türkiye, Mart 2022

Premature thelarche: is it related to nutrition., Konferans, 2nd International Eurasian Congress of Social Pediatrics, Türkiye, Kasım 2020

Vitamin D: Present and future, Konferans, Milli Pediatri Derneği/5th Italian-Turkish Pediatric Symposium. November 1, 2019, Kyrenia, Kıbrıs (Kktc), Kasım 2019

Obesity management in children and adolescents. Master clinician case discussion, Çalıştay, European Association for the Study of obesity (EASO) National COMs Teaching Course, Türkiye, Eylül 2018

Obesity management in children and adolescents. Master clinician case discussion., Çalıştay, European Association for the Study of obesity (EASO) Training the East Mediterranean Trainers in the Prevention and management of obesity. National COMs Teaching Course, Türkiye, Eylül 2018

Neuroendocrin regulation of energy homeostasis and hypothalamic obesity in children, Konferans, Türkiye, Temmuz 2018

Evaluation of short children, Çalıştay, IVth.Novo Nordisk BANE Unit "Summit for Growth" Meeting, Dubai, 2016., Birleşik Arap Emirlikleri, Mayıs 2016

Hypothalamic Obesity – Mechanisms and treatment options. Overview, Çalıştay, 4th International multidisciplinary Postgraduate Course on Childhood Craniopharyngioma, April 7-10, 2016, Bad Zwischenahn, Germany, Almanya, Nisan 2016

Challenges in GH treatment across various indications: Turner Syndrome and SGA, Konferans, NordiUp Meeting, Tehran, 13-11-2015, İran, Kasım 2015

Case discussion of a girl with Turner Syndrome, Konferans, Growth Hormone Case Council by Novo Nordisk" Barcelona, Spain October 2, 2015, İspanya, Ekim 2015

A Controversial Case discussion of Precocious Puberty, Çalıştay, 29. European Society for Pediatric Endocrinology ESPE Summer School, Barcelona, , İspanya, Eylül 2015

Addressing puberty delay in growth disorders, Çalıştay, 3rd Novo Nordisk BANE Unit "Summit for Growth" Meeting, Türkiye, Mayıs 2015

Trends, challenges and efforts for better management of Diabetes in a transitional society, Konferans, IIIrd ISPAD Course and Conference. Future Diabetes. Varna, Bulgaria , April 24-26, 2015, Türkiye, Nisan 2015

A challenging patient with diabetes (Meet the expert session), Konferans, IIIrd International Society of Pediatric and Adolescent Diabetes Course and Conference. Future Diabetes. Varna, Bulgaria , April 24-26, 2015, Bulgaristan, Nisan 2015

Nutritional Vitamin D deficiency: Recent evidence. Definitions, clinical implications and therapeutic options, Seminer, Turkish National Pediatric Society and American Academy of Pediatrics Joint meeting Pediatrik Disease Course. 2014, Antalya , Türkiye, Ekim 2014

A controversial case of Diabetes, Çalıştay, 28. European Society for Pediatric Endocrinology ESPE Summer School, Dublin, Ireland 2014 , İrlanda, Eylül 2014

Addressing puberty delay in growth disorders, Seminer, EXEMED, Excellence in medical education. Advances in diagnosis and treatment of growth disorders symposium, Türkiye, Mayıs 2014

Growth hormone Transition to Adulthood: in whom? And How?, Çalıştay, 2nd. Novo Nordisk BANE Unit "Summit for Growth" Meeting, Türkiye, Nisan 2014

Optimized use of GH in short statured children, Konferans, 6th Iranian Congress on update in Pediatric Endocrinology and Metabolism, İran, Haziran 2013

Evaluation of worrisome growth, Çalıştay, 1.st Novo Nordisk BANE Unit "Summit for Growth" Meeting , Türkiye, Mayıs 2013

Epidemic of obesity and type 2 diabetes in children: implications for the future, Konferans, Excellence in Diabetes Meeting, Türkiye, Şubat 2013

Disorders of Puberty, Konferans, Saudi Pediatric Association Meeting, 47th Jeddah Pediatric Seminar, Suudi Arabistan, Şubat 2013

Type 2 diabetes in children, Çalıştay, 27. European Society for Pediatric Endocrinology ESPE Summer School , Almanya, Eylül 2012

Meet the Expert Session. Management of Central Precocious Puberty in girls. , Konferans, 51st European Society for Pediatric Endocrinology Meeting, Leipzig, Germany, Almanya, Eylül 2012

Impact of Growth hormone treatment in Turner Syndrome and SGA, Konferans, Norditropin Growth hormone Summit-Iraq, Türkiye, Aralık 2011

Understanding everyday difficulties of treating patients from 0 to 17 with type 1 diabetes including pump therapy, Konferans, Lilly Regional Diabetes Summit, Türkiye, Kasım 2011

Current status of childhood obesity and associated comorbidities in Turkey. Childhood obesity Task Force Meeting. , Konferans, 18.th European Congress of Obesity, Türkiye, Mayıs 2011

Thyroid health in Pregnancy Symposium. Acibadem University/Neonatal hypothyroid Screening for Congenital hypothyroidism in Turkey, Seminer, Acibadem Mehmet Ali Aydınlar Üniversitesi, Türkiye, Nisan 2011

Treatment of Hirsutism in Nonclassical Congenital Adrenal Hyperplasia, Konferans, Androgen Excesss & PCOS Society Update Meeting-Pathogenesis, Diagnosis and Treatment of Hirsutism, Türkiye, Mayıs 2010

Complications and metabolic risks of obesity in childhood and adolescence. , Konferans, 37th PanHellenic Congress of Endocrinology, and 1st Meditarrenean Endocrine day. Athens, Greece, 2010, Yunanistan, Nisan 2010

Rickets in children: Nature versus Nurture, Konferans, 53. Milli Pediatri Derneği/1.Turkish-Egyptian Pediatric Symposium., Türkiye, Ekim 2009

Meet the expert session. Nutritional Rickets, Konferans, 8th.Joint Meeting of LWPES/ European Society for Pediatric Endocrinology, New York, USA, 2009., Amerika Birleşik Devletleri, Eylül 2009

Preoperative assessment of a child with craniopharyngeoma, Konferans, European Society for Pediatric Neurosurgery (ESPN) /Postgraduate course: Consensus conference on Craniopharyngeomas in children, Türkiye, Nisan 2007

Definition and clinical approach to growth retardation in children, Konferans, 29. Congress of union of middle eastern

and mediterranean pediatric societies , Türkiye, Eylül 2005

Implications for physical Activity and obesity: From mechanism to management, , Seminer, European Association for Study of obesity (EASO)/ 14. European congress of Obesity satellite symposium, Türkiye, Haziran 2005

Short stature: A view of a Pediatric Endocrinologist , Konferans, Europediatrics 2003, Prague, , Çek Cumhuriyeti, Ekim 2003

Nutritional Rickets in Developing Countries. A consensus symposium on Vitamin D supplementation in children and adolescents. , Konferans, 6th Joint Meeting of LWPES/ESPE, Kanada, Temmuz 2001

## Burslar

Personalized approach to non-syndromic childhood obesity usiig multi-omics disease signature, Diğer Uluslararası Organizasyonlar, 2017 - Devam Ediyor

SBAG-2327 (100S079), (Metilfenidat tedavisinin çocuklarda büyüme ve IGF sistemi üzerine etkileri),2000, TÜBİTAK, 2000 - 2020

SBAG (108S133) (Doğumsal adrenal hiperplazili çocuklarda CYP21 mutasyonlarının araştırılması),2008, TÜBİTAK, 2008 - 2008

Rickets in the Middle-east, Diğer Uluslararası Organizasyonlar, 2004 - 2006

## Ödüller

Bereket A., Outstanding Clinician ödülü, European Society For Pediatric Endocrinology, Ekim 2022

Demircioğlu S., Güran T., Seven Menevşe T., Bereket A., Sözel Bildiri Üçüncülük Ödülü-Adrenokortikal hormon profilleri: KAH dışı primer adrenal yetmezlikte moleküler etiyolojiyi öngörebilir mi?, Xxv. Ulusal Pediatrik Endokrinoloji & Diyabet Kongresi , Ekim 2021

Sürekli Ö., Tamer S. A., Demircioğlu S., Güran T., Bereket A., Yegen B., Dr Tolga Köroğlu Özel Ödülü, 8. Marmara Pediatri Kongresi, Şubat 2021

Güran T., Demircioğlu S., Bereket A., Kaygusuz S. B., Eltan M., Tutar E., Gürpınar Tosun B., Seven Menevşe T., Poster Bildiri Birincilik Ödülü-Parsiyel Pankreatik Agenezi ve Sendromik Hipopituitarizm, Xxiv. Ulusal Pediatrik Endokrinoloji Ve Diyabet Çevrim İçi Kongresi, Kasım 2020

Güran T., Demircioğlu S., Bereket A., Haklar G., Sözlü Bildiri Birincilik Ödülü Steroid 11β-hidroksilaz eksikliği olan 100 çocuk hastanın klinik bulgularının genetik ve adrenokortikal hormon profili ile ilişkisinin değerlendirilmesi, Xxiv. Ulusal Pediatrik Endokrinoloji Ve Diyabet Çevrim İçi Kongresi, Kasım 2020

Güran T., Demircioğlu S., Bereket A., Eltan M., Gürpınar Tosun B., Kısa Sözel Bildiri Üçüncülük Ödülü- Biallelik PPP2R3C mutasyonları 46, XX ve 46, XY gonadal disgeneziye yol açar, Xxiv. Ulusal Pediatrik Endokrinoloji Ve Diyabet Çevrim İçi Kongresi, Kasım 2020

Haliloğlu B., Hysenaj G., Atay Z., Güran T., Abalı S., Demircioğlu S., Ellard S., Bereket A., En iyi ikinci sözel sunum, Ulusal Pediatrik Endokrinoloji Kongresi Bilimsel Kurul, Ekim 2013

Bereket A., TUBA GEBİP Seçkin Genç Bilimci Ödülü, Türkiye Bilimler Akademisi, Haziran 2001

## Akademi Dışı Deneyim

Hacettepe Üniversitesi