

## Prof.Dr. ABDULLAH BEREKET

### Kişisel Bilgiler

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### 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 dalı'nı 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

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

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

### I. Personalized approach to childhood obesity: Lessons from gut microbiota and omics studies.

#### Narrative review and insights from the 29th European childhood obesity congress

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### II. Dysgenesis and dysfunction of pancreas and pituitary due to FOXA2 gene defects.

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- V. **Cranial MRI abnormalities and long-term follow-up of the lesions in 770 girls with Central Precocious Puberty.**  
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- VII. **Does Genotype–Phenotype Correlation Exist in Vitamin D-Dependent Rickets Type IA: Report of 13 New Cases and Review of the Literature**  
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- XIII. **Management of Systemic Hypersensitivity Reactions to Gonadotropin-Releasing Hormone Analogues during Treatment of Central Precocious Puberty**  
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- XIV. **Nationwide Turkish Cohort Study of Hypophosphatemic Rickets**

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- XV. **Clinical Significance of Hypophosphatasemia in Children**  
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- XVI. **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**  
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- XVII. **Once-Weekly somapacitan vs daily GH in children with GH deficiency: Results from a randomized phase 2 trial**  
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- XVIII. **Fibroblast Growth Factor-23 and Matrix Extracellular Phosphoglycoprotein Levels in Healthy Children and, Pregnant and Puerperal Women**  
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- XXII. **Low DHEAS Concentration in a Girl Presenting with Short Stature and Premature Pubarche: A Novel PAPSS2 Gene Mutation**  
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- XXV. **Adrenocortical carcinoma in atypical Beckwith-Wiedemann syndrome due to loss of methylation at imprinting control region 2**  
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Bereket A.

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- XXIX. **Simplifying the interpretation of steroid metabolome data by a machine-learning approach**  
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- XXXIII. **Rare Causes of Osteogenesis Imperfecta are Common in Consanguineous Pedigrees**  
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- XXXIV. **A rare cause of hypophosphatemia: Raine Syndrome**  
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- 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**  
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- XXXVII. **How to Approach Systemic Hypersensitivity reactions to Gonadotropin Releasing Hormone Analogues during treatment of Central Precocious Puberty**  
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- XXXVIII. **Genotype and Phenotype Characterization of Turkish Patients with Vitamin D Dependent Rickets Type IA**  
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- XLIII. Exome Sequencing of a Primary Ovarian Insufficiency Cohort Reveals Common Molecular Etiologies for a Spectrum of Disease**  
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- XLV. Comparison of Treatment Regimens in Management of Severe Hypercalcemia Due to Vitamin D Intoxication in Children**  
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- XV. **Relation of serum IGF-1 and IGFBP3 levels with acute exacerbation in cystic fibrosis**  
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- XVI. **Persistent Mullerian duct syndrome: Rare But Important Aetiology of an Inguinal Hernia and Cryptorchidism in Boys**  
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- XVII. **An 18 Month Old Boy with Hypoglycemic Convulsion and Obesity Due to POMC Deficiency**  
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- XVIII. **Isolated Congenital Central Hypothyroidism due to a Novel Mutation in TSH Beta Subunit Gene**  
KIRKGÖZ T., ÖZHAN B., ÇETİN G. O. , KAYGUSUZ S. B. , DEMİRCİOĞLU S., BEREKET A., GÜRAN T.  
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- XIX. **Recurrent Hypoglycemia in a Preschooler Girl with Overgrowth: Isolated ACTH-Deficiency with a Novel TPIT Mutation**  
YAVAŞ ABALI Z., YEŞİL G., KIRKGOZ T., KAYGUSUZ S. B. , DEMİRCİOĞLU S., BEREKET A., GÜRAN T.  
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- XX. **Evaluation of molecular characteristics and steroid metabolomics in a large cohort of children with 3β-Hydroxysteroid Dehydrogenase 2 deficiency.**  
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- XXI. **Çok Merkezli Olarak Hipofosfatemik Riketsli Olguların Değerlendirilmesi**  
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- XXII. **Relation of serum IGF-1 and IGFBP3 levels with acute exacerbation in cystic fibrosis**  
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- XXIII. **Neuroendocrin Regulation of Energy Homeostasis and Hypothalamic Obesity in Children**  
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- XXIV. **Clinical characteristics of Turkish children and adolescents with type 2 diabetes**  
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- XXV. **17OH-pregnenolone seems a major drive of androgen excess in patient with 11 beta hydroxylase deficiency**  
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- XXVI. **SIMULTANEOUS PROFILING OF 17 STEROID HORMONES USING LIQUID CHROMATOGRAPHY TANDEM MASS SPECTROMETRY IN NEWBORN AND EARLY INFANCY.**

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- XXVIII. **CLINICAL AND LABORATORY CHARACTERISTICS OF HYPERPROLACTINEMIC CHILDREN AND ADOLESCENTS: NATIONAL SURVEY.**  
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- XXXIII. **Etiologic Distribution and Characteristics of Patients with Short Stature in a Pediatric Endocrinology Clinic**  
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- XXXIV. **Frequency of Recessive Osteogenesis Imperfecta in a Turkish Cohort and Genetic Causes**  
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- XXXV. **Reconsideration of Mid Parental Height Calculation**  
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- XXXVI. **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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- XLII. **Boy Kısaliğı Olan Hastalarda Özellikler ve Etiyolojik Dağılım Bir Çocuk Endokrinoloji Kliniğı Verileri**  
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- XLIII. **Nonklasik Konjenital Adrenal Hiperplazi Hastalarının Genotip Ve Fenotip Özellikleri**  
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- XLIV. **Merkezi Yenidogan Tarama Programi ile Tani Almis Konjenital Hipotiroidili Vakalarimizin İzlemi**  
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- XLV. **Otozomal Resesif Osteogenezis İmprefekta Populasyonumuzdaki Sıklığı Ve Genetik Nedenleri**  
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- XLVI. **Friedreich s Ataxia Presenting with Diabetes Mellitusin an Adolescent**  
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- XLVII. **Aetiolojik Spektrum ve Klinik Özellikleri 129 Çocukta Gonadotropin Bağımsız Erken Puberte Bir Ulusal Kohort Çalışması**  
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- L. **Hereditary Vitamin D Resistant Rickets Report of Four Cases with Successful Use of Intermittent Intravenous Calcium Via Peripheral Route**  
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- LI. **Central or Primary Hypothyroidism How to Differentiate in Patients with Low T4 but Mildly Elevated TSH Levels**  
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- LII. **Factors Effecting Response to Growth Hormone Treatment in Children with Turner Syndrome**  
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- LIV. **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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- LVI. **Identification of novel dentin matrix protein-1 (DMP1) mutations in two unrelated kindreds with autosomal recessive hypophosphatemia**  
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- LVII. **Detection of Y chromosomal material in patients with a 45,X karyotype by PCR method**  
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## Desteklenen Projeler

Boran P., İpar 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

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## Bilimsel Kuruluşlardaki Üyelikler / Görevler

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## Etkinlik Organizasyonlarındaki Görevler

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## Atıflar

Toplam Atıf Sayısı (WOS):2877  
h-indeksi (WOS):28

## Davetli Konuşmalar

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