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Eğitim Bilgileri

Post Doktora, The University Of Michigan, Medical School, Pediatric Nephrology, Amerika Birleşik Devletleri 2004 - 2005

Doktora, Hacettepe Üniversitesi, Tıp Fakültesi, Çocuk Nefrolojisi , Türkiye 2000 - 2002

Tıpta Uzmanlık, Hacettepe Üniversitesi, Tıp Fakültesi, Çocuk Sağlığı Ve Hastalıkları, Türkiye 1994 - 2000

Lisans, Ege Üniversitesi, Tıp Fakültesi, Türkiye 1988 - 1994

Yabancı Diller

İngilizce, B2 Orta Üstü

Araştırma Alanları

Tıp, Sağlık Bilimleri, Dahili Tıp Bilimleri, Çocuk Sağlığı ve Hastalıkları, Pediatrik Nefroloji

Akademik Unvanlar / Görevler

Doç.Dr., Hacettepe Üniversitesi, Tıp Fakültesi (ingilizce), Dahili Tıp Bilimleri Bölümü, 2006 - Devam Ediyor

Yrd.Doç.Dr., Hacettepe Üniversitesi, Tıp Fakültesi (ingilizce), Dahili Tıp Bilimleri Bölümü, 2005 - 2006

Verdiği Dersler

SODYUM VE SU DENGESİ VE BOZUKLUKLARI (DÖNEM III, TÜRKÇE TIP), Ön Lisans, 2016 - 2017

AKUT BÖBREK YETMEZLİĞİ (DÖNEM III, TÜRKÇE TIP), Ön Lisans, 2016 - 2017

SIVI-ELEKTROLİT (DÖNEM IV, PDÖ), Ön Lisans, 2016 - 2017

DEHİDRATASYON VE TEDAVİSİ (DÖNEM IV, TÜRKÇE TIP), Ön Lisans, 2016 - 2017

SODYUM VE SU DENGESİ BOZUKLUKLARI (DÖNEM III, İNGİLİZCE TIP), Ön Lisans, 2016 - 2017

AKUT VE KRONİK BÖBREK YETMEZLİĞİNE YAKLAŞIM (DÖNEM IV), Ön Lisans, 2016 - 2017

POTASYUM DENGESİ VE BOZUKLUKLARI (DÖNEM III, TÜRKÇE TIP), Ön Lisans, 2016 - 2017

Akut Böbrek Yetmezliği, Lisans, 2016 - 2017

EKSTREMİTE MUAYENESİ VE KAN BASINCI ÖLÇÜMÜ (DÖNEM IV), Ön Lisans, 2016 - 2017

DEHİDRATASYON VE TEDAVİSİ (DÖNEM IV, İNGİLİZCE TIP), Ön Lisans, 2016 - 2017

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İDRAR KAÇIRAN ÇOCUĞA YAKLAŞIM, Ön Lisans, 2016 - 2017

SCI, SSCI ve AHCI İndekslerine Giren Dergilerde Yayınlanan Makaleler

- I. **Management of congenital nephrotic syndrome: consensus recommendations of the ERKNet-ESPN Working Group**
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- II. **Determinants of outcomes in chronic pediatric peritoneal dialysis: a single center experience**
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- III. **Transplantation in pediatric aHUS within the era of eculizumab therapy**
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- IV. **Genetic aspects of congenital nephrotic syndrome: a consensus statement from the ERKNet-ESPN inherited glomerulopathy working group**
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- V. **A homozygousHOXA11variation as a potential novel cause of autosomal recessive congenital anomalies of the kidney and urinary tract**
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- VI. **Renal Biopsy Prognostic Findings in Children With Atypical Hemolytic Uremic Syndrome**
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- VII. **Clinical characteristics of children with congenital anomalies of the kidney and urinary tract and predictive factors of chronic kidney disease**
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- VIII. **Predictors for the use of herbal and dietary supplements in children and adolescents with kidney and urinary tract diseases**
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- IX. **Cystinosis beyond kidneys: gastrointestinal system and muscle involvement**
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- X. **CD80 expression and infiltrating regulatory T cells in idiopathic nephrotic syndrome of childhood**
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- XI. **Surgical management of renovascular hypertension in children and young adults: a 13-year experience**
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- XII. **Long-term follow-up results of patients with ADCK4 mutations who have been diagnosed in the asymptomatic period: effects of early initiation of CoQ10 supplementation**
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- XIV. **Persistent hypoglycemic attacks during hemodialysis sessions in an infant with congenital nephrotic syndrome: Questions and Answers**
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- XV. **An immunohistochemical approach to detect oncogenic CTNNB1 mutations in primary neoplastic tissues**
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- XVI. **BK virus associated nephropathy and severe pneumonia in a kidney transplanted adolescent with Schimke immune-osseous-dysplasia**
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- XVIII. **MYCETOMA OF THE URINARY TRACT IN AN INFANT WITH HORSESHOE KIDNEY: USEFULNESS OF MECHANICAL REMOVAL AND AMPHOTERICIN-B IRRIGATION**
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- XIX. **EARLY USE OF ECULIZUMAB IN A CASE OF STEC-HUS BEFORE THE DEVELOPMENT OF CLINICAL SYMPTOMS**
BAYRAM M. T. , DEMİR B. K. , SOYLU A., KAVUKÇU S., ÖZALTIN F.
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- XX. **LONG-TERM FOLLOW-UP RESULTS OF PATIENTS WITH ADCK4 MUTATIONS WHO HAVE BEEN DIAGNOSED IN ASYMPTOMATIC PERIOD: EFFECTS OF EARLY INITIATION OF COQ10 SUPPLEMENTATION**
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- XXII. **TURKISH ATYPICAL HEMOLYTIC UREMIC SYNDROME REGISTRY: ECULIZUMAB TREATMENT IN AHUS PATIENTS**
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- XXIII. **CLINICAL AND PATHOLOGIC CHARACTERISTICS OF GENETICALLY CONFIRMED ALPORT SYNDROME PATIENTS**
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- XXIV. **LUPUS NEPHRITIS: LONG TERM FOLLOW-UP AND EFFECT OF TREATMENT ON GROWTH**

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- XXVII. **CLINICAL AND MUTATIONAL SPECTRUM OF CHILDREN WITH AUTOSOMAL RECESSIVE AND
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- XXVIII. **ADRENAL INSUFFICIENCY IN AN INFANT WITH CONGENITAL NEPHROTIC SYNDROME: NEPHROTIC
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- XXIX. **GASTRIC DUPLICATION CYST IN AN INFANT WITH FINNISH-TYPE CONGENITAL NEPHROTIC
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- XXX. **Extra-Renal manifestations of atypical hemolytic uremic syndrome in children**
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- XXXI. **Effect of IGF-1 and HGF induced bone marrow mesenchymal stem cells on focal segmental
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- XXXII. **Atypical Hemolytic Uremic Syndrome in Children Aged < 2 Years**
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- XXXIII. **Nephropathic Cystinosis Mimicking Bartter Syndrome A Novel Mutation**
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- XXXIV. **Mutations in KEOPS-complex genes cause nephrotic syndrome with primary microcephaly**
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- XXXV. **Long-Term Outcome of Steroid-Resistant Nephrotic Syndrome in Children**
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- XXXVI. **USE OF HERBAL AND DIETARY SUPPLEMENTS IN CHILDREN WITH KIDNEY AND URINARY TRACT
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- XXXVII. **DOES C.3979 G > A/P.VAL1327MET VARIANT OF COL4A4 HAS ANY PATHOGENIC EFFECT IN**

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- LI. **Loss of diacylglycerol kinase epsilon in mice causes endothelial distress and impairs glomerular Cox-2 and PGE(2) production**
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