

Doç. Dr. BORA GÜLHAN

Kişisel Bilgiler

E-posta: bora.gulhan@hacettepe.edu.tr

Web: <https://avesis.hacettepe.edu.tr/bora.gulhan>

Posta Adresi: boragulhan@yahoo.com

Uluslararası Araştırmacı ID'leri

ORCID: 0000-0003-0236-5786

Yoksis Araştırmacı ID: 170066

Eğitim Bilgileri

Tıpta Uzmanlık, Hacettepe Üniversitesi, Tıp Fakültesi, Çocuk Sağlığı Ve Hastalıkları, Türkiye 2004 - Devam Ediyor

Tıpta Yandal Uzmanlık, Hacettepe Üniversitesi, Tıp Fakültesi, Çocuk Sağlığı Ve Hastalıkları, Türkiye 2009 - 2013

Yabancı Diller

İngilizce, C1 İleri

Araştırma Alanları

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

Akademik Unvanlar / Görevler

Doç. Dr., Hacettepe Üniversitesi, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2016 - Devam Ediyor

Yrd. Doç. Dr., Hacettepe Üniversitesi, Tıp Fakültesi (İngilizce), Dahili Tıp Bilimleri Bölümü, 2015 - 2016

Uzman, Hacettepe Üniversitesi, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2013 - 2015

Uzman, Hacettepe Üniversitesi, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2009 - 2013

Araştırma Görevlisi, Hacettepe Üniversitesi, Tıp Fakültesi (İngilizce), Dahili Tıp Bilimleri Bölümü, 2004 - 2009

SCI, SSCI ve AHCI İndekslerine Giren Dergilerde Yayınlanan Makaleler

- Kidney transplantation in children and adolescents with C3 glomerulopathy or immune complex membranoproliferative glomerulonephritis: a real-world study within the CERTAIN research network**
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- IV. **Variable phenotype and genotype of pediatric patients with HNF1B nephropathy.**
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- V. **Acute kidney injury in children with moderate-severe COVID-19 and multisystem inflammatory syndrome in children: a referral center experience**
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- VI. **Pediatric kidney care experience after the 2023 Turkey/Syria earthquake.**
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- VII. **Management of pediatric hemolytic uremic syndrome**
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- VIII. **Omic Studies on In Vitro Cystinosis Model: siRNA-Mediated CTNS Gene Silencing in HK-2 Cells**
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- IX. **An unusual cause of diarrhea in a child with nephrotic syndrome: Questions.**
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- XI. **Long-term kidney follow-up after pediatric acute kidney support therapy for children less than 15 kg**
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- XII. **The outcomes of renin-angiotensin-aldosterone system inhibition and immunosuppressive therapy in children with X-linked Alport syndrome**
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- XIV. **Metabolomic Analyses to Identify Candidate Biomarkers of Cystinosis**
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- XV. **The Clinical and Mutational Spectrum of 69 Turkish Children with Autosomal Recessive or Autosomal Dominant Polycystic Kidney Disease: A Multicenter Retrospective Cohort Study**
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- XVI. **Case Report: Severe McCune–Albright syndrome presenting with neonatal Cushing syndrome: navigating through clinical obstacles**
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- XVII. **Hearing Loss Related to Gene Mutations in Distal Renal Tubular Acidosis**
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- XVIII. **CANDIDATE BIOMARKER(S) FOR CYSTINOSIS WITH OMIC-BASED TECHNOLOGY: FROM LABORATORY TO BED-SIDE**
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- XIX. **ACUTE KIDNEY INJURY ASSOCIATED WITH COVID-19 AND MULTISYSTEM INFLAMMATORY SYNDROME IN CHILDREN (MIS-C)**
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- XX. **ACUTE HEMODIALYSIS EXPERIENCE IN PEDIATRIC PATIENTS WEIGHING LESS THAN 15 KG**
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- XXI. **COVID-19 VACCINE-RELATED SIDE EFFECTS AMONG ADOLESCENTS WITH CHRONIC KIDNEY CONDITIONS: A SINGLE-CENTER EXPERIENCE**
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- XXIII. **EFFECTS OF SIROLIMUS ON RENAL FUNCTIONS AND GROWTH IN PEDIATRIC RENAL TRANSPLANT RECIPIENTS**
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- XXIV. **CLINICAL COURSE OF ADOLESCENT ONSET ATYPICAL HEMOLYTIC UREMIC SYNDROME: A STUDY OF TURKISH AHUS REGISTRY**
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- XXV. **Oral Coenzyme Q10 supplementation leads to better preservation of kidney function in steroid-resistant nephrotic syndrome due to primary Coenzyme Q10 deficiency**
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- XXVI. **Variation of the clinical spectrum and genotype-phenotype associations in Coenzyme Q10 deficiency associated glomerulopathy**

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XXXI. **Glomerulonephritis with crescents in childhood; etiologies and significance of M2 macrophages**

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XXXIII. **CLINICAL AND MUTATIONAL SPECTRUM OF CHILDREN WITH AUTOSOMAL RECESSIVE AND AUTOSOMAL DOMINANT POLYCYSTIC KIDNEY DISEASE**

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XXXIV. **EFFECTS OF RAAS INHIBITION AND IMMUNOSUPPRESSIVE THERAPY IN PEDIATRIC PATIENTS WITH X-LINKED ALPORT SYNDROME**

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- XXXIX. **TAKAYASU ARTERITIS WITH RENAL ARTERY INVOLVEMENT IN CHILDREN: 12 YEARS EXPERIENCE OF A TERTIARY CENTER**
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- XL. **Fludrocortisone is the salvage treatment in cases with calcineurin inhibitor related hyperkalemia**
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- XLI. **Neonatal McCune Albright Syndrome Presenting with Diabetes Mellitus**
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- XLII. **Hedera helix L: a possible cause of severe acute tubulointerstitial nephritis in an infant**
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- XLIII. **Clinical practice recommendations for recurrence of focal and segmental glomerulosclerosis/steroid-resistant nephrotic syndrome**
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- XLIV. **CLINICAL AND MUTATIONAL SPECTRUM OF CHILDREN WITH AUTOSOMAL RECESSIVE AND AUTOSOMAL DOMINANT POLYCYSTIC KIDNEY DISEASE**
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- XLV. **EARLY URINARY SYSTEM COMPLICATIONS IN CHILDREN WITH HEMATOPOETIC STEM CELL TRANSPLANTATION**
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- XLVI. **Transplantation in pediatric aHUS within the era of eculizumab therapy**
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- XLVII. **Aetiology, course and treatment of acute tubulointerstitial nephritis in paediatric patients: a cross-sectional web-based survey**
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- XLVIII. **Predictors for the use of herbal and dietary supplements in children and adolescents with kidney and urinary tract diseases**
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- XLIX. **Acute kidney injury in a patient with COVID-19: Answers**
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- L. **Determinants of outcomes in chronic pediatric peritoneal dialysis: a single center experience**
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- LI. **COL4A3 mutation is an independent risk factor for poor prognosis in children with Alport syndrome**
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- LII. **Clinical characteristics of children with congenital anomalies of the kidney and urinary tract and predictive factors of chronic kidney disease**
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- LIII. **Cystinosis beyond kidneys: gastrointestinal system and muscle involvement**
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- LIV. **CD80 expression and infiltrating regulatory T cells in idiopathic nephrotic syndrome of childhood**
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- LV. **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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- LVI. **Rituximab for Children With Difficult-to-Treat Nephrotic Syndrome: Its Effects on Disease Progression and Growth**
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