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Education Information

Expertise In Medicine, Hacettepe University, Tıp Fakültesi, Çocuk Sağlığı Ve Hastalıkları, Turkey 2004 - Continues
Post Doctorate of Medicine, Hacettepe University, Tıp Fakültesi, Çocuk Sağlığı Ve Hastalıkları, Turkey 2009 - 2013

Foreign Languages

English, C1 Advanced

Research Areas

Medicine, Health Sciences, Internal Medicine Sciences, Child Health and Diseases, Pediatric Nephrology

Academic Titles / Tasks

Associate Professor, Hacettepe University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2016 - Continues

Assistant Professor, Hacettepe University, Tıp Fakültesi (İngilizce), Dahili Tıp Bilimleri Bölümü, 2015 - 2016

Expert, Hacettepe University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2013 - 2015

Expert, Hacettepe University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2009 - 2013

Research Assistant, Hacettepe University, Tıp Fakültesi (İngilizce), Dahili Tıp Bilimleri Bölümü, 2004 - 2009

Published journal articles indexed by SCI, SSCI, and AHCI

I. Adolescence-onset atypical hemolytic uremic syndrome: is it different from infant-onset?

Celegen K., Gulhan B., Fidan K., Yuksel S., Yilmaz N., Yilmaz A. C., Demircioğlu Kılıç B., Gokce I., Kavaz Tufan A., Kalyoncu M., et al.

Clinical and experimental nephrology, 2024 (SCI-Expanded)

II. Variable phenotype and genotype of pediatric patients with HNF1B nephropathy.

Gülhan B., Ekici O., Dursun İ., Göknar N., Yuksel S., Alaygut D., Özçakar Z. B., Nalçacıoğlu H., Demircioğlu Kılıç B., Söylemezoglu O., et al.

Clinical nephrology, 2024 (SCI-Expanded)

III. Acute kidney injury in children with moderate-severe COVID-19 and multisystem inflammatory syndrome in children: a referral center experience

- TAŞTEMEL ÖZTÜRK T., DÜZOVA A., OYGAR P. D., BALTU D., ÖZCİLİNGİR P., LAÇİNEL GÜRLEVİK S., KURT ŞÜKÜR E. D., AYKAN H. H., ÖZEN S., ERTUĞRUL İ., et al.
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- IV. **Pediatric kidney care experience after the 2023 Turkey/Syria earthquake.**
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- V. **Management of pediatric hemolytic uremic syndrome**
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Turkish Journal of Pediatrics, vol.66, no.1, pp.1-16, 2024 (SCI-Expanded)
- VI. **Omic Studies on In Vitro Cystinosis Model: siRNA-Mediated CTNS Gene Silencing in HK-2 Cells**
Baysal İ., YABANOĞLU ÇİFTÇİ S., NEMUTLU E., EYLEM C. C., Gök-Topak E. D., ULUBAYRAM K., KIR S., GÜLHAN B., UÇAR G., ÖZALTIN F., et al.
Laboratory investigation; a journal of technical methods and pathology, vol.104, no.1, pp.100287, 2024 (SCI-Expanded)
- VII. **An unusual cause of diarrhea in a child with nephrotic syndrome: Questions.**
Baltu D., Kurt Sukur E. D., Gumus E., Tastemel Ozturk T., Ergen Y. M., Demirtas D., Gülhan B., Ozaltin F., Orhan D., Özén H., et al.
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- VIII. **An unusual cause of diarrhea in a child with nephrotic syndrome: Answers.**
Baltu D., Kurt Sukur E. D., Gumus E., Tastemel Ozturk T., Ergen Y. M., Demirtas D., Gülhan B., Ozaltin F., Orhan D., Özén H., et al.
Pediatric nephrology (Berlin, Germany), vol.38, no.12, pp.3977-3981, 2023 (SCI-Expanded)
- IX. **Long-term kidney follow-up after pediatric acute kidney support therapy for children less than 15 kg**
GÜLÇEK Ö. N., GÜLHAN B., KESİCİ S., KURT ŞÜKÜR E. D., HAYRAN K. M., ÖZALTIN F., DÜZOVA A., BAYRAKCİ B., TOPALOĞLU R.
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- X. **The outcomes of renin-angiotensin-aldosterone system inhibition and immunosuppressive therapy in children with X-linked Alport syndrome**
ÖZDEMİR E. G., GÜLHAN B., ŞÜKÜR E. D. K., Atayar E., ATAN R., DURSUN İ., ÖZÇAKAR Z. B., SAYGILI S. K., SOYLU A., SÖYLEMEZOĞLU O., et al.
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- XI. **Metabolomic Analyses to Identify Candidate Biomarkers of Cystinosis**
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- XII. **Predictors of kidney complications and analysis of hypertension in children with allogeneic hematopoietic stem cell transplantation**
Gurbanov A., GÜLHAN B., KUŞKONMAZ B. B., OKUR F. V., ÖZALTIN F., DÜZOVA A., ÇETİNKAYA F. D., TOPALOĞLU R.
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- XIII. **The Clinical and Mutational Spectrum of 69 Turkish Children with Autosomal Recessive or Autosomal Dominant Polycystic Kidney Disease: A Multicenter Retrospective Cohort Study**
Tutal O., GÜLHAN B., Atayar E., Yuksel S., ÖZÇAKAR Z. B., Soylemezoglu O., SAYGILI S. K., ÇALIŞKAN S., Inozu M., Baskin E., et al.
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- XIV. **Hearing Loss Related to Gene Mutations in Distal Renal Tubular Acidosis**
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Audiology & neuro-otology, vol.28, no.5, pp.350-359, 2023 (SCI-Expanded)

- XV. **Case Report: Severe McCune-Albright syndrome presenting with neonatal Cushing syndrome: navigating through clinical obstacles**
ÜNSAL Y., GÖZMEN O., USER İ. R., Hizarcıoglu H., GÜLHAN B., EKİNCİ S., KARAGÖZ T., ÖZÖN Z. A., GÖNC E. N.
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- XVI. **ACUTE KIDNEY INJURY ASSOCIATED WITH COVID-19 AND MULTISYSTEM INFLAMMATORY SYNDROME IN CHILDREN (MIS-C)**
TAŞTEMEL ÖZTÜRK T., DÜZOVA A., OYGAR P. D., BALTU D., Ozcilingir P., LAÇİNEL GÜRLEVİK S., Kurtsukur E. D.,
AYKAN H. H., ÖZEN S., ERTUĞRUL İ., et al.
PEDIATRIC NEPHROLOGY, vol.37, no.11, pp.2804, 2022 (SCI-Expanded)
- XVII. **CLINICAL COURSE OF ADOLESCENT ONSET ATYPICAL HEMOLYTIC UREMIC SYNDROME: A STUDY OF TURKISH AHUS REGISTRY**
Celegen K., GÜLHAN B., FİDAN H. K., YÜKSEL S., Yilmaz N., Yilmaz A. C., KILIÇ B. D., GÖKCE İ., KAVAZ TUFAN A.,
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- XVIII. **EFFECTS OF SIROLIMUS ON RENAL FUNCTIONS AND GROWTH IN PEDIATRIC RENAL TRANSPLANT RECIPIENTS**
TAŞ N., GÜLHAN B., Ozdemir G., Ozturk T. T., BALTU D., KURT ŞÜKÜR E. D., ORHAN D., ÖZALTIN F., DÜZOVA A.,
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- XIX. **CANDIDATE BIOMARKER(S) FOR CYSTINOSIS WITH OMIC-BASED TECHNOLOGY: FROM LABORATORY TO BED-SIDE**
NEMUTLU E., Eylem C. C., GÜLHAN B., BAYSAL İ., Yabanoglu-Ciftci S., KIR S., ULUBAYRAM K., UÇAR G., SEZERMAN O. U., ÖZALTIN F., et al.
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- XX. **COVID-19 VACCINE-RELATED SIDE EFFECTS AMONG ADOLESCENTS WITH CHRONIC KIDNEY CONDITIONS: A SINGLE-CENTER EXPERIENCE**
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- XXI. **ACUTE HEMODIALYSIS EXPERIENCE IN PEDIATRIC PATIENTS WEIGHING LESS THAN 15 KG**
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- XXII. **COVID-19 IN CHILDREN WITH CHRONIC KIDNEY DISEASE; DOES IT DIFFER MUCH?**
BALTU D., KURT ŞÜKÜR E. D., TAŞTEMEL ÖZTÜRK T., GÜLHAN B., ÖZALTIN F., DÜZOVA A., TOPALOĞLU R.
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- XXIII. **Variation of the clinical spectrum and genotype-phenotype associations in Coenzyme Q10 deficiency associated glomerulopathy**
Drovandi S., Lipska-Zietkiewicz B. S., ÖZALTIN F., Emma F., GÜLHAN B., Boyer O., Trautmann A., Zietkiewicz S., Xu H.,
Shen Q., et al.
KIDNEY INTERNATIONAL, vol.102, no.3, pp.592-603, 2022 (SCI-Expanded)
- XXIV. **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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- XXV. **A broad clinical spectrum of PLC epsilon 1-related kidney disease and intrafamilial variability**
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- XXVI. **Apparent mineralocorticoid excess: A diagnosis beyond classical causes of severe hypertension in a child**

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- XXVII. **Long-term renal survival of paediatric patients with lupus nephritis**
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- XXVIII. **Eculizumab treatment and discontinuation in pediatric patients with atypical hemolytic uremic syndrome: a multicentric retrospective study**
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- XXIX. **Glomerulonephritis with crescents in childhood; etiologies and significance of M2 macrophages**
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- XXX. **Could plasma based therapies still be considered in selected cases with atypical hemolytic uremic syndrome?**
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- XXXI. **CHARACTERISTICS AND OUTCOME OF BK VIRUS INFECTION IN PEDIATRIC RENAL TRANSPLANT RECIPIENT**
KURT ŞÜKÜR E. D., Ozdemir G., TAŞTEMEL ÖZTÜRK T., BALTU D., GÜLHAN B., ÖZALTIN F., DÜZOVA A., TOPALOĞLU R.
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- XXXII. **AUDIOLOGICAL FINDINGS IN DISTAL RENAL TUBULAR ACIDOSIS**
Ay E., GÜRSES E., Arslan F., GÜLHAN B., Alniacik A., DÜZOVA A., BAJİN M. D., SENNAROĞLU L., GENÇ G. A., ÖZALTIN F., et al.
PEDIATRIC NEPHROLOGY, vol.36, no.10, pp.3428, 2021 (SCI-Expanded)
- XXXIII. **PRIMARY COQ10 DEFICIENCY: CLINICAL SPECTRUM AND GENOTYPE-PHENOTYPE CORRELATIONS**
Drovandi S., Lipska-zietkiewicz B. S., ÖZALTIN F., Emma F., GÜLHAN B., Boyer O., Trautmann A., Shen Q., Rao J., Riedhammer K. M., et al.
PEDIATRIC NEPHROLOGY, vol.36, no.10, pp.3445-3446, 2021 (SCI-Expanded)
- XXXIV. **EFFECTS OF RAAS INHIBITION AND IMMUNOSUPPRESSIVE THERAPY IN PEDIATRIC PATIENTS WITH X-LINKED ALPORT SYNDROME**
Ozdemir G., GÜLHAN B., KURT ŞÜKÜR E. D., Atayar E., DURSUN İ., ÖZÇAKAR Z. B., Saygili S., SOYLU A., Soylemezoglu O., Yilmaz A., et al.
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- XXXV. **TAKAYASU ARTERITIS WITH RENAL ARTERY INVOLVEMENT IN CHILDREN: 12 YEARS EXPERIENCE OF A TERTIARY CENTER**
TAŞTEMEL ÖZTÜRK T., BALTU D., KURT ŞÜKÜR E. D., GÜLHAN B., ÖZEN S., DÜZOVA A., ÖZALTIN F., TOPALOĞLU R.
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- XXXVI. **LONG TERM FOLLOW UP IN RENAL SCARRING OF URINARY TRACT INFECTION: ALBUMINURIA, DIASTOLIC BLOOD PRESSURE**
BALTU D., VOLKAN SALANCI B., GÜLHAN B., TAŞTEMEL ÖZTÜRK T., KURT ŞÜKÜR E. D., ÖZALTIN F., DÜZOVA A., TOPALOĞLU R.
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- XXXVII. **CLINICAL AND MUTATIONAL SPECTRUM OF CHILDREN WITH AUTOSOMAL RECESSIVE AND AUTOSOMAL DOMINANT POLYCYSTIC KIDNEY DISEASE**
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- XXXVIII. Neonatal McCune Albright Syndrome Presenting with Diabetes Mellitus**
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- XXXIX. Fludrocortisone is the salvage treatment in cases with calcineurin inhibitor related hyperkalemia**
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- XL. Hedera helix L: a possible cause of severe acute tubulointerstitial nephritis in an infant**
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- XLII. Transplantation in pediatric aHUS within the era of eculizumab therapy**
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- XLIII. Clinical practice recommendations for recurrence of focal and segmental glomerulosclerosis/steroid-resistant nephrotic syndrome**
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- XLIV. EARLY URINARY SYSTEM COMPLICATIONS IN CHILDREN WITH HEMATOPOETIC STEM CELL TRANSPLANTATION**
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- XLV. Aetiology, course and treatment of acute tubulointerstitial nephritis in paediatric patients: a cross-sectional web-based survey**
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- XLVI. Predictors for the use of herbal and dietary supplements in children and adolescents with kidney and urinary tract diseases**
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- XLVIII. Determinants of outcomes in chronic pediatric peritoneal dialysis: a single center experience**
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- XLIX. COL4A3 mutation is an independent risk factor for poor prognosis in children with Alport syndrome**
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- L. Clinical characteristics of children with congenital anomalies of the kidney and urinary tract and**

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- L.I. Cystinosis beyond kidneys: gastrointestinal system and muscle involvement**
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- L.II. CD80 expression and infiltrating regulatory T cells in idiopathic nephrotic syndrome of childhood**
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- L.III. 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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- L.IV. Rituximab for Children With Difficult-to-Treat Nephrotic Syndrome: Its Effects on Disease Progression and Growth**
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- LV. Do not Miss Rare and Treatable Cause of Early-Onset Hemolytic Uremic Syndrome: Cobalamin C Deficiency**
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- L.VI. BK virus associated nephropathy and severe pneumonia in a kidney transplanted adolescent with Schimke immune-osseous-dysplasia**
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- L.VII. CLINICAL AND PATHOLOGIC CHARACTERISTICS OF GENETICALLY CONFIRMED ALPORT SYNDROME PATIENTS**
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- L.VIII. CLINICAL AND MUTATIONAL SPECTRUM OF CHILDREN WITH AUTOSOMAL RECESSIVE AND AUTOSOMAL DOMINANT POLYCYSTIC KIDNEY DISEASE**
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