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Öğrenim Bilgisi

Post Doktora 2004 - 2005	The University of Michigan, Medical School, Pediatric Nephrology, Amerika Birleşik Devletleri
Doktora 2000 - 2002	Hacettepe Üniversitesi, Tıp Fakültesi, Çocuk Nefrolojisi, Türkiye
Tıpta Uzmanlık 1994 - 2000	Hacettepe Üniversitesi, Tıp Fakültesi, Çocuk Sağlığı Ve Hastalıkları, Türkiye
Lisans 1988 - 1994	Ege Üniversitesi, Tıp Fakültesi, Türkiye

Yabancı Diller

İngilizce, B2 Orta Üstü

Akademik Unvanlar / Görevler

Prof. Dr. 2018 - Devam Ediyor	Hacettepe Üniversitesi, Tıp Fakültesi (İngilizce), Dahili Tıp Bilimleri Bölümü
Doç. Dr. 2006 - 2018	Hacettepe Üniversitesi, Tıp Fakültesi (İngilizce), Dahili Tıp Bilimleri Bölümü
Yrd. Doç. Dr. 2005 - 2006	Hacettepe Üniversitesi, Tıp Fakültesi (İngilizce), Dahili Tıp Bilimleri Bölümü

Desteklenen Projeler

- ÖZALTIN F., TOPALOĞLU R., TÜRKOĞLU Ö., ATAYAR E., GÜLHAN B., Yükseköğretim Kurumları Destekli Proje,

Çocukluk Çağı Otozomal Dominant ve Otozomal Resesif Polikistik Böbrek Hastalıklarının Genetik ve Klinik Özelliklerinin Araştırılması, 2019 - 2021

2. ÖZALTIN F., TOPALOĞLU R., EROĞLU İ., NEMUTLU E., ÖZBAY H. S., YABANOĞLU ÇİFTÇİ S., Yükseköğretim Kurumları Destekli Proje, CoQ10 Nefropatisinde Koenzim Q10 İçerikli Nanoterapötiklerin Tedavi Etkiliğinin Araştırılması ve İn Vitro Modelde Mitokondriyal Fonksiyonlarının Karşılaştırılması, 2018 - 2021
3. ÖZALTIN F., BEŞBAŞ N., Yükseköğretim Kurumları Destekli Proje, Atipik Hemolitik Üremik Sendromlu Hastalarda Kompleman Düzenleyici Genlerdeki Mutasyonların Saptanması, 2011 - 2016

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1. **Calcineurin inhibitor-related hyperkalemia is caused by hyporeninemic hypoaldosteronism and fludrocortisone is an effective treatment: Report of a case series and review of the literature**
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4. **Acute kidney injury in children with moderate-severe COVID-19 and multisystem inflammatory syndrome in children: a referral center experience**
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6. **Management of pediatric hemolytic uremic syndrome**
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7. **Genotype/phenotype relationship in mild congenital nephrotic syndrome**
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8. **A novel homozygous missense variant in TBC1D31 in a consanguineous family with congenital anomalies of the kidney and urinary tract (CAKUT)**
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9. **Long-term kidney follow-up after pediatric acute kidney support therapy for children less than 15 kg**
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10. **COVID-19 associated thrombotic microangiopathy**
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11. **Outcomes of steroid-resistant nephrotic syndrome in children not treated with intensified immunosuppression**
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12. **A rare cause of nephrotic syndrome-sphingosine-1-phosphate lyase (SGPL1) deficiency: 6 cases and a review of the literature**
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13. **HOXA11 is another monogenic cause of congenital anomalies of the kidney and urinary tract**
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17. **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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18. **Response to Dr. Spizzirri et al**
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23. **Long-term renal survival of paediatric patients with lupus nephritis**
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24. **Eculizumab treatment and discontinuation in pediatric patients with atypical hemolytic uremic syndrome: a multicentric retrospective study**
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25. **Clinically Different Presentations of Family Members With thSame Homozygote Diacylglycerol Kinase Epsilon Mutation: Case Report**
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26. **Mitochondria-targeted CoQ(10) loaded PLGA-b-PEG-TTP nanoparticles: Their effects on mitochondrial functions of COQ8B(-/-) HK-2 cells**
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27. **A splice site mutation in the TSEN2 causes a new syndrome with craniofacial and central nervous system malformations, and atypical hemolytic uremic syndrome**
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28. **Mitigation of portal fibrosis and cholestatic liver disease in ANKS6-deficient livers by macrophage depletion**
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29. **Glomerulonephritis with crescents in childhood; etiologies and significance of M2 macrophages**
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30. **Could plasma based therapies still be considered in selected cases with atypical hemolytic uremic syndrome?**
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31. **An international cohort study spanning five decades assessed outcomes of nephropathic cystinosis**
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32. **AGTR1-RELATED RENAL TUBULAR DYSGENESES MAY NOT BE FATAL**
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33. **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.
PEDIATRIC NEPHROLOGY, cilt.36, sa.10, ss.3441, 2021 (SCI-Expanded)
34. **PRIMARY COQ10 DEFICIENCY: CLINICAL SPECTRUM AND GENOTYPE-PHENOTYPE CORRELATIONS**

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35. **CHARACTERISTICS AND OUTCOME OF BK VIRUS INFECTION IN PEDIATRIC RENAL TRANSPLANT RECIPIENT**

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

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

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42. **A HOMOZYGOUS HOXA11 VARIATION AS A POTENTIAL NOVEL CAUSE OF AUTOSOMAL RECESSIVE CONGENITAL ANOMALIES OF THE KIDNEY AND URINARY TRACT**

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43. **DGKE MUTATION IN A CHILD TREATED WITH ECULIZUMAB**

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49. **AGTR1-related Renal Tubular Dysgeneses May Not Be Fatal**
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56. **COL4A3 mutation is an independent risk factor for poor prognosis in children with Alport syndrome**
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