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Learning Knowledge

Post Doctorate 2004 - 2005	The University of Michigan, Medical School, Pediatric Nephrology, United States Of America
Doctorate 2000 - 2002	Hacettepe University, Tıp Fakültesi, Çocuk Nefrolojisi , Turkey
Expertise In Medicine 1994 - 2000	Hacettepe University, Tıp Fakültesi, Çocuk Sağlığı Ve Hastalıkları, Turkey
Undergraduate 1988 - 1994	Ege University, Faculty Of Medicine, Turkey

Foreign Languages

English, B2 Upper Intermediate

Academic Titles / Tasks

Professor 2018 - Continues	Hacettepe University, Tıp Fakültesi (İngilizce), Dahili Tıp Bilimleri Bölümü
Associate Professor 2006 - 2018	Hacettepe University, Tıp Fakültesi (İngilizce), Dahili Tıp Bilimleri Bölümü
Assistant Professor 2005 - 2006	Hacettepe University, Tıp Fakültesi (İngilizce), Dahili Tıp Bilimleri Bölümü

Supported Projects

- ÖZALTIN F., TOPALOĞLU R., TÜRKOĞLU Ö., ATAYAR E., GÜLHAN B., Project Supported by Higher Education

Institutions, 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., Project Supported by Higher Education Institutions, CoQ10 Nefropatisinde Koenzim Q10 İçerikli Nanoterapötiklerin Tedavi Etkiliğinin Araştırılması ve İn Vitro Modelde Mitokondriyal Fonksiyonlarının Karşılaştırılması, 2018 - 2021
3. ÖZALTIN F., BEŞBAŞ N., Project Supported by Higher Education Institutions, Atipik Hemolitik Üremik Sendromlu Hastalarda Kompleman Düzenleyici Genlerdeki Mutasyonların Saptanması, 2011 - 2016

Published journal articles indexed by SCI, SSCI, and AHCI

1. **Steroid-Resistant Nephrotic Syndrome due to NPHS2 Variants Is Not Associated With Posttransplant Recurrence**
Kachmar J., Boyer O., Lipska-Ziętkiewicz B., Morinière V., Gribouval O., Heidet L., Balasz-Chmielewska I., Benetti E., Cloarec S., Csaicsich D., et al.
Kidney International Reports, vol.9, no.4, pp.973-981, 2024 (SCI-Expanded)
2. **Implication of transcription factor FOXD2 dysfunction in syndromic congenital anomalies of the kidney and urinary tract (CAKUT)**
Riedhammer K. M., Nguyen T. T., KOŞUKCU C., Calzada-Wack J., Li Y., Assia Batzir N., SAYGILI S. K., Wimmers V., Kim G., Chrysanthou M., et al.
Kidney International, vol.105, no.4, pp.844-864, 2024 (SCI-Expanded)
3. **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., ÖZÇİLİNGİR P., LAÇİNEL GÜRLEVİK S., KURT ŞÜKÜR E. D., AYKAN H. H., ÖZEN S., ERTUĞRUL İ., et al.
Pediatric Nephrology, vol.39, no.3, pp.867-877, 2024 (SCI-Expanded)
4. **Management of pediatric hemolytic uremic syndrome**
GÜLHAN B., ÖZALTIN F., Fidan K., ÖZÇAKAR Z. B., Söylemezoğlu O.
Turkish Journal of Pediatrics, vol.66, no.1, pp.1-16, 2024 (SCI-Expanded)
5. **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)
6. **A novel homozygous missense variant in TBC1D31 in a consanguineous family with congenital anomalies of the kidney and urinary tract (CAKUT)**
SAYGILI S. K., KOŞUKCU C., BAŞTUĞ T., DOĞAN EKİCİ A. İ., YILMAZ E. M., Kalyoncu A. U., Ağbaş A., CANPOLAT N., Çalıskan S., ÖZALTIN F.
Clinical Genetics, vol.104, no.6, pp.679-685, 2023 (SCI-Expanded)
7. **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., BAYRAKCI B., TOPALOĞLU R.
Pediatric Nephrology, vol.38, no.11, pp.3811-3821, 2023 (SCI-Expanded)
8. **COVID-19 associated thrombotic microangiopathy**
YILMAZ E. M., Cebi M. N., Karahan I., SAYGILI S. K., Gulmez R., Demirgan E. B., Durak C., AYĞÜN F., ÖZALTIN F., Caliskan S., et al.
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9. **Outcomes of steroid-resistant nephrotic syndrome in children not treated with intensified immunosuppression**
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- A., ÇALIŞKAN S., Saeed B., et al.
PEDIATRIC NEPHROLOGY, vol.38, no.5, pp.1499-1511, 2023 (SCI-Expanded)
10. **A rare cause of nephrotic syndrome-sphingosine-1-phosphate lyase (SGPL1) deficiency: 6 cases and a review of the literature**
TAŞTEMEL ÖZTÜRK T., CANPOLAT N., SAYGILI S. K., BAYRAKÇI U. S., Soylemezoglu O., ÖZALTIN F., TOPALOĞLU R.
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 11. **HOXA11 is another monogenic cause of congenital anomalies of the kidney and urinary tract**
SAYGILI S. K., ÇALIŞKAN S., ÖZALTIN F.
PEDIATRIC NEPHROLOGY, vol.38, no.3, pp.933-934, 2023 (SCI-Expanded)
 12. **Metabolomic Analyses to Identify Candidate Biomarkers of Cystinosis**
NEMUTLU E., ÖZALTIN F., YABANOĞLU ÇİFTÇİ S., GÜLHAN B., Eylem C. C., BAYSAL İ., Gök-Topak E. D., ULUBAYRAM K., SEZERMAN O. U., UÇAR G., et al.
International Journal of Molecular Sciences, vol.24, no.3, 2023 (SCI-Expanded)
 13. **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.
PEDIATRIC NEPHROLOGY, vol.38, no.2, pp.461-469, 2023 (SCI-Expanded)
 14. **Hearing Loss Related to Gene Mutations in Distal Renal Tubular Acidosis**
Ay E., GÜRSES E., ASLAN F., GÜLHAN B., Alniacik A., DÜZOVA A., BAJİN M. D., SENNAROĞLU L., GENÇ G. A., ÖZALTIN F., et al.
Audiology & neuro-otology, vol.28, no.5, pp.350-359, 2023 (SCI-Expanded)
 15. **Hemoglobin cast nephropathy: a rare but serious complication of hemolysis in a pediatric patient**
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Turkish Journal of Pediatrics, vol.65, no.5, pp.874-880, 2023 (SCI-Expanded)
 16. **Response to Dr. Spizzirri et al**
TAŞTEMEL ÖZTÜRK T., ÖZALTIN F., TOPALOĞLU R.
Pediatric Nephrology, vol.38, no.1, pp.309-310, 2023 (SCI-Expanded)
 17. **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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 18. **Variation of the clinical spectrum and genotype-phenotype associations in Coenzyme Q10 deficiency associated glomerulopathy**
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 19. **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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 20. **A broad clinical spectrum of PLC epsilon 1-related kidney disease and intrafamilial variability**
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 21. **Long-term renal survival of paediatric patients with lupus nephritis**
Demir S., GÜLHAN B., ÖZEN S., Çelëgen K., BATU AKAL E. D., Taş N., ORHAN D., BİLGİNER Y., DÜZOVA A., ÖZALTIN F., et al.
Nephrology Dialysis Transplantation, vol.37, no.6, pp.1069-1077, 2022 (SCI-Expanded)
 22. **Eculizumab treatment and discontinuation in pediatric patients with atypical hemolytic uremic**

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JOURNAL OF NEPHROLOGY, vol.35, no.4, pp.1213-1222, 2022 (SCI-Expanded)

23. **Clinically Different Presentations of Family Members With thSame Homozygote Diacylglycerol Kinase Epsilon Mutation: Case Report**
Çelikkaya E, Güngör T, Karakaya D, Çakıcı E. K, Yazılıtaş F, ÖZALTIN F, Bülbül M.
Experimental and Clinical Transplantation, vol.20, no.5, pp.45-48, 2022 (SCI-Expanded)
24. **Mitochondria-targeted CoQ(10) loaded PLGA-b-PEG-TTP nanoparticles: Their effects on mitochondrial functions of COQ8B(-/-) HK-2 cells**
Sena Ozbay H, Yabanoglu-Ciftci S, Baysal I, Gultekinoglu M, Can Eylem C, Ulubayram K, Nemutlu E, Topaloglu R, Ozaltin F.
EUROPEAN JOURNAL OF PHARMACEUTICS AND BIOPHARMACEUTICS, vol.173, pp.22-33, 2022 (SCI-Expanded)
25. **A splice site mutation in the TSEN2 causes a new syndrome with craniofacial and central nervous system malformations, and atypical hemolytic uremic syndrome**
CANPOLAT N, Liu D, Atayar E, SAYGILI S. K, Kara N. S, Westfall T. A, Ding Q, Brown B. J, Braun T. A, Slusarski D, et al.
CLINICAL GENETICS, vol.101, pp.346-358, 2022 (SCI-Expanded)
26. **Mitigation of portal fibrosis and cholestatic liver disease in ANKS6-deficient livers by macrophage depletion**
Airik M, McCourt B, TAŞTEMEL ÖZTÜRK T, Huynh A. B, Zhang X, Tometich J. T, TOPALOĞLU R, ÖZEN H, ORHAN D, Nejak-Bowen K, et al.
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27. **Glomerulonephritis with crescents in childhood; etiologies and significance of M2 macrophages**
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TURKISH JOURNAL OF PEDIATRICS, vol.64, no.1, pp.59-68, 2022 (SCI-Expanded)
28. **An international cohort study spanning five decades assessed outcomes of nephropathic cystinosis**
Emma F, van't Hoff W, Hohenfellner K, TOPALOĞLU R, Greco M, Ariceta G, Bettini C, Bockenbauer D, Veys K, Pape L, et al.
KIDNEY INTERNATIONAL, vol.100, no.5, pp.1112-1123, 2021 (SCI-Expanded)
29. **Could plasma based therapies still be considered in selected cases with atypical hemolytic uremic syndrome?**
ÖZLÜ S. G., GÜLHAN B., Aydog O, Atayar E, Delibas A, Parmaksiz G, Ozdogan E. B., ÇOMAK E., Tasdemir M., Acar B., et al.
TURKISH JOURNAL OF PEDIATRICS, vol.63, no.6, pp.986-993, 2021 (SCI-Expanded)
30. **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)
31. **CLINICAL AND MUTATIONAL SPECTRUM OF CHILDREN WITH AUTOSOMAL RECESSIVE AND AUTOSOMAL DOMINANT POLYCYSTIC KIDNEY DISEASE**
Tutal O, GÜLHAN B, Atayar E, YÜKSEL S, Ozcakar B, Soylemezoglu O, Saygili S, Inozu M, Baskin E, DÜZOVA A, et al.
PEDIATRIC NEPHROLOGY, vol.36, no.10, pp.3421-3422, 2021 (SCI-Expanded)
32. **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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33. **DGKE MUTATION IN A CHILD TREATED WITH ECULIZUMAB**

- Tekcan D., Nalcacioglu H., Karadag S. I. K., Onal H. G., ÖZALTIN F., AYDOĞ Ö.
PEDIATRIC NEPHROLOGY, vol.36, no.10, pp.3403, 2021 (SCI-Expanded)
34. **AGTR1-RELATED RENAL TUBULAR DYSGENESES MAY NOT BE FATAL**
Demirgan E. B., Saygili S., CANPOLAT N., Sever L., Kilicaslan I., Taylan D., ÇALIŞKAN S., ÖZALTIN F.
PEDIATRIC NEPHROLOGY, vol.36, no.10, pp.3455, 2021 (SCI-Expanded)
35. **INFANTILE NEPHROTIC SYNDROME WITH PLCE1 MUTATION; TREATMENT OF CYCLOSPORINE-3-YEAR FOLLOW-UP**
Tekcan D., Nalcacioglu H., Onal H. G., ÖZALTIN F., AYDOĞ Ö.
PEDIATRIC NEPHROLOGY, vol.36, no.10, pp.3475-3476, 2021 (SCI-Expanded)
36. **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.
PEDIATRIC NEPHROLOGY, vol.36, no.10, pp.3434, 2021 (SCI-Expanded)
37. **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)
38. **A HOMOZYGOUS HOXA11 VARIATION AS A POTENTIAL NOVEL CAUSE OF AUTOSOMAL RECESSIVE CONGENITAL ANOMALIES OF THE KIDNEY AND URINARY TRACT**
Saygili S., Atayar E., CANPOLAT N., ELİÇEVİK M., KURUĞOĞLU S., Sever L., ÇALIŞKAN S., ÖZALTIN F.
PEDIATRIC NEPHROLOGY, vol.36, no.10, pp.3404-3405, 2021 (SCI-Expanded)
39. **NEPHROTIC SYNDROME WITH MUTATION IN SPHINGOSINE-1-PHOSPHATE LYASE: 6 CASES**
TAŞTEMEL ÖZTÜRK T., CANPOLAT N., Saygili S., BAYRAKCI U. S., Soylemezoglu O., ÖZALTIN F., TOPALOĞLU R.
PEDIATRIC NEPHROLOGY, vol.36, no.10, pp.3476, 2021 (SCI-Expanded)
40. **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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41. **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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42. **Outcome of diacylglycerol kinase epsilon-mediated hemolytic uremic syndrome in an infant**
KOYUN M., KAYA AKSOY G., ÇOMAK E., ÖZALTIN F., AKMAN S.
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43. **CLINICAL AND MUTATIONAL SPECTRUM OF CHILDREN WITH AUTOSOMAL RECESSIVE AND AUTOSOMAL DOMINANT POLYCYSTIC KIDNEY DISEASE**
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44. **Transplantation in pediatric aHUS within the era of eculizumab therapy**
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45. **EARLY URINARY SYSTEM COMPLICATIONS IN CHILDREN WITH HEMATOPOETIC STEM CELL TRANSPLANTATION**
Gurbanov A., GÜLHAN B., KUŞKONMAZ B. B., OKUR F. V., ÇETİNKAYA F. D., Ozdemir G., TAŞ N., Celegen K., ÖZALTIN F., DÜZOVA A., et al.
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46. **Management of congenital nephrotic syndrome: consensus recommendations of the ERKNet-ESPN Working Group**
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47. **AGTR1-related Renal Tubular Dysgeneses May Not Be Fatal**
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48. **Predictors for the use of herbal and dietary supplements in children and adolescents with kidney and urinary tract diseases**
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49. **Acute kidney injury in a patient with COVID-19: Answers**
TAŞTEMEL ÖZTÜRK T., BALTU D., KURT ŞÜKÜR E. D., ÖZSÜREKÇİ Y., Gucer S., BAŞARAN H. Ö., GÜLHAN B., ÖZALTIN F., DÜZOVA A., TOPALOĞLU R.
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51. **Determinants of outcomes in chronic pediatric peritoneal dialysis: a single center experience**
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52. **Genetic aspects of congenital nephrotic syndrome: a consensus statement from the ERKNet-ESPN inherited glomerulopathy working group**
Lipska-Zietkiewicz B. S., ÖZALTIN F., Holtta T., Bockenhauer D., Berody S., Levtchenko E., Vivarelli M., Webb H., Haffner D., Schaefer F., et al.
EUROPEAN JOURNAL OF HUMAN GENETICS, vol.28, no.10, pp.1368-1378, 2020 (SCI-Expanded)
53. **Renal Biopsy Prognostic Findings in Children With Atypical Hemolytic Uremic Syndrome**
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54. **COL4A3 mutation is an independent risk factor for poor prognosis in children with Alport syndrome**
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55. **A homozygousHOXA11variation as a potential novel cause of autosomal recessive congenital anomalies of the kidney and urinary tract**
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56. **Clinical characteristics of children with congenital anomalies of the kidney and urinary tract and predictive factors of chronic kidney disease**
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57. **Cystinosis beyond kidneys: gastrointestinal system and muscle involvement**
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58. **CD80 expression and infiltrating regulatory T cells in idiopathic nephrotic syndrome of childhood**
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1. **CLINICAL CHARACTERISTICS AND OUTCOME OF BK VIRUS INFECTION IN PEDIATRIC RENAL TRANSPLANT RECIPIENTS**

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9. **CLINICAL AND PATHOLOGIC CHARACTERISTICS OF GENETICALLY CONFIRMED ALPORT SYNDROME PATIENTS**
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10. **TURKISH ATYPICAL HEMOLYTIC UREMIC SYNDROME REGISTRY: ECULIZUMAB TREATMENT IN AHUS PATIENTS**
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11. **MYCETOMA OF THE URINARY TRACT IN AN INFANTWITH HORSESHOE KIDNEY: USEFULNESS OFMECHANICAL REMOVAL AND AMPHOTERICIN-BIRRIGATION**
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12. **Lupus nephritis: Long term follow-up and effect of treatment on growth**
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13. **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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14. **Clinical and Mutational Spectrum of Children with Autosomal Recessive and Autosomal Dominant**

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15. **ACUTE HEMODIALYSIS EXPERIENCE IN CHILDREN WEIGHING LESS THAN 15 KG**
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16. **Effect of IGF-1 and HGF Induced Bone Marrow Mesenchymal Stem Cells on Focal Segmental Glomerulosclerosis in Sprague-Dawley Rats: A pilot study**
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18. Ulusal Vasküler ve Endovasküler Cerrahi Kongresi 9. Ulusal Fleboloji Kongresi, Girne, Cyprus (Kktc), 9 - 12 December 2017
20. **Çocuk ve genç erişkinlerde renovasküler hipertansiyonda cerrahi tedaviler**
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22. **USE OF HERBAL AND DIETARY SUPPLEMENTS IN CHILDREN WITH KIDNEY AND URINARY TRACT DISEASE**
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25. **Does C.3979 G A/P.VAL1327MET variant of COL4A4 has any pathogenic effect in Turkish patients with Alport Syndrome ?**
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30. **NEXT GENERATION GENE PANEL SCREENING IN STEROID-RESISTANT NEPHROTIC SYNDROME**
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32. **NORMAL 25-HYDROXYVITAMIN D LEVELS ARE ASSOCIATED WITH LESS PROTEINURIA AND ATTENUATE RENAL FAILURE PROGRESSION IN CHILDREN WITH CHRONIC KIDNEY DISEASE**
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23. ULUSAL NEONATOLOJİ KONGRESİ, Antalya, Turkey, 19 - 22 April 2015
34. **CONVERSION TO SIROLIMUS IN PEDIATRIC RENAL TRANSPLANT RECIPIENTS**
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Courses

- SODYUM VE SU DENGESİ VE BOZUKLUKLARI (DÖNEM III, TÜRKÇE TIP), Associate Degree, 2016 - 2017
 AKUT BÖBREK YETMEZLİĞİ (DÖNEM III, TÜRKÇE TIP), Associate Degree, 2016 - 2017
 SIVI-ELEKTROLİT (DÖNEM IV, PDÖ), Associate Degree, 2016 - 2017
 DEHİDRATASYON VE TEDAVİSİ (DÖNEM IV, TÜRKÇE TIP), Associate Degree, 2016 - 2017
 SODYUM VE SU DENGESİ BOZUKLUKLARI (DÖNEM III, İNGİLİZCE TIP), Associate Degree, 2016 - 2017
 AKUT VE KRONİK BÖBREK YETMEZLİĞİNE YAKLAŞIM (DÖNEM IV), Associate Degree, 2016 - 2017

POTASYUM DENGESİ VE BOZUKLUKLARI (DÖNEM III, TÜRKÇE TIP), Associate Degree, 2016 - 2017
Akut Böbrek Yetmezliği, Undergraduate, 2016 - 2017
EKSTREMİTE MUAYENESİ VE KAN BASINCI ÖLÇÜMÜ (DÖNEM IV), Associate Degree, 2016 - 2017
DEHİDRATASYON VE TEDAVİSİ (DÖNEM IV, İNGİLİZCE TIP), Associate Degree, 2016 - 2017
POTASYUM DENGESİ VE BOZUKLUKLARI (DÖNEM III, İNGİLİZCE TIP), Associate Degree, 2016 - 2017
İDRAR KAÇIRAN ÇOCUĞA YAKLAŞIM, Associate Degree, 2016 - 2017

Advising Theses

Özaltın F., Çocukluk çağı otozomal dominant ve otozomal resesif polikistik böbrek hastalıklarının klinik ve genetik özelliklerinin araştırılması, Expertise In Medicine, Ö.TUTAL(Student), 2021
Özaltın F., ADCK4 mutasyonu saptanan hastaların uzun dönem izlem sonuçları, Expertise In Medicine, M.ATMACA(Student), 2017

Metrics

Publication: 232
Citation (WoS): 3571
Citation (Scopus): 4053
H-Index (WoS): 31
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Research Areas

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

Non Academic Experience

Hacettepe Üniversitesi Tıp Fakültesi