

Prof. FATİH ÖZALTIN

Personal Information

Office Phone: [+90 312 305 2645](tel:+903123052645)

Office Phone: [+90 312 305 1246](tel:+903123051246)

Email: fozaltin@hacettepe.edu.tr

Web: <https://www.fatihozaltin.com>

Education Information

Post Doctorate, The University Of Michigan, Medical School, Pediatric Nephrology, United States Of America 2004 - 2005

Doctorate, Hacettepe Üniversitesi, Tıp Fakültesi, Çocuk Nefrolojisi, Turkey 2000 - 2002

Expertise In Medicine, Hacettepe Üniversitesi, Tıp Fakültesi, Çocuk Sağlığı Ve Hastalıkları, Turkey 1994 - 2000

Under Graduate, Ege Üniversitesi, Tıp Fakültesi, Turkey 1988 - 1994

Foreign Languages

English, B2 Upper Intermediate

Research Areas

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

Academic Titles / Tasks

Associate Professor, Hacettepe Üniversitesi, Tıp Fakültesi (ingilizce), Dahili Tıp Bilimleri Bölümü, 2006 - Continues

Assistant Professor, Hacettepe Üniversitesi, Tıp Fakültesi (ingilizce), Dahili Tıp Bilimleri Bölümü, 2005 - 2006

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, Under Graduate, 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

Articles Published in Journals That Entered SCI, SSCI and AHCI Indexes

- I. **Management of congenital nephrotic syndrome: consensus recommendations of the ERKNet-ESPN Working Group**
Boyer O., Schaefer F., Haffner D., Bockenbauer D., Holttä T., Berody S., Webb H., Heselden M., Lipska-Zietkiewicz B. S., ÖZALTIN F., et al.
NATURE REVIEWS NEPHROLOGY, 2021 (Journal Indexed in SCI)
- II. **Determinants of outcomes in chronic pediatric peritoneal dialysis: a single center experience**
Asi T., DÜZÖVA A., DOĞAN H. S. , Karakurt G., BAHADIR Ö. F. , Bozaci A. C. , GÜLHAN B., ÖZALTIN F., AKI F. T. , TEKGÜL S., et al.
TURKISH JOURNAL OF PEDIATRICS, vol.62, no.6, pp.940-948, 2020 (Journal Indexed in SCI)
- III. **Transplantation in pediatric aHUS within the era of eculizumab therapy**
ÖZÇAKAR Z. B. , ÖZALTIN F., GÜLHAN B., ÇOMAK E., Parmaksiz G., Baskin E., TOPALOĞLU R., Kasap Demir B., Canpolat N., Yuruk Yildirim Z., et al.
PEDIATRIC TRANSPLANTATION, 2020 (Journal Indexed in SCI)
- IV. **Genetic aspects of congenital nephrotic syndrome: a consensus statement from the ERKNet-ESPN inherited glomerulopathy working group**
Lipska-Zietkiewicz B. S. , ÖZALTIN F., Holttä T., Bockenbauer D., Berody S., Levchenko E., Vivarelli M., Webb H., Haffner D., Schaefer F., et al.
EUROPEAN JOURNAL OF HUMAN GENETICS, vol.28, no.10, pp.1368-1378, 2020 (Journal Indexed in SCI)
- V. **A homozygousHOXA11variation 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., Caliskan S., ÖZALTIN F.
CLINICAL GENETICS, vol.98, no.4, pp.390-395, 2020 (Journal Indexed in SCI)
- VI. **Renal Biopsy Prognostic Findings in Children With Atypical Hemolytic Uremic Syndrome**
YÜKSEL S., GÖNÜL İ. I. , Canpolat N., GÖKCE İ., Ozlu S. G. , ÖZÇAKAR Z. B. , ÖZALTIN F., SÖYLEMEZOĞLU O.
PEDIATRIC AND DEVELOPMENTAL PATHOLOGY, vol.23, no.5, pp.362-371, 2020 (Journal Indexed in SCI)
- VII. **Clinical characteristics of children with congenital anomalies of the kidney and urinary tract and predictive factors of chronic kidney disease**
Cetinkaya P. G. , GÜLHAN B., DÜZÖVA A., Besbas N., HAYRAN K. M. , TOPALOĞLU R., ÖZALTIN F.
TURKISH JOURNAL OF PEDIATRICS, vol.62, no.5, pp.746-755, 2020 (Journal Indexed in SCI)
- VIII. **Predictors for the use of herbal and dietary supplements in children and adolescents with kidney and urinary tract diseases**
Ozturk T. T. , KANBUR N., ÖZMERT E. N. , GÜLHAN B., ÖZALTIN F., TOPALOĞLU R., DÜZÖVA A.
EUROPEAN JOURNAL OF PEDIATRICS, 2020 (Journal Indexed in SCI)
- IX. **Cystinosis beyond kidneys: gastrointestinal system and muscle involvement**
TOPALOĞLU R., GÜLTEKİNGİL KESER A., GÜLHAN B., ÖZALTIN F., DEMİR H., Ciftci T., DEMİR N., TEMUÇİN Ç. M. , YÜCE A., Akhan O.
BMC GASTROENTEROLOGY, vol.20, no.1, 2020 (Journal Indexed in SCI)
- X. **CD80 expression and infiltrating regulatory T cells in idiopathic nephrotic syndrome of childhood**
Eroglu F. K. , ORHAN D., Inozu M., DÜZÖVA A., GÜLHAN B., ÖZALTIN F., TOPALOĞLU R.
PEDIATRICS INTERNATIONAL, vol.61, no.12, pp.1250-1256, 2019 (Journal Indexed in SCI)
- XI. **Surgical management of renovascular hypertension in children and young adults: a 13-year experience**
Peker O., AKI F. T. , Kumbasar U., GÜVENER M., YILMAZ M., DOĞAN R., ÖZALTIN F., DÜZÖVA A., TOPALOĞLU R., PEYNİRCİOĞLU B., et al.
INTERACTIVE CARDIOVASCULAR AND THORACIC SURGERY, vol.29, no.5, pp.746-752, 2019 (Journal Indexed in SCI)
- 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**
ATMACA M., GÜLHAN B., Atayar E., KARABAY BAYAZIT A., CANDAN C., ARICI M., TOPALOĞLU R., ÖZALTIN F.
TURKISH JOURNAL OF PEDIATRICS, vol.61, no.5, pp.657-663, 2019 (Journal Indexed in SCI)

- XIII. **Rituximab for Children With Difficult-to-Treat Nephrotic Syndrome: Its Effects on Disease Progression and Growth**
TOPALOĞLU R., GÜLHAN B., Celegen K., Inozu M., Hayran M., Duzoya A., ÖZALTIN F.
FRONTIERS IN PEDIATRICS, vol.7, 2019 (Journal Indexed in SCI)
- XIV. **Persistent hypoglycemic attacks during hemodialysis sessions in an infant with congenital nephrotic syndrome: Questions and Answers**
Saygili S., CANPOLAT N., Sever L., ÇALIŞKAN S., Atayar E., ÖZALTIN F.
PEDIATRIC NEPHROLOGY, vol.34, no.1, pp.75-79, 2019 (Journal Indexed in SCI)
- XV. **An immunohistochemical approach to detect oncogenic CTNNB1 mutations in primary neoplastic tissues**
AKYOL A., GUNER G., OZSEKER H. S. , IŞIK A., Atci O., UZUN S., ATAYAR E., ÖZALTIN F., GEDIKOĞLU G., SÖKMENSÜER C., et al.
LABORATORY INVESTIGATION, vol.99, no.1, pp.128-137, 2019 (Journal Indexed in SCI)
- XVI. **BK virus associated nephropathy and severe pneumonia in a kidney transplanted adolescent with Schimke immune-osseous-dysplasia**
DÜZOVA A., GÜLHAN B., TOPALOĞLU R., ÖZALTIN F., CENGİZ A. B. , Yetimakman A. F. , Dogru D., Gucer S., Besbas N.
TURKISH JOURNAL OF PEDIATRICS, vol.61, no.1, pp.111-116, 2019 (Journal Indexed in SCI)
- XVII. **Response to Early Coenzyme Q10 Supplementation Is not Sustained in CoQ10 Deficiency Caused by CoQ2 Mutation**
Eroglu F. K. , ÖZALTIN F., Gonc N., Nalcacioglu H., Ozcakar Z. B. , YALNIZOĞLU D., Gucer S., ORHAN D., Eminoglu F. T. , GÖÇMEN R., et al.
PEDIATRIC NEUROLOGY, vol.88, pp.71-74, 2018 (Journal Indexed in SCI)
- XVIII. **MYCETOMA OF THE URINARY TRACT IN AN INFANT WITH HORSESHOE KIDNEY: USEFULNESS OF MECHANICAL REMOVAL AND AMPHOTERICIN-B IRRIGATION**
OĞUZ B., Celegen K., ÇİFTÇİ A. Ö. , ÜNAL E., ŞENOCAK M. E. , ÖZKALE YAVUZ Ö. , TAŞ N., GÜLHAN B., ÖZALTIN F., TOPALOĞLU R., et al.
PEDIATRIC NEPHROLOGY, vol.33, no.10, pp.1984, 2018 (Journal Indexed in SCI)
- 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.
PEDIATRIC NEPHROLOGY, vol.33, no.10, pp.1845, 2018 (Journal Indexed in SCI)
- 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**
ATMACA M., GÜLHAN B., KARABAY BAYAZIT A., CANDAN C., ARICI M., TOPALOĞLU R., ÖZALTIN F.
PEDIATRIC NEPHROLOGY, vol.33, no.10, pp.1821-1822, 2018 (Journal Indexed in SCI)
- XXI. **Familial Mediterranean fever patients homozygous for E148Q variant may have milder disease**
TOPALOĞLU R., BATU AKAL E. D. , Yildiz C., Korkmaz E., ÖZEN S., BEŞBAŞ N., ÖZALTIN F.
INTERNATIONAL JOURNAL OF RHEUMATIC DISEASES, vol.21, no.10, pp.1857-1862, 2018 (Journal Indexed in SCI)
- XXII. **TURKISH ATYPICAL HEMOLYTIC UREMIC SYNDROME REGISTRY: ECULIZUMAB TREATMENT IN AHUS PATIENTS**
Baskin E., Canpolat N., Gulleroglu K., Yilmaz A., MELEK E., Yuksel S., GÜLHAN B., KALYONCU M., Parmaksiz G., Ozcakar B., et al.
PEDIATRIC NEPHROLOGY, vol.33, no.10, pp.1832, 2018 (Journal Indexed in SCI)
- XXIII. **CLINICAL AND PATHOLOGIC CHARACTERISTICS OF GENETICALLY CONFIRMED ALPORT SYNDROME PATIENTS**
TOPALOĞLU R., Ozdemir G., GÜLHAN B., Atayar E., DÜZOVA A., SOYLU A., Ozcakar B., SÖYLEMEZOĞLU O., ÖZALTIN F.
PEDIATRIC NEPHROLOGY, vol.33, no.10, pp.1826-1827, 2018 (Journal Indexed in SCI)
- XXIV. **LUPUS NEPHRITIS: LONG TERM FOLLOW-UP AND EFFECT OF TREATMENT ON GROWTH**
TOPALOĞLU R., TAŞ N., Celegen K., Ozdemir G., GÜLHAN B., DÜZOVA A., ÖZALTIN F., ORHAN D.

- PEDIATRIC NEPHROLOGY, vol.33, no.10, pp.1947, 2018 (Journal Indexed in SCI)
- XXV. **ACUTE HEMODIALYSIS EXPERIENCE IN CHILDREN WEIGHING LESS THAN 15 KG**
TAŞ N., GÜLHAN B., Celegen K., Ozdemir G., DÜZOVA A., ÖZALTIN F., TOPALOĞLU R.
PEDIATRIC NEPHROLOGY, vol.33, no.10, pp.1893, 2018 (Journal Indexed in SCI)
- XXVI. **GASTROINTESTINAL SYSTEM INVOLVEMENT IN ATYPICAL HEMOLYTIC UREMIC SYNDROME**
Fidan K., Yildirim Z. Y., Goknar N., GÜLHAN B., Gulleroglu K., Ozcakar Z. B., Baskin E., Hayran M., ÖZALTIN F., Soylemezoglu O.
PEDIATRIC NEPHROLOGY, vol.33, no.10, pp.1843, 2018 (Journal Indexed in SCI)
- XXVII. **CLINICAL AND MUTATIONAL SPECTRUM OF CHILDREN WITH AUTOSOMAL RECESSIVE AND AUTOSOMAL DOMINANT POLYCYSTIC KIDNEY DISEASE**
TÜRKOĞLU Ö., GÜLHAN B., YÜKSEL S., Caliskan S., DÜZOVA A., ÇAKAR N., SÖYLEMEZOĞLU O., TOPALOĞLU R., ÖZALTIN F.
PEDIATRIC NEPHROLOGY, vol.33, no.10, pp.1814, 2018 (Journal Indexed in SCI)
- XXVIII. **ADRENAL INSUFFICIENCY IN AN INFANT WITH CONGENITAL NEPHROTIC SYNDROME: NEPHROTIC SYNDROME-14**
Saygili S., Canpolat N., Caliskan S., Atayar E., Sever L., ÖZALTIN F.
PEDIATRIC NEPHROLOGY, vol.33, no.10, pp.1929, 2018 (Journal Indexed in SCI)
- XXIX. **GASTRIC DUPLICATION CYST IN AN INFANT WITH FINNISH-TYPE CONGENITAL NEPHROTIC SYNDROME: CONCURRENCE OR COINCIDENCE?**
Eroglu T. G. F. K., Can G. G., Cakici E. K., Yazilitas F., Celikkaya E., ÖZALTIN F., Bulbul M.
PEDIATRIC NEPHROLOGY, vol.33, no.10, pp.1931-1932, 2018 (Journal Indexed in SCI)
- XXX. **Extra-Renal manifestations of atypical hemolytic uremic syndrome in children**
Fidan K., Goknar N., GÜLHAN B., MELEK E., Yildirim Z. Y., Baskin E., HAYRAN K. M., Gulleroglu K., Ozcakar Z. B., ÖZALTIN F., et al.
PEDIATRIC NEPHROLOGY, vol.33, no.8, pp.1395-1403, 2018 (Journal Indexed in SCI)
- XXXI. **Effect of IGF-1 and HGF induced bone marrow mesenchymal stem cells on focal segmental glomerulosclerosis in Sprague-Dawley rats: a pilot study**
ŞAHAN Ö., KORKMAZ E., ONBAŞILAR İ., Gucer S., KAYMAZ F. F., ÖZALTIN F., ÖZCAN A.
FEBS OPEN BIO, vol.8, pp.152, 2018 (Journal Indexed in SCI)
- XXXII. **Atypical Hemolytic Uremic Syndrome in Children Aged < 2 Years**
ÇAKAR N., ÖZÇAKAR Z. B., ÖZALTIN F., KOYUN M., ACAR B. Ç., BAHAT ÖZDOĞAN E., GÜLHAN B., Korkmaz E., Yurt A., Yilmaz S., et al.
NEPHRON, vol.139, no.3, pp.211-218, 2018 (Journal Indexed in SCI)
- XXXIII. **Nephropathic Cystinosis Mimicking Bartter Syndrome A Novel Mutation**
Bastug F., Nalcacioglu H., ÖZALTIN F., Korkmaz E., Yel S.
IRANIAN JOURNAL OF KIDNEY DISEASES, vol.12, no.1, pp.61-63, 2018 (Journal Indexed in SCI)
- XXXIV. **Mutations in KEOPS-complex genes cause nephrotic syndrome with primary microcephaly**
Braun D. A., Rao J., Mollet G., Schapiro D., Dageron M., Tan W., Gribouval O., Boyer O., Revy P., Jobst-Schwan T., et al.
NATURE GENETICS, vol.49, no.10, pp.1529-1541, 2017 (Journal Indexed in SCI)
- XXXV. **Long-Term Outcome of Steroid-Resistant Nephrotic Syndrome in Children**
Trautmann A., Schnaidt S., Lipska-Zietkiewicz B. S., Bodria M., ÖZALTIN F., Emma F., Anarat A., Melk A., Azocar M., Oh J., et al.
JOURNAL OF THE AMERICAN SOCIETY OF NEPHROLOGY, vol.28, no.10, pp.3055-3065, 2017 (Journal Indexed in SCI)
- XXXVI. **USE OF HERBAL AND DIETARY SUPPLEMENTS IN CHILDREN WITH KIDNEY AND URINARY TRACT DISEASE**
TAŞTEMEL ÖZTÜRK T., Ozmert E., KANBUR N., GÜLHAN B., ÖZALTIN F., TOPALOĞLU R., DÜZOVA A.
PEDIATRIC NEPHROLOGY, vol.32, no.9, pp.1675, 2017 (Journal Indexed in SCI)
- XXXVII. **DOES C.3979 G > A/P.VAL1327MET VARIANT OF COL4A4 HAS ANY PATHOGENIC EFFECT IN TURKISH PATIENTS WITH ALPORT SYNDROME?**

- YILDIZ N., ATA P., ALPAY H., GÖKCE İ., ÖZALTIN F.
PEDIATRIC NEPHROLOGY, vol.32, no.9, pp.1773, 2017 (Journal Indexed in SCI)
- XXXVIII. **CLINICAL CHARACTERISTICS OF PATIENTS WITH MTORI CONVERSION AND EFFECTS ON RENAL OUTCOME**
TAŞ N., GÜLHAN B., Inozu M., Celegen K., DÜZOVA A., ÖZALTIN F., TOPALOĞLU R.
PEDIATRIC NEPHROLOGY, vol.32, no.9, pp.1803, 2017 (Journal Indexed in SCI)
- XXXIX. **GENETIC AND CLINICAL CHARACTERISTICS OF PATIENTS WITH C3 GLOMERULOPATHY**
TOPALOĞLU R., GÜLHAN B., Korkmaz E., DÜZOVA A., ÖZALTIN F.
PEDIATRIC NEPHROLOGY, vol.32, no.9, pp.1669, 2017 (Journal Indexed in SCI)
- XL. **RISK FACTORS FOR CARDIOVASCULAR COMORBIDITIES IN CHILDREN WITH STEROID-SENSITIVE AND STEROID-RESISTANT NEPHROTIC SYNDROME**
Eroglu F. K. , TOPALOĞLU R., KARAGÖZ T., OĞUZ B., ÖZALTIN F., GÜLHAN B., DÜZOVA A.
PEDIATRIC NEPHROLOGY, vol.32, no.9, pp.1661, 2017 (Journal Indexed in SCI)
- XLI. **IS RAAS BLOCKADE EFFECTIVE IN NPHS2 GLOMERULOPATHY?**
Trautmann A., ÖZALTIN F., Zurowksa A., Helena H., Saeed B., Azocar M., Yilmaz A., Anarat A., Balat A., Caliskan S., et al.
PEDIATRIC NEPHROLOGY, vol.32, no.9, pp.1661, 2017 (Journal Indexed in SCI)
- XLII. **LONG TERM RESULTS OF RITUXIMAB IN A TERTIARY REFERRAL CENTER FOR DIFFICULT-TO-TREAT NEPHROTIC SYNDROME**
Celegen K., GÜLHAN B., Inozu M., TAŞ N., ÖZALTIN F., DÜZOVA A., TOPALOĞLU R.
PEDIATRIC NEPHROLOGY, vol.32, no.9, pp.1769-1770, 2017 (Journal Indexed in SCI)
- XLIII. **Follow-up results of patients with ADCK4 mutations and the efficacy of CoQ10 treatment**
Atmaca M., GÜLHAN B., KORKMAZ E., INOZU M., Soylemezoglu O., CANDAN C., Bayazit A. K. , Elmaci A. M. , Parmaksiz G., DÜZOVA A., et al.
PEDIATRIC NEPHROLOGY, vol.32, no.8, pp.1369-1375, 2017 (Journal Indexed in SCI)
- XLIV. **Low renal but high extrarenal phenotype variability in Schimke immuno-osseous dysplasia**
Lipska-Zietkiewicz B. S. , Gellermann J., Boyer O., Gribouval O., Zietkiewicz S., Kari J. A. , Shalaby M. A. , ÖZALTIN F., Dusek J., Melk A., et al.
PLOS ONE, vol.12, no.8, 2017 (Journal Indexed in SCI)
- XLV. **Tocilizumab treatment in childhood Takayasu arteritis: Case series of four patients and systematic review of the literature**
Batu E. D. , Sonmez H. E. , HAZIROLAN T., ÖZALTIN F., BİLGİNER Y., ÖZEN S.
SEMINARS IN ARTHRITIS AND RHEUMATISM, vol.46, no.4, pp.529-535, 2017 (Journal Indexed in SCI)
- XLVI. **Turkish pediatric atypical hemolytic uremic syndrome registry: initial analysis of 146 patients**
BESBAS N., GULHAN B., SOYLEMEZOGLU O., OZCAKAR Z. B. , KORKMAZ E., HAYRAN M., Ozaltin F.
BMC NEPHROLOGY, vol.18, 2017 (Journal Indexed in SCI)
- XLVII. **Timing of renal replacement therapy does not influence survival and growth in children with congenital nephrotic syndrome caused by mutations in NPHS1: data from the ESPN/ERA-EDTA Registry**
Holta T., Bonthuis M., Van Stralen K. J. , Bjerre A., TOPALOĞLU R., ÖZALTIN F., Holmberg C., Harambat J., Jager K. J. , Schaefer F., et al.
PEDIATRIC NEPHROLOGY, vol.31, no.12, pp.2317-2325, 2016 (Journal Indexed in SCI)
- XLVIII. **First-Line, Early and Long-Term Eculizumab Therapy in Atypical Hemolytic Uremic Syndrome: A Case Series in Pediatric Patients**
YÜKSEL S., Evrengul H., ÖZÇAKAR Z. B. , Becerir T., Yalcin N., Korkmaz E., ÖZALTIN F.
PEDIATRIC DRUGS, vol.18, no.6, pp.413-420, 2016 (Journal Indexed in SCI)
- XLIX. **Genetic and clinical features of cryopyrin-associated periodic syndromes in Turkish children**
EROGLU F. K. , Kasapcopur O., BESBAS N., ÖZALTIN F., BİLGİNER Y., Barut K., MENSA-VILARO A., Nakagawa K., Heike T., Nishikomori R., et al.
CLINICAL AND EXPERIMENTAL RHEUMATOLOGY, vol.34, no.6, 2016 (Journal Indexed in SCI)
- L. **Turkish Atypical Hemolytic Uremic Syndrome Registry: Evaluation of 146 Patients**

Besbas N., Soylemezoglu O., GÜLHAN B., Ozcazar Z. B., KORKMAZ E., HAYRAN K. M., ÖZALTIN F.
PEDIATRIC NEPHROLOGY, vol.31, no.10, pp.1884, 2016 (Journal Indexed in SCI)

- LI. **Loss of diacylglycerol kinase epsilon in mice causes endothelial distress and impairs glomerular Cox-2 and PGE(2) production**
Zhu J., Chaki M., Lu D., Ren C., Wang S., Rauhauser A., Li B., Zimmerman S., Jun B., Du Y., et al.
AMERICAN JOURNAL OF PHYSIOLOGY-RENAL PHYSIOLOGY, vol.310, no.9, 2016 (Journal Indexed in SCI)
- LII. **Mutations in nuclear pore genes NUP93, NUP205 and XPO5 cause steroid-resistant nephrotic syndrome**
Braun D. A., Sadowski C. E., Kohl S., Lovric S., Astrinidis S. A., Pabst W. L., Gee H. Y., Ashraf S., Lawson J. A., Shril S., et al.
NATURE GENETICS, vol.48, no.4, pp.457-467, 2016 (Journal Indexed in SCI)
- LIII. **Whole exome sequencing identifies causative mutations in the majority of consanguineous or familial cases with childhood-onset increased renal echogenicity**
Braun D. A., Schueler M., Halbritter J., Gee H. Y., Porath J. D., Lawson J. A., Airik R., Shril S., Allen S. J., Stein D., et al.
KIDNEY INTERNATIONAL, vol.89, no.2, pp.468-475, 2016 (Journal Indexed in SCI)
- LIV. **Normal 25-Hydroxyvitamin D Levels Are Associated with Less Proteinuria and Attenuate Renal Failure Progression in Children with CKD**
Shroff R., Aitkenhead H., Costa N., Trivelli A., Litwin M., Picca S., Anarat A., Sallay P., ÖZALTIN F., Zurowska A., et al.
JOURNAL OF THE AMERICAN SOCIETY OF NEPHROLOGY, vol.27, no.1, pp.314-322, 2016 (Journal Indexed in SCI)
- LV. **ADCK4-Associated Glomerulopathy Causes Adolescence-Onset FSGS**
Korkmaz E., Lipska-Zietkiewicz B. S., Boyer O., Gribouval O., Fourrage C., Tabatabaei M., Schnaidt S., Gucer S., Kaymaz F., ARICI M., et al.
JOURNAL OF THE AMERICAN SOCIETY OF NEPHROLOGY, vol.27, no.1, pp.63-68, 2016 (Journal Indexed in SCI)
- LVI. **Lupus in a patient with cystinosis: is it drug induced?**
Eroglu F. K., BESBAS N., ÖZALTIN F., TOPALOĞLU R., ÖZEN S.
LUPUS, vol.24, no.13, pp.1452-1454, 2015 (Journal Indexed in SCI)
- LVII. **MCP1 2518 A/G polymorphism affects progression of childhood focal segmental glomerulosclerosis**
Besbas N., KALYONCU M., Cil O., ÖZGÜL R. K., Bakkaloglu A., ÖZALTIN F.
RENAL FAILURE, vol.37, no.9, pp.1435-1439, 2015 (Journal Indexed in SCI)
- LVIII. **NEPHROPATHIC CYSTINOSIS MIMICKING BARTTER SYNDROME: NOVEL MUTATION**
BaStuG F., Yel S., ÖZALTIN F., Korkmaz E., Uytun S., DÜŞÜNSEL R.
PEDIATRIC NEPHROLOGY, vol.30, no.9, pp.1681, 2015 (Journal Indexed in SCI)
- LIX. **SEVERE PNEUMONIA ASSOCIATED WITH BK VIRUS NEPHROPATHY IN A KIDNEY TRANSPLANTED ADOLESCENT WITH SCHIMKE IMMUNO-OSSEOUS-DYSPLASIA**
DÜZOVA A., GÜLHAN B., ÖZALTIN F., TOPALOĞLU R., BİLGİNER Y., BEŞBAŞ N.
PEDIATRIC NEPHROLOGY, vol.30, no.9, pp.1619-1620, 2015 (Journal Indexed in SCI)
- LX. **CHARACTERISTICS OF RENAL TRANSPLANT CHILDREN WITH CHRONIC ALLOGRAFT NEPHROPATHY: EXPERIENCE OF A TERTIARY REFERRAL CENTER**
GÜLHAN B., INOZU M., BİLGİNER Y., ÖZALTIN F., DÜZOVA A., ÖZEN S., TOPALOĞLU R., AKI F. T., BeSbaS N.
PEDIATRIC NEPHROLOGY, vol.30, no.9, pp.1721, 2015 (Journal Indexed in SCI)
- LXI. **ATYPICAL HEMOLYTIC UREMIC SYNDROME UNRESPONSIVE TO ECULIZUMAB THERAPY**
Inozu M., BODUR İ., DÜZOVA A., BİLGİNER Y., ÖZALTIN F., BEŞBAŞ N.
PEDIATRIC NEPHROLOGY, vol.30, no.9, pp.1648, 2015 (Journal Indexed in SCI)
- LXII. **RESULTS OF TURKISH MULTICENTRIC NATIONAL CYSTINOSIS REGISTRY**
TOPALOĞLU R., GÜLHAN B., ÖZALTIN F., BODUR İ., BEŞBAŞ N., Dursun H., Yilmaz A., GÜRGÖZE M. K., GÖKCE İ., Akinci N., et al.
PEDIATRIC NEPHROLOGY, vol.30, no.9, pp.1668-1669, 2015 (Journal Indexed in SCI)
- LXIII. **RITUXIMAB EXPERIENCE OF A TERTIARY REFERRAL CENTER FOR DIFFICULT-TO-TREAT NEPHROTIC SYNDROME**
GÜLHAN B., TOPALOĞLU R., ÖZALTIN F., DÜZOVA A., EROGLU F. K., Bodur I., BİLGİNER Y., ÖZEN S., BesbaS N.
PEDIATRIC NEPHROLOGY, vol.30, no.9, pp.1692, 2015 (Journal Indexed in SCI)

- LXIV. RESPONSE TO INTENSIFIED IMMUNOSUPPRESSIVE THERAPY AND IDENTIFICATION OF GENETIC DISEASE ARE HIGHLY PREDICTIVE OF LONG-TERM RENAL OUTCOME IN CHILDREN WITH STEROID RESISTANT NEPHROTIC SYNDROME (SRNS)**
Trautmann A., Schnaidt S., Ghiggeri G. M. , ÖZALTIN F., Saeed B., Drozd D., Caliskan S., Anarat A., Oh J., Bogdanovic R., et al.
PEDIATRIC NEPHROLOGY, vol.30, no.9, pp.1556-1557, 2015 (Journal Indexed in SCI)
- LXV. RISK FACTORS FOR POST-TRANSPLANT RECURRENCE OF STEROID RESISTANT NEPHROTIC SYNDROME (SRNS): RESULTS FROM THE PODONET REGISTRY**
Trautmann A., Ghiggeri G. M. , Azocar M., Remuzzi G., ÖZALTIN F., Melk A., Jankauskiene A., Peco-antic A., Gellermann J., Emma F., et al.
PEDIATRIC NEPHROLOGY, vol.30, no.9, pp.1557-1558, 2015 (Journal Indexed in SCI)
- LXVI. Genetic abnormalities and prognosis in patients with congenital and infantile nephrotic syndrome**
Cil O., Besbas N., DÜZOVA A., TOPALOĞLU R., PECO-ANTIC A., KORKMAZ E., ÖZALTIN F.
PEDIATRIC NEPHROLOGY, vol.30, no.8, pp.1279-1287, 2015 (Journal Indexed in SCI)
- LXVII. Thrombophilic Risk Factors And The Efficiency Of Prophylactic Anticoagulation Therapy In Children Who Underwent Renal Transplantation**
GÜLHAN B., TAVIL B., DÜZOVA A., ÖZALTIN F., ÖZEN S., TOPALOĞLU R., BİLGİNER Y., GÜMRÜK F., BEŞBAŞ N.
PEDIATRIC NEPHROLOGY, vol.29, no.9, pp.1694, 2014 (Journal Indexed in SCI)
- LXVIII. Mutations In Anks6 Cause A Nephronophthisis-like Phenotype With End Stage Renal Disease**
Taskiran E., Korkmaz E., Gucer S., KOŞUKCU C., Kaymaz F., Koyunlar C., Bryda E. C. , Chaki M., Lu D. D. , Vadnagara K., et al.
PEDIATRIC NEPHROLOGY, vol.29, no.9, pp.1820, 2014 (Journal Indexed in SCI)
- LXIX. A novel CFHR5 mutation associated with C3 glomerulonephritis in a Turkish girl**
BEŞBAŞ N., GÜLHAN B., GÜÇER K. Ş. , Korkmaz E., ÖZALTIN F.
JOURNAL OF NEPHROLOGY, vol.27, no.4, pp.457-460, 2014 (Journal Indexed in SCI)
- LXX. Diagnostic validity of colchicine in patients with Familial Mediterranean fever**
ÖZALTIN F., BİLGİNER Y., GÜLHAN B., Bajin I., Erdogan O., HAYRAN K. M. , Yilmaz E., ÖZEN S.
CLINICAL RHEUMATOLOGY, vol.33, no.7, pp.969-974, 2014 (Journal Indexed in SCI)
- LXXI. Mutations in EMP2 Cause Childhood-Onset Nephrotic Syndrome**
Gee H. Y. , Ashraf S., Wan X., Vega-Warner V., Esteve-Rudd J., Lovric S., Fang H., Hurd T. W. , Sadowski C. E. , Allen S. J. , et al.
AMERICAN JOURNAL OF HUMAN GENETICS, vol.94, no.6, pp.884-890, 2014 (Journal Indexed in SCI)
- LXXII. Post-transplant hypertension in pediatric kidney transplant recipients**
GÜLHAN B., TOPALOĞLU R., KARABULUT E., ÖZALTIN F., AKI F. T. , BİLGİNER Y., Besbas N.
PEDIATRIC NEPHROLOGY, vol.29, no.6, pp.1075-1080, 2014 (Journal Indexed in SCI)
- LXXIII. Primary coenzyme Q(10) (CoQ(10)) deficiencies and related nephropathies**
ÖZALTIN F.
PEDIATRIC NEPHROLOGY, vol.29, no.6, pp.961-969, 2014 (Journal Indexed in SCI)
- LXXIV. Genotype-phenotype associations in WT1 glomerulopathy**
Lipska-Zietkiewicz B. S. , Ranchin B., Iatropoulos P., Gellermann J., Melk A., ÖZALTIN F., Caridi G., Seeman T., Tory K., Jankauskiene A., et al.
KIDNEY INTERNATIONAL, vol.85, no.5, pp.1169-1178, 2014 (Journal Indexed in SCI)
- LXXV. NPHS2 Mutations in Steroid-Resistant Nephrotic Syndrome: A Mutation Update and the Associated Phenotypic Spectrum**
Bouchireb K., Boyer O., Gribouval O., Nevo F., Huynh-Cong E., Moriniere V., Campait R., Ars E., Brackman D., Dantal J., et al.
HUMAN MUTATION, vol.35, no.2, pp.178-186, 2014 (Journal Indexed in SCI)
- LXXVI. Proteinuria in Frasier Syndrome**
Peco-Antic A., ÖZALTIN F., Parezanovic V., Milosevski-Lomic G., Zdravkovic V.
SRPSKI ARHIV ZA CELOKUPNO LEKARSTVO, vol.141, pp.685-688, 2013 (Journal Indexed in SCI)
- LXXVII. The Effect Of Fibroblast Growth Factor 23 On Left Ventricular Function In Peritoneal Dialysis**

Children

Hacihamdioglu D. O. , BEŞBAŞ N., ALEHAN D., OĞUZ B., ÖZALTIN F., DÜZOVA A., ÖZEN S., TOPALOĞLU R., Gök F.
PEDIATRIC NEPHROLOGY, vol.28, no.8, pp.1643, 2013 (Journal Indexed in SCI)

LXXVIII. **CONTRIBUTORY RISK FACTORS FOR DEVELOPMENT OF THROMBOSIS IN CHILDREN WITH NEPHROTIC SYNDROME**

KaraEroglu F., Tavil B., ÖZALTIN F., BEŞBAŞ N., ÖZEN S., ÇETİN M., GÜMRÜK F., TOPALOĞLU R.
PEDIATRIC NEPHROLOGY, vol.28, no.8, pp.1584, 2013 (Journal Indexed in SCI)

LXXIX. **Genetic screening in adolescents with steroid-resistant nephrotic syndrome**

Lipska-Zietkiewicz B. S. , Iatropoulos P., Maranta R., Caridi G., ÖZALTIN F., Anarat A., Balat A., Gellermann J., Trautmann A., Erdogan O., et al.
KIDNEY INTERNATIONAL, vol.84, no.1, pp.206-213, 2013 (Journal Indexed in SCI)

LXXX. **DGKE Variants Cause a Glomerular Microangiopathy That Mimics Membranoproliferative GN**

ÖZALTIN F., Li B., Rauhauser A., An S., Soylemezoglu O., Gonul I. I. , Taskiran E. Z. , Ibsirlioglu T., Korkmaz E., BİLGİNER Y., et al.

JOURNAL OF THE AMERICAN SOCIETY OF NEPHROLOGY, vol.24, no.3, pp.377-384, 2013 (Journal Indexed in SCI)

LXXXI. **Neonatal onset atypical hemolytic uremic syndrome successfully treated with eculizumab**

Besbas N., Gulhan B., Karpman D., Topaloglu R., Duzova A., Korkmaz E., ÖZALTIN F.
PEDIATRIC NEPHROLOGY, vol.28, no.1, pp.155-158, 2013 (Journal Indexed in SCI)

LXXXII. **Circulating suPAR in Two Cohorts of Primary FSGS**

Wei C., Trachtman H., Li J., Dong C., Friedman A. L. , Gassman J. J. , McMahan J. L. , Radeva M., Heil K. M. , Trautmann A., et al.

JOURNAL OF THE AMERICAN SOCIETY OF NEPHROLOGY, vol.23, no.12, pp.2051-2059, 2012 (Journal Indexed in SCI)

LXXXIII. **WT1 screening in nephrotic syndrome - lessons from PodoNet**

Lipska-Zietkiewicz B. S. , Iatropoulos P., Ranchin B., Dusek J., ÖZALTIN F., Melk A., Gellermann J., Jankauskiene A., Peco-Antic A., Zurowska A., et al.

PEDIATRIC NEPHROLOGY, vol.27, no.9, pp.1617-1618, 2012 (Journal Indexed in SCI)

LXXXIV. **PERITONEAL DIALYSIS IN CHILDREN UNDER TWO YEARS OF AGE**

GÜLHAN B., DÜZOVA A., ÖZALTIN F., TOPALOĞLU R., ÖZEN S., BİLGİNER Y., Tayfur A. C. , Yildiz C., BEŞBAŞ N.
PEDIATRIC NEPHROLOGY, vol.27, no.9, pp.1799-1800, 2012 (Journal Indexed in SCI)

LXXXV. **Rituximab in steroid-dependent and resistant nephrotic syndrome patients**

TOPALOĞLU R., GÜLHAN B., ÖZALTIN F., DÜZOVA A., BEŞBAŞ N.

PEDIATRIC NEPHROLOGY, vol.27, no.9, pp.1710-1711, 2012 (Journal Indexed in SCI)

LXXXVI. **ATYPICAL HUS AND ECULIZUMAB TREATMENT: EXPERIENCE OF A TERTIARY CENTER**

BEŞBAŞ N., GÜLHAN B., ÖZEN S., TOPALOĞLU R., DÜZOVA A., Yildiz C., ÖZALTIN F.

PEDIATRIC NEPHROLOGY, vol.27, no.9, pp.1630, 2012 (Journal Indexed in SCI)

LXXXVII. **Takayasu arteritis in a 4-year-old girl: case report and brief overview of the pediatric literature**

Aypar E., Celebi-Tayfur A., Keser M., Odabas D., ÖZALTIN F., PAKSOY Y., ÖZEN S.

TURKISH JOURNAL OF PEDIATRICS, vol.54, no.5, pp.536-539, 2012 (Journal Indexed in SCI)

LXXXVIII. **Response to Intensified Immunosuppressive Therapy Predicts Long-Term Prognosis in Steroid Resistant Nephrotic Syndrome (SRNS)**

Trautmann A., ÖZALTIN F., Bodria M., Anarat A., Saeed B., Azocar M., Gellermann J., Drozd D., Bogdanovic R., Krmar R., et al.

PEDIATRIC NEPHROLOGY, vol.27, no.9, pp.1639-1640, 2012 (Journal Indexed in SCI)

LXXXIX. **THE BENEFITS OF CYCLOSPORINE TREATMENT OF THE PATIENT WITH NPHS2 MUTATION**

Hacihamdioglu D. O. , ÖZALTIN F., Zeybek C., Kalman S., Demirkaya E., Gök F.

PEDIATRIC NEPHROLOGY, vol.27, no.9, pp.1703-1704, 2012 (Journal Indexed in SCI)

XC. **C1q deficiency: identification of a novel missense mutation and treatment with fresh frozen plasma**

TOPALOĞLU R., TASKIRAN E. Z. , TAN Ç., Erman B., ÖZALTIN F., SANAL O.

CLINICAL RHEUMATOLOGY, vol.31, no.7, pp.1123-1126, 2012 (Journal Indexed in SCI)

XCI. **Genetic basis of cystinosis in Turkish patients: a single-center experience**

- TOPALOĞLU R., Vilboux T., COŞKUN T., ÖZALTIN F., Tinloy B., Gunay-Aygun M., BAKKALOĞLU A., BEŞBAŞ N., van den Heuvel L., Kleta R., et al.
PEDIATRIC NEPHROLOGY, vol.27, no.1, pp.115-121, 2012 (Journal Indexed in SCI)
- XCII. **NPHS2 gene mutation in an Iranian family with familial steroid-resistant nephrotic syndrome**
Ameli S., Mazaheri M., Zare-Shahabadi A., ÖZALTIN F., Asgarian F., Monajemzadeh M., Bazargani B., Ataei N., Hajezadeh N., Madani A., et al.
NEFROLOGIA, vol.32, no.5, pp.674-676, 2012 (Journal Indexed in SCI)
- XCIII. **OUTCOME OF NEPHROPATHIC CYSTINOSIS IN PEDIATRIC RENAL TRANSPLANTATION: A SINGLE CENTRE EXPERIENCE**
TOPALOĞLU R., BİLGİNER Y., ÖZALTIN F., DÜZOVA A., Aki T., ÖZEN S., BAKKALOĞLU A., BEŞBAŞ N.
PEDIATRIC TRANSPLANTATION, vol.15, pp.82, 2011 (Journal Indexed in SCI)
- XCIV. **Disruption of PTPRO Causes Childhood-Onset Nephrotic Syndrome**
ÖZALTIN F., Ibsirlioglu T., Taskiran E. Z. , Baydar D. E. , Kaymaz F., Buyukcelik M., Kilic B. D. , Balat A., Iatropoulos P., Asan E., et al.
AMERICAN JOURNAL OF HUMAN GENETICS, vol.89, no.1, pp.139-147, 2011 (Journal Indexed in SCI)
- XCv. **MYO1E Mutations and Childhood Familial Focal Segmental Glomerulosclerosis**
Mele C., Iatropoulos P., Donadelli R., Calabria A., Maranta R., Cassis P., Buelli S., Tomasoni S., Piras R., Krendel M., et al.
NEW ENGLAND JOURNAL OF MEDICINE, vol.365, no.4, pp.295-306, 2011 (Journal Indexed in SCI)
- XCVI. **Respiratory-chain deficiency presenting as diffuse mesangial sclerosis with NPHS3 mutation**
Baskin E., Bayrakci U. S. , Alehan F., Ozdemir H., Oner A., Horvath R., Vega-Warner V., Hildebrandt F., ÖZALTIN F.
PEDIATRIC NEPHROLOGY, vol.26, no.7, pp.1157-1161, 2011 (Journal Indexed in SCI)
- XCvII. **COQ6 mutations in human patients produce nephrotic syndrome with sensorineural deafness**
Heeringa S. F. , Chernin G., Chaki M., Zhou W., Sloan A. J. , Ji Z., Xie L. X. , Salviati L., Hurd T. W. , Vega-Warner V., et al.
JOURNAL OF CLINICAL INVESTIGATION, vol.121, no.5, pp.2013-2024, 2011 (Journal Indexed in SCI)
- XCvIII. **Follow-Up of Patients With Juvenile Nephronophthisis After Renal Transplantation: A Single Center Experience**
Tayfur A. C. , Besbas N., BİLGİNER Y., ÖZALTIN F., DÜZOVA A., Bakkaloglu M., AKI F. T. , ÖZEN S., TOPALOĞLU R., Bakkaloglu A.
TRANSPLANTATION PROCEEDINGS, vol.43, no.3, pp.847-849, 2011 (Journal Indexed in SCI)
- XCIX. **Spondyloenchondrodysplasia with Systemic Lupus Erythematosus: a report of three cases**
BİLGİNER Y., ALANAY Y., DÜZOVA A., TOPALOĞLU R., Superti-Furga A., ÖZEN S., ÖZALTIN F., BEŞBAŞ N.
CLINICAL AND EXPERIMENTAL RHEUMATOLOGY, vol.29, no.2, pp.430, 2011 (Journal Indexed in SCI)
- C. **Three cases of a rare disease, congenital chloride diarrhea, summons up the variation in the clinical course and significance of early diagnosis and adequate treatment in the prevention of intellectual disability**
Gurakan F., Baysoy G., Wedenoja S., Uslu N., ÖZEN H., ÖZALTIN F., Hoglund P.
TURKISH JOURNAL OF PEDIATRICS, vol.53, no.2, pp.194-198, 2011 (Journal Indexed in SCI)
- CI. **Macrophage Activating Syndrome complicating Wegener Granulomatosis: treatment with plasma exchange**
Eroglu F. K. , GÜLHAN B., ÜNAL S., ÖZALTIN F., ORHAN D., ÖZEN S.
CLINICAL AND EXPERIMENTAL RHEUMATOLOGY, vol.29, no.2, pp.463, 2011 (Journal Indexed in SCI)
- CII. **The bone and mineral disorder of children undergoing chronic peritoneal dialysis**
Borzych D., Rees L., Ha I. S. , Chua A., Valles P. G. , Lipka M., Zambrano P., Ahlenstiel T., BAKKALOĞLU EZGÜ S. A. , Spizzirri A. P. , et al.
KIDNEY INTERNATIONAL, vol.78, no.12, pp.1295-1304, 2010 (Journal Indexed in SCI)
- CIII. **Neuroendocrine immune system in familial Mediterranean fever**
TOPALOĞLU R., BİLGİNER Y., ALİKAŞİFOĞLU A., ÖZALTIN F., BEŞBAŞ N., ÖZEN S., BAKKALOĞLU A.
TURKISH JOURNAL OF PEDIATRICS, vol.52, no.6, pp.588-593, 2010 (Journal Indexed in SCI)
- CIV. **Genotype/Phenotype Correlation in Nephrotic Syndrome Caused by WT1 Mutations**
Chernin G., Vega-Warner V., Schoeb D. S. , Heeringa S. F. , Ovunc B., Saisawat P., Cleper R., ÖZALTIN F., Hildebrandt

F.

CLINICAL JOURNAL OF THE AMERICAN SOCIETY OF NEPHROLOGY, vol.5, no.9, pp.1655-1662, 2010 (Journal Indexed in SCI)

- CV. **New syndrome - Situs inversus totalis with cystic dysplasia of kidneys, pancreas and bowing**
Balci S., ÖZALTIN F., Bostanoglu S.
CLINICAL DYSMORPHOLOGY, vol.19, no.3, pp.173-174, 2010 (Journal Indexed in SCI)
- CVI. **Risk factors in community-acquired urinary tract infections caused by ESBL-producing bacteria in children**
TOPALOĞLU R., Er I., Dogan B. G. , BİLGİNER Y., ÖZALTIN F., Besbas N., ÖZEN S., Bakkaloglu A., GÜR D.
PEDIATRIC NEPHROLOGY, vol.25, no.5, pp.919-925, 2010 (Journal Indexed in SCI)
- CVII. **Clinical course of primary focal segmental glomerulosclerosis (FSGS) in Turkish children: a report from the Turkish Pediatric Nephrology FSGS Study Group**
BEŞBAŞ N., ÖZALTIN F., Emre S., Anarat A., ALPAY H., BAKKALOĞLU A., Baskin E., BUYAN N., DÖNMEZ O., DÜŞÜNSEL R., et al.
TURKISH JOURNAL OF PEDIATRICS, vol.52, no.3, pp.255-261, 2010 (Journal Indexed in SCI)
- CVIII. **A Case Report of Thrombocytopenia-associated Multiple Organ Failure Secondary to Salmonella enterica Serotype Typhi Infection in a Pediatric Patient: Successful Treatment With Plasma Exchange**
Yildirim I., CEYHAN M., BAYRAKÇI B., Uysal M., Kuskonmaz B. B. , ÖZALTIN F.
THERAPEUTIC APHERESIS AND DIALYSIS, vol.14, no.2, pp.226-229, 2010 (Journal Indexed in SCI)
- CIX. **Toll-like receptors 2 and 4 cell surface expression reflects endotoxin tolerance in Henoch-Schönlein purpura.**
CANPINAR H., ÖZALTIN F., BILGINER Y., BAKKALOĞLU A., Ozen S.
The Turkish journal of pediatrics, vol.52, pp.22-7, 2010 (Journal Indexed in SCI Expanded)
- CX. **A novel CLCN7 mutation resulting in a most severe form of autosomal recessive osteopetrosis**
BEŞBAŞ N., Draaken M., Ludwig M., DEREN Ö., ORHAN D., BİLGİNER Y., ÖZALTIN F.
EUROPEAN JOURNAL OF PEDIATRICS, vol.168, no.12, pp.1449-1454, 2009 (Journal Indexed in SCI)
- CXI. **Treatment of severe Henoch-Schonlein nephritis: justifying more immunosuppression**
Altugan F. S. , ÖZEN S., Aktay-Ayaz N., Gucer S., TOPALOĞLU R., DÜZOVA A., ÖZALTIN F., Besbas N.
TURKISH JOURNAL OF PEDIATRICS, vol.51, no.6, pp.551-555, 2009 (Journal Indexed in SCI)
- CXII. **Thirteen novel NPHS1 mutations in a large cohort of children with congenital nephrotic syndrome**
Heeringa S. F. , Vlangos C. N. , Chernin G., Hinkes B., Gbadegesin R., Liu J., Hoskins B. E. , ÖZALTIN F., Hildebrandt F.
NEPHROLOGY DIALYSIS TRANSPLANTATION, vol.23, no.11, pp.3527-3533, 2008 (Journal Indexed in SCI)
- CXIII. **Evaluation of intima media thickness of the common and internal carotid arteries with inflammatory markers in familial Mediterranean fever as possible predictors for atherosclerosis**
BİLGİNER Y., ÖZALTIN F., Basaran C., DÜZOVA A., Besbas N., TOPALOĞLU R., ÖZEN S., Bakkaloglu A.
RHEUMATOLOGY INTERNATIONAL, vol.28, no.12, pp.1211-1216, 2008 (Journal Indexed in SCI)
- CXIV. **Mutations in PLCE1 are a major cause of isolated diffuse mesangial sclerosis (IDMS)**
Gbadegesin R., Hinkes B. G. , Hoskins B. E. , Vlangos C. N. , Heeringa S. F. , Liu J., Loirat C., ÖZALTIN F., Hashmi S., Ulmer F., et al.
NEPHROLOGY DIALYSIS TRANSPLANTATION, vol.23, no.4, pp.1291-1297, 2008 (Journal Indexed in SCI)
- CXV. **Eye involvement in children with primary focal segmental glomerulosclerosis**
ÖZALTIN F., Heeringa S., Poyraz C. E. , BİLGİNER Y., KADAYIFÇILAR S., BEŞBAŞ N., TOPALOĞLU R., ÖZEN S., Hildebrandt F., BAKKALOĞLU A.
PEDIATRIC NEPHROLOGY, vol.23, no.3, pp.421-427, 2008 (Journal Indexed in SCI)
- CXVI. **Right atrial thrombosis complicating renal transplantation in a child**
BİLGİNER Y., ÖZALTIN F., DÜZOVA A., Erdogan I., AKI F. T. , DEMİRCİN M., Bakkaloglu M., Bakkaloglu A.
PEDIATRIC TRANSPLANTATION, vol.12, no.2, pp.251-255, 2008 (Journal Indexed in SCI)
- CXVII. **Specific podocin mutations correlate with age of onset in steroid-resistant nephrotic syndrome**
Hinkes B., Vlangos C., Heeringa S., Mucha B., Gbadegesin R., Liu J., Hasselbacher K., ÖZALTIN F., Hildebrandt F.
JOURNAL OF THE AMERICAN SOCIETY OF NEPHROLOGY, vol.19, no.2, pp.365-371, 2008 (Journal Indexed in SCI)
- CXVIII. **Triple immunosuppression with tacrolimus in pediatric renal transplantation: Single-center**

experience

DÜZOVA A., AKI F. T. , Bakkaloglu A., Besbas N., TOPALOĞLU R., ÖZEN S., ÖZALTIN F., BİLGİNER Y., Demirkaya E., Bakkaloglu M.

TRANSPLANTATION PROCEEDINGS, vol.40, no.1, pp.132-134, 2008 (Journal Indexed in SCI)

CXIX. Cerebral sinovenous thrombosis in a child with steroid sensitive nephrotic syndrome

Isik Balci Y., Tavil B., Fidan G., Ozaltin F.

EUROPEAN JOURNAL OF PEDIATRICS, vol.166, no.7, pp.757-758, 2007 (Journal Indexed in SCI)

CXX. Nephrotic syndrome in the first year of life: Two thirds of cases are caused by mutations in 4 genes (NPHS1, NPHS2, WT1, and LAMB2)

Hinkes B. G. , Mucha B., Vlangos C. N. , Gbadegesin R., Liu J., Hasselbachera K., Hangan D., Ozaltin F., Zenker M., Hildebrandt F.

PEDIATRICS, vol.119, no.4, 2007 (Journal Indexed in SCI)

CXXI. Childhood vasculitides in Turkey: a nationwide survey

Ozen S., Bakkaloglu A., Dusunsel R., Soylemezoglu O., Ozaltin F., Poyrazoglu H., KASAPÇOPUR Ö., Ozkaya O., Yalcinkaya F., Balat A., et al.

CLINICAL RHEUMATOLOGY, vol.26, no.2, pp.196-200, 2007 (Journal Indexed in SCI)

CXXII. Novel OCRL1 mutations in patients with the phenotype of dent disease

Utsch B., Boekenkamp A., Benz M. R. , Besbas N., Doetsch J., Franke I., Freund S., Gok F., Hoppe B., Karle S., et al.

AMERICAN JOURNAL OF KIDNEY DISEASES, vol.48, no.6, pp.942-954, 2006 (Journal Indexed in SCI)

CXXIII. Positional cloning uncovers mutations in PLCE1 responsible for a nephrotic syndrome variant that may be reversible

Hinkes B., Wiggins R. C. , Gbadegesin R., Vlangos C. N. , Seelow D., Nuernberg G., Garg P., Verma R., Chaib H., Hoskins B. E. , et al.

NATURE GENETICS, vol.38, no.12, pp.1397-1405, 2006 (Journal Indexed in SCI)

CXXIV. Serum IgD concentrations in patients with ataxia telangiectasia and with selective IgA deficiency

SANAL O., Ozaltin F., TEZCAN İ., ERSOY F.

INTERNATIONAL ARCHIVES OF ALLERGY AND IMMUNOLOGY, vol.116, no.3, pp.246, 1998 (Journal Indexed in SCI)

Articles Published in Other Journals

I. Clinicopathological and immunohistological features in childhood IgA nephropathy: a single-centre experience

TOPALOĞLU R., Orhan D., BİLGİNER Y., KARABULUT E., ÖZALTIN F., DÜZOVA A., KALE G., BEŞBAŞ N.

CLINICAL KIDNEY JOURNAL, vol.6, no.2, pp.169-175, 2013 (Journal Indexed in ESCI)

Refereed Congress / Symposium Publications in Proceedings

I. CLINICAL CHARACTERISTICS OF PATIENTS WITH GENETICALLY CONFIRMED ALPORT SYNDROME

Ozdemir G., GÜLHAN B., Atayar E., Canpolat N., SÖYLEMEZOĞLU O., ÖZÇAKAR Z. B. , Eroglu F. K. , Candan C., Demir B. K. , SOYLU A., et al.

57th ERA-EDTA Congress, ELECTR NETWORK, 6 - 09 June 2020, vol.35, pp.359

II. ANALYSIS OF LATE RENAL COMPLICATIONS AND RISK FACTORS IN CHILDREN WITH HEMATOPOIETIC STEM CELL TRANSPLANTATION

Gurbanov A., GÜLHAN B., KUŞKONMAZ B. B. , OKUR F. V. , Cetinkaya D. U. , Ozdemir G., Tas N., Celegen K., ÖZALTIN F., DÜZOVA A., et al.

57th ERA-EDTA Congress, ELECTR NETWORK, 6 - 09 June 2020, vol.35, pp.2140

III. EARLY URINARY SYSTEM COMPLICATIONS IN CHILDREN WITH HEMATOPOIETIC STEM CELL TRANSPLANTATION

Gurbanov A., GÜLHAN B., KUŞKONMAZ B. B. , OKUR F. V. , Cetinkaya D. U. , Ozdemir G., Tas N., Celegen K., ÖZALTIN

F., DÜZOVA A., et al.

57th ERA-EDTA Congress, ELECTR NETWORK, 6 - 09 June 2020, vol.35, pp.2123

- IV. **Hematopoetik Kök Hücre Nakli Yapılan Çocuk Hastalarda Erken Dönem Üriner Sistem Komplikasyonları**
Gurbanov A., GÜLHAN B., KUŞKONMAZ B. B. , OKUR F. V. , Uçkan Çetinkaya D., Özdemir G., Taş N., Çelegen K., ÖZALTIN F., DÜZOVA A., et al.
10. Çocuk Nefrolojisi Kongresi, Turkey, 1 - 04 May 2019
- V. **Hematopoetik Kök Hücre Nakli Yapılan Çocuk Hastalarda Geç Dönem Renal Komplikasyonların Ve RiskFaktörlerinin Analizi**
Gurbanov A., GÜLHAN B., KUŞKONMAZ B. B. , OKUR F. V. , Uçkan Çetinkaya D., Özdemir G., TAŞ N., Çelegen K., ÖZALTIN F., DÜZOVA A., et al.
10. Çocuk Nefrolojisi Kongresi, Turkey, 1 - 04 May 2019
- VI. **Çocukluk Çağı Preemptif Ve Preemptif Olmayan Böbrek Nakli Hastalarında Nakil Sonrası İlk Bir Yıldaki GreftFonksiyonlarının Değerlendirilmesi**
TAŞ N., AKI F. T. , GÜLHAN B., ÖZALTIN F., DÜZOVA A., TEKGÜL S., TOPALOĞLU R.
10. Çocuk Nefrolojisi Kongresi, Turkey, 1 - 04 May 2019
- VII. **A REFRACTORY CLINICAL CASE OF LUPUS NEPHRITIS WHO UNDERWENT BONE MARROW TRANSPLANTATION,**
TAŞ N., DEMİR S., ÇAĞDAŞ AYVAZ D. N. , ÖZALTIN F., KUŞKONMAZ B. B. , TEZCAN F. İ. , ÖZEN S., ORHAN D., TOPALOĞLU R.
European Society for Paediatric Nephrology, 51th Annual Meeting, 3 - 06 October 2018
- VIII. **PEPTIDE BIOMARKER SIGNATURES IN STEROID-RESISTANT NEPHROTIC SYNDROME**
Zurbig P., ÖZALTIN F., Anarat A., Paripovic D., Yilmaz A., Caliskan S., Jankauskiene A., Trautmann A., Mischak H., Schaefer F.
55th Congress of the European-Renal-Association (ERA) and European-Dialysis-and-Transplantation-Association (EDTA), Copenhagen, Denmark, 24 - 27 May 2018, vol.33
- IX. **Çocuk ve genç erişkinlerde renovasküler hipertansiyonda cerrahi tedaviler.**
PEKER R. O. , AKI F. T. , KUMBASAR U., GÜVENER M., YILMAZ M., ÖZALTIN F., DÜZOVA A., TOPALOĞLU R., PEYNİRCİOĞLU B., DEMİRCİN M.
18. Ulusal Vasküler ve Endovasküler Cerrahi Kongresi 9. Ulusal Fleboloji Kongresi, Girne, Cyprus (Kktc), 9 - 12 December 2017
- X. **Çocuk ve Genç Erişkinlerde Renovasküler Hipertansiyonda Cerrahi Tedavi**
PEKER R. O. , AKI F. T. , KUMBASAR U., GÜVENER M., YILMAZ M., DOĞAN R., ÖZALTIN F., DÜZOVA A., DEMİRCİN M., TOPALOĞLU R., et al.
18. Ulusal Vasküler ve Endovasküler Cerrahi Kongresi, Turkey, 9 - 12 November 2017
- XI. **Risk factors for cardiovascular comorbidities in children with steroid-sensitive and steroid-resistant nephrotic syndrome.**
Kara Eroğlu F., TOPALOĞLU R., KARAGÖZ T., OĞUZ B., ÖZALTIN F., GÜLHAN B., DÜZOVA A.
50th Anniversary Meeting of the ESPN, 6 - 09 September 2017, vol.32, pp.1661
- XII. **OC-7 Cresentic glomerulonephritis (CGN) in childhood classification of aetiology and clinicopathological importance of cd163 positive (M2) macrophages**
KAYKI G., ORHAN D., ÖZALTIN F., TALİM B., DÜZOVA A., AKÇAÖREN z., TOPALOĞLU R., GÜÇER K. Ş.
Oral Communications, 7 - 10 June 2017
- XIII. **Tocilizumab Treatment in childhood takayasu arteritis case series of four patients and systematic review of the literature**
BATU E. D. , SÖNMEZ H. E. , HAZIROLAN T., ÖZALTIN F., BİLGİNER Y., ÖZEN S.
PRES 2016 Genova, 28 September - 01 October 2016
- XIV. **NEXT GENERATION GENE PANEL SCREENING IN STEROID-RESISTANT NEPHROTIC SYNDROME**
Lipska-Zietkiewicz B. S. , Boyer O., Gribouval O., Tabatabaei M., Fourrage C., Nischke P., Bole-Feysot C., Rotthier A., ÖZALTIN F., Noris M., et al.
53rd ERA-EDTA Congress, Vienna, Austria, 21 - 24 May 2016, vol.31, pp.353

- XV. **SMARCAL1 SCREENING IN NEPHROTIC SYNDROME - LESSONS FROM PODONET**
Lipska-Zietkiewicz B. S. , Gellermann J., Boyer O., Shalaby M. A. , ÖZALTIN F., Dusek J., Melk A., KARABAY BAYAZIT A., Massella L., Schaefer F.
53rd ERA-EDTA Congress, Vienna, Austria, 21 - 24 May 2016, vol.31, pp.353-354
- XVI. **NORMAL 25-HYDROXYVITAMIN D LEVELS ARE ASSOCIATED WITH LESS PROTEINURIA AND ATTENUATE RENAL FAILURE PROGRESSION IN CHILDREN WITH CHRONIC KIDNEY DISEASE**
Shroff R., Trivelli A., Litwin M., Picca S., Anarat A., Sallay P., ÖZALTIN F., Zurowska A., Jankauskiene A., Montini G., et al.
52nd Congress of the European-Renal-Association-European-Dialysis-and-Transplant-Association, London, Canada, 28 - 31 May 2015, vol.30
- XVII. **CONVERSION TO SIROLIMUS IN PEDIATRIC RENAL TRANSPLANT RECIPIENTS**
TOPALOĞLU R., GÜLHAN B., BİLGİNER Y., Tayfur A. C. , Yildiz C., ÖZALTIN F., DÜZOVA A., ÖZEN S., Aki T., BEŞBAŞ N.
50th European-Renal-Association - European-Dialysis-and-Transplant-Association Congress, İstanbul, Turkey, 18 - 21 May 2013, vol.28, pp.294
- XVIII. **Eye involvement in children with primary FSGS**
BİLGİNER Y., Poyraz C. E. , ÖZALTIN F., KADAYIFÇILAR S., BEŞBAŞ N., Topaloglu R., ÖZEN S., Hildebrandt F., BAKKALOĞLU A.
44th ERA-EDTA Congress, Barcelona, Spain, 22 - 24 June 2007, vol.22, pp.292
- XIX. **Influence of monocyte chemoattractant protein-1-2518A > G polymorphism on childhood focal segmental glomerulosclerosis**
Kalyoncu M., Besbas N., Ozgul K., Ozaltin F., Duzova A., Topaloglu R., Ozen S., Ozguc M., Bakkaloglu A.
43rd ERA-EDTA Congress, Glasgow, England, 15 - 18 July 2006, vol.21, pp.328

Supported Projects

ÖZALTIN F., BEŞBAŞ N., Project Supported by Higher Education Institutions, Atipik Hemolitik Üremik Sendromlu Hastalarda Kompleman Düzenleyici Genlerdeki Mutasyonların Saptanması, 2011 - 2016

Citations

Total Citations (WOS):2605

h-index (WOS):26