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Eğitim Bilgileri

Tıpta Uzmanlık, Dokuz Eylül Üniversitesi, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, Türkiye 2002 - 2011

Tıpta Uzmanlık, Dokuz Eylül Üniversitesi, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, Türkiye 1997 - 2001

Tıpta Uzmanlık, Dokuz Eylül Üniversitesi, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, Türkiye 1991 - 1997

Lisans, Hacettepe Üniversitesi, Tıp Fakültesi, Tıp Pr. (İngilizce), Türkiye 1984 - 1991

Yabancı Diller

İngilizce, C1 İleri

Araştırma Alanları

Sağlık Bilimleri

Akademik Unvanlar / Görevler

Prof. Dr., Dokuz Eylül Üniversitesi, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 1991 - Devam Ediyor

SCI, SSCI ve AHCI İndekslerine Giren Dergilerde Yayınlanan Makaleler

- Causes and management of urinary system problems in children on long-term home invasive mechanical ventilation**
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Pediatric Pulmonology, 2024 (SCI-Expanded)
- AN INFANT WITH ALDOSTERONE SYNTHASE DEFICIENCY WHO WAS ERRONEOUSLY TREATED AS BARTTER SYNDROME**
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- IV. **CLINICAL AND GENETIC ANALYSES OF 16 CHILDREN WITH NEPHRONOPHTISIS- RELATED CILIOPATHY**
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- V. **SURVIVAL BEYOND THREE YEARS OF AGE WITH MILD CHRONIC RENAL DISEASE IN A CHILD WITH RENAL TUBULAR DYSGENESIS DUE TO ACE MUTATION**
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- VI. **REVERSIBLE RENAL IMPAIREMENT DUE TO ACQUIRED HYPOTHYROIDISM IN A CHILD WITH ADCK3 MUTATION**
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- VII. **ISOLATED NON-ORTHOSTATIC PERSISTENT PROTEINURIA IN A CHILD: DO NOT OVERLOOK CUBN VARIANTS**
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- VIII. **NEPHROCALCINOSIS: THE PUZZLE IS ALMOST COMPLETE WITH WHOLE EXOME SEQUENCING**
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- IX. **The outcomes of renin-angiotensin-aldosterone system inhibition and immunosuppressive therapy in children with X-linked Alport syndrome**
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- X. **Unexpected finding in kidney biopsy of a child with nephrotic proteinuria: Questions**
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- XV. **RELATIONSHIP BETWEEN THE RENAL FUNCTION AND THE RENAL PELVIS ANTEROPOSTERIOR DIAMETER IN CHILDREN WITH URETEROPELVIC JUNCTION OBSTRUCTION**
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- XIX. **Late onset Bartter syndrome: Bartter syndrome type 2 presenting with isolated nephrocalcinosis and high parathyroid hormone levels mimicking primary hyperparathyroidism**
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- XX. **Comparison of clinical, pathological and long-term renal outcomes of children with Henoch-Schonlein purpura nephritis and IgA nephropathy**
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- XXX. **The role of L-carnitine in treatment of a murine model of asthma**
 Uzuner N., Kavukcu S., Yilmaz O., Ozkal S., Isekel H., Karaman O., Soylu A., Kargi A.

- ACTA MEDICA OKAYAMA, cilt.56, sa.6, ss.295-301, 2002 (SCI-Expanded)
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- XXXV. **The clinical value of urinary N-acetyl-beta-D-glucosaminidase levels in childhood age group**
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- XXXVI. **Systemic lupus erythematosus presenting with normocomplementemic urticarial vasculitis in a 4-year-old girl**
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- XL. **Effect of socioeconomic status on the blood pressure in children living in a developing country**
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- XLI. **Hypercalciuria preceding IgA nephropathy in a child with haematuria**
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- XLII. **The role of vitamin A in preventing renal scarring secondary to pyelonephritis**
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- XLIII. **Effect of seasonal changes on the cyclosporine A blood levels in renal transplant recipients during childhood**
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- XLIV. **Clinical quiz. Fanconi syndrome.**
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- XLV. **Serum vitamin A and beta-carotene concentrations and renal scarring in urinary tract infections**
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- XLVI. **Evaluation of the resistance induction in enteric flora in children caused by oral ampicillin plus**

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Diğer Dergilerde Yayınlanan Makaleler

I. Demographics, clinical, laboratory findings and treatment results of pediatric patients with IgA Vasculitis: Single-center experience

AÇARI C., BAYRAM M., yıldız g., kavukcu s., Soylu A.

Pamukkale Tıp Dergisi, cilt.16, sa.1, ss.73-80, 2023 (Scopus)

II. Effect Of MEFV Variants On The Presentation And Clinical Course Of Henoch-Schonlein Purpura In Children?

AÇARI C., TORUN BAYRAM M., YILDIZ G., KAVUKÇU S., SOYLU A.

Dokuz Eylül Üniversitesi Tıp Fakültesi Dergisi, cilt.36, sa.3, 2022 (Hakemli Dergi)

III. Mutation Analysis of the AGXT Gene in Combined Liver-Kidney and Isolated Liver Transplanted Children for Primary Hyperoxaluria Type 1: a Single Center Experience

TÜRKMEN M. A., ALAYGUT D., Ağilkaya S., BAYRAM M. T., DEMİR B., SOYLU A., KAVUKÇU S., CİNGÖZ S.

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IV. Mutation Analysis of the AGXT Gene in Combined Liver-Kidney and Isolated Liver Transplanted Children for Primary Hyperoxaluria Type 1

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Desteklenen Projeler

GİRAY BOZKAYA Ö., SOYLU A., ÜLGENALP A., KOCABEY M., Yükseköğretim Kurumları Destekli Proje, MEFV Geninde Mutasyon Saptanamayan veya Tek Mutasyon Saptanan Ailesel Akdeniz Ateşi Olgularında Farklı Gen Varyantlarının Araştırılması, 2020 - 2021

SOYLU A., Yükseköğretim Kurumları Destekli Proje, Proteinüri ile seyreden böbrek hastalıklarında aşırı yüksek hesaplanan kreatinin klirensinin, glomerül filtrasyon hızında gerçek bir artışı temsil edip etmediğinin ve uzun vadede böbrek fonksiyonlarına etkisinin araştırılması, 2018 - 2019

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