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Tıpta Uzmanlık, Hacettepe Üniversitesi, Çocuk Sağlığı Ve Hastalıkları, Türkiye 1981 - 1986

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Araştırma Alanları

Tıp, Sağlık Bilimleri, Dahili Tıp Bilimleri, Çocuk Sağlığı ve Hastalıkları, Pediatrik Hematoloji

Akademik Unvanlar / Görevler

Prof. Dr., Hacettepe Üniversitesi, Tıp Fakültesi (Türkçe), Dahili Tıp Bilimleri Bölümü, 1981 - Devam Ediyor

Akademik İdari Deneyim

Hacettepe Üniversitesi, Tıp Fakültesi, 2016 - Devam Ediyor

SCI, SSCI ve AHCI İndekslerine Giren Dergilerde Yayınlanan Makaleler

- I. Alu-Mediated Deletion of *FANCA* in Turkish Families With Fanconi Anemia: Evidence of a Founder Effect**
DURMAZ ÖZDİNÇ C. D., GÜMRÜK F., Celkan T., Unal S., ÇETİNKAYA A.
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- II. Single-center experience of four cases with iron-refractory iron deficiency anemia (IRIDA)**
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- III. Outcome of the Modified St. Jude Total XV Protocol in Turkish Children with Newly Diagnosed Acute Lymphoblastic Leukemia: A Single-Center Retrospective Analysis**
YILMAZ H., AYTAÇ EYÜPOĞLU Ş. S., KUŞKONMAZ B. B., ÇETİNKAYA F. D., Unal S., GÜMRÜK F.
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- IV. A rare case of Klippel-Trenaunay syndrome presenting with chronic myeloid leukemia**
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- V. **First report of t(1;9)(q21;q34) in Fanconi anemia as a preceding chromosomal aberration before leukemia development**
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- VII. **Usage of Plasma Presepsin, C-Reactive Protein, Procalcitonin and Proadrenomedullin to Predict Bacteremia in Febrile Neutropenia of Pediatric Hematological Malignancy Patients**
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- VIII. **Loss of C2orf69 defines a fatal autoinflammatory syndrome in humans and zebrafish that evokes a glycogen-storage-associated mitochondriopathy**
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- IX. **Proerythroblast Cells of Diamond-Blackfan Anemia Patients With RPS19 and CECR1 Mutations Have Similar Transcriptomic Signature**
KARAOŞMANOĞLU B., KURŞUNEL M. A., ÇETİNKAYA F. D., GÜMRÜK F., ESENDAĞLI G., ÜNAL Ş., Taskiran E. Z.
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- X. **Hemophagocytosis in bone marrow aspirates in multisystem inflammatory syndrome in children**
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- XI. **Transplacental hemophilia A and prophylactic treatment with intravenous immunoglobulin and recombinant factor VIIa in the newborn period: a case report**
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- XII. **Acute promyelocytic leukemia in a child with reticulin fibrosis**
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- XIII. **Central nervous system lesions in Fanconi anemia: Experience from a research center for Fanconi anemia patients**
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- XIV. **Comment on: Clinical, cytogenetic, and molecular analyses of 17 neonates with transient abnormal myelopoiesis and nonconstitutional trisomy 21**
AKSU T., GÜMRÜK F., ÜNAL Ş.
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- XV. **Hemophagocytic Lymphohistiocytosis in Patients With Primary Immunodeficiency**
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- XVI. **Three patients with glucose-6 phosphatase catalytic subunit 3 deficiency**
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- XVII. **A rare form of congenital neutropenia: VPS45 deficiency**
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- XVIII. **Autoinflammation in addition to combined immunodeficiency: SLC29A3 gene defect**
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- XIX. **Hb H Disease Diagnosed During Adolescent Pregnancy**
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- XX. **The remarkable response to ponatinib therapy in a child with blastic phase of chronic myeloid leukemia**
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- XXI. **A Monogenic Disease with a Variety of Phenotypes: Deficiency of Adenosine Deaminase 2**
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- XXII. **Rare Cytogenetic Anomalies in Two Pediatric Patients with Acute Leukemia**
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- XXIII. **A Spectrum of Clinical Findings from ALPS to CVID: Several Novel LRBA Defects.**
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- XXIV. **Comparison of ferrous sulfate, polymaltose complex and iron-zinc in iron deficiency anemia**
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- XXV. **Fanconi anemia: a single center experience of a large cohort**
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- XXVI. **Infant lymphoblastic leukemia: a single centers 10 year experience**
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- XXVIII. **Deferasirox in children with transfusion-dependent thalassemia or sickle cell anemia: A large cohort real-life experience from Turkey (REACH-THEM)**
Antmen B., KARAKAŞ Z., Yesilipek M. A., KÜPESİZ O. A., Sasmaz I., Uygun V., Kurtoglu E., Oktay G., Aydogan G., Akin M., et al.
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- XXIX. **Vacuolization in myeloid and erythroid precursors in a child with menkes disease Menkes hastalıklı bir çocukta myeloid ve eritroid öncüllerde vaküolizasyon**
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- XXX. **Comparison of different types of twin pregnancies in terms of obstetric and perinatal outcomes: association of vanished twins with methylenetetrahydrofolate reductase (MTHFR) polymorphism(s)**
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- XXXI. **Methylenetetrahydrofolate Reductase Polymorphisms and Pregnancy Outcome**
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- XXXII. **Bone Marrow Mesenchymal Stem Cells Carrying FANCD2 Mutation Differ from the Other Fanconi Anemia Complementation Groups in Terms of TGF-beta 1 Production**
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- XXXIII. **Mechanism for survival of homozygous nonsense mutations in the tumor suppressor gene BRCA1**
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- XXXIV. **Assessment of Peripheral Neuropathy in Patients with β -Thalassemia via Electrophysiological Study: Reevaluation in the Era of Iron Chelators**
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- XXXV. **The clinical and laboratory evaluation of familial hemophagocytic lymphohistiocytosis and the importance of hepatic and spinal cord involvement: a single center experience**
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- XXXVI. **Molecular genetic analysis of the F11 gene in 14 Turkish patients with factor XI deficiency: identification of novel and recurrent mutations and their inheritance within families**
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- XXXVII. **Tyrosine Kinase Inhibitor Responses in Pediatric CML Patients: Hacettepe Experience**
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- XXXVIII. **Human Bocavirus: Can It Trigger Hemophagocytic Lymphohistiocytosis?**
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- XXXIX. **Heavy metal levels in patients with ineffective erythropoiesis**
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- XL. **Liver transplantation from a deceased donor with beta-thalassemia intermedia is not contraindicated: A case report**
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- XLI. **The questioning for routine monthly monitoring of proteinuria in patients with β -thalassemia on deferasirox chelation**
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- XLII. **Macrophage activation syndrome in children with systemic juvenile idiopathic arthritis and systemic lupus erythematosus**
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- XLIII. **Hypereosinophilic Syndrome: Hacettepe Experience**
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- XLIV. **Severe Hypercalcemia in a Child With Acute Lymphoblastic Leukemia Relapse: Successful Management With Combination of Calcitonin and Bisphosphonate**
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- XLV. **Sorafenib-induced Posterior Reversible Encephalopathy Syndrome in a Child With FLT3-ITD-positive Acute Myeloid Leukemia**
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- XLVI. **Successful Outcome With Fludarabine-Based Conditioning Regimen for Hematopoietic Stem Cell Transplantation From Related Donor in Fanconi Anemia: A Single Center Experience From Turkey**

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- XLVII. **The genetic basis of asymptomatic codon 8 frame-shift (HBB: C25_26delAA) β 0-thalassaemia homozygotes**
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- XLVIII. **Effects of blood transfusion on cytokine profile and pulmonary function in patients with thalassemia major**
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- XLIX. **Foetal and neonatal intracranial haemorrhage in term newborn infants: Hacettepe University experience**
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- LI. **Hereditary Elliptocytosis with Pyropoikilocytosis**
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- LII. **Neonates born to mothers with immune thrombocytopenic purpura: a single-center experience of 20 years**
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- LIII. **The Feasibility of Magnetic Resonance Imaging for Quantification of Liver, Pancreas, Spleen, Vertebral Bone Marrow, and Renal Cortex R2*and Proton Density Fat Fraction in Transfusion-Related Iron Overload**
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- LIV. **Serum lipids in Turkish patients with β -thalassemia major and β -thalassemia minor Türk β -talasemi majör ve β -talasemi minör hastalarının serum lipidleri**
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- LV. **Immune Thrombocytopenic Purpura During Maintenance Phase of Acute Lymphoblastic Leukemia: A Rare Coexistence Requiring a High Degree of Suspicion, a Case Report and Review of the Literature.**
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- LVI. **OSTEOPOROSIS IN PATIENTS WITH ACUTE LYMPHOBLASTIC LEUKEMIA TREATED ON MODIFIED ST JUDE TOTAL XV THERAPY**
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- LVIII. **Renal transplantation experience in a patient with factor V Leiden homozygous, MTHFR C677T heterozygous, and PAI heterozygous mutation**
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- LX. **Case series of thromboembolic complications in childhood nephrotic syndrome: Hacettepe experience.**
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- LXI. **Risk of thrombosis in a cohort of pediatric acute lymphoblastic leukemia patients**
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- LXII. **Basal cell carcinoma after treatment of childhood acute lymphoblastic leukemia and concise review of the literature.**
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- LXIII. **Iron chelation with deferasirox in a patient with de-novo ferroportin mutation.**
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- LXIV. **Acute Megakaryoblastic Leukemia with t(1;22) Mimicking Neuroblastoma in an Infant.**
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- LXV. **The hematological and molecular spectrum of α -thalassemias in Turkey: The hacettepe experience Türkiye’de alfa talasemilerin hematolojik ve moleküler spektrumu: Hacettepe deneyimi**
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- LXVII. **Molecular Analyses of Pyruvate Kinase Deficient Turkish Patients from a Single Center.**
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- LXVIII. **The first report of a homozygous codons 9/10 (+T) β -thalassemia mutation in a Turkish patient.**
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Kitap & Kitap Bölümleri

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I. TİP III VON WİLLEBRAND HASTALIĞI OLAN ÇOCUKLARDA UZUN SÜRELİ PROFİLAKSİ PROFİLAKSİ VERELİM Mİ? VERMEYELİM Mİ?

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Desteklenen Projeler

GÜMRÜK F., BAYHAN T., ÜNAL Ş., ÇETİN M., Yükseköğretim Kurumları Destekli Proje, Rotasyonel tromboelastografi ROTEM cihazı kullanılarak çocuklarda santral venöz kateterlerde tromboz gelişimi riskinin öngörülebilirliğinin belirlenmesi, 2015 - 2020

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