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Kişisel Bilgiler

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Eğitim Bilgileri

Tıpta Yandal Uzmanlık, Hacettepe Üniversitesi, Çocuk Sağlığı Ve Hastalıkları Hematoloji Bilim Dalı, Türkiye 1989 - 1994

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

Lisans, Hacettepe Üniversitesi, Tıp Fakültesi, Türkiye 1975 - 1981

Araştırma Alanları

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

Mesleki Deneyim

Bölüm Başkanı, Hacettepe Üniversitesi, Tıp Fakültesi, 2016 - Devam Ediyor

SCI, SSCI ve AHCI İndekslerine Giren Dergilerde Yayınlanan Makaleler

- I. **Central nervous system lesions in Fanconi anemia: Experience from a research center for Fanconi anemia patients**
AKSU T., GÜMRÜK F., Bayhan T., Coskun C., Oguz K. K. , ÜNAL Ş.
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- II. **Hemophagocytic Lymphohistiocytosis in Patients With Primary Immunodeficiency**
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- IV. **Comment on: Clinical, cytogenetic, and molecular analyses of 17 neonates with transient abnormal myelopoiesis and nonconstitutional trisomy 21**
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- VI. **A rare form of congenital neutropenia: VPS45 deficiency**
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- XII. **Infant lymphoblastic leukemia: a single centers 10 year experience**
Yaman-Bajin I., Aytac S., Kuskonmaz B. B. , Ucan-Cetinkaya D., ÜNAL Ş., GÜMRÜK F., Cetin M.
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- XIII. **Deferasirox in children with transfusion-dependent thalassemia or sickle cell anemia: A large cohort real-life experience from Turkey (REACH-THEM)**
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- XIV. **Methylenetetrahydrofolate Reductase Polymorphisms and Pregnancy Outcome**
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- XV. **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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- XVI. **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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- XVII. **Human Bocavirus: Can It Trigger Hemophagocytic Lymphohistiocytosis?**
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- XVIII. **Tyrosine Kinase Inhibitor Responses in Pediatric CML Patients: Hacettepe Experience**
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- XIX. **Liver transplantation from a deceased donor with beta-thalassemia intermedia is not contraindicated: A case report**
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- XX. **Macrophage activation syndrome in children with systemic juvenile idiopathic arthritis and systemic lupus erythematosus**
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- XXII. **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-**

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- XXIV. **OSTEOPOROSIS IN PATIENTS WITH ACUTE LYMPHOBLASTIC LEUKEMIA TREATED ON MODIFIED ST JUDE TOTAL XV THERAPY**
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- XXVI. **Risk of thrombosis in a cohort of pediatric acute lymphoblastic leukemia patients**
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- XXVII. **THE OUTCOME OF PEDIATRIC ACUTE LYMPHOBLASTIC LEUKEMIA IN PATIENTS WHO PRESENTED WITH HYPERLEUKOCYTOSIS**
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- XXVIII. **Pediatric Myelodysplastic Syndrome: Experience from a Single Center**
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- XXIX. **the Effects of Deferasirox on Iron in Pituitary, Pancreas and Thyroid Glands: An Observational Case-Control Study**
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- XXX. **PEDIATRIC MYELODYSPLASTIC SYNDROME: EXPERIENCE FROM A SINGLE CENTER**
ÇETİN M., ÜNAL S., BAYHAN T., AYTAÇ S. A. , KUŞKONMAZ B. B. , TAVİL B., UÇKAN-CETİNKAYA D., TUNCER M., GÜMRÜK F.
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- XXXI. **Thrombophilic Risk Factors And The Efficiency Of Prophylactic Anticoagulation Therapy In Children Who Underwent Renal Transplantation**
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- XXXII. **An infant presented with deficiencies of vitamin K dependent factors due to an inherited novel mutation in GGCX gene**
ÜNAL S., KHANİYEV S., WATZKA M., OLDENBURG J., GÜMRÜK F.
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- XXXIV. **Striking Hematological Abnormalities in Patients With Microcephalic Osteodysplastic Primordial Dwarfism Type II (MOPD II): A Potential Role of Pericentrin in Hematopoiesis**
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- XXXV. **IMPROVED OUTCOME IN HIGH RISK GROUP PATIENTS WITH HIGH DOSE METHYLPREDNISOLONE DURING INDUCTION OF MODIFIED ST JUDE TOTAL XIII A**

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- XXXVI. **CONTRIBUTORY RISK FACTORS FOR DEVELOPMENT OF THROMBOSIS IN CHILDREN WITH NEPHROTIC SYNDROME**
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- XXXVIII. **Double trouble: Duchenne muscular dystrophy and hemophilia**
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- XLI. **Prenatal Diagnosis of Hemoglobinopathies in Hacettepe University, Turkey**
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- XLIV. **MEASLES, MUMPS, AND RUBELLA ANTIBODY STATUS AND RESPONSE TO IMMUNIZATION IN CHILDREN AFTER THERAPY FOR ACUTE LYMPHOBLASTIC LEUKEMIA**
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- XLVIII. **Chronic Recurrent Multifocal Osteomyelitis as the First Presentation of Acute Lymphoblastic Leukemia in a 2-year-old Boy**
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- L. Wilms Tumor, AML and Medulloblastoma in a Child With Cancer Prone Syndrome of Total Premature Chromatid Separation and Fanconi Anemia**
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- LI. PARVOVIRUS B19-INDUCED PERSISTENT PURE RED CELL APLASIA IN A CHILD WITH T-CELL IMMUNODEFICIENCY**
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- LII. Hyperimmunoglobulinemia D and periodic fever syndrome; treatment with etanercept and follow-up**
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- LIII. Congenital coagulation factor deficiencies: Hacettepe experience**
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- LIV. Pearson syndrome associated with hemophagocytic syndrome in a child**
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- LV. CD-34 selected hematopoietic stem cell transplantation from HLA identical family members for fanconi anemia**
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- LVI. Interleukin-6 (IL-6), tumor necrosis factor-alpha (TNF-alpha) levels and IL-6, TNF-Polymorphisms in children with thrombosis**
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- LIX. Haematological findings in children with inborn errors of metabolism**
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- LX. Molecular characterization of Turkish patients with pyrimidine 5 ' nucleotidase-I deficiency**
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Diğer Dergilerde Yayınlanan Makaleler

- I. Infant Acute Lymphoblastic Leukemia with Atypical Presentation**

YAMAN BAJİN H. İ. , YILDIZ Y., akın ş., AYTAÇ EYÜPOĞLU Ş. S. , ÜNAL CANGÜL Ş., KUŞKONMAZ B. B. , Cetin M., SİVRİ H. S. , GÜMRÜK F.

Acta Medica, cilt.50, sa.4, ss.57-59, 2019 (Diğer Kurumların Hakemli Dergileri)

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