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Eğitim Bilgileri

Tıpta Yandal Uzmanlık, Hacettepe Üniversitesi, Çocuk Hematolojisi Ve Onkolojisi, Türkiye 2005 - 2010

Tıpta Uzmanlık, Hacettepe Üniversitesi, Çocuk Sağlığı Ve Hastalıkları, Türkiye 1999 - 2005

Lisans, Hacettepe Üniversitesi, Tıp Fakültesi (İngilizce), Türkiye 1993 - 1999

Yabancı Diller

İngilizce, C1 İleri

Araştırma Alanları

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

Akademik Unvanlar / Görevler

Prof. Dr., Hacettepe Üniversitesi, Tıp Fakültesi (İngilizce), Dahili Tıp Bilimleri Bölümü, 2017 - Devam Ediyor

Doç. Dr., Hacettepe Üniversitesi, Tıp Fakültesi (İngilizce), 2011 - 2017

Verdiği Dersler

BİTİRME PROJESİ - II, Lisans, 2017 - 2018, 2016 - 2017

LİSANS ARAŞTIRMA PROJESİ, Lisans, 2017 - 2018

MEKATRONİK, Lisans, 2017 - 2018, 2016 - 2017

MAKİNE ELEMANLARI TASARIMI, Lisans, 2017 - 2018

BİTİRME PROJESİ - I, Lisans, 2017 - 2018

ARAÇ PARÇA TASARIMI, Lisans, 2017 - 2018

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- II. **Central nervous system lesions in Fanconi anemia: Experience from a research center for Fanconi anemia patients**
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- V. **TURKEY CONGENITAL THROMBOTIC THROMBOCYTOPENIC PURPURA REGISTRATION SYSTEM RESULTS**
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- VIII. **Outcomes of Eltrombopag Treatment and Development of Iron Deficiency in Children with Immune Thrombocytopenia in Turkey**
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- IX. **Characterization of Two Cases of Congenital Dyserythropoietic Anemia Type I Shed Light on the Uncharacterized C15orf41 Protein.**
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- X. **Refugee children with beta-thalassemia in Turkey: Overview of demographic, socioeconomic, and medical characteristics.**
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- XI. **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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- XII. **Both Granulocytic and Non-Granulocytic Blood Cells Are Affected in Patients with Severe Congenital Neutropenia and Their Non-Neutropenic Family Members: An Evaluation of Morphology, Function, and Cell Death**
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- XVI. **MECOM-associated syndrome: a heterogeneous inherited bone marrow failure syndrome with amegakaryocytic thrombocytopenia.**
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- XVII. **Assessment of Peripheral Neuropathy in Patients with β -Thalassemia via Electrophysiological Study: Reevaluation in the Era of Iron Chelators**
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- XVIII. **A National Registry of Thalassemia in Turkey: Demographic and Disease Characteristics of Patients, Achievements, and Challenges in Prevention.**
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- XIX. **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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- XX. **Acute myeloid leukemia in a child with dedicator of cytokinesis 8 (DOCK8) deficiency**
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- XXV. **Predictors of Suboptimal Follow-up in Pediatric Cancer Survivors.**
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- XXVII. **Hypereosinophilic Syndrome: Hacettepe Experience**
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- XXVIII. **THE GLYCOCALYX AND TRAUMA: A REVIEW**
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- XXX. **Sorafenib-induced Posterior Reversible Encephalopathy Syndrome in a Child With FLT3-ITD-positive Acute Myeloid Leukemia**
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- XXXII. **Foetal and neonatal intracranial haemorrhage in term newborn infants: Hacettepe University experience**
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- XXXVI. **Neonates born to mothers with immune thrombocytopenic purpura: a single-center experience of 20 years**
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- XL. **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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- XLI. **An Infant With Congenital Leukemia Cutis and AML-M5 With MLL Gene Rearrangement.**
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- XLII. **Biochemical markers of glucose metabolism may be used to estimate the degree and progression of iron overload in the liver and pancreas of patients with β -thalassemia major.**
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- XLIII. **Case series of thromboembolic complications in childhood nephrotic syndrome: Hacettepe experience.**
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- XLVII. **Molecular Analyses of Pyruvate Kinase Deficient Turkish Patients from a Single Center.**
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- LII. **Retrospective cohort study of 205 cases with congenital dyserythropoietic anemia type II: definition of clinical and molecular spectrum and identification of new diagnostic scores.**
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- LVI. **Lysosomal vesicles, giant granules, and erythrophagocytosis in chédiak-higashi syndrome.**
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- LVII. **A Novel Mutation in Protein C Gene (PROC) Causing Severe Phenotype in Neonatal Period**
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- LVIII. **Congenital thrombotic thrombocytopenic purpura with novel mutations in three unrelated Turkish children.**
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- LIX. **Kindlin-3-independent adhesion of neutrophils from patients with leukocyte adhesion deficiency type III**
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- LX. **Atypical combined immunodeficiency due to Artemis defect: a case presenting as hyperimmunoglobulin M syndrome and with LGLL.**
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- LXI. **Nocturnal enuresis in sickle cell disease and thalassemia major: associated factors in a clinical sample.**
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- LXII. **Recurrent macrophage activation syndrome associated with heterozygous perforin W374X gene mutation in a child with systemic juvenile idiopathic arthritis.**
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