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Eğitim Bilgileri

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Lisans, Hacettepe Üniversitesi, Tıp Fakültesi (ingilizce), 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 (ingilizce), Dahili Tıp Bilimleri Bölümü, 2017 - Devam Ediyor

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

Verdiği Dersler

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

LİSANS ARAŞTIRMA PROJESİ, Lisans, 2017 - 2018

MEKATRONİK, Lisans, 2016 - 2017, 2017 - 2018

MAKİNE ELEMANLARI TASARIMI, Lisans, 2017 - 2018

BİTİRME PROJESİ - I, Lisans, 2017 - 2018

ARAÇ PARÇA TASARIMI, Lisans, 2017 - 2018

Özel Konular, Yüksek Lisans, 2017 - 2018

MEKATRONİK TASARIM, Lisans, 2017 - 2018

LİSANS ÜSTÜ DERS DANIŞMANLIĞI, Lisans, 2016 - 2017

DESİGN OF MACHİNE ELEMENTS, Lisans, 2016 - 2017

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SCI, SSCI ve AHCI İndekslerine Giren Dergilerde Yayınlanan Makaleler

- I. **Modelling the future of HIV in Turkey: disease implications of improving prevention, diagnosis and treatment**
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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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- IV. **TURKEY CONGENITAL THROMBOTIC THROMBOCYTOPENIC PURPURA REGISTRATION SYSTEM RESULTS**
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- V. **Outcomes of Eltrombopag Treatment and Development of Iron Deficiency in Children with Immune Thrombocytopenia in Turkey**
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- VI. **Rare Cytogenetic Anomalies in Two Pediatric Patients with Acute Leukemia**
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- VII. **A Monogenic Disease with a Variety of Phenotypes: Deficiency of Adenosine Deaminase 2.**
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- VIII. **Vacuolization in Myeloid and Erythroid Precursors in a Child with Menkes Disease**
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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. **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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- XII. **Molecular approaches to diagnose Diamond-Blackfan anemia: The EuroDBA experience.**
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- XIII. **Mechanism for survival of homozygous nonsense mutations in the tumor suppressor gene BRCA1**
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- XIV. **Multi-gene panel testing improves diagnosis and management of patients with hereditary anemias.**
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- XV. **A National Registry of Thalassemia in Turkey: Demographic and Disease Characteristics of Patients, Achievements, and Challenges in Prevention.**
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- XVI. **Assessment of Peripheral Neuropathy in Patients with β -Thalassemia via Electrophysiological Study: Reevaluation in the Era of Iron Chelators.**
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- XVII. **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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- XVIII. **Acute myeloid leukemia in a child with dedicator of cytokinesis 8 (DOCK8) deficiency**
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- XIX. **Mutations in signal recognition particle SRP54 cause syndromic neutropenia with Shwachman-Diamond-like features.**
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- XX. **Heavy metal levels in patients with ineffective erythropoiesis.**
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- XXI. **The questioning for routine monthly monitoring of proteinuria in patients with β -thalassemia on deferasirox chelation.**
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- XXII. **Rapid lung magnetic resonance imaging in children with pulmonary infection: reply to Sodhi et al.**
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- XXIII. **Predictors of Suboptimal Follow-up in Pediatric Cancer Survivors.**
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- XXIV. **Magnetic resonance imaging of pulmonary infection in immunocompromised children: comparison with multidetector computed tomography.**
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- XXV. **Hypereosinophilic Syndrome: Hacettepe Experience**
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- XXVI. **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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- XXVII. **Sorafenib-induced Posterior Reversible Encephalopathy Syndrome in a Child With FLT3-ITD-positive Acute Myeloid Leukemia**
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- XXVIII. **THE GLYCOLALYX AND TRAUMA: A REVIEW.**
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- XXIX. **Serum Lipids in Turkish Patients with β -Thalassemia Major and β -Thalassemia Minor.**
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- XXX. **The genetic basis of asymptomatic codon 8 frame-shift (HBB:c25_26delAA) $\beta(0)$ -thalassaemia homozygotes.**
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- XXXI. **Foetal and neonatal intracranial haemorrhage in term newborn infants: Hacettepe University experience**
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- XXXII. **Effects of blood transfusion on cytokine profile and pulmonary function in patients with thalassemia major.**
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- XXXIII. **The European Hematology Association Roadmap for European Hematology Research: a consensus document.**
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- XXXIV. **Neonates born to mothers with immune thrombocytopenic purpura: a single-center experience of 20 years**
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- XXXV. **Targeted high-throughput sequencing for genetic diagnostics of hemophagocytic lymphohistiocytosis.**
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- XXXVI. **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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- XXXVII. **Spectrum of Atypical Clinical Presentations in Patients with Biallelic PRF1 Missense Mutations.**
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- XXXVIII. **Transcobalamin II Deficiency in Four Cases with Novel Mutations.**
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- XXXIX. **An Infant With Congenital Leukemia Cutis and AML-M5 With MLL Gene Rearrangement.**
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- XL. **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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- XLI. **Case series of thromboembolic complications in childhood nephrotic syndrome: Hacettepe experience**
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- XLII. **The Hematological and Molecular Spectrum of α -Thalassemias in Turkey: The Hacettepe Experience.**
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- XLIII. **Basal cell carcinoma after treatment of childhood acute lymphoblastic leukemia and concise review of the literature.**
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- XLV. **The first report of a homozygous codons 9/10 (+T) β -thalassemia mutation in a Turkish patient.**
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- XLVI. **Molecular Analyses of Pyruvate Kinase Deficient Turkish Patients from a Single Center.**
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- XLVII. **Acute megakaryoblastic leukemia with t(1;22) mimicking neuroblastoma in an infant Nöroblastomu taklit eden t(1;22) pozitif akut megakaryoblastik lösemili bir süt Çocuğu**
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- XLVIII. **Fanconi's Anemia Effect or Sickle Cell Anemia Effect: That is the Question.**
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- XLIX. **Mean Platelet Volume can Predict Cerebrovascular Events in Patients with Sickle Cell Anemia.**
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- L. **Bleeding and non-bleeding phenotypes in patients with GGX gene mutations.**
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- LI. **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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- LII. **Successful hematopoietic stem cell transplantation in a patient with congenital dyserythropoietic anemia type II**
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- LIII. **The relationship between hematological findings and coronary artery aneurysm in kawasaki disease.**
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- LIV. **Lysosomal vesicles, giant granules, and erythrophagocytosis in chédiak-higashi syndrome.**
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- LV. **A novel mutation in protein C gene (PROC) causing severe phenotype in neonatal period.**
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- LVI. **Congenital thrombotic thrombocytopenic purpura with novel mutations in three unrelated Turkish children.**
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- LX. **Recurrent macrophage activation syndrome associated with heterozygous perforin w374x gene mutation in a child with systemic juvenile idiopathic arthritis**
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- LXI. **THROMBOPHILIC RISK FACTORS AND THE EFFICIENCY OF PROPHYLACTIC ANTICOAGULATION THERAPY IN CHILDREN WHO UNDERWENT RENAL TRANSPLANTATION**
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- LXIII. **Comparison of primary human cytotoxic T-cell and natural killer cell responses reveal similar molecular requirements for lytic granule exocytosis but differences in cytokine production.**
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- LXIV. **Medical management of moyamoya disease and recurrent stroke in an infant with Majewski osteodysplastic primordial dwarfism type II (MOPD II).**
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