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Education Information

Post Doctorate of Medicine, Erciyes University, Tıp Fakültesi, Dahili Tıp Bil., Turkey 2015 - 2018

Expertise In Medicine, Erciyes University, Tıp Fakültesi, Dahili Tıp Bil., Turkey 2007 - 2012

Foreign Languages

English, C1 Advanced

Research Areas

Health Sciences, Medicine, Internal Medicine Sciences, Child Health and Diseases, Pediatric Hematology

Published journal articles indexed by SCI, SSCI, and AHCI

- I. **Identification of the molecular etiology in rare congenital hemolytic anemias using next-generation sequencing with exome-based copy number variant analysis**
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- II. **Pyruvate kinase deficiency in 29 Turkish patients with two novel intronic variants**
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- III. **COVID-19 disease in children and adolescents following allogeneic hematopoietic stem cell transplantation: A report from the Turkish pediatric bone marrow transplantation study group**
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- V. **A Novel Biallelic LCK Variant Resulting in Profound T-Cell Immune Deficiency and Review of the Literature**

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- VI. **Defective Treg generation and increased type 3 immune response in leukocyte adhesion deficiency 1**
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- VII. **Central nervous system thrombosis in pediatric acute lymphoblastic leukemia in Turkey: A multicenter study**
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- VIII. **Monogenic early-onset lymphoproliferation and autoimmunity: Natural history of STAT3 gain-of-function syndrome.**
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- IX. **Antibody Response against Vaccine Antigens in Children after TCRαβ-Depleted Haploidentical Stem Cell Transplantation: Is It Similar to That in Recipients with Fully Matched Donors?**
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- X. **Immunodeficiency associated with a novel functionally defective variant of *SLC19A1* benefits from folinic acid treatment**
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- XII. **Treatment of Infantile Fibrosarcoma in the Era of Targeted Therapies**
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- XIII. **Aceruloplasminemia presenting with microcytic anemia in a Turkish boy due to a novel pathogenic variant**
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- XIV. **Assessment of extracorporeal photopheresis related cell damage.**
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- XV. **Evaluation of primary care physicians' approaches to hemophilia and bleeding disorders: a questionnaire survey**
Samur B. M., Samur T. G., Çiflikli F., Özcan A., Gök V., Soykan R., Soytürk F., Kılıç Ö., Kandur M., Kandemir R., et al.
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- XVI. **Characterization of cord blood CD3⁺ TCRVα7.2⁺ CD161^{high} T and innate lymphoid cells in the pregnancies with gestational diabetes, morbidly adherent placenta, and pregnancy hypertension diseases.**
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- XVII. **A novel missense mutation outside the DNAJ domain of DNAJC21 is associated with Shwachman-**

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- XVIII. **Castleman Disease: A Multicenter Case Series from Turkey.**
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- XIX. **Social exclusion and behavior problems in adolescents with cancer and healthy counterparts**
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- XX. **A teenager boy with a novel variant of Sitosterolemia presented with pancytopenia.**
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- XXI. **A nonsense mutation in DIAPH1 gene presents with major T cell defects**
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- XXII. **Common Variable Immunodeficiency, Autoimmune Hemolytic Anemia, and Pancytopenia Associated With a Defect in IKAROS**
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- XXIII. **Comprehensive Mutation Analysis of the RAS/RAF/MEK/ERK Pathway in Paediatric Leukaemia and Significant Inferences**
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- XXV. **Different Clinical Presentation of 3 Children With Familial Hemophagocytic Lymphohistiocytosis With 2 Novel Mutations.**
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- XXVI. **ILC3 deficiency and generalized ILC abnormalities in DOCK8-deficient patients**
Eken A., Cansever M., Okus F. Z., Erdem S., Nain E., Azizoglu Z. B., Haliloglu Y., Karakukcu M., Ozcan A., Devecioglu O., et al.
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- XXVII. **Congenital afibrinogenemia in a 4-year-old girl complicated with acute lymphoblastic leukemia**
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- XXVIII. **Twenty children with non-Wilms renal tumors from a reference center in Central Anatolia, Turkey**
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- XXIX. **Invasive Fungal Infections in Children with Acute Lymphoblastic Leukemia: Experience from a Reference University Hospital in Cappadocia**
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- XXX. **CD137 deficiency causes immune dysregulation with predisposition to lymphomagenesis**
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- XXXI. **Two Different Endocrine Cancer, One Disease; DICER-1 Mutation**
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- XXXII. **A rare cause of vomiting in an adolescent: gastric Burkitt's lymphoma**
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- XXXIII. **The relationship between the prognosis of children with acute arterial stroke and polymorphisms of CDKN2B, HDAC9, NINJ2, NAA25 genes**
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- XXXIV. **Propranolol treatment for chylothorax due to diffuse lymphangiomatosis**
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- XXXV. **Genetic Deficiency and Biochemical Inhibition of ITK Affect Human Th17, Treg, and Innate Lymphoid Cells**
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- XXXVI. **A Rare Case of Activated Phosphoinositide 3-Kinase Delta Syndrome (APDS) Presenting With Hemophagocytosis Complicated With Hodgkin Lymphoma.**
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- XXXVII. **A mummy emerges from the grave: Scurvy confounding the clinical presentation of a child with Fanconi anemia**
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- XXXVIII. **Scurvy: A rare cause of arthritis in a child with neurologic disorder.**
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- XXXIX. **Vena Cava Superior Syndrome in Children with Mediastinal Tumors: Single Center Experience**
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- XL. **Tumour Lysis Syndrome in Children with Hematologic Malignancies: Single Center Experience**
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- XLI. **DOES EVEN LOW DOSE ENALAPRIL CAUSE LIFE THREATENING ACUTE KIDNEY INJURY?**
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- XLII. **Cerebral Sinus Venous Thrombosis and Prothrombotic Risk Factors in Children: A Single-Center Experience From Turkey**
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- XLIII. **ALLOGENEIC BONE MARROW TRANSPLANTATION IN THREE CASES WITH DOCK 8 DEFICIENCY**
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- XLIV. **The relationship between hematological parameters and prognosis of children with acute ischemic stroke**

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- XLV. **Use of a chlorhexidine-impregnated dressing reduced catheter-related bloodstream infections caused by Gram-positive microorganisms**
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- XLVI. **A case of congenital fibrinogen deficiency complicated with acute lymphoblastic leukemia**
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- XLVII. **Interstitial deletion 5p14.1-p15.2 and 5q14.3-q23.2 in a patient with clubfoot, blepharophimosis, arthrogyriposis, and multiple congenital abnormalities**
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- XLVIII. **A Case of Familial Hemophagocytic Lymphohistiocytosis Type 4 With Involvement of the Central Nervous System Complicated With Infarct**
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- L. **Neuroblastoma in a Child With Wolf-Hirschhorn Syndrome**
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- L.I. **Evaluation of vitamin D prophylaxis in 3-36-month-old infants and children.**
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- L.II. **Functional Selectivity in Cytokine Signaling Revealed Through a Pathogenic EPO Mutation**
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- L.III. **Clinical Toxicity of Acute Overdoses With L-Thyroxine in Children**
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- L.IV. **The First Reported Case of Down Syndrome Co-existing with Glycogen Storage Disease**
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- L.V. **Fibromuscular Dysplasia Complicated With Cerebral Stroke in a Child With Congenital Dyserythropoietic Anemia Type II**
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- L.VI. **THE ROLE OF HLA TYPING TO DIAGNOSE MATERNAL ENGRAFTMENT SYNDROME IN PATIENTS WITH SEVERE COMBINED IMMUNE DEFICIENCY**
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- L.VII. **Comparing the Efficacy of 7%, 3% and 0.9% Saline in Moderate to Severe Bronchiolitis in Infants**
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- LVIII. Combination of two different homozygote mutations in Pompe disease**
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- LIX. Effect of parenterally L-arginine supplementation on the respiratory distress syndrome in preterm newborns**
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- LX. Late presentation of congenital diaphragmatic hernia**
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- LXI. Dystonia of neck and oculogyric dystonic reaction due to sertraline use**
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- LXII. A rare cause of recurrent respiratory tract infection: isolated absence of the right pulmonary artery**
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- LXIII. Mycophenolate mofetil-induced pseudotumor cerebri in a boy with autoimmune lymphoproliferative disease**
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- I. Comparing Pediatric and Adult Primary Chest Wall Tumors**
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- II. The influence of RANKL/osteoprotegerin on the prognosis of childhood acute lymphoblastic leukemia**
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- III. Wilms Tumor Experience of 30 years in the Single Center, Central Anatolia Region of Turkey**
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- IV. Single-center experience of childhood Hodgkin lymphoma treated without radiotherapy**
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- V. The spectrum of underlying diseases in children with autoimmune hemolytic anemia**
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- VI. Transplantation for ultra high-risk neuroblastoma patients: Effect of tandem autologous stem cell transplantation**
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