

## Assoc. Prof. ALPER ÖZCAN

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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. **Clinical spectrum of primary hemophagocytic lymphohistiocytosis: experience of reference centers in Central and Southeast Anatolia.**  
Akyol Ş., Yılmaz E., Tokgöz H., Karaman K., Pekpak E., Özcan A., Şi Mşek A., Arslan B., Ören A. C., Gökçeli H. S., et al.  
Annals of hematology, 2024 (SCI-Expanded)
- II. **DIAPH1-Deficiency is Associated with Major T, NK and ILC Defects in Humans.**  
Azizoglu Z. B., Babayeva R., Haskologlu Z. S., Acar M. B., Ayaz-Guner S., Okus F. Z., Alsavaf M. B., Can S., Basaran K. E., Canatan M. F., et al.  
Journal of clinical immunology, vol.44, no.8, pp.175, 2024 (SCI-Expanded)
- III. **Identification of the molecular etiology in rare congenital hemolytic anemias using next-generation sequencing with exome-based copy number variant analysis**  
IŞIK E., AYDINOK Y., Albayrak C., Durmus B., Karakas Z., Orhan M. F., SARPER N., Aydin S., ÜNAL S., Oymak Y., et al.  
EUROPEAN JOURNAL OF HAEMATOLOGY, vol.113, no.1, pp.82-89, 2024 (SCI-Expanded)
- IV. **Pyruvate kinase deficiency in 29 Turkish patients with two novel intronic variants**  
GÖK V., LEBLEBİSATAN G., Gokcebay D. G., GÜLER S., Dogan M. E., Bozdogan S. T., Yozgat A. K., ÖZCAN A., Sahinoglu E. P., Tokgoz H., et al.  
BRITISH JOURNAL OF HAEMATOLOGY, vol.205, no.1, pp.236-242, 2024 (SCI-Expanded)
- V. **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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- VI. **Alantolactone ameliorates graft versus host disease in mice**  
Odabas G. P., Aslan K., Suna P. A., Kendirli P. K., Erdem Ş., Çakır M., Özcan A., Yılmaz E., Karakukcu M., Donmez-Altuntas H., et al.  
INTERNATIONAL IMMUNOPHARMACOLOGY, vol.128, 2024 (SCI-Expanded)
- VII. **A Novel Biallelic LCK Variant Resulting in Profound T-Cell Immune Deficiency and Review of the Literature**  
Lanz A., Erdem S., ÖZCAN A., CEYLANER G., CANSEVER M., CEYLANER S., Conca R., Magg T., Acuto O., Latour S., et al.  
Journal of Clinical Immunology, vol.44, no.1, 2024 (SCI-Expanded)
- VIII. **A case of autoimmune lymphoproliferative syndrome with a novel de novo FAS variant**  
ÖZÇELİK F., ASLAN K., Gök V., Ari M. B., ÖZCAN A., EKEN A., Ünal E., ÖZKUL Y., DÜNDAR M.  
Pediatric Hematology and Oncology, vol.41, no.4, pp.301-309, 2024 (SCI-Expanded)
- IX. **Treatment of Infantile Fibrosarcoma in the Era of Targeted Therapies**  
Yildirim U. M., Kebudi R., Asarcikli F., Sozmen B. O. F. L. A. Z., ÜNAL E., ÖZCAN A., Zulfikar B.  
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- X. **Central nervous system thrombosis in pediatric acute lymphoblastic leukemia in Turkey: A multicenter study**  
Guzelkucuk Z., Karapınar D. Y., Gelen S. A., Tokgoz H., ÖZCAN A., Ay Y., BAHADIR A., Ozbek N. Y., Oren A. C., Ayhan A. C., et al.  
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- XI. **Defective Treg generation and increased type 3 immune response in leukocyte adhesion deficiency 1**  
Erdem S., Haskoloğlu Z. Ş., Haliloglu Y., Celikzencir H., Arik E., Keskin O., Eltan S. B., Yücel E., Canatan H., Avçılar H., et al.  
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- XII. **Monogenic early-onset lymphoproliferation and autoimmunity: Natural history of STAT3 gain-of-function syndrome.**  
Leiding J. W., Vogel T. P., Santarlas V. G. J., Mhaskar R., Smith M. R., Carisey A., Vargas-Hernandez A., Silva-Carmona M., Heeg M., Rensing-Ehl A., et al.  
The Journal of allergy and clinical immunology, vol.151, no.4, pp.1081-1095, 2023 (SCI-Expanded)
- XIII. **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?**  
Kondolot M., Yılmaz E., Sahin N. E., Ozcan A., Kaynar L., Unal E., Karakukcu M.  
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- XIV. **Haploidentical Antibody Response against Vaccine Antigens in Children after TCRab-Depleted Haploidentical Stem Cell Transplantation: Is It Similar to That in Recipients with Fully Matched Donors?**  
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- XV. **Immunodeficiency associated with a novel functionally defective variant of *SLC19A1* benefits from folinic acid treatment**  
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- XVI. **Aceruloplasminemia presenting with microcytic anemia in a Turkish boy due to a novel pathogenic variant**  
Gok V., Ozcan A., Ozer S., Karaman F., Aykutlu E., Yılmaz E., Karakukcu M., Bisgin A., Unal E.  
PEDIATRIC HEMATOLOGY AND ONCOLOGY, vol.40, no.7, pp.673-681, 2023 (SCI-Expanded)
- XVII. **Assessment of extracorporeal photopheresis related cell damage.**  
Samur B. M., KARAKÜKCÜ Ç., ÖZCAN A., ÜNAL E., YILMAZ E., KARAKÜKCÜ M.

Transfusion and apheresis science : official journal of the World Apheresis Association : official journal of the European Society for Haemapheresis, vol.61, no.6, pp.1034-72, 2022 (SCI-Expanded)

- XVIII. **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. BLOOD COAGULATION & FIBRINOLYSIS, vol.33, no.7, pp.381-388, 2022 (SCI-Expanded)
- XIX. **Characterization of cord blood CD3<sup>+</sup>TCRV $\alpha$ 7.2<sup>+</sup>CD161<sup>high</sup> T and innate lymphoid cells in the pregnancies with gestational diabetes, morbidly adherent placenta, and pregnancy hypertension diseases**  
Haliloglu Y., Ozcan A., Erdem S., Azizoglu Z. B., Bicer A., Ozarslan Ö. Y., Kilic Ö., Okus F. Z., Demir F., Canatan H., et al. AMERICAN JOURNAL OF REPRODUCTIVE IMMUNOLOGY, vol.88, no.1, 2022 (SCI-Expanded)
- XX. **A novel missense mutation outside the DNAJ domain of DNAJC21 is associated with Shwachman-Diamond syndrome**  
Alsavaf M. B., Verboon J. M., DOĞAN M. E., Azizoglu Z. B., Okus F. Z., ÖZCAN A., DÜNDAR M., EKEN A., ALTUNTAŞ H., Sankaran V. G., et al. BRITISH JOURNAL OF HAEMATOLOGY, vol.197, no.6, 2022 (SCI-Expanded)
- XXI. **Castleman Disease: A Multicenter Case Series from Turkey.**  
Gündüz E., Kırkızlar H. O., Ümit E. G., Karaman Gülsaran S., Özkocaman V., Özkalemkaş F., Candar Ö., Elverdi T., Küçükyurt S., Paydaş S., et al. Turkish journal of haematology : official journal of Turkish Society of Haematology, vol.39, pp.130-135, 2022 (SCI-Expanded)
- XXII. **Social exclusion and behavior problems in adolescents with cancer and healthy counterparts**  
SEZER EFE Y., ÖZBEY H., CANER N., ERDEM E., Kuzucu E. G., KARAKÜKCÜ M., PATIROĞLU T., ÜNAL E., YILMAZ E., ÖZCAN A. JOURNAL OF PEDIATRIC NURSING-NURSING CARE OF CHILDREN & FAMILIES, vol.64, 2022 (SCI-Expanded)
- XXIII. **A teenager boy with a novel variant of Sitosterolemia presented with pancytopenia.**  
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- XXIV. **A nonsense mutation in DIAPH1 gene presents with major T cell defects**  
Azizoglu Z. B., Okus F. Z., ÖZCAN A., Sawaf B., KÖSE M., CANÖZ Ö., GÜMÜŞ H., Ceylaner S., PATIROĞLU T., ÜNAL E., et al. EUROPEAN JOURNAL OF IMMUNOLOGY, vol.51, pp.296, 2021 (SCI-Expanded)
- XXV. **Common Variable Immunodeficiency, Autoimmune Hemolytic Anemia, and Pancytopenia Associated With a Defect in IKAROS**  
Yilmaz E., Kuehn H., Odakir E., Niemela J., Ozcan A., Eken A., Rohlf M., Cansever M., Gok V., Aydin F., et al. JOURNAL OF PEDIATRIC HEMATOLOGY ONCOLOGY, no.3, 2021 (SCI-Expanded)
- XXVI. **Comprehensive Mutation Analysis of the RAS/RAF/MEK/ERK Pathway in Paediatric Leukaemia and Significant Inferences**  
Akin-Bali D. F., AKTAŞ S. H., ÖZCAN A., YILMAZ E., AYDIN F., GÖK V., ÜNAL E., KARAKÜKCÜ M. HONG KONG JOURNAL OF PAEDIATRICS, vol.26, no.2, pp.75-87, 2021 (SCI-Expanded)
- XXVII. **Hepatitis-associated aplastic anemia in pediatric patients: single center experience.**  
Altay D., Yilmaz E., Özcan A., Karakükçü M., Ünal E., Arslan D. Transfusion and apheresis science : official journal of the World Apheresis Association : official journal of the European Society for Haemapheresis, vol.59, 2020 (SCI-Expanded)
- XXVIII. **Different Clinical Presentation of 3 Children With Familial Hemophagocytic Lymphohistiocytosis With 2 Novel Mutations.**  
Akyol S., Ozcan A., Sekine T., Chiang S., Yilmaz E., Karakurkcu M., Patiroglu T., Bryceson Y., Unal E. Journal of pediatric hematology/oncology, vol.42, 2020 (SCI-Expanded)
- XXIX. **ILC3 deficiency and generalized ILC abnormalities in DOCK8-deficient patients**  
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- XXX. **Congenital afibrinogenemia in a 4-year-old girl complicated with acute lymphoblastic leukemia**  
ÖZCAN A., Samur B., AKYOL Ş., Erdogmus N. A., PATIROĞLU T., KARAKÜKCÜ M., ÜNAL E.  
TURKISH JOURNAL OF PEDIATRICS, vol.62, no.2, pp.289-292, 2020 (SCI-Expanded)
- XXXI. **Twenty children with non-Wilms renal tumors from a reference center in Central Anatolia, Turkey**  
Ünal E., Yilmaz E., Özcan A., Işık B., Karakükcü M., Turan C., Akgün H., Öztürk F., Coşkun A., Özdemir M., et al.  
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- XXXII. **Invasive Fungal Infections in Children with Acute Lymphoblastic Leukemia: Experience from a Reference University Hospital in Cappadocia**  
YILMAZ E., Erdogmus A., ÖZCAN A., GÖRKEM S. B., Ceylan O., DENİZ K., ÜNAL E., ATALAY M. A., KARAKÜKCÜ M., KOÇ A. N., et al.  
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- XXXIII. **CD137 deficiency causes immune dysregulation with predisposition to lymphomagenesis**  
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- XXXIV. **Two Different Endocrine Cancer, One Disease; DICER-1 Mutation**  
Tatlı Z. U., Direk G., ÖZCAN A., HATİPOĞLU N., KENDİRCİ M., Kurtoglu S.  
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- XXXV. **A rare cause of vomiting in an adolescent: gastric Burkitt's lymphoma**  
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TURKISH JOURNAL OF PEDIATRICS, vol.61, no.3, pp.431-435, 2019 (SCI-Expanded)
- XXXVI. **Propranolol treatment for chylothorax due to diffuse lymphangiomas**  
HANGÜL M., KÖSE M., ÖZCAN A., ÜNAL E.  
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- XXXVII. **Genetic Deficiency and Biochemical Inhibition of ITK Affect Human Th17, Treg, and Innate Lymphoid Cells**  
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- XXXVIII. **The relationship between the prognosis of children with acute arterial stroke and polymorphisms of CDKN2B, HDAC9, NINJ2, NAA25 genes**  
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- XXXIX. **A Rare Case of Activated Phosphoinositide 3-Kinase Delta Syndrome (APDS) Presenting With Hemophagocytosis Complicated With Hodgkin Lymphoma.**  
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- XL. **A mummy emerges from the grave: Scurvy confounding the clinical presentation of a child with Fanconi anemia**  
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AMERICAN JOURNAL OF HEMATOLOGY, vol.94, no.4, pp.506-507, 2019 (SCI-Expanded)
- XLI. **Scurvy: A rare cause of arthritis in a child with neurologic disorder.**  
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European journal of rheumatology, vol.5, pp.283-284, 2018 (SCI-Expanded)
- XLII. **Tumour Lysis Syndrome in Children with Hematologic Malignancies: Single Center Experience**  
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- XLIII. **Vena Cava Superior Syndrome in Children with Mediastinal Tumors: Single Center Experience**  
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- XLIV. **DOES EVEN LOW DOSE ENALAPRIL CAUSE LIFE THREATENING ACUTE KIDNEY INJURY?**  
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- XLV. **Cerebral Sinus Venous Thrombosis and Prothrombotic Risk Factors in Children: A Single-Center Experience From Turkey**  
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- XLVI. **ALLOGENEIC BONE MARROW TRANSPLANTATION IN THREE CASES WITH DOCK 8 DEFICIENCY**  
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- XLVII. **The relationship between hematological parameters and prognosis of children with acute ischemic stroke**  
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- XLVIII. **Use of a chlorhexidine-impregnated dressing reduced catheter-related bloodstream infections caused by Gram-positive microorganisms**  
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- XLIX. **A case of congenital fibrinogen deficiency complicated with acute lymphoblastic leukemia**  
ÖZCAN A., SAMUR M. B., ERDOĞMUŞ A., PATIROĞLU T., KARAKÜKCÜ M., ÜNAL E.  
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- L. **Interstitial deletion 5p14.1-p15.2 and 5q14.3-q23.2 in a patient with clubfoot, blepharophimosis, arthrogyposis, and multiple congenital abnormalities**  
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AMERICAN JOURNAL OF MEDICAL GENETICS PART A, vol.173, no.10, pp.2798-2802, 2017 (SCI-Expanded)
- LI. **A Case of Familial Hemophagocytic Lymphohistiocytosis Type 4 With Involvement of the Central Nervous System Complicated With Infarct**  
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- LII. **INHERITED PROTHROMBOTIC RISK FACTORS IN TURKISH CHILDREN WITH HEREDITARY ANGIOEDEMA. SINGLE CENTER EXPERIENCE**  
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- LIII. **Neuroblastoma in a Child With Wolf-Hirschhorn Syndrome**  
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- LIV. **Evaluation of vitamin D prophylaxis in 3-36-month-old infants and children.**  
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- LV. **Functional Selectivity in Cytokine Signaling Revealed Through a Pathogenic EPO Mutation**  
Kim A. R., Ulirsch J. C., WILMES S., ÜNAL E., Moraga I., KARAKÜKCÜ M., YUAN D., KAZEROUNIAN S., Abdulhay N. J., King D. S., et al.  
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- LVI. **Clinical Toxicity of Acute Overdoses With L-Thyroxine in Children**  
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- LVII. **The First Reported Case of Down Syndrome Co-existing with Glycogen Storage Disease**  
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- LVIII. Fibromuscular Dysplasia Complicated With Cerebral Stroke in a Child With Congenital Dyserythropoietic Anemia Type II**  
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- LIX. THE ROLE OF HLA TYPING TO DIAGNOSE MATERNAL ENGRAFTMENT SYNDROME IN PATIENTS WITH SEVERE COMBINED IMMUNE DEFICIENCY**  
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- LX. Combination of two different homozygote mutations in Pompe disease**  
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- LXI. Comparing the Efficacy of 7%, 3% and 0.9% Saline in Moderate to Severe Bronchiolitis in Infants**  
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- LXII. Effect of parenterally L-arginine supplementation on the respiratory distress syndrome in preterm newborns**  
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- LXIII. Late presentation of congenital diaphragmatic hernia**  
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- LXIV. Dystonia of neck and oculogyric dystonic reaction due to sertraline use**  
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- LXV. A rare cause of recurrent respiratory tract infection: isolated absence of the right pulmonary artery**  
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- LXVI. Mycophenolate mofetil-induced pseudotumor cerebri in a boy with autoimmune lymphoproliferative disease**  
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## Articles Published in Other Journals

- I. **30 Years of Wilms Tumor Experience at One Center in Türkiye's Central Anatolia Region**  
AKYOL Ş., ODABAŞ G. P., ÖZCAN A., YILMAZ E., KARAMAN Z. F., ÖZTÜRK F., Akgun H., DOĞAN A. B., EROĞLU C., Unal E., et al.  
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## Metrics

Publication: 205

Citation (WoS): 417

Citation (Scopus): 466

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H-Index (Scopus): 9

## Non Academic Experience

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ENERJİ VE TABİİ KAYNAKLAR BAKANLIĞI