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Araştırma Alanları

Pediatrik Hematoloji, Pediatrik Onkoloji

SCI, SSCI ve AHCI İndekslerine Giren Dergilerde Yayınlanan Makaleler

I. **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.
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II. **A case of autoimmune lymphoproliferative syndrome with a novel de novo FAS variant**
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III. **Immunodeficiency associated with a novel functionally defective variant of SLC19A1 benefits from folinic acid treatment**
Gök V., Erdem Ş., Haliloğlu Y., Bişgin A., Belkaya S., Başaran K. E., Canatan M. F., Özcan A., Yılmaz E., Acıpayam C., et al.
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IV. **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., Yilmaz E., Karakukcu M., Bisgin A., Unal E.
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V. **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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VI. **A teenager boy with a novel variant of Sitosterolemia presented with pancytopenia.**
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VII. **Evolution and long-term outcomes of combined immunodeficiency due to CARMIL2 deficiency**
Kocamış B., Baser D., Akcam B., Danielson J., Eltan S. B., Haliloglu Y., Sefer A. P., Babayeva R., Akgun G., Charbonnier L., et al.

ALLERGY, cilt.77, sa.3, ss.1004-1019, 2022 (SCI-Expanded)

VIII. **NOVEL VARIANTS IN ETV6 AND FYB1 GENES: TWO CHILDREN WITH RARE INHERITED THROMBOCYTOPENIA**
Gok V., Karaca M., Ozcan A., Yilmaz E., Bozdogan S. T., Bisgin A., Karakukcu M., Unal E.
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IX. **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., Rohlfs M., Cansever M., Gok V., Aydin F., et al.
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X. **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.
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XI. **HEMOPHILIA INTERNATIONAL ACCREDITATION TRAINING PLAN IN ERCIYES UNIVERSITY PEDIATRIC HEMATOLOGY CLINIC**
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XII. **A CHILD WHO DEVELOPED LUPUS ANTICOAGULANT HYPOPROTHROMBINEMIA SYNDROME AFTER VIRAL GASTROENTERITIS**
Gok V., Simsek N., Aydin F., Ozcan A., Yilmaz E., Unal E., Karakukcu M., Patiroglu T.
HAEMOPHILIA, ss.148-149, 2021 (SCI-Expanded)

XIII. **The Spectrum of Underlying Diseases in Children with Torticollis**
Tumturk A., Gorkem S. B., Ozmansur E. N., Peduk Y., Arslan U., Gok V., Dogan H., Cetin B. Ş., Sahin A., Gumus H., et al.
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Gok V., Isik E., Yilmaz E., Aydin F., Ozcan A., Unal E., Karakukcu M., Atik T., Patiroglu T.
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Diğer Dergilerde Yayınlanan Makaleler

I. **Rare Coagulation Factor Deficiencies: Multicenter Experience With 188 Cases**
GÖK V., Sahinoglu E. P., Tokgoz H., Mutlu F. T., Acipayam C., Karaman K., Tuncel D. A., Oren A. C., Simsek A., Arslan B., et al.
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II. **Changes in Body Mass Index and Height Among Survivors of Childhood Acute Lymphoblastic Leukemia: A Single Institutional Experience**
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III. **A Fatal Case of Familial Hemophagocytic Lymphohistiocytosis Associated with Fusarium Infection and Rare Mutation**
DÜNDAR M. A., ORAK F., ACIPAYAM C., Aslan K., Gök V., ÇETİN B. Ş., ÜNAL E., KLEIN C., AKYILDIZ B.
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Akyol Ş., Göl D. K., Yılmaz E., Karaman Z. F., Özcan A., Küçük A., Gök V., Aydın F., Per H., Karakükcü M., et al.
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V. **Triosephosphate isomerase deficiency in an infant**
Karaca M., Gök V., Sarı Ü. A., Dündar M. A., Aydın F., Özcan A., Yılmaz E., Canpolat M., Ceylaner S., Kardaş F., et al.
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VI. **The Significance of soluble CD40 and CD40 ligand levels in childhood acute lymphoblastic leukemia patients**
Gök V., Özcan A., Yılmaz E., Karakükcü M., Patıroğlu T., Özdemir M. A., Ünal E.
Journal of Translational and Practical Medicine, cilt.1, sa.2, ss.49-56, 2022 (Hakemli Dergi)

VII. **Comprehensive approach to hemophilia**
Gök V., Ünal E.
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VIII. **Etiological Factors of Opsoclonus Myoclonus Ataxia Syndrome: A Single Center Experience with Eight Children**
Gök V., Gümüş G., Durmuş H., Ünal E., Gümüş H., Karakükcü M., Bayram A., Per H.
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IX. **Type 2B Von Willebrand Disease Mimicking Autoimmune Thrombocytopenia in the Neonatal Period**
GÖK V., IŞIK E., YILMAZ E., AYDIN F., ÖZCAN A., ÜNAL E., KARAKÜKCÜ M., ATİK T., PATIROĞLU T.
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X. **Two Siblings Followed Up for Hereditary Multiple Exostoses**
Erol M., Yigit O., Adanir O., Toksoz M., Narin H., GÖK V., Borakay D., Konya M.
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Hakemli Kongre / Sempozyum Bildiri Kitaplarında Yer Alan Yayınlar

I. **SEIZURE AS A RARE SIDE EFFECT OF L-ASPARAGINASE INFUSION IN A LEUKEMIC CHILD WITH DOWN SYNDROME**
GÖK V., AYDIN F., ÖZCAN A., YILMAZ E., ACER H., GÖRKEM S. B., Erdoğmuş N. A., ÜNAL E., KARAKÜKCÜ M., PATIROĞLU T.
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II. **COMBINATION OF TWO RARE GENETIC DISEASES OF FANCONI APLASTIC ANEMIA AND 46,X,DEL(X)(Q23) IN A TURKISH GIRL**
GÖK V., DOĞAN M. E., YILMAZ E., ÖZCAN A., TORUN Y., AYDIN F., PER H., ÜNAL E., KARAKÜKCÜ M., PATIROĞLU T.
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III. **Afibrinogenemia Experience: Multicenter Study**
TUNCEL D. A., GÖK V., PEKPAK E., TOKGÖZ H., ACIPAYAM C., YILMAZ E., ÖREN A. C., ÖZCAN A., KARAKÜKCÜ M., ÜNAL E., et al.
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Metrikler

Yayın: 32
Atıf (WoS): 27

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