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Kişisel Bilgiler

Uluslararası Araştırmacı ID'leri

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Eğitim Bilgileri

Tıpta Yandal Uzmanlık, Çukurova Üniversitesi, Çocuk Sağlığı Ve Hastalıkları , Çocuk Romatoloji, Türkiye 2016 - Devam Ediyor

Tıpta Uzmanlık, Mersin Üniversitesi, Çocuk Sağlığı Ve Hastalıkları , Türkiye 2004 - 2010

Lisans, Van Yüzüncü Yıl Üniversitesi, Tıp Fakültesi, Türkiye 1997 - 2003

Yabancı Diller

İngilizce, B2 Orta Üstü

Yaptığı Tezler

Tıpta Uzmanlık, TNF- α -308, IL-10-1082 gen polimorfizmlerinin romatizmal kalp hastalığının şiddeti ve duyarlılığı ile ilişkisi, Mersin Üniversitesi, Çocuk Sağlığı Ve Hastalıkları, Çocuk Sağlığı Ve Hastalıkları Anabilim Dalı, 2010

Araştırma Alanları

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

Akademik Unvanlar / Görevler

Uzman, Çukurova Üniversitesi, Tıp Fakültesi, Çocuk Romatoloji, 2016 - Devam Ediyor

SCI, SSCI ve AHCI İndekslerine Giren Dergilerde Yayınlanan Makaleler

- Differentiating children with familial Mediterranean fever from other recurrent fever syndromes: The utility of new Eurofever/PRINTO classification criteria**
KIŞLA EKİNCİ R. M., Balci S., Erol A. H., Karagoz D., ALTINTAŞ D. U., BIŞGİN A.
ARCHIVES OF RHEUMATOLOGY, cilt.36, sa.4, ss.493-498, 2021 (SCI-Expanded)
- Clinical features of childhood uveitis at a tertiary referral center in Southern Turkey**
Esen E., Sızmaz S., Balci S., Kışla Ekinci R. M., Demircan N.
INTERNATIONAL OPHTHALMOLOGY, cilt.41, sa.6, ss.2073-2081, 2021 (SCI-Expanded)
- Camptodactyly-Arthropathy-Coxa Vara-Pericarditis Syndrome Resembling Juvenile Idiopathic Arthritis: A Single-Center Experience from Southern Turkey**

KIŞLA EKİNCİ R. M., Balci S., Dogan H., Ceylaner S., Varan C., ALINÇ ERDEM S., Coban F., BİŞGİN A.
MOLECULAR SYNDROMOLOGY, cilt.12, sa.2, ss.112-117, 2021 (SCI-Expanded)

- IV. **Altered expression of apoptosis-related, circulating cell-free miRNAs in children with familial Mediterranean fever: a cross-sectional study**
Karpuzoglu E. M., KIŞLA EKİNCİ R. M., Balci S., BİŞGİN A., YILMAZ M.
RHEUMATOLOGY INTERNATIONAL, cilt.41, sa.1, ss.103-111, 2021 (SCI-Expanded)
- V. **Twenty-Year Experience of a Single Referral Center on Pediatric Familial Mediterranean Fever What Has Changed Over the Last Decade?**
KIŞLA EKİNCİ R. M., Balci S., Dogruel D., ALTINTAŞ D. U., YILMAZ M.
JCR-JOURNAL OF CLINICAL RHEUMATOLOGY, cilt.27, sa.1, ss.18-24, 2021 (SCI-Expanded)
- VI. **Evaluation of different classification criteria in children with Behcet disease: results from a single referral center**
KIŞLA EKİNCİ R. M., ESEN E., Erol A. H., SIZMAZ S., Karagoz D., ALTINTAŞ D. U., Balci S.
EXPERT REVIEW OF CLINICAL IMMUNOLOGY, cilt.16, sa.11, ss.1093-1097, 2020 (SCI-Expanded)
- VII. **Clinical manifestations and outcomes of 420 children with Henoch Schonlein Purpura from a single referral center from Turkey: A three-year experience**
KIŞLA EKİNCİ R. M., Balci S., MELEK E., KARABAY BAYAZIT A., DOĞRUDEL D., ALTINTAŞ D. U., YILMAZ M.
MODERN RHEUMATOLOGY, cilt.30, sa.6, ss.1039-1046, 2020 (SCI-Expanded)
- VIII. **Retrospective Analysis of the Factors Affecting Growth Parameters in Turkish Children With Systemic Lupus Erythematosus**
Balci S., KIŞLA EKİNCİ R. M., MELEK E., KARABAY BAYAZIT A., Dogruel D., ALTINTAŞ D. U., YILMAZ M.
ARCHIVES OF RHEUMATOLOGY, cilt.35, sa.3, ss.357-365, 2020 (SCI-Expanded)
- IX. **Different clinical manifestations of three prime repair exonuclease 1 mutation: A case series**
İNCECİK F., Balci S., KIŞLA EKİNCİ R. M., Herguner O. M., BİŞGİN A., YILMAZ M.
ANNALS OF INDIAN ACADEMY OF NEUROLOGY, cilt.23, sa.5, ss.699-703, 2020 (SCI-Expanded)
- X. **Growth Parameters of Turkish Children With an Autoinflammatory Disease Before and After Canakinumab Treatment**
Balci S., KIŞLA EKİNCİ R. M., Dogruel D., ALTINTAŞ D. U., YILMAZ M.
INDIAN PEDIATRICS, cilt.57, sa.7, ss.637-640, 2020 (SCI-Expanded)
- XI. **Pre-Pulseless Takayasu Arteritis in a Child Represented With Prolonged Fever of Unknown Origin and Successful Management With Concomitant Mycophenolate Mofetil and Infliximab**
KIŞLA EKİNCİ R. M., Balci S., Piskin F. C., Varan C., ALINÇ ERDEM S., YILMAZ M.
ARCHIVES OF RHEUMATOLOGY, cilt.35, sa.2, ss.278-282, 2020 (SCI-Expanded)
- XII. **Phenotypic variability in two patients with tumor necrosis factor receptor associated periodic fever syndrome emphasizes a rare manifestation: Immunoglobulin A nephropathy**
Balci S., KIŞLA EKİNCİ R. M., MELEK E., ATMIŞ B., BİŞGİN A., YILMAZ M.
EUROPEAN JOURNAL OF MEDICAL GENETICS, cilt.63, sa.4, 2020 (SCI-Expanded)
- XIII. **Unicentric Castleman Disease Mimicking an Autoinflammatory Disorder: A Diagnostic Challenge in a Pediatric Patient With Recurrent Fever**
KIŞLA EKİNCİ R. M., Balci S., ERGİN M., KÜPELİ S., BAYRAM İ., YILMAZ M., KOCABAŞ E.
JOURNAL OF PEDIATRIC HEMATOLOGY ONCOLOGY, cilt.42, sa.3, ss.204-207, 2020 (SCI-Expanded)
- XIV. **Baricitinib experience on STING-associated vasculopathy with onset in infancy: A representative case from Turkey**
Balci S., KIŞLA EKİNCİ R. M., de Jesus A. A., Goldbach-Mansky R., YILMAZ M.
CLINICAL IMMUNOLOGY, cilt.212, ss.113-116, 2020 (SCI-Expanded)
- XV. **Deficiency of adenosine deaminase 2: a case series revealing clinical manifestations, genotypes and treatment outcomes from Turkey**
KIŞLA EKİNCİ R. M., Balci S., Hershfield M., BİŞGİN A., DOĞRUDEL D., ALTINTAŞ D. U., YILMAZ M.
RHEUMATOLOGY, cilt.59, sa.1, ss.254-256, 2020 (SCI-Expanded)
- XVI. **Canakinumab in Children with Familial Mediterranean Fever: A Single-Center, Retrospective Analysis**
KIŞLA EKİNCİ R. M., Balci S., DOĞRUDEL D., ALTINTAŞ D. U., YILMAZ M.

- PEDIATRIC DRUGS, cilt.21, sa.5, ss.389-395, 2019 (SCI-Expanded)
- XVII. **A homozygote novel L451W mutation in CECR1 gene causes deficiency of adenosine deaminase 2 in a pediatric patient representing with chronic lymphoproliferation and cytopenia**
EKINCI R. M. K., Balci S., BİŞGİN A., ŞAŞMAZ I., LEBLEBİSATAN G., İNCECİK F., YILMAZ M.
PEDIATRIC HEMATOLOGY AND ONCOLOGY, cilt.36, sa.6, ss.376-381, 2019 (SCI-Expanded)
- XVIII. **Disease Severity and Genotype Affect Physical Growth in Children With Familial Mediterranean Fever**
KIŞLA EKİNCİ R. M., Balci S., Akay E., DOĞRUDEL D., ALTINTAŞ D. U., YILMAZ M.
ARCHIVES OF RHEUMATOLOGY, cilt.34, sa.3, ss.288-293, 2019 (SCI-Expanded)
- XIX. **Etanercept for the Treatment of Chronic Arthritis Related to Chronic Granulomatous Disease: A Case**
Balci S., KIŞLA EKİNCİ R. M., SERBES M., DOĞRUDEL D., ALTINTAŞ D. U., YILMAZ M.
PEDIATRIC ALLERGY IMMUNOLOGY AND PULMONOLOGY, cilt.32, sa.3, ss.131-134, 2019 (SCI-Expanded)
- XX. **Hyperimmunoglobulinemia D syndrome with recurrent perianal abscess successfully treated with canakinumab**
KIŞLA EKİNCİ R. M., Balci S., BİŞGİN A., TÜMGÖR G., DOĞRUDEL D., YILMAZ M.
SCOTTISH MEDICAL JOURNAL, cilt.64, sa.3, ss.103-107, 2019 (SCI-Expanded)
- XXI. **Do practical laboratory indices predict the outcomes of children with Henoch-Schonlein purpura?**
EKINCI R. M. K., Balci S., GOKAY S. S., YILMAZ H. L., DOĞRUDEL D., ALTINTAŞ D. U., YILMAZ M.
POSTGRADUATE MEDICINE, cilt.131, sa.4, ss.295-298, 2019 (SCI-Expanded)
- XXII. **Juvenile systemic lupus erythematosus: a single-center experience from southern Turkey**
Balci S., Ekinci R., Bayazit A., Melek E., Dogruel D., Altintas D. U., Yilmaz M.
CLINICAL RHEUMATOLOGY, cilt.38, sa.5, ss.1459-1468, 2019 (SCI-Expanded)
- XXIII. **Recurrent Henoch Schonlein purpura without renal involvement successfully treated with methotrexate**
KIŞLA EKİNCİ R. M., Balci S., SERBES M., Aycin G. D., DOĞRUDEL D., ALTINTAŞ D. U., YILMAZ M.
SCOTTISH MEDICAL JOURNAL, cilt.64, sa.2, ss.74-77, 2019 (SCI-Expanded)
- XXIV. **Autoimmune Manifestations in Heterozygote Type I Complement 2 Deficiency: A Child Eventually Diagnosed With Systemic Lupus Erythematosus**
KIŞLA EKİNCİ R. M., Balci S., BİŞGİN A., ATMIŞ B., DOĞRUDEL D., YILMAZ M.
ARCHIVES OF RHEUMATOLOGY, cilt.34, sa.1, ss.96-99, 2019 (SCI-Expanded)
- XXV. **Frequency of functional gastrointestinal disorders in children with familial Mediterranean fever**
Ekinci R. M., Balci S., AKAY E., TUMGOR G., DOGRUEL D., Altintas D. U., Yilmaz M.
CLINICAL RHEUMATOLOGY, cilt.38, sa.3, ss.921-926, 2019 (SCI-Expanded)
- XXVI. **Hyperphosphatemic Familial Tumoral Calcinosis in Two Siblings with a Novel Mutation in *GALNT3* Gene: Experience from Southern Turkey**
Kışla E., Gürbüz F., BALCI S., Bişgin A., TAŞTAN M., Yüksel B., Yılmaz M.
Journal of clinical research in pediatric endocrinology, cilt.11, ss.94-99, 2019 (SCI-Expanded)
- XXVII. **Is Henoch-Schonlein purpura a susceptibility factor for functional gastrointestinal disorders in children?**
KIŞLA EKİNCİ R. M., Balci S., Mart O. O., TÜMGÖR G., Yavuz S., Celik H., DOĞRUDEL D., ALTINTAŞ D. U., YILMAZ M.
RHEUMATOLOGY INTERNATIONAL, cilt.39, sa.2, ss.317-322, 2019 (SCI-Expanded)
- XXVIII. **MEW gene variants in children with Henoch-Schonlein purpura and association with clinical manifestations: a single-center Mediterranean experience**
Ekinci R. M., Balci S., Bisgin A., Atmis B., Dogruel D., Altintas D. U., Yilmaz M.
POSTGRADUATE MEDICINE, cilt.131, sa.1, ss.68-72, 2019 (SCI-Expanded)
- XXIX. **Renal Amyloidosis in Deficiency of Adenosine Deaminase 2: Successful Experience With Canakinumab**
KIŞLA EKİNCİ R. M., Balci S., BİŞGİN A., Hershfield M., ATMIŞ B., DOĞRUDEL D., YILMAZ M.
PEDIATRICS, cilt.142, sa.5, 2018 (SCI-Expanded)
- XXX. **Decreased serum vitamin B-12 and vitamin D levels affect sleep quality in children with familial Mediterranean fever**

Ekinci R., Balci S., Serbes M., Dogruel D., Altintas D. U., Yilmaz M.
RHEUMATOLOGY INTERNATIONAL, cilt.38, sa.1, ss.83-87, 2018 (SCI-Expanded)

XXXI. A homozygote TREX1 mutation in two siblings with different phenotypes: Chilblains and cerebral vasculitis.

Kisla E., Balci S., Bisgin A., Altintas D. U., Yilmaz M.
European journal of medical genetics, cilt.60, ss.690-694, 2017 (SCI-Expanded)

XXXII. The Influence of Concomitant Disorders on Disease Severity of Familial Mediterranean Fever in Children.

Kişla E., Balci S., Ufuk A., Yilmaz M.
Archives of rheumatology, cilt.33, ss.282-287, 2017 (SCI-Expanded)

Diğer Dergilerde Yayınlanan Makaleler

- I. **Complement C2 polymorphisms in children with Henoch Schonlein Purpura**
KIŞLA EKİNCİ R. M., Balci S., ATMIŞ B., KARABAY BAYAZIT A., DOĞRUDEL D., ALTINTAŞ D. U., BIŞGIN A.
CUKUROVA MEDICAL JOURNAL, cilt.45, sa.1, ss.89-95, 2020 (ESCI)
- II. **The frequency of avascular necrosis in juvenile systemic lupus erythematosus**
Balci S., Kişla Ekinci R. M., Piskin F. C., Melek E., Atmış B., Doğruel D., Altıntaş D. U., Karabay Bayazıt A.
CUKUROVA MEDICAL JOURNAL, cilt.45, ss.200-207, 2020 (ESCI)
- III. **Growth parameters of Turkish children with juvenile idiopathic arthritis**
Balci S., Calkan M., Dilek S. O., Dogruel D., Altintas D. U., KIŞLA EKİNCİ R. M.
CUKUROVA MEDICAL JOURNAL, cilt.45, sa.2, ss.495-501, 2020 (ESCI)
- IV. **Does selective IgA deficiency have a good prognostic role on juvenile dermatomyositis? a case report**
Balci S., KIŞLA EKİNCİ R. M., DOĞRUDEL D., SERBES M., ALTINTAŞ D. U., YILMAZ M.
CUKUROVA MEDICAL JOURNAL, cilt.44, sa.4, ss.1511-1514, 2019 (ESCI)
- V. **Symptomatic multifocal avascular necrosis in an adolescent with neuropsychiatric systemic lupus erythematosus**
Ekinci R. M., Balci S., ÇELİK G., DOĞRUDEL D., ALTINTAŞ D. U., YILMAZ M.
Reumatologia, cilt.57, sa.3, ss.182-187, 2019 (Scopus)
- VI. **Çocukluk çağı Miks Konnektif Doku Hastalığı: tek merkez deneyimi**
KIŞLA EKİNCİ R. M., BALCI S., ALTINTAŞ D. U., YILMAZ M.
CUKUROVA MEDICAL JOURNAL, cilt.43, sa.2, ss.1, 2018 (Hakemli Dergi)
- VII. **Juvenile mixed connective tissue disease: case series**
KIŞLA EKİNCİ R. M., Balci S., Altintas D. U., YILMAZ M.
CUKUROVA MEDICAL JOURNAL, cilt.43, sa.1, ss.235-240, 2018 (ESCI)
- VIII. **Juvenile idiopathic inflammatory myopathy: single center data**
Balci S., Ekinci R. M. K., ALTINTAŞ D. U., YILMAZ M.
CUKUROVA MEDICAL JOURNAL, cilt.43, sa.3, ss.685-691, 2018 (ESCI)

Hakemli Kongre / Sempozyum Bildiri Kitaplarında Yer Alan Yayınlar

- I. **Genotype-phenotype correlation of patients with familial mediterranean fever in childhood: Experience of a single center**
DOĞRUDEL D., KIŞLA EKİNCİ R. M., BALCI S., YILMAZ M., ALTINTAŞ D. U.
9. International Congress of Familial Mediterranean Fever and Systemic Auto Inflammatory Diseases, Kıbrıs (Kkctc),
4 - 07 Mayıs 2017

Desteklenen Projeler

KIŞLA EKİNCİ R. M., BIŞGİN A., ALTINTAŞ D. U., YILMAZ M., BALCI S., DOĞRUEL D., Yükseköğretim Kurumları Destekli Proje, Henoch Schönlein Purpuralı hastalarda C2 gen polimorfizm sıklığı ve klinik bulgulara etkisi, 2018 - 2019

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Atf (Scopus): 166

H-İndeks (WoS): 6

H-İndeks (Scopus): 8