

## Personal Information

### International Researcher IDs

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## Education Information

Post Doctorate of Medicine, Cukurova University, Çocuk Sağlığı Ve Hastalıkları , Çocuk Romatoloji, Turkey 2016 - Continues

Expertise In Medicine, Mersin University, Çocuk Sağlığı Ve Hastalıkları , Turkey 2004 - 2010

Undergraduate, Van Yüzüncü Yil University, Tıp Fakültesi, Turkey 1997 - 2003

## Foreign Languages

English, B2 Upper Intermediate

## Dissertations

Expertise In Medicine, TNF- $\alpha$ -308, IL-10-1082 gen polimorfizmlerinin romatizmal kalp hastalığının şiddeti ve duyarlılığı ile ilişkisi, Mersin University, Çocuk Sağlığı Ve Hastalıkları, Çocuk Sağlığı Ve Hastalıkları Anabilim Dalı, 2010

## Research Areas

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

## Academic Titles / Tasks

Expert, Cukurova University, Tıp Fakültesi, Çocuk Romatoloji, 2016 - Continues

## Published journal articles indexed by SCI, SSCI, and AHCI

- I. 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., BİŞGİN A.  
ARCHIVES OF RHEUMATOLOGY, vol.36, no.4, pp.493-498, 2021 (SCI-Expanded)
- II. 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, vol.41, no.6, pp.2073-2081, 2021 (SCI-Expanded)
- III. 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, vol.12, no.2, pp.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, vol.41, no.1, pp.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, vol.27, no.1, pp.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, vol.16, no.11, pp.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., DOĞRUEL D., ALTINTAŞ D. U., YILMAZ M.  
MODERN RHEUMATOLOGY, vol.30, no.6, pp.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, vol.35, no.3, pp.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, vol.23, no.5, pp.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, vol.57, no.7, pp.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, vol.35, no.2, pp.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, vol.63, no.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, vol.42, no.3, pp.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, vol.212, pp.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., DOĞRUEL D., ALTINTAŞ D. U., YILMAZ M.  
RHEUMATOLOGY, vol.59, no.1, pp.254-256, 2020 (SCI-Expanded)
- XVI. **Canakinumab in Children with Familial Mediterranean Fever: A Single-Center, Retrospective Analysis**  
KIŞLA EKİNCİ R. M., Balci S., DOĞRUEL D., ALTINTAŞ D. U., YILMAZ M.

PEDIATRIC DRUGS, vol.21, no.5, pp.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, vol.36, no.6, pp.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, vol.34, no.3, pp.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, vol.32, no.3, pp.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, vol.64, no.3, pp.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, vol.131, no.4, pp.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, vol.38, no.5, pp.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, vol.64, no.2, pp.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, vol.34, no.1, pp.96-99, 2019 (SCI-Expanded)
- XXV. **Frequency of functional gastrointestinal disorders in children with familial Mediterranean fever**  
Ekinci R. M., Balci S., AKAY E., Tumor G., DOGRUEL D., Altintas D. U., Yilmaz M.  
CLINICAL RHEUMATOLOGY, vol.38, no.3, pp.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, vol.11, pp.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, vol.39, no.2, pp.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, vol.131, no.1, pp.68-72, 2019 (SCI-Expanded)
- XXIX. **Renal Amyloidosis in Deficiency of Adenosine Deaminase 2: Successful Experience With Canakinumab**  
Kışla Ekinci R. M., Balci S., Bişgin A., Hershfield M., Atmiş B., Doğruel D., Yılmaz M.  
PEDIATRICS, vol.142, no.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, vol.38, no.1, pp.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, vol.60, pp.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, vol.33, pp.282-287, 2017 (SCI-Expanded)

## Articles Published in Other Journals

- I. **The adherence of parents regarding epinephrine auto-injector use in anaphylaxis management: a real-life study**  
Serbes M., Sasihuseyinoglu A. S., ÖZCAN D., Kisla Ekinci R. M., BALCI S., Altintas D. U.  
Cukurova Medical Journal, vol.48, no.1, pp.234-242, 2023 (ESCI)
- II. **Retrospective analyzes of adverse events during biologic agents in children with juvenile idiopathic arthritis from a single center in Turkey**  
BALCI ÇETİNKAYA S., DEMİR İ., SERBES M., ÖZCAN D., ALTINTAŞ D. U., KIŞLA EKİNCİ R. M.  
RHEUMATOLOGIA, vol.58, no.6, pp.367-374, 2020 (Scopus)
- III. **Frequency of functional gastrointestinal disorders in children with juvenile idiopathic arthritis**  
BALCI ÇETİNKAYA S., KIŞLA EKİNCİ R. M., DEMİR İ., SERBES M., ÖZCAN D., ALTINTAŞ D. U., TÜMGÖR G.  
CUKUROVA MEDICAL JOURNAL, vol.45, no.4, pp.1768-1774, 2020 (ESCI)
- IV. **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 Bayazit A.  
CUKUROVA MEDICAL JOURNAL, vol.45, pp.200-207, 2020 (ESCI)
- V. **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, vol.45, no.2, pp.495-501, 2020 (ESCI)
- VI. **Complement C2 polymorphisms in children with Henoch Schonlein Purpura**  
KIŞLA EKİNCİ R. M., Balci S., ATMIŞ B., KARABAY BAYAZIT A., DOĞRUEL D., ALTINTAŞ D. U., BİŞGİN A.  
CUKUROVA MEDICAL JOURNAL, vol.45, no.1, pp.89-95, 2020 (ESCI)
- VII. **Symptomatic multifocal avascular necrosis in an adolescent with neuropsychiatric systemic lupus erythematosus**  
Ekinci R. M., Balci S., ÇELİK G., DOĞRUEL D., ALTINTAŞ D. U., YILMAZ M.  
Reumatologia, vol.57, no.3, pp.182-187, 2019 (Scopus)
- VIII. **Does selective IgA deficiency have a good prognostic role on juvenile dermatomyositis? a case report**  
Balci S., KIŞLA EKİNCİ R. M., DOĞRUEL D., SERBES M., ALTINTAŞ D. U., YILMAZ M.  
CUKUROVA MEDICAL JOURNAL, vol.44, no.4, pp.1511-1514, 2019 (ESCI)
- IX. **Çocukluk çağı Miks Konnektif Doku Hastalığı: tek merkez deneyimi**  
KIŞLA EKİNCİ R. M., BALCI S., ALTINTAŞ D. U., YILMAZ M.  
CUKUROVA MEDICAL JOURNAL, vol.43, no.2, pp.1, 2018 (Peer-Reviewed Journal)
- X. **Juvenile mixed connective tissue disease: case series**  
KIŞLA EKİNCİ R. M., Balci S., Altintas D. U., YILMAZ M.  
CUKUROVA MEDICAL JOURNAL, vol.43, no.1, pp.235-240, 2018 (ESCI)
- XI. **Juvenile idiopathic inflammatory myopathy: single center data**  
Balci S., Ekinci R. M. K., ALTINTAŞ D. U., YILMAZ M.  
CUKUROVA MEDICAL JOURNAL, vol.43, no.3, pp.685-691, 2018 (ESCI)

## Books & Book Chapters

### I. Juvenil Dermatomiyozi

KIŞLA EKİNCİ R. M., BALCI ÇETİNKAYA S.

in: Klinik Romatoloji, Kısacık Bünyamin, Bes Cemal, Koca Süleyman Serdar, Kasapçopur Özgür, Editor, EMA TIP KİTABEVİ, pp.768-777, 2023

## Refereed Congress / Symposium Publications in Proceedings

### I. Genotype-phenotype correlation of patients with familial mediterranean fever in childhood: Experience of a single center

DOĞRUEL D., KIŞLA EKİNCİ R. M., BALCI S., YILMAZ M., ALTINTAŞ D. U.

9. International Congress of Familial Mediterranean Fever and Systemic Auto Inflammatory Diseases, Cyprus (Kktc), 4 - 07 May 2017

## Supported Projects

KIŞLA EKİNCİ R. M., BIŞGİN A., ALTINTAŞ D. U., YILMAZ M., BALCI S., DOĞRUEL D., Project Supported by Higher Education Institutions, Henoch Schönlein Purpuralı hastalarda C2 gen polimorfizm sıklığı ve klinik bulgulara etkisi, 2018 - 2019

## Metrics

Publication: 45

Citation (WoS): 112

Citation (Scopus): 148

H-Index (WoS): 6

H-Index (Scopus): 7

## Non Academic Experience

KAYSERİ ŞEHİR EĞİTİM ve ARAŞTIRMA HASTANESİ

FORUM YAŞAM HASTANESİ MERSİN

CITY HOSPİTAL MERSİN

MEDICAL PARK TARSUS HASTANESİ

MURADIYE DEVLET HASTANESİ