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

- I. Clinical and molecular evaluation of MEFV gene variants in the Turkish population: a study by the National Genetics Consortium  
DÜNDAR M., FAHRİOĞLU U., Yıldız S. H., Bakır-Gungor B., TEMEL Ş. G., AKIN H., ARTAN S., CORA T., ŞAHİN F. İ., DURSUN A., et al.  
FUNCTIONAL & INTEGRATIVE GENOMICS, vol.22, no.3, pp.291-315, 2022 (SCI-Expanded)
- II. The cumulative effects of MEFV gene polymorphisms and mutations in patients with inflammatory bowel diseases  
AĞİN M., BİŞGİN A., YILMAZ M., KONT ÖZHAN A., MAMMADOV F., TÜMGÖR G.  
JOURNAL OF THE PAKISTAN MEDICAL ASSOCIATION, vol.71, no.2, pp.479-483, 2021 (SCI-Expanded)
- III. 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)
- IV. Twenty-Year Experience of a Single Referral Center on Pediatric Familial Mediterranean Fever What Has Changed Over the Last Decade?  
KİSLA 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)
- V. Clinical manifestations and outcomes of 420 children with Henoch Schonlein Purpura from a single referral center from Turkey: A three-year experience  
KİSLA EKİNCİ R. M., Balci S., MELEK E., KARABAY BAYAZIT A., DOĞRU EL D., ALTINTAŞ D. U., YILMAZ M.  
MODERN RHEUMATOLOGY, vol.30, no.6, pp.1039-1046, 2020 (SCI-Expanded)
- VI. 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)
- VII. 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)
- VIII. Pre-Pulseless Takayasu Arteritis in a Child Represented With Prolonged Fever of Unknown Origin and Successful Management With Concomitant Mycophenolate Mofetil and Infliximab  
KİSLA EKİNCİ R. M., Balci S., Piskin F. C., Varan C., ALİNÇ ERDEM S., YILMAZ M.  
ARCHIVES OF RHEUMATOLOGY, vol.35, no.2, pp.278-282, 2020 (SCI-Expanded)
- IX. Unicentric Castleman Disease Mimicking an Autoinflammatory Disorder: A Diagnostic Challenge in a Pediatric Patient With Recurrent Fever  
KİSLA 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)
- X. Phenotypic variability in two patients with tumor necrosis factor receptor associated periodic fever

- syndrome emphasizes a rare manifestation: Immunoglobulin A nephropathy**  
 Balci S., KİŞLA EKİNCİ R. M., MELEK E., ATMIŞ B., BİŞGİN A., YILMAZ M.  
 EUROPEAN JOURNAL OF MEDICAL GENETICS, vol.63, no.4, 2020 (SCI-Expanded)
- XI. Baricitinib experience on STING-associated vasculopathy with onset in infancy: A representative case from Turkey**  
 Balci S., KİŞLA EKİNCİ R. M., de Jesus A. A., Goldbach-Mansky R., YILMAZ M.  
 CLINICAL IMMUNOLOGY, vol.212, pp.113-116, 2020 (SCI-Expanded)
- XII. Deficiency of adenosine deaminase 2: a case series revealing clinical manifestations, genotypes and treatment outcomes from Turkey**  
 KİŞLA EKİNCİ R. M., Balci S., Hershfield M., BİŞGİN A., DOĞRUCEL D., ALTINTAŞ D. U., YILMAZ M.  
 RHEUMATOLOGY, vol.59, no.1, pp.254-256, 2020 (SCI-Expanded)
- XIII. Abatacept as a Long-Term Targeted Therapy for LRBA Deficiency**  
 Kiykim A., Öğülür İ., Dursun E., Charbonnier L. M., Nain E., Çekiç Ş., Doğruel D., Karaca N. E., Çögürlü M. T., Bilir O. A., et al.  
 JOURNAL OF ALLERGY AND CLINICAL IMMUNOLOGY-IN PRACTICE, vol.7, pp.2790-2815, 2019 (SCI-Expanded)
- XIV. Canakinumab in Children with Familial Mediterranean Fever: A Single-Center, Retrospective Analysis**  
 KİŞLA EKİNCİ R. M., Balci S., DOĞRUCEL D., ALTINTAŞ D. U., YILMAZ M.  
 PEDIATRIC DRUGS, vol.21, no.5, pp.389-395, 2019 (SCI-Expanded)
- XV. Etanercept for the Treatment of Chronic Arthritis Related to Chronic Granulomatous Disease: A Case**  
 Balci S., KİŞLA EKİNCİ R. M., SERBES M., DOĞRUCEL D., ALTINTAŞ D. U., YILMAZ M.  
 PEDIATRIC ALLERGY IMMUNOLOGY AND PULMONOLOGY, vol.32, no.3, pp.131-134, 2019 (SCI-Expanded)
- XVI. Disease Severity and Genotype Affect Physical Growth in Children With Familial Mediterranean Fever**  
 KİŞLA EKİNCİ R. M., Balci S., Akay E., DOĞRUCEL D., ALTINTAŞ D. U., YILMAZ M.  
 ARCHIVES OF RHEUMATOLOGY, vol.34, no.3, pp.288-293, 2019 (SCI-Expanded)
- XVII. Hyperimmunoglobulinemia D syndrome with recurrent perianal abscess successfully treated with canakinumab**  
 KİŞLA EKİNCİ R. M., Balci S., BİŞGİN A., TÜMGÖR G., DOĞRUCEL D., YILMAZ M.  
 SCOTTISH MEDICAL JOURNAL, vol.64, no.3, pp.103-107, 2019 (SCI-Expanded)
- XVIII. Unusual and early onset IPEX syndrome: a case report**  
 DOĞRUCEL D., GÜRBÜZ F., TURAN İ., ALTINTAŞ D. U., YILMAZ M., YÜKSEL B.  
 TURKISH JOURNAL OF PEDIATRICS, vol.61, no.4, pp.580-584, 2019 (SCI-Expanded)
- XIX. Two years of newborn screening for cystic fibrosis in Turkey: Cukurova experience**  
 Sasihuseyinoglu A. S., ALTINTAŞ D. U., BİŞGİN A., DOĞRUCEL D., YILMAZ M., SERBES M.  
 TURKISH JOURNAL OF PEDIATRICS, vol.61, no.4, pp.505-512, 2019 (SCI-Expanded)
- XX. Concomitance of Familial Mediterranean Fever and Gitelman syndrome in an adolescent**  
 ATMIŞ B., KİŞLA EKİNCİ R. M., MELEK E., BİŞGİN A., YILMAZ M., Anarat A., KARABAY BAYAZIT A.  
 TURKISH JOURNAL OF PEDIATRICS, vol.61, no.3, pp.444-448, 2019 (SCI-Expanded)
- XXI. Recurrent Henoch Schonlein purpura without renal involvement successfully treated with methotrexate**  
 KİŞLA EKİNCİ R. M., Balci S., SERBES M., Aycin G. D., DOĞRUCEL D., ALTINTAŞ D. U., YILMAZ M.  
 SCOTTISH MEDICAL JOURNAL, vol.64, no.2, pp.74-77, 2019 (SCI-Expanded)
- XXII. Clinical findings and genetic analysis of the patients with IL-12R beta 1 deficiency from southeast Turkey**  
 DOĞRUCEL D., ÖZGÜR GÜNDEŞLİOĞLU Ö., YILMAZ M., Alabaz D., ALTINTAŞ D. U., KOCABAŞ E.  
 TURKISH JOURNAL OF PEDIATRICS, vol.61, no.2, pp.174-179, 2019 (SCI-Expanded)
- XXIII. Evaluation of high resolution computed tomography findings of cystic fibrosis**  
 Sasihuseyinoglu A. S., ALTINTAŞ D. U., SOYUPAK S., DOĞRUCEL D., YILMAZ M., SERBES M., Duyuler G.  
 KOREAN JOURNAL OF INTERNAL MEDICINE, vol.34, no.2, pp.335-343, 2019 (SCI-Expanded)
- XXIV. Autoimmune Manifestations in Heterozygote Type I Complement 2 Deficiency: A Child Eventually Diagnosed With Systemic Lupus Erythematosus**

- KİŞLA EKİNCİ R. M., Balci S., BİŞGİN A., ATMIŞ B., DOĞRUCL D., YILMAZ M.  
ARCHIVES OF RHEUMATOLOGY, vol.34, no.1, pp.96-99, 2019 (SCI-Expanded)
- XXV. Is Henoch-Schonlein purpura a susceptibility factor for functional gastrointestinal disorders in children?  
KİŞLA EKİNCİ R. M., Balci S., Mart O. O., TÜMGÖR G., Yavuz S., Celik H., DOĞRUCL D., ALTINTAŞ D. U., YILMAZ M.  
RHEUMATOLOGY INTERNATIONAL, vol.39, no.2, pp.317-322, 2019 (SCI-Expanded)
- XXVI. Dendritic cell activation is blunted in patients with coronary artery disease and diabetes mellitus  
Yetkin G. I., YÜCEL A. A., TEKİN İ. Ö., YILMAZ M., Atalay H., Yetkin E.  
JOURNAL OF DIABETES AND ITS COMPLICATIONS, vol.33, no.2, pp.134-139, 2019 (SCI-Expanded)
- XXVII. Renal Amyloidosis in Deficiency of Adenosine Deaminase 2: Successful Experience With Canakinumab  
Kişla Ekinci R. M., Balci S., Bişgin A., Hershfield M., Atmiş B., Doğruel D., Yılmaz M.  
PEDIATRICS, vol.142, no.5, 2018 (SCI-Expanded)
- XXVIII. Coexistence of 2 rare autosomal recessively inherited disorders manifesting with immune deficiency; IL-12 receptor beta 1 and biotinidase deficiencies  
DOĞRUCL D., BULUT F. D., YILMAZ M., Onenli-Mungan N., ALTINTAŞ D. U.  
TURKISH JOURNAL OF PEDIATRICS, vol.60, no.5, pp.584-587, 2018 (SCI-Expanded)
- XXIX. Squamous Cell Carcinoma With Hyper-IgE Syndrome: A Case Report  
Sasihuseyinoglu A. S., YILMAZ M., ALTINTAŞ D. U., DOĞRUCL D., SERBES M., Uguz A. H., Kiroglu M.  
JOURNAL OF PEDIATRIC HEMATOLOGY ONCOLOGY, vol.40, no.6, 2018 (SCI-Expanded)
- XXX. Ataxia-Telangiectasia Clinical and Laboratory Features: Single Center Results  
Sasihuseyinoglu A. S., YILMAZ M., BİŞGİN A., DOĞRUCL D., ALTINTAŞ D. U., Duyuler G., SERBES M.  
PEDIATRIC ALLERGY IMMUNOLOGY AND PULMONOLOGY, vol.31, no.1, pp.9-14, 2018 (SCI-Expanded)
- XXXI. Hematopoietic stem cell transplantation from unrelated donors in children with DOCK8 deficiency  
Uygun D. F. K., Uygun V., REİSLİ İ., KELEŞ S., Ozen A., YILMAZ M., Sayar E. H., Daloglu H., Ozturkmen S. I., Caki S., et al.  
PEDIATRIC TRANSPLANTATION, vol.21, no.7, 2017 (SCI-Expanded)
- XXXII. The extended clinical phenotype of 64 patients with dedicator of cytokinesis 8 deficiency  
Engelhardt K. R., Gertz M. E., Keles S., Schaeffer A. A., Sigmund E. C., Glockner C., Saghafi S., Pourpak Z., Ceja R., Sassi A., et al.  
JOURNAL OF ALLERGY AND CLINICAL IMMUNOLOGY, vol.136, no.2, pp.402-412, 2015 (SCI-Expanded)
- XXXIII. Griscelli syndrome type 3-like phenotype with MYO-5A exon-F deletion  
YILMAZ M., Cagdas D., Grandin V., ALTINTAŞ D. U., Tezcan I., de Saint Basile G., Sanal O.  
PEDIATRIC ALLERGY AND IMMUNOLOGY, vol.25, no.8, pp.817-819, 2014 (SCI-Expanded)
- XXXIV. A Novel Mutation in Leukocyte Adhesion Deficiency Type II/CDGIIc  
Cagdas D., YILMAZ M., KANDEMİR N., Tezcan I., Etzioni A., Sanal O.  
JOURNAL OF CLINICAL IMMUNOLOGY, vol.34, no.8, pp.1009-1014, 2014 (SCI-Expanded)
- XXXV. AUTOIMMUNE HEPATITIS AND SCLERODERMA IN A PATIENT WITH X-LINKED AGAMMAGLOBULINAEMIA  
YILMAZ M., BİNGÖL KARAKOÇ G., ALTINTAŞ D. U., Kotil A.  
JOURNAL OF CLINICAL IMMUNOLOGY, vol.34, no.6, pp.723, 2014 (SCI-Expanded)
- XXXVI. EFFECTS OF ANTIDOTAL THERAPY ON TESTIS TISSUE IN ORGANOPHOSPHATE POISONING  
GUMUSAY U., SEBE A., SATAR D. A., AY M. O., Yilmaz M., METE U. Ö.  
ACTA MEDICA MEDITERRANEA, vol.30, no.2, pp.435-439, 2014 (SCI-Expanded)
- XXXVII. Correlation between nasal eosinophils and nasal airflows in children with asthma and/or rhinitis monosensitised to house dust mites  
Yukselen A., Kendirli S. G., YILMAZ M., ALTINTAŞ D. U., BİNGÖL KARAKOÇ G.  
ALLERGOLOGIA ET IMMUNOPATHOLOGIA, vol.42, no.1, pp.50-55, 2014 (SCI-Expanded)
- XXXVIII. Clinical, functional, and genetic characterization of chronic granulomatous disease in 89 Turkish patients  
KÖKER M. Y., Camcioglu Y., van Leeuwen K., KILIÇ S. Ş., Barlan I., YILMAZ M., Metin A., de Boer M., AVCILAR H., PATIROĞLU T., et al.

- JOURNAL OF ALLERGY AND CLINICAL IMMUNOLOGY, vol.132, no.5, pp.1156-1168, 2013 (SCI-Expanded)
- XXXIX. **Evaluation of efficacy of immunotherapy in children with asthma monosensitized to Alternaria**  
 Kilic M., ALTINTAŞ D. U., YILMAZ M., BİNGÖL KARAKOÇ G., Burgut R., Guneser-Kendirli S.  
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- XL. **Temptamy Preaxial Brachydactyly Syndrome Is Caused by Loss-of-Function Mutations in Chondroitin Synthase 1, a Potential Target of BMP Signaling**  
 LI Y., LAUE K., TEMTAMY S., AGLAN M., KOTAN L. D., YIGIT G., CANAN H., PAWLIK B., NUERNBERG G., WAKELING E. L., et al.  
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- XLI. **Exhaled Breath Condensate MMP-9 Levels in Children With Bronchiectasis**  
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 PEDIATRIC PULMONOLOGY, vol.44, no.10, pp.1010-1016, 2009 (SCI-Expanded)
- XLII. **The clinical and genetical features of 124 children with Familial Mediterranean fever: experience of a single tertiary center**  
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- XLIII. **Defects along the T(H)17 differentiation pathway underlie genetically distinct forms of the hyper IgE syndrome**  
 Al Khatib S., Keles S., Garcia-Lioret M., Koc-Aydiner E. K., Reisli I., ARTAÇ H., Camcioglu Y., Cokugras H., Somer A., KÜTÜKÇÜLER N., et al.  
 JOURNAL OF ALLERGY AND CLINICAL IMMUNOLOGY, vol.124, no.2, pp.342-348, 2009 (SCI-Expanded)
- XLIV. **Exposure to house dust endotoxin and allergic sensitization in allergic and nonallergic children living in Adana, Turkey**  
 YILMAZ M., ALTINTAŞ D. U., BİNGÖL KARAKOÇ G., Inal A., Kilic M., Sutolukta Z., Guneser-Kendirli S.  
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- XLV. **Effect of outdoor fungus concentrations on symptom severity of children with asthma and/or rhinitis monosensitized to molds**  
 Inal A., BİNGÖL KARAKOÇ G., ALTINTAŞ D. U., Pinar M., Ceter T., YILMAZ M., Kendirli S. G.  
 ASIAN PACIFIC JOURNAL OF ALLERGY AND IMMUNOLOGY, vol.26, no.1, pp.11-17, 2008 (SCI-Expanded)

### Articles Published in Other Journals

- I. **Does selective IgA deficiency have a good prognostic role on juvenile dermatomyositis? a case report**  
 Balci S., KİŞLA EKİNCİ R. M., DOĞRU EL D., SERBES M., ALTINTAŞ D. U., YILMAZ M.  
 CUKUROVA MEDICAL JOURNAL, vol.44, no.4, pp.1511-1514, 2019 (ESCI)
- II. **Term ve geç preterm bebeklerde kordon klempeme zamanının lenfosit alt grupları üzerine etkisi**  
 BAHAR N., SATAR M., YILMAZ M., BÜYÜKKURT S., OZLU F., YAPICIOĞLU YILDIZDAŞ H., YAMAN A.  
 TURK PEDIATRI ARSIVI, vol.53, no.4, pp.214-221, 2018 (Scopus)
- III. **A 6-Month-Old Boy with Reddish, Scaly Skin: Netherton Syndrome**  
 BULUT F. D., KOR D., Yilmaz B. S., YILMAZ M., ALTINTAŞ D. U., Ceylaner S., Kilavuz S., Mungan N. O.  
 JOURNAL OF PEDIATRIC RESEARCH, vol.5, no.1, pp.54-56, 2018 (ESCI)
- IV. **T helper, cytotoxic T, and natural killer T cell profiles and their association with clinical prognosis in children with sickle cell anemia**  
 ATMIŞ B., KILINÇ Y., YILMAZ M., Atmis A., Karagun B. S., ŞAŞMAZ H. İ.  
 CUKUROVA MEDICAL JOURNAL, vol.43, no.4, pp.1002-1007, 2018 (ESCI)
- V. **Juvenile mixed connective tissue disease: case series**  
 KİŞLA EKİNCİ R. M., Balci S., Altintas D. U., YILMAZ M.  
 CUKUROVA MEDICAL JOURNAL, vol.43, no.1, pp.235-240, 2018 (ESCI)
- VI. **Effects of physical exercise on clinical and functional parameters in children with asthma**

- DOĞRUEL D., ALTINTAŞ D. U., YILMAZ M.  
CUKUROVA MEDICAL JOURNAL, vol.43, no.2, pp.457-462, 2018 (ESCI)
- VII. **Distribution of Allergen Sensitization in Childhood with the Skin Test**  
Sasihuseyinoglu A. S., Kont Ozhan A., SERBES M., Duyuler G. A., BINGOL G., YILMAZ M., ALTINTAŞ D. U.  
ASIM ALLERJI IMMUNOLOJİ, vol.15, no.1, pp.43-48, 2017 (ESCI)
- VIII. **Yenidoğan döneminde sağlık bakımı ile ilişkili enfeksiyon geçirmiş ve geniş spektrumlu antibiyotik kullanmış olmanın erken çocukluk döneminde gözlenen astma, allerjik rinit ve atopik dermatit üzerine etkisi**  
YAPICIOĞLU YILDIZDAŞ H., ÖZCAN A., YILMAZ M., SERTDEMİR Y.  
CUKUROVA MEDICAL JOURNAL, vol.42, no.1, pp.132-139, 2017 (Peer-Reviewed Journal)
- IX. **Effect of healthcare associated infections and broad spectrum antibiotic use in newborn period on development of asthma, allergic rhinitis and atopic dermatitis in early childhood**  
YILDIZDAS H. Y., OEZCAN A., Sertdemir Y., Yilmaz M.  
CUKUROVA MEDICAL JOURNAL, vol.42, no.1, pp.132-139, 2017 (ESCI)

### Refereed Congress / Symposium Publications in Proceedings

- I. **The evaluation of rare and low-frequency genetic variants in common variable immune deficiency (CVID) patients together with CV/RVCD (common variant/rare variant, common disease) hypothesis for final interpretation**  
BİŞGİN A., SÖNMEZLER Ö., BOGA İ., YILMAZ M.  
6. Uluslararası Katılımlı Erciyes Tıp Tibbi Genetik Kongresi, Turkey, 16 September 2021
- II. **geç preterm ve term bebeklerde kordon klempleme zamanının lenfosit alt gruplarına etkisi**  
Bahar N., Satar M., Yılmaz M., Büyükkurt S., Ozlu F., Yapıcıoğlu Yıldızdaş H.  
UNEKO-26, İskele, Cyprus (Kktc), 14 - 18 April 2018
- III. **Nöroihtiyotik Sendromlar: Bir Vaka Serisi**  
İNCECİK F., HERGÜNER M. Ö., GÜNGÖR S., YILMAZ M., BEŞEN Ş., GÜL MERT G.  
19.Uluslararası Çocuk Nörolojisi Kongresi, Turkey, 19 - 23 April 2017
- IV. **AUTOIMMUNE HEPATITIS AND SCLERODERMA IN A PATIENT WITH X-LINKED AGAMMAGLOBULINAEMIA**  
YILMAZ M., BİNGÖL KARAKOÇ G., ALTINTAŞ D. U., Kotil A.  
100th J Project Meeting, Antalya, Turkey, 12 - 14 March 2014, vol.34, pp.723

### Metrics

Publication: 58  
Citation (WoS): 483  
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H-Index (Scopus): 9

### Non Academic Experience

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