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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., Bakir-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**
AĞIN 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?**
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)
- V. **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)
- 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**
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)
- IX. **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)
- X. **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)

- XI. **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)
- XII. **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, 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., Çöğürü M. T., Bilir O. A., et al.
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- XIV. **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, 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., 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)
- XVI. **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)
- XVII. **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)
- XVIII. **Unusual and early onset IPEX syndrome: a case report**
DOĞRUDEL 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., DOĞRUDEL D., YILMAZ M., SERBES M.
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- XX. **Concomitance of Familial Mediterranean Fever and Gitelman syndrome in an adolescent**
ATMIŞ B., KIŞLA EKİNCİ R. M., MELEK E., BİŞGİN A., YILMAZ M., Anarat A., KARABAY BAYAZIT A.
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- XXI. **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)
- XXII. **Clinical findings and genetic analysis of the patients with IL-12R beta 1 deficiency from southeast Turkey**
DOĞRUDEL 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., DOĞRUDEL 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**

- KIŞLA EKİNCİ R. M., Balci S., BİŞGİN A., ATMIŞ B., DOĞRUEL 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?**
KIŞLA EKİNCİ R. M., Balci S., Mart O. O., TÜMGÖR G., Yavuz S., Celik H., DOĞRUEL 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**
Kışla Ekinci R. M., Balci S., Bişgin A., Hershfield M., Atmiş B., Doğruel D., Yılmaz M.
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- XXVIII. **Coexistence of 2 rare autosomal recessively inherited disorders manifesting with immune deficiency; IL-12 receptor beta 1 and biotinidase deficiencies**
DOĞRUEL 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., DOĞRUEL D., SERBES M., Uguz A. H., Kiroglu M.
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- XXX. **Ataxia-Telangiectasia Clinical and Laboratory Features: Single Center Results**
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- 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.
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- 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., Glocker C., Saghafi S., Pourpak Z., Ceja R., Sassi A., et al.
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- XXXIII. **Griselli 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.
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- XXXIV. **A Novel Mutation in Leukocyte Adhesion Deficiency Type II/CD11c**
Cagdas D., YILMAZ M., KANDEMİR N., Tezcan I., Etzioni A., Sanal O.
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- 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 monosensitized 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**
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- XL. **Temtamy Preaxial Brachydactyly Syndrome Is Caused by Loss-of-Function Mutations in Chondroitin Synthase 1, a Potential Target of BMP Signaling**
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- XLI. **Exhaled Breath Condensate MMP-9 Levels in Children With Bronchiectasis**
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- XLII. **The clinical and genetical features of 124 children with Familial Mediterranean fever: experience of a single tertiary center**
Inal A., YILMAZ M., Kendirli S. G., ALTINTAŞ D. U., BİNGÖL KARAKOÇ G.
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- XLIII. **Defects along the T(H)17 differentiation pathway underlie genetically distinct forms of the hyper IgE syndrome**
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- XLIV. **Exposure to house dust endotoxin and allergic sensitization in allergic and nonallergic children living in Adana, Turkey**
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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.
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Articles Published in Other Journals

- I. **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)
- II. **Term ve geç preterm bebeklerde kordon klempleme zamanının lenfosit alt grupları üzerine etkisi**
BAHAR N., SATAR M., YILMAZ M., BÜYÜKKURT S., OZLU F., YAPICIOĞLU YILDIZDAŞ H., YAMAN A.
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- 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.
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- 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.
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- V. **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)
- 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 ALLERJİ 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 klemleme 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.Ulusal 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

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Non Academic Experience

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