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Kişisel Bilgiler

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Eğitim Bilgileri

Doktora, Marmara Üniversitesi, Sağlık Bilimleri Enstitüsü, Tıbbi Biyoloji Ve Genetik Anabilim Dalı, Türkiye 1996 - 2000

Yüksek Lisans, University of London-Imperial College of Science, Technology and Medicine, Biyokimya ve Genetik, İnsan Genetiği, İngiltere 1994 - 1995

Lisans, University of Nottingham, Doğa Bilimleri, Genetik, İngiltere 1991 - 1994

Yabancı Diller

İngilizce, C2 Ustalık

Yaptığı Tezler

Yüksek Lisans, A Linkage Study of NMDAR1 Receptors and Schizophrenia, Imperial College of Science, Technology and Medicine, 2005

Doktora, Dikkat eksikliği aşırı hareketlilik bozukluğunun dopamin genleri ile ilişkisi, Marmara Üniversitesi, Sağlık Bilimleri Enstitüsü, Tıbbi Biyoloji Ve Genetik Anabilim Dalı, 2000

Araştırma Alanları

Sağlık Bilimleri, Yaşam Bilimleri

Akademik Unvanlar / Görevler

Prof.Dr., Acibadem Mehmet Ali Aydınlar Üniversitesi, Fen-Edebiyat Fakültesi, Moleküler Biyoloji Ve Genetik Bölümü, 2021 - Devam Ediyor

Prof.Dr., İstanbul Teknik Üniversitesi, Fen-Edebiyat, Moleküler Biyoloji Ve Genetik, 2009 - 2021

Doç.Dr., İstanbul Teknik Üniversitesi, Fen-Edebiyat, Moleküler Biyoloji Ve Genetik, 2009 - 2016

Yrd.Doç.Dr., İstanbul Teknik Üniversitesi, Fen-Edebiyat, Moleküler Biyoloji Ve Genetik, 2005 - 2010

Öğretim Görevlisi Dr., İstanbul Teknik Üniversitesi, Fen-Edebiyat, Moleküler Biyoloji Ve Genetik, 2004 - 2005

Öğretim Görevlisi Dr., Boğaziçi Üniversitesi, Fen-Edebiyat Fakültesi, Moleküler Biyoloji Ve Genetik Bölümü, 2001 - 2003

Akademik İdari Deneyim

Bölüm Başkanı, Acibadem Mehmet Ali Aydınlar Üniversitesi, Fen-Edebiyat Fakültesi, Moleküler Biyoloji Ve Genetik Bölümü, 2021 - Devam Ediyor

Bölüm Başkan Yardımcısı, İstanbul Teknik Üniversitesi, Fen-Edebiyat, 2009 - 2013

Erasmus Koordinatörü, İstanbul Teknik Üniversitesi, Fen-Edebiyat, 2007 - 2013

Verdiği Dersler

Genetics, Lisans, 2021 - 2022

Yönetilen Tezler

Tahir Turanlı E., VARIANT PATHOGENICITY PREDICTION TOOL BASED ON PROTEIN-PROTEIN INTERACTIONS AND THE EFFECTS OF VARIANTS ON 3-DIMENSIONAL PROTEIN STRUCTURE:A MODEL FOR MONOGENIC AUTOINFLAMMATORY DISORDERS, Yüksek Lisans, A.ALPER(Öğrenci), 2021

Tahir Turanlı E., The relationship between urinary exosomal miRNA levels and renal outcome in type 2 diabetic nephropathy patients, Yüksek Lisans, M.SEYİT(Öğrenci), 2021

Tahir Turanlı E., Disease gene identification using linkage and exome analysis, Yüksek Lisans, D.YAVUZ(Öğrenci), 2020

Tahir Turanlı E., Disease gene identification using linkage and exome analyses, Doktora, İ.KARACAN(Öğrenci), 2019

Tahir Turanlı E., Investigation of MEFV gene expression and pyrin levels in familial mediterranean fever and behçet syndrome, Yüksek Lisans, M.ÇİFTÇİ(Öğrenci), 2019

Tahir Turanlı E., Differential expression of proteins in active and inactive phases of Behçet's syndrome, Doktora, K.ASLI(Öğrenci), 2019

Tahir Turanlı E., Investigation of novel genes in autosomal dominant Behçet syndrome, Yüksek Lisans, G.TURAN(Öğrenci), 2019

Tahir Turanlı E., Expression analysis of hla-b gene in sporadic behcet syndrome patients, Yüksek Lisans, E.KIZILTEPE(Öğrenci), 2018

Tahir Turanlı E., TRANSCRIPTION ANALYSIS OF HLA-B GENE IN SPORADIC BEHCET SYNDROME PATIENTS, Yüksek Lisans, E.Kızıltepe(Öğrenci), 2017

Tahir Turanlı E., Investigation of the possible effect of intragenic MEFV gene CpG island methylation on mRNA transcription and pyrin localization, Doktora, G.ERDEM(Öğrenci), 2017

Tahir Turanlı E., Analysis of mefv variations, expression and pyrin levels in familial mediterranean fever disease, Yüksek Lisans, N.SEVİNÇ(Öğrenci), 2016

Tahir Turanlı E., A genetic analysis of autoinflammatory diseases, Yüksek Lisans, A.BALAMİR(Öğrenci), 2016

Tahir Turanlı E., Investigation of notch signalling pathway proteins as potential biomarkers for differentiating multiple sclerosis subtypes, Yüksek Lisans, Z.ÖZTÜRK(Öğrenci), 2016

Tahir Turanlı E., MEFV mutation analysis in familial Mediterranean fever, gout and adult-onset still's diseases, Yüksek Lisans, S.GILA(Öğrenci), 2015

Tahir Turanlı E., Analysis of histone modifications in familial Mediterranean fever patients using chromatin immunoprecipitation sequencing assay, Yüksek Lisans, B.FİDAN(Öğrenci), 2015

Tahir Turanlı E., Genetic and epigenetic analyses of hla-b5 gene in familial behçet syndrome pedigrees, Yüksek Lisans, P.KÖPRÜLÜ(Öğrenci), 2015

Tahir Turanlı E., Investigation of molecular pathways and biomarkers in Multiple sclerosis clinical subtypes, Doktora, T.AVŞAR(Öğrenci), 2015

Tahir Turanlı E., A linkage analysis and a genome-wide association study on familial multiple sclerosis, Yüksek Lisans, E.EVEREST(Öğrenci), 2015

Tahir Turanlı E., A study on the localization of alternative MEFV transcripts in neutrophil-like cells, Yüksek Lisans, Ş.ERDEMİR(Öğrenci), 2013

Tahir Turanlı E., Analysis of MEFV gene alternatively spliced transcripts expression patterns in cell culture models, Yüksek Lisans, İ.ABACI(Öğrenci), 2013

SCI, SSCI ve AHCI İndekslerine Giren Dergilerde Yayınlanan Makaleler

- I. **Investigation of multiple sclerosis-related pathways through the integration of genomic and proteomic data**
Everest E., Ulgen E., UYGUNOĞLU U., TÜTÜNCÜ M., SAİP S., SEZERMAN O. U. , SİVA A., TAHİR TURANLI E.
PEERJ, cilt.9, 2021 (SCI İndekslerine Giren Dergi)
- II. **LACC1 deficiency links juvenile arthritis with autophagy and metabolism in macrophages**
Omarjee O., Mathieu A., Quiniou G., Moreews M., Ainouze M., Frachette C., Melki I., Dumaine C., Gerfaud-Valentin M., Duquesne A., et al.
JOURNAL OF EXPERIMENTAL MEDICINE, cilt.218, sa.3, 2021 (SCI İndekslerine Giren Dergi)
- III. **Preparation and in vitro characterization of monoclonal antibody ranibizumab conjugated magnetic nanoparticles for ocular drug delivery**
Ayata N., SEZER A. D. , Bucak S., Turanli E.
BRAZILIAN JOURNAL OF PHARMACEUTICAL SCIENCES, cilt.56, 2020 (SCI İndekslerine Giren Dergi)
- IV. **A 9.5-year-old boy with recurrent neurological manifestations and severe hypertension, treated initially for polyarteritis nodosa, was subsequently diagnosed with adenosine deaminase type 2 deficiency (DADA2) which responded to anti-TNF-alpha**
Sahin S., Adrovic A., BARUT K., Baran S., Turanli E., CANPOLAT N., KIZILKILIÇ O., Ozkaya O., KASAPÇOPUR Ö.
PAEDIATRICS AND INTERNATIONAL CHILD HEALTH, cilt.40, sa.1, ss.65-68, 2020 (SCI İndekslerine Giren Dergi)
- V. **Peripheral blood mononuclear cell proteome profile in Behcet's syndrome**
Aydin A. K. , Ozguler Y., Ucar D., KASAP M., AKPINAR G., SEYAHİ E., Turanli E.
RHEUMATOLOGY INTERNATIONAL, cilt.40, sa.1, ss.65-74, 2020 (SCI İndekslerine Giren Dergi)
- VI. **Atypical phenotype of an old disease or typical phenotype of a new disease: deficiency of adenosine deaminase 2**
Cakan M., Aktay-Ayaz N., Karadag S. G. , Tahir-Turanli E., Stafstrom K., Bainter W., Geha R. S. , Chou J.
TURKISH JOURNAL OF PEDIATRICS, cilt.61, sa.3, ss.413-417, 2019 (SCI İndekslerine Giren Dergi)
- VII. **Diagnostic utility of a targeted next-generation sequencing gene panel in the clinical suspicion of systemic autoinflammatory diseases: a multi-center study**
Karacan I., Balamir A., Ugurlu S., Aydin A. K. , Everest E., Zor S., Onen M. O. , Dasdemir S., Ozkaya O., Sozeri B., et al.
RHEUMATOLOGY INTERNATIONAL, cilt.39, sa.5, ss.911-919, 2019 (SCI İndekslerine Giren Dergi)
- VIII. **Spectrum of the neurologic manifestations in childhood-onset cryopyrin-associated periodic syndrome**
Kilic H., Sahin S., Duman C., Adrovic A., BARUT K., Turanli E., YILDIRIM S. R. , KIZILKILIÇ O., KASAPÇOPUR Ö., Saltik S.
EUROPEAN JOURNAL OF PAEDIATRIC NEUROLOGY, cilt.23, sa.3, ss.466-472, 2019 (SCI İndekslerine Giren Dergi)
- IX. **Correction to: Diagnostic utility of a targeted next-generation sequencing gene panel in the clinical suspicion of systemic autoinflammatory diseases: a multi-center study.**
Karacan İ., Balamir A., Uğurlu S., Aydin A. K. , Everest E., Zor S., Önen M. Ö. , Daşdemir S., Özkaya O., Sözeri B., et al.
Rheumatology international, cilt.39, ss.921, 2019 (SCI İndekslerine Giren Dergi)
- X. **A Novel ATP6V0A2 Mutation Causing Recessive Cutis Laxa with Unusual Manifestations of Bleeding Diathesis and Defective Wound Healing**
Karacan I., Kucukkaya R. D. , Karakus F. N. , Solakoglu S., Tolun A., Hancer V. S. , Turanli E.
TURKISH JOURNAL OF HEMATOLOGY, cilt.36, sa.1, ss.29-36, 2019 (SCI İndekslerine Giren Dergi)
- XI. **C3 glomerulopathy in NLRP12-related autoinflammatory disorder: case-based review**
BAŞARAN H. Ö. , Uncu N., Cakar N., Turanli E., Kiremitci S., Aydin F., BAYRAKCI U. S.
RHEUMATOLOGY INTERNATIONAL, cilt.38, sa.8, ss.1571-1576, 2018 (SCI İndekslerine Giren Dergi)
- XII. **LACC1 Gene Defects in Familial Form of Juvenile Arthritis**
Karacan I., Ugurlu S., Sahin S., Everest E., KASAPÇOPUR Ö., Tolun A., Ozdogan H., Turanli E.
JOURNAL OF RHEUMATOLOGY, cilt.45, sa.5, ss.726-728, 2018 (SCI İndekslerine Giren Dergi)
- XIII. **Familial Mediterranean fever in childhood: a single-center experience**
BARUT K., Sahin S., ADROVIC YILDIZ A., Sinoplu A. B. , Yucel G., Pamuk G., Aydin A. K. , Dasdemir S., Turanli E., BUYRU A. N. , et al.

- RHEUMATOLOGY INTERNATIONAL, cilt.38, sa.1, ss.67-74, 2018 (SCI İndekslerine Giren Dergi)
- XIV. **Clinical, imaging and genotypical features of three deceased and five surviving cases with ADA2 deficiency**
Sahin S., ADROVIC YILDIZ A., BARUT K., Ugurlu S., Turanli E., Ozdogan H., KASAPÇOPUR Ö.
RHEUMATOLOGY INTERNATIONAL, cilt.38, sa.1, ss.129-136, 2018 (SCI İndekslerine Giren Dergi)
- XV. **Other autoinflammatory disease genes in an FMF-prevalent population: a homozygous MVK mutation and a novel heterozygous TNFRSF1A mutation in two different Turkish families with clinical FMF.**
Karacan İ., Uğurlu S., Tolun A., Tahir Turanlı E., Ozdogan H.
Clinical and experimental rheumatology, ss.75-81, 2017 (SCI Expanded İndekslerine Giren Dergi)
- XVI. **Alternatively spliced MEFV transcript lacking exon 2 and its protein isoform pyrin-2d implies an epigenetic regulation of the gene in inflammatory cell culture models**
Erdem G. C. , Erdemir S., Abaci I., Aydin A. K. K. , Everest E., Turanli E.
GENETICS AND MOLECULAR BIOLOGY, cilt.40, sa.3, ss.688-697, 2017 (SCI İndekslerine Giren Dergi)
- XVII. **Integrated Genomic and Proteomic Analyses of Multiple Sclerosis**
Everest E., Uygunoglu U., TÛTÛNCÛ M., SAİP S., Sezerman O. U. , Siva A., Turanli E.
NEUROLOGY, cilt.88, 2017 (SCI İndekslerine Giren Dergi)
- XVIII. **A NOVEL Missense Mvk mutation in a Family with Familial Mediterranean Fever-like Disease**
Karacan I., UĞURLU S., Tolun A., Turanli E., Ozdogan H.
ARTHRITIS & RHEUMATOLOGY, cilt.68, 2016 (SCI İndekslerine Giren Dergi)
- XIX. **Nanomedicine**
Turanli E., Everest E.
LOW-DIMENSIONAL AND NANOSTRUCTURED MATERIALS AND DEVICES: PROPERTIES, SYNTHESIS, CHARACTERIZATION, MODELLING AND APPLICATIONS, ss.579-587, 2016 (SCI İndekslerine Giren Dergi)
- XX. **Methylation Analysis of HLA-B Locus in Familial Behcet Syndrome**
Turanli E., Koprulu P., UĞURLU S., Yazici H., Seyahi E.
ARTHRITIS & RHEUMATOLOGY, cilt.67, 2015 (SCI İndekslerine Giren Dergi)
- XXI. **CSF Proteomics Identifies Specific and Shared Pathways for Multiple Sclerosis Clinical Subtypes**
Avsar T., Durasi I. M. , Uygunoglu U., Tutuncu M., Demirci N. O. , Saip S., Sezerman O. U. , Siva A., Tahir Turanlı E.
PLOS ONE, cilt.10, sa.5, 2015 (SCI İndekslerine Giren Dergi)
- XXII. **Protein biomarkers for multiple sclerosis: semi-quantitative analysis of cerebrospinal fluid candidate protein biomarkers in different forms of multiple sclerosis**
Avsar T., Korkmaz D., TÛTÛNCÛ M., Demirci N. O. , SAİP S., Kamaşak M. E. , SİVA A., Turanli E.
MULTIPLE SCLEROSIS JOURNAL, cilt.18, sa.8, ss.1081-1091, 2012 (SCI İndekslerine Giren Dergi)
- XXIII. **Analysis of MEFV exon methylation and expression patterns in familial Mediterranean fever**
Kirectepe A. K. , KASAPÇOPUR Ö., Arisoy N., Erdem G. C. , HATEMİ G., Ozdogan H., Turanli E.
BMC MEDICAL GENETICS, cilt.12, 2011 (SCI İndekslerine Giren Dergi)
- XXIV. **Increased expression of exon 2 deleted MEFV transcript in familial Mediterranean fever patients**
Kirectepe A. K. , Erdem G. C. , Senturk N., Arisoy N., Hatemi G., Ozdogan H., KASAPÇOPUR Ö., Turanli E.
INTERNATIONAL JOURNAL OF IMMUNOGENETICS, cilt.38, sa.4, ss.327-329, 2011 (SCI İndekslerine Giren Dergi)
- XXV. **The prevalence of Behçet's syndrome, familial Mediterranean fever, HLA-B51 and MEFV gene mutations among ethnic Armenians living in Istanbul, Turkey.**
Seyahi E., Tahir Turanlı E., Mangan M. S. , Celikyapi G., Oktay V., Cevirgen D., Kuzuoglu D., Ozoglu S., Yazici H.
Clinical and experimental rheumatology, cilt.28, 2010 (SCI Expanded İndekslerine Giren Dergi)
- XXVI. **A twin study in Behçet's syndrome.**
Masatlioglu S., Seyahi E., Tahir Turanlı E., Fresko I., Gogus F., Senates E., Oguz Savran F., Yazici H.
Clinical and experimental rheumatology, cilt.28, 2010 (SCI Expanded İndekslerine Giren Dergi)
- XXVII. **A TWIN STUDY IN BEHCET'S SYNDROME**
Masatlioglu S., Seyahi E., Turanli E., FRESKO İ., Gogus F., Senates E., Savran F. O. , Yazici H.
CLINICAL AND EXPERIMENTAL RHEUMATOLOGY, cilt.28, sa.4, 2010 (SCI İndekslerine Giren Dergi)
- XXVIII. **Common MEFV mutations and polymorphisms in an elderly population: an association with E148Q polymorphism and rheumatoid factor levels.**

- Turanli E., Beger T., Erdinçler D., Curgunlu A., Karaman S., Karaca E., Dasdemir S., Bolayirli M., Yazici H.
Clinical and experimental rheumatology, cilt.27, ss.340-3, 2009 (SCI Expanded İndekslerine Giren Dergi)
- XXIX. **Analysis of the dopamine beta hydroxylase gene in Gilles de la Tourette syndrome**
Ozbay F., Wigg K. G. , Turanli E., Asherson P., Yazgan Y., Sandor P., Barr C. L.
AMERICAN JOURNAL OF MEDICAL GENETICS PART B-NEUROPSYCHIATRIC GENETICS, sa.6, ss.673-677, 2006 (SCI İndekslerine Giren Dergi)
- XXX. **Stable transmission and expression of the hepatitis B virus total genome in hybrid transgenic mice until F10 generation**
Bagis H., Arat S., Mercan H., Aktoprakligil D., Caner M., Turanli E., Baysal K., Turgut G., Sekmen S., Cirakoglu B.
JOURNAL OF EXPERIMENTAL ZOOLOGY PART A-ECOLOGICAL AND INTEGRATIVE PHYSIOLOGY, sa.5, ss.420-427, 2006 (SCI İndekslerine Giren Dergi)
- XXXI. **Functional effects of a tandem duplication polymorphism in the 5'flanking region of the DRD4 gene**
D'Souza U., Russ C., Tahir E., Mill J., McGuffin P., Asherson P., Craig I.
BIOLOGICAL PSYCHIATRY, cilt.56, sa.9, ss.691-697, 2004 (SCI İndekslerine Giren Dergi)
- XXXII. **Joint analysis of the DRD5 marker concludes association with attention-deficit/hyperactivity disorder confined to the predominantly inattentive and combined subtypes**
Lowe N., Kirley A., Hawi Z., Sham P., Wickham H., Kratochvil C., Smith S., Lee S., Levy F., Kent L., et al.
AMERICAN JOURNAL OF HUMAN GENETICS, cilt.74, sa.2, ss.348-356, 2004 (SCI İndekslerine Giren Dergi)
- XXXIII. **Association analysis of MAOA and COMT with neuroticism assessed by peers**
Eley T., Tahir E., Angleitner A., Harriss K., McClay J., Plomin R., Riemann R., Spinath F., Craig I.
AMERICAN JOURNAL OF MEDICAL GENETICS PART B-NEUROPSYCHIATRIC GENETICS, sa.1, ss.90-96, 2003 (SCI İndekslerine Giren Dergi)
- XXXIV. **Evaluation of the genes for the adrenergic receptors alpha 2A and alpha 1C and Gilles de la Tourette syndrome**
Xu C., Ozbay F., Wigg K., Shulman R., Tahir E., Yazgan Y., Sandor P., Barr C.
AMERICAN JOURNAL OF MEDICAL GENETICS PART B-NEUROPSYCHIATRIC GENETICS, sa.1, ss.54-59, 2003 (SCI İndekslerine Giren Dergi)
- XXXV. **Association study of a dopamine transporter polymorphism and attention deficit hyperactivity disorder in UK and Turkish samples**
Curran S., Mill J., Tahir E., Kent L., Richards S., Gould A., Hockett L., Sharp J., Batten C., Fernando S., et al.
MOLECULAR PSYCHIATRY, cilt.6, sa.4, ss.425-428, 2001 (SCI İndekslerine Giren Dergi)
- XXXVI. **Association and linkage of DRD4 and DRD5 with attention deficit hyperactivity disorder (ADHD) in a sample of Turkish children**
Tahir E., Yazgan Y., Cirakoglu B., Ozbay F., Waldman I., Asherson P.
MOLECULAR PSYCHIATRY, cilt.5, sa.4, ss.396-404, 2000 (SCI İndekslerine Giren Dergi)

Diğer Dergilerde Yayınlanan Makaleler

- I. **Chemerin rs17173608 Gene Polymorphism is not Associated with Type 2 Diabetes Mellitus: a Cross-sectional Study.**
Olt S., Öznas O., Bağış H., Turanli E.
Folia medica, cilt.61, ss.69-75, 2019 (Diğer Kurumların Hakemli Dergileri)
- II. **Quality assurance of genetic laboratories and the EBTNA practice certification, a simple standardization assurance system for a laboratory network**
Precone V., DÜNDAR M., Beccari T., Turanli E., Cecchin S., Marceddu G., Manara E., Bertelli M.
EUROBIOTECH JOURNAL, cilt.2, sa.4, ss.215-258, 2018 (ESCI İndekslerine Giren Dergi)
- III. **Role of genetics in pediatric rheumatology**
Turanli E., Everest E., Balamir A., Aydin A. K. , KASAPÇOPUR Ö.
TURK PEDIATRI ARSIVI-TURKISH ARCHIVES OF PEDIATRICS, cilt.52, sa.3, ss.113-121, 2017 (ESCI İndekslerine Giren Dergi)

Kitap & Kitap Bölümleri

- I. **Behçet Hastalığı: Genetik ve Epigenetik Araştırmalar**
Tahir Turanlı E., Kireçtepe Aydın A., Özkılınç Önen M.
Behçet Hastalığı, Emire Seyahi, Editör, Türkiye Klinikleri Yayınevi, Ankara, ss.21-28, 2020
- II. **Nanomedicine**
TAHİR TURANLI E., EVEREST E.
Low Dimensional and Nanostructured Materials and Devices, , Editör, Springer International Publishing, ss.579-587, 2016

Hakemli Kongre / Sempozyum Bildiri Kitaplarında Yer Alan Yayınlar

- I. **Increased Burden of Susceptibility Variants in Familial Multiple Sclerosis**
Everest E., UYGUNOĞLU U., TÜTÜNCÜ M., SAİP S., Duman T., Riley B., Siva A., Turanlı E.
Annual Meeting of the American-Academy-of-Neurology, Toronto, Kanada, 25 Nisan - 01 Mayıs 2020, cilt.94
- II. **Prospective Analysis of Cerebrospinal Fluid Protein Levels Reveals Candidate Biomarkers for Disability Outcome in Multiple Sclerosis**
Siva A., Everest E., UYGUNOĞLU U., Yavuz D., Bulbul A., TÜTÜNCÜ M., SAİP S., Avsar T., Turanlı E.
Annual Meeting of the American-Academy-of-Neurology, Toronto, Kanada, 25 Nisan - 01 Mayıs 2020, cilt.94
- III. **Genetic screening in the clinical suspicion of autoinflammatory diseases**
Turanlı E.
European Biotechnology Congress, Valencia, İspanya, 11 - 13 Nisan 2019, cilt.305
- IV. **Investigation of Peripheral Blood Mononuclear Cells (PBMC) Proteome Profile in Behçet's Disease**
Aydın A. F. , Ozguler Y., UÇAR D., Seyahi E., Yazici H., Turanlı E.
51st Conference of the European-Society-of-Human-Genetics (ESHG) in conjunction with the European Meeting on Psychosocial Aspects of Genetics (EMPAG), Milan, İtalya, 16 - 19 Haziran 2018, cilt.27, ss.201-202
- V. **Candidate Gene Search For Autosomal Dominant Behçet's Disease Through Whole Exome Sequencing**
UĞURLU S., Turan G., Karacan I., Dasdemir S., Seven M., Duz M. B. , Ozdogan H., Tolun A., Turanlı E.
51st Conference of the European-Society-of-Human-Genetics (ESHG) in conjunction with the European Meeting on Psychosocial Aspects of Genetics (EMPAG), Milan, İtalya, 16 - 19 Haziran 2018, cilt.27, ss.936
- VI. **URINARY EXOSOMAL MIRNA LEVELS REVEAL NEW POTENTIAL BIOMARKERS IN TYPE 2 DIABETIC NEPHROPATHY AND FSGS**
Trabulus S., Zor M. S. , Turanlı E., Dincer T., Alagoz S., Mese M., Yilmaz E., SEYAHİ N.
56th Congress of the European-Renal-Association (ERA)-European-Dialysis-and-Transplant-Association (EDTA) - Burden, Access and Disparities in Kidney Disease, Budapest, Macaristan, 13 - 16 Haziran 2019, cilt.34, ss.214
- VII. **Lack of Low-Frequency Complete-Penetrance Coding Variants Responsible from Familial Multiple Sclerosis**
Siva A., Everest E., UYGUNOĞLU U., TÜTÜNCÜ M., SAİP S., Duman T., Turanlı E.
71st Annual Meeting of the American-Academy-of-Neurology (AAN), Pennsylvania, Amerika Birleşik Devletleri, 4 - 10 Mayıs 2019, cilt.92
- VIII. **LACC1 mutations in familial form of juvenile idiopathic arthritis**
Karacan I., Ugurlu S., Sahin S., KASAPÇOPUR Ö., Tolun A., Ozdogan H., Turanlı E.
50th European-Society-of-Human-Genetics (ESHG) Conference, Copenhagen, Danimarka, 27 - 30 Mayıs 2017, cilt.26, ss.915
- IX. **Whole genome linkage and exome sequencing analyses in an autosomal recessive Takayasu arteritis family**
Karacan I., Turanlı E., Tolun A., Seyahi E.
50th European-Society-of-Human-Genetics (ESHG) Conference, Copenhagen, Danimarka, 27 - 30 Mayıs 2017, cilt.26, ss.323

- X. **WHOLE GENOME LINKAGE AND EXOME SEQUENCING ANALYSES IN TAKAYASU ARTERITIS FAMILIES**
Turanli E., Karacan I., Esatlioglu S. N., Sahin S., KASAPÇOPUR Ö., Tolun A., Seyahi E.
Congress of the European-League-Against-Rheumatism (EULAR), Amsterdam, Hollanda, 13 - 16 Haziran 2018, cilt.77, ss.231-232
- XI. **NADİR BİR OLGU: ADENOZİN DEAMİNAZ-2 EKSİKLİĞİ**
AYDIN M., BABAZADA K., TAŞDEMİR E., ŞAHİN S., ADROVIC A., TURAN O., BARUT K., AKSOY S., SAYGILI S. K., KURUĞOĞLU S., et al.
54. TÜRK PEDIATRİ KONGRESİ, Türkiye, 6 - 10 Mayıs 2018
- XII. **NLRP12 varyasyonu taşıyan bir çocuk olguda otoinflamatuvar hastalık ve C3 Glomerulopati birlikteliği.**
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