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Education Information

Doctorate, Cumhuriyet Üniversitesi, Tıp, Temel Tı Bilimler Bölümü, Sağlık Bilimler Enstitüsü, Tıbbi Biyoloji Ve Genetik- Tıbbi Genetik Bilim Dalı, Turkey 1989 - 1992

Postgraduate, Cumhuriyet Üniversitesi, Tıp, Temel Tı Bilimler Bölümü, Sağlık Bilimler Enstitüsü, Tıbbi Biyoloji Ve Genetik- Tıbbi Genetik Bilim Dalı, Turkey 1987 - 1989

Undergraduate, Hacettepe University, Fen Fakültesi, Biyoloji, Turkey 1981 - 1986

Foreign Languages

English, C1 Advanced

Dissertations

Doctorate, "5-Azodeoksisitidinin (baz analogu) insan 11,16 ve X kromozomları üzerindeki etkisinin C ve bromodeoksi uridin (BrdU) bant teknikleriyle araştırılması", Cumhuriyet Üniversitesi, Temel Tıp Bilimleri, Tıbbi Biyoloji Ve Genetik Anabilim Dalı - Tıbbi Genetik Bilim Dalı, 1992

Postgraduate, "Populasyonda kromozom polimorfizm sıklığı ve beraberindeki genetik stigmatlar", Cumhuriyet Üniversitesi, Temel Tıp Bilimleri, Tıbbi Biyoloji Ve Genetik- Tıbbi Genetik Bilim Dalı, 1989

Research Areas

Medicine, Health Sciences, Internal Medicine Sciences, Medical Genetics

Academic Titles / Tasks

Professor, Çanakkale Onsekiz Mart University, Tıp Fakültesi, Dahili Tıp Bilimleri, 2010 - Continues

Professor, Cumhuriyet Üniversitesi, Tıp , Dahili Tıp Bilimleri, 2010 - 2010

Associate Professor, Cumhuriyet Üniversitesi, Tıp , Dahili Tıp Bilimleri, 2003 - 2006

Associate Professor, Cumhuriyet Üniversitesi, Tıp , Temel Tıp Bilimleri Bölümü, 2000 - 2003

Assistant Professor, Cumhuriyet Üniversitesi, Tıp , Temel Tıp Bilimleri Bölümü, 1997 - 2000

Lecturer PhD, Cumhuriyet Üniversitesi, Tıp , Temel Tıp Bilimleri Bölümü, 1993 - 1997

Research Assistant, Cumhuriyet Üniversitesi, Tıp , Temel Tıp Bilimleri Bölümü, 1987 - 1993

ADVISING 1 NESSES

- ÖZDEMİR Ö., Çoklu ligasyonla prob amlifikasyonu (MLPA) yönteminin prenatal tanıdaki yeri ve önemi, Postgraduate, E.ARI(Student), 2015
- ÖZDEMİR Ö., Tekrarlayan gebelik kaybı olan çiftlerde sayısal ve yapısal kromozom aberasyonlarının FISH yöntemi ile ileri düzeyde araştırılması, Postgraduate, B.UYSAL(Student), 2014
- ÖZDEMİR Ö., Abortus etiolojisinde embriyonal ve parental genetik faktörlerin değerlendirilmesi, Expertise In Medicine, D.A.(Student), 2013
- ÖZDEMİR Ö., Populasyonda Faktör V Leiden (G1619A) ve Faktör V H1299R (R2) gen polimorfizmi ve allel sıklığı, Expertise In Medicine, D.ÖZEN(Student), 2009
- ÖZDEMİR Ö., Sivas popülasyonunda anjiyotensin convreting enzim (ACE) gen polimorfizmi ve allel sıklığı, Expertise In Medicine, D.E.(Student), 2008
- ÖZDEMİR Ö., Populasyonda metilentetrahidrofolat redüktaz (MTHFR) gen polimorfizmi ve allel sıklığı, Expertise In Medicine, D.KOÇAK(Student), 2008
- ÖZDEMİR Ö., İyonize radyasyon, MR ve CT'ye maruz bırakılan rat kemikliği stem hücrelerinde mikroçekirdek sıklığı, Postgraduate, E.GÜL(Student), 2005
- ÖZDEMİR Ö., 3-Metilkolantren ile indüklenip metil- di- t-bütülfenol ile promote edilen rat akciğer dokusunda Ki-RAS exon-2 mutasyonları", Doctorate, F.POLAT(Student), 2004

Articles Published in Journals That Entered SCI, SSCI and AHCI Indexes

- I. **The high frequency of chromosomal copy number variations and candidate genes in epilepsy patients ***
ALBUZ B., ÖZDEMİR Ö., SILAN F.
CLINICAL NEUROLOGY AND NEUROSURGERY, vol.202, 2021 (Journal Indexed in SCI)
- II. **The Analysis of GJB2, GJB3, and GJB6 Gene Mutations in Patients with Hereditary Non-Syndromic Hearing Loss Living in Sivas**
KÜÇÜK KURTULGAN H., ALTUNTAŞ E. E. , YILDIRIM M. E. , ÖZDEMİR Ö., BAĞCI B., Sezgin I.
JOURNAL OF INTERNATIONAL ADVANCED OTOLOGY, vol.15, no.3, pp.373-378, 2019 (Journal Indexed in SCI)
- III. **Blau syndrome with a rare mutation in exon 9 of NOD2 gene**
Velickovic J., SILAN F., Bir F. D. , SILAN C., ALBUZ B., ÖZDEMİR Ö.
AUTOIMMUNITY, vol.52, pp.256-263, 2019 (Journal Indexed in SCI)
- IV. **Prevalence of MEFV gene mutations in a large cohort of patients with suspected familial Mediterranean fever in Central Anatolia**
YILDIRIM M. E. , KÜÇÜK KURTULGAN H., ÖZDEMİR Ö., KILIÇGÜN H., Aydemir D. S. , Baser B., Sezgin I.
ANNALS OF SAUDI MEDICINE, vol.39, no.6, pp.382-387, 2019 (Journal Indexed in SCI)
- V. **The Genomics of Arthrogyrosis, a Complex Trait: Candidate Genes and Further Evidence for Oligogenic Inheritance**
Pehlivan D., Bayram Y., Gunes N., Akdemir Z. C. , Shukla A., Bierhals T., TABAKCI B., Sahin Y., Gezdirici A., Fatih J. M. , et al.
AMERICAN JOURNAL OF HUMAN GENETICS, vol.105, no.1, pp.132-150, 2019 (Journal Indexed in SCI)
- VI. **Evaluation of the Association between Lithium Treatment and GSK-3 beta Polymorphism in Bipolar Disorder Patients**
ALTINBAŞ K., Yesilbas D., Ince B., Cansiz A., SILAN F., ÖZDEMİR Ö., Guloksuz S.
TURK PSIKIYATRI DERGISI, vol.29, no.2, pp.73-78, 2018 (Journal Indexed in SSCI)
- VII. **The prevalence of human papillomavirus (HPV) genotypes detected by PCR in women with normal and abnormal cervico-vaginal cytology**
BEYAZIT F., SILAN F., Gencer M., Aydin B., Paksoy B., ÜNSAL M. A. , ÖZDEMİR Ö.
GINEKOLOGIA POLSKA, vol.89, no.2, pp.62-67, 2018 (Journal Indexed in SCI)
- VIII. **Possible association between germline methylenetetrahydrofolate reductase gene polymorphisms and psoriasis risk in a Turkish population**
KILIÇ S., ÖZDEMİR Ö., SILAN F., IŞIK S., YILDIZ Ö., KARAAGACLI D., SILAN C., OGRETMEN Z.

Clinical and Experimental Dermatology, vol.42, no.1, pp.8-13, 2017 (Journal Indexed in SCI)

- IX. **THE MEFV GENE PATHOGENIC VARIANTS AND PHENOTYPE-GENOTYPE CORRELATION IN CHILDREN WITH FAMILIAL MEDITERRANEAN FEVER IN THE CANAKKALE POPULATION**
BATTAL F., SILAN F., TOPALOĞLU N., AYLANÇ H., YILDIRIM Ş., Binnetoglu K. F. , TEKİN M., KAYMAZ N., ÖZDEMİR Ö.
BALKAN JOURNAL OF MEDICAL GENETICS, vol.19, no.2, pp.23-28, 2016 (Journal Indexed in SCI)
- X. **Prevalence and mutations of β -thalassemia trait and abnormal hemoglobins in premarital screening in Çanakkale province, Turkey.**
Uludağ A., UYSAL A., Uludağ A., Ertekin Y. H. , Tekin M., KÜTÜK B., Silan F., Özdemir Ö.
Balkan journal of medical genetics : BJMG, vol.19, pp.29-34, 2016 (Journal Indexed in SCI)
- XI. **Associations of fractalkine receptor (CX3CR1) and CCR5 gene variants with hypertension, diabetes and atherosclerosis in chronic renal failure patients undergoing hemodialysis**
BAĞCI B., Bağci G., Huzmeli C., Sezgin I., ÖZDEMİR Ö.
INTERNATIONAL UROLOGY AND NEPHROLOGY, vol.48, no.7, pp.1163-1170, 2016 (Journal Indexed in SCI)
- XII. **The CYP4502D6*4 and*6 alleles are the molecular genetic markers for drug response: implications in colchicine non-responder FMF patients**
Yalcintepe S., ÖZDEMİR Ö., SILAN C., Ozen F., ULUDAĞ A., CANDAN F., SILAN F.
EUROPEAN JOURNAL OF DRUG METABOLISM AND PHARMACOKINETICS, vol.41, no.3, pp.281-286, 2016 (Journal Indexed in SCI)
- XIII. **Vitamin D Receptor Gene BSMI, FOKI, APAI, and TAQI Polymorphisms and the Risk of Atopic Dermatitis**
KILIÇ S., SILAN F., Hiz M. M. , IŞIK S., OGRET MEN Z., ÖZDEMİR Ö.
JOURNAL OF INVESTIGATIONAL ALLERGOLOGY AND CLINICAL IMMUNOLOGY, vol.26, no.2, pp.106-110, 2016 (Journal Indexed in SCI)
- XIV. **KRAS, BRAF oncogene mutations and tissue specific promoter hypermethylation of tumor suppressor SFRP2, DAPK1, MGMT, HIC1 and p16 genes in colorectal cancer patients**
BAĞCI B., SARI M., KARADAYI K., Turan M., ÖZDEMİR Ö., Bağci G.
CANCER BIOMARKERS, vol.17, no.2, pp.133-143, 2016 (Journal Indexed in SCI)
- XV. **The relationship between C-reactive protein rs3091244 polymorphism and ankylosing spondylitis**
Akbal A., REŞORLU H., Gokmen F., Savas Y., ZATERİ C., Sargin B., BOZKURT E., SILAN F., ÖZDEMİR Ö.
INTERNATIONAL JOURNAL OF RHEUMATIC DISEASES, vol.19, no.1, pp.43-48, 2016 (Journal Indexed in SCI)
- XVI. **Hyperimmunoglobulin D Syndrome: Case Report**
Sen H., SILAN F., Binnetoglu E., Gunes F., Akurut C., ULUDAĞ A., ÖZDEMİR Ö.
ARCHIVES OF RHEUMATOLOGY, vol.30, no.3, pp.244-246, 2015 (Journal Indexed in SCI)
- XVII. **Aldosterone Synthase CYP11B2 Gene Promoter Polymorphism in a Turkish Population With Chronic Kidney Disease**
YILMAZ M., Sari I., BAĞCI B., Gumus E., ÖZDEMİR Ö.
IRANIAN JOURNAL OF KIDNEY DISEASES, vol.9, no.3, pp.209-214, 2015 (Journal Indexed in SCI)
- XVIII. **The protective effect of MCP-1-2518 A > G promoter polymorphism in Turkish chronic renal failure patients requiring long-term hemodialysis**
BAĞCI B., Bağci G., CANDAN F., ÖZDEMİR Ö., Sezgin I.
INTERNATIONAL UROLOGY AND NEPHROLOGY, vol.47, no.3, pp.551-556, 2015 (Journal Indexed in SCI)
- XIX. **BcII-RFLP profiles for serum amyloid A1 and mutated MEFV gene prevalence in chronic renal failure patients requiring long-term hemodialysis**
Ozdemir O., Kayatas M., Cetinkaya S., YILDIRIM M. E. , SILAN F., Kurtulgan H. K. , Koksall B., Urfali M., CANDAN F.
RENAL FAILURE, vol.37, no.2, pp.292-296, 2015 (Journal Indexed in SCI)
- XX. **Is the HLA B27 genotype a risk factor for psoriatic arthritis and psoriasis vulgaris?**
Ogretmen Z., Hiz M. M. , SILAN F., Kosar S., ÖZDEMİR Ö.
TURKDERM-TURKISH ARCHIVES OF DERMATOLOGY AND VENEROLOGY, vol.48, no.3, pp.131-134, 2014 (Journal Indexed in SCI)
- XXI. **Effects of the Chemokine Receptor 5 (CCR5)-Delta32 Mutation on Hepatitis C Virus-Specific Immune Responses and Liver Tissue Pathology in HCV Infected Patients**

- Yilmaz A., Alagozlu H., Ozdemir O., Arici S.
HEPATITIS MONTHLY, vol.14, no.7, 2014 (Journal Indexed in SCI)
- XXII. **Association of endothelial nitric oxide synthase Glu298Asp gene polymorphism in psoriasis cases with hypertension.**
Ogretmen Z., Hiz M. M. , SILAN F., ULUDAĞ A., ÖZDEMİR Ö.
Annals of Saudi medicine, vol.34, no.4, pp.340-5, 2014 (Journal Indexed in SCI Expanded)
- XXIII. **Association Between Inherited Thrombophilia and Impaired Right Ventricular Function in Deep Vein Thrombosis Without Symptomatic Pulmonary Embolism**
AŞGÜN H. F. , KIRILMAZ B., Saygi S., Ozturk O., SILAN F., KARATAĞ O., Kosar S., ÖZDEMİR Ö.
CLINICAL AND APPLIED THROMBOSIS-HEMOSTASIS, vol.20, no.3, pp.270-277, 2014 (Journal Indexed in SCI)
- XXIV. **The prevalence of Familial Mediterranean Fever common gene mutations in patients with simple febrile seizures**
OZEN F., Kocak N., Kelekci S., Yildirim I. H. , HACIMUTO G., ÖZDEMİR Ö.
EUROPEAN REVIEW FOR MEDICAL AND PHARMACOLOGICAL SCIENCES, vol.18, no.5, pp.657-660, 2014 (Journal Indexed in SCI)
- XXV. **Relationship Between Response to Colchicine Treatment and MDR1 Polymorphism in Familial Mediterranean Fever Patients**
ULUDAĞ A., SILAN C., Atik S., Akurut C., Uludag A., SILAN F., ÖZDEMİR Ö.
GENETIC TESTING AND MOLECULAR BIOMARKERS, vol.18, no.2, pp.73-76, 2014 (Journal Indexed in SCI)
- XXVI. **Endothelial function and germ-line ACE I/D, eNOS and PAI-1 gene profiles in patients with coronary slow flow in the Canakkale population: multiple thrombophilic gene profiles in coronary slow flow**
GAZİ E., Temiz A., Altun B., BARUTÇU A., SILAN F., Colkesen Y., ÖZDEMİR Ö.
CARDIOVASCULAR JOURNAL OF AFRICA, vol.25, no.1, pp.9-14, 2014 (Journal Indexed in SCI)
- XXVII. **Methylenetetrahydrofolate Reductase Gene Germ-Line C677T and A1298C SNPs are Associated with Colorectal Cancer Risk in the Turkish Population**
Ozen F., Sen M., Ozdemir O.
ASIAN PACIFIC JOURNAL OF CANCER PREVENTION, vol.15, no.18, pp.7731-7735, 2014 (Journal Indexed in SCI)
- XXVIII. **Intercellular Adhesion Molecule-1 K469E and Angiotensinogen T207M Polymorphisms in Coronary Slow Flow**
GAZİ E., BARUTÇU A., Altun B., Temiz A., Bekler A., Urfali M., SILAN F., Colkesen Y., ÖZDEMİR Ö.
MEDICAL PRINCIPLES AND PRACTICE, vol.23, no.4, pp.346-350, 2014 (Journal Indexed in SCI)
- XXIX. **Association Between ABCB1 (MDR1) Gene Polymorphism and Unresponsiveness Combined Therapy in Chronic Hepatitis C virus**
Timucin M., Alagozlu H., ÖZDEMİR S., Ozdemir O.
HEPATITIS MONTHLY, vol.13, no.4, 2013 (Journal Indexed in SCI)
- XXX. **The Proto-Oncogene KRAS and BRAF Profiles and Some Clinical Characteristics in Colorectal Cancer in the Turkish Population**
OZEN F., ÖZDEMİR S., ZEMHERİ E., HACIMUTO G., SILAN F., ÖZDEMİR Ö.
GENETIC TESTING AND MOLECULAR BIOMARKERS, vol.17, no.2, pp.135-139, 2013 (Journal Indexed in SCI)
- XXXI. **Prevalence of Common YMDD Motif Mutations in Long Term Treated Chronic HBV Infections in a Turkish Population**
Alagozlu H., ÖZDEMİR Ö., Koksall B., Yilmaz A., Coskun M.
ASIAN PACIFIC JOURNAL OF CANCER PREVENTION, vol.14, no.9, pp.5489-5494, 2013 (Journal Indexed in SCI)
- XXXII. **Possible Roles of the Xenobiotic Transporter P-glycoproteins Encoded by the MDR1 3435 C > T Gene Polymorphism in Differentiated Thyroid Cancers**
ÖZDEMİR S., ULUDAĞ A., SILAN F., Atik S. Y. , Turgut B., ÖZDEMİR Ö.
ASIAN PACIFIC JOURNAL OF CANCER PREVENTION, vol.14, no.5, pp.3213-3217, 2013 (Journal Indexed in SCI)
- XXXIII. **DOUBLE TRANSLOCATION: AN INTERESTING FAMILY HISTORY**
UYSAL A. O. , ULUDAĞ A., SILAN F., Ercelen N., Zafer C., ÖZDEMİR Ö.
BALKAN JOURNAL OF MEDICAL GENETICS, vol.16, no.1, pp.77-80, 2013 (Journal Indexed in SCI)
- XXXIV. **Combined Germline Variations of Thrombophilic Genes Promote Genesis of Lung Cancer**

- Ozen F., POLAT F., ARSLAN S., Ozdemir O.
ASIAN PACIFIC JOURNAL OF CANCER PREVENTION, vol.14, no.9, pp.5449-5454, 2013 (Journal Indexed in SCI)
- XXXV. **The effect of 3-methylcholanthrene and butylated hydroxytoluene on glycogen levels of liver, muscle, testis, and tumor tissues of rats**
POLAT F., DERE E., Gul E., YELKUVAN İ., ÖZDEMİR Ö., BİNGÖL G.
TURKISH JOURNAL OF BIOLOGY, vol.37, no.1, pp.33-38, 2013 (Journal Indexed in SCI)
- XXXVI. **The prevalence of VKORC1 1639 G > A and CYP2C9*2*3 genotypes in patients that requiring anticoagulant therapy in Turkish population**
SILAN C., DOĞAN Ö. T. , Silan F., Kukulguven F. M. , AŞGÜN H. F. , Ozdemir S., ULUDAĞ A., Atik S., Gungor B., AKDUR S., et al.
MOLECULAR BIOLOGY REPORTS, vol.39, no.12, pp.11017-11022, 2012 (Journal Indexed in SCI)
- XXXVII. **Increased T-Allele Frequency of 677 C > T Polymorphism in the Methylenetetrahydrofolate Reductase Gene in Differentiated Thyroid Carcinoma**
ÖZDEMİR S., SILAN F., HASBEK Z., ULUDAĞ A., Atik S., Erselcan T., ÖZDEMİR Ö.
GENETIC TESTING AND MOLECULAR BIOMARKERS, vol.16, no.7, pp.780-784, 2012 (Journal Indexed in SCI)
- XXXVIII. **Alterations in Promoter Methylation Status of Tumor Suppressor HIC1, SFRP2, and DAPK1 Genes in Prostate Carcinomas**
Kilinc D., Ozdemir O., ÖZDEMİR S., KORĞALI E., Koksall B., USLU A., Gultekin Y. E.
DNA AND CELL BIOLOGY, vol.31, no.5, pp.826-832, 2012 (Journal Indexed in SCI)
- XXXIX. **Recurrent Pregnancy Loss and Its Relation to Combined Parental Thrombophilic Gene Mutations**
ÖZDEMİR Ö., Yenicesu G. I. , SILAN F., Koksall B., Atik S., Ozen F., Gol M., ÇETİN A.
GENETIC TESTING AND MOLECULAR BIOMARKERS, vol.16, no.4, pp.279-286, 2012 (Journal Indexed in SCI)
- XL. **Combined point mutations in codon 12 and 13 of KRAS oncogene in prostate carcinomas**
SILAN F., Gultekin Y., Atik S., Kilinc D., ALAN C., Yildiz F., ULUDAĞ A., ÖZDEMİR Ö.
MOLECULAR BIOLOGY REPORTS, vol.39, no.2, pp.1595-1599, 2012 (Journal Indexed in SCI)
- XLI. **Combined Effect of Factor V Leiden, MTHFR, and Angiotensin-Converting Enzyme (Insertion/Deletion) Gene Mutations in Hypertensive Adult Individuals: A Population-Based Study from Sivas and Canakkale, Turkey**
DEMİREL Y., Dogan S., ULUDAĞ A., SILAN C., Atik S., SILAN F., ÖZDEMİR Ö.
GENETIC TESTING AND MOLECULAR BIOMARKERS, vol.15, no.11, pp.785-791, 2011 (Journal Indexed in SCI)
- XLII. **Increased T allele frequency in MTHFR C677T gene in thyroid carcinoma**
ÖZDEMİR S., SILAN F., Erselcan T., ULUDAĞ A., Ustun F., Colak A., Atik S., ÖZDEMİR Ö.
CHROMOSOME RESEARCH, vol.19, 2011 (Journal Indexed in SCI)
- XLIII. **Prevalence of known mutations in the MEFV gene in a population screening with high rate of carriers**
Ozdemir O., Sezgin I., Kurtulgan H. K. , CANDAN F., Koksall B., Sumer H., Icagasioglu D., USLU A., Yildiz F., ARSLAN S., et al.
MOLECULAR BIOLOGY REPORTS, vol.38, no.5, pp.3195-3200, 2011 (Journal Indexed in SCI)
- XLIV. **The endemic RTL80V/I and RTM204V/I YMDD mutation profiles in a case of chronic hepatitis B**
ÖZDEMİR Ö., Alagozlu H., Timucin M., ÖZDEMİR S., KORKMAZ M., Koksall B., Ozen O.
CHROMOSOME RESEARCH, vol.19, 2011 (Journal Indexed in SCI)
- XLV. **In Vivo Evaluation of the Genotoxic Effects of Gonadotropins on Rat Reticulocytes**
Duran B., Koc O., OZDEMIRCI S., Topcuoglu A., ÖZDEMİR Ö.
CURRENT THERAPEUTIC RESEARCH-CLINICAL AND EXPERIMENTAL, vol.72, no.2, pp.60-70, 2011 (Journal Indexed in SCI)
- XLVI. **Association between ABCB1 (MDR1) Gene 3435 C > T Polymorphism and Colchicine Unresponsiveness of FMF Patients**
Ozen F., SILAN C., ULUDAĞ A., CANDAN F., SILAN F., Ozdemir S., Atik S., ÖZDEMİR Ö.
RENAL FAILURE, vol.33, no.9, pp.899-903, 2011 (Journal Indexed in SCI)
- XLVII. **GJB2 35delG and Mitochondrial A1555G Mutations and Etiology of Deafness at the Gelibolu School for the Deaf in Turkey**

- SILAN F., GÜÇLÜ O., KADIOĞLU L. E. , SILAN C., ATIK S., ULUDAĞ A., DEMIRAY A., ÖZDEMİR Ö., Derekoy F. S.
JOURNAL OF INTERNATIONAL ADVANCED OTOLOGY, vol.7, no.3, pp.361-371, 2011 (Journal Indexed in SCI)
- XLVIII. **CCR2 Polymorphism in Chronic Renal Failure Patients Requiring Long-Term Hemodialysis**
Sezgin I., Koksall B., Bagci G., Kurtulgan H. K. , ÖZDEMİR Ö.
INTERNAL MEDICINE, vol.50, no.21, pp.2457-2461, 2011 (Journal Indexed in SCI)
- XLIX. **Frequencies of micronuclei (MNi), nucleoplasmic bridges (NPBs), and nuclear buds (NBUDs) in farmers exposed to pesticides in Canakkale, Turkey**
Coskun M., Coskun M., ÇAYIR A., ÖZDEMİR Ö.
ENVIRONMENT INTERNATIONAL, vol.37, no.1, pp.93-96, 2011 (Journal Indexed in SCI)
- L. **Frequency dependence of conductivity in intrinsic amorphous silicon carbide film, assessed through admittance measurement of metal insulator semiconductor structure**
OZDEMIR O., ATILGAN İ., AKAOĞLU B., Sel K., KATIRCIÖĞLU B.
THIN SOLID FILMS, vol.497, pp.149-156, 2006 (Journal Indexed in SCI)
- LI. **Iodine-131 treatment and chromosomal damage: in vivo dose-effect relationship**
ERSELCAN T., SUNGU S., Ozdemir S., TURGUT B., DOĞAN D., OZDEMIR O.
EUROPEAN JOURNAL OF NUCLEAR MEDICINE AND MOLECULAR IMAGING, vol.31, no.5, pp.676-684, 2004 (Journal Indexed in SCI)

Articles Published in Other Journals

- I. **Diagnostic outcomes for genetic testing of 54 genes in pregnancy loss using array CGH method: A two-year retrospective study Gebelik kayıplarında 54 genin array CGH methoduyla yapılan tanısal sonuçları: İki yıllık retrospektif çalışma**
PAKSOY B., ÖZDEMİR Ö., SILAN F.
Jinekoloji-Obstetrik ve Neonatoloji Tıp Dergisi, vol.17, no.4, 2020 (Other Refereed National Journals)
- II. **Tedaviyi Etkileyen Tarama Testleri**
KARAKAYA T., SILAN F., ÖZDEMİR Ö.
Türkiye Klinikleri Sağlık Bilimleri Dergisi, 2020 (Other Refereed National Journals)
- III. **Is BCL11B a potential candidate gene for the diffuse cutaneous mastocytosis: A case report**
SILAN F., ALBUZ B., Bourouba R., ÖZTÜRK M., YILDIZ O., ÖZDEMİR Ö.
Cumhuriyet Medical Journal, vol.42, no.3, 2020 (Other Refereed National Journals)
- IV. **MEME KANSERLİ HASTALAR VE BİRİNCİ DERECEYAKINLARINDA PERİFERAL TRANSKRİPTOMPROFİLLERİNİN ARAŞTIRILMASI**
ERDİŞ E., YÜCEL B., ÖZDEMİR Ö.
Bozok Tıp Derg, vol.9, no.4, pp.27-31, 2019 (Other Refereed National Journals)
- V. **Warfarin Resistance: A Case Report**
GÖNLÜGÜR U., GÖNLÜGÜR U., ÖZDEMİR Ö., SILAN F.
EURASIAN JOURNAL OF EMERGENCY MEDICINE, vol.18, no.1, pp.61-63, 2019 (Journal Indexed in ESCI)
- VI. **The diagnostic accuracy of non-invasive fetal RhD genotyping by using cell-free fetal DNA in maternal plasma. (Maternal plazmadaki hücre dışı serbest fetal DNA kullanılarak girişimsel olmayan fetal RhD genotiplemesinin tanısal doğruluğu)**
Akurut Ç., SILAN F., YALÇINTEPE S., ÖZDEMİR Ö.
FAMILY PRACTICE AND PALLIATIVE CARE, vol.4, no.1, pp.1-6, 2019 (Other Refereed National Journals)
- VII. **Prenatal diagnosis of aneuploidies and microdeletion/duplication in amniotic fluid and fetal aborted material by QF-PCR and MLPA analysis.**
Ari E., ÖZDEMİR Ö., Djurovic J., SILAN F.
Biomed Genet Genomics, vol.3, no.1, pp.1-6, 2018 (Refereed Journals of Other Institutions)
- VIII. **The clinical, cytogenetics and molecular characterization of inverted duplication/deletion of chromosome 8p in a boy with mental and motor retardation: Genotype-phenotype correlation in a case report**

SILAN F., BOUROUBA R., KARAKAYA T., YILDIZ O., PAKSOY B., URFALI M., ÖZDEMİR Ö.

Egyptian Journal of Medical Human Genetics, vol.19, no.4, pp.437-441, 2018 (Refereed Journals of Other Institutions)

- IX. **Evaluation of the physical and biological dosimetry methods in iodine-131-treated patients**
Özdal A., ERSELCAN T., ÖZDEMİR Ö., ÖZGÜVEN Y., Silov G., Erdoğan Z.
World Journal of Nuclear Medicine, vol.17, no.4, pp.253-260, 2018 (Refereed Journals of Other Institutions)
- X. **The frequency of toll-like receptor 4 gene polymorphism in ankylosing spondylitis and its relationship between disease activity.**
Sargin B., AKBAL A., REŞORLU H., SAVAŞ Y., ZATERİ C., SILAN F., ÖZDEMİR Ö.
The European Research Journal, 2017 (Refereed Journals of Other Institutions)
- XI. **THE FREQUENCY OF TOLL-LIKE RECEPTOR 4 GENE POLYMORPHISM IN ANKYLOSING SPONDYLITIS AND ITS RELATIONSHIP BETWEEN DISEASE ACTIVITY**
SARGIN B., AKBAL A., REŞORLU H., SAVAŞ Y., ZATERİ C., SILAN F., ÖZDEMİR Ö.
THE EUROPEAN RESEARCH JOURNAL, 2017 (Refereed Journals of Other Institutions)
- XII. **Kanser Etiyolojisinde Tetikleyici Moleküler Mekanizmalar**
ÖZDEMİR Ö., KURU B., PAKSOY B., SILAN F.
Türkiye Klinikleri J Med Genet-Special Topics, vol.2, no.2, pp.74-87, 2017 (Other Refereed National Journals)
- XIII. **Günümüzde Kanser Tanısında Kullanılan Geçerli ve Güvenilir Moleküler Tetkikler**
Yalcintepe S., ÖZDEMİR Ö., Guler Z., SILAN F.
Türkiye Klinikleri Journal of Medical Genetics-Special Topics, vol.2, no.2, pp.113-122, 2017 (Other Refereed National Journals)
- XIV. **Çok Parametrel ve Çoklu Bilinmeyenli Kanser**
ÖZDEMİR Ö.
Türkiye Klinikleri Journal of Medical Genetics-Special Topics, vol.2, no.2, pp.60-68, 2017 (Other Refereed National Journals)
- XV. **Kanser Etiyolojisinde Tetikleyici Moleküler Mekanizmalar**
ÖZDEMİR Ö., Kuru B., PAKSOY B., SILAN F.
Türkiye Klinikleri Journal of Medical Genetics-Special Topics, vol.2, no.2, pp.74-87, 2017 (Other Refereed National Journals)
- XVI. **Macular and choroidal thickness of children with Familial Mediterranean Fever gene mutation**
BATTAL F., AYLANÇ H., YILDIRIM Ş., YELİZ E., SILAN F., ÖZDEMİR Ö.
Family Practice and Palliative Care, pp.23-28, 2017 (Refereed Journals of Other Institutions)
- XVII. **The GJB2 gene mutation profiles in hearing impaired patients from Western Turkey, Canakkale**
SILAN F., Kankaya D., KARAKAYA T., PAKSOY B., TÜRÜNZ V., ÖZDEMİR Ö.
Biomedical Genetics and Genomics, vol.2, no.2, pp.1-5, 2017 (Refereed Journals of Other Institutions)
- XVIII. **A mosaic infertile case of isodicentric Y-chromosome with duplicated SRY, SHOX and deleted AZF locus**
ÖZDEMİR Ö., PAKSOY B., SILAN F.
Biomedical Genetics and Genomics, vol.2, no.1, pp.1-3, 2017 (Refereed Journals of Other Institutions)
- XIX. **Fatma Silan, Duygu Kankaya, Taner Karakaya, Baris Paksoy, Volkan Turunz and Ozturk Ozdemir. The GJB2 gene mutation profiles in hearing impaired patients from Western Turkey, Canakkale.**
SILAN F., Kankaya D., KARAKAYA T., PAKSOY B., Turunz V., ÖZDEMİR Ö.
Biomedical Genetics and Genomics, vol.2, no.2, pp.1-5, 2017 (Refereed Journals of Other Institutions)
- XX. **Hypermethylated promoter profiles for tumour suppressor APC p53 MSH6 and MGMT genes in CRC tumours**
SİK E., ÖZDEMİR Ö., KÜÇÜK KURTULGAN H., URFALI M., ŞEN M., SILAN F.
Journal of Biomedical Research, vol.2, no.6, pp.41-47, 2016 (Refereed Journals of Other Institutions)
- XXI. **A mental and motor retarded dysmorphic case with heterozygous 1p36 deletion. Comparable results from cytogenetic MicroArray CGH FISH and MLPA techniques**
SILAN F., YILDIZ O., URFALI M., GÜLER Z., ÖZDEMİR Ö.
Merit Research Journal of Medicine and Medical Sciences, 2016 (Refereed Journals of Other Institutions)

- XXII. **9qh liđi Molar Gebelik için bir risk faktörü mü**
ÇAKIR GÜNGÖR A. N. , SILAN F., KILINÇ N., GENCER M., ÖZDEMİR Ö.
Jinekoloji-Obstetrik ve Neonatoloji Tıp Dergisi, pp.14-15, 2016 (Refereed Journals of Other Institutions)
- XXIII. **308G/A and 238G/A polymorphisms in the TNF-alpha gene may not contribute to the risk of arthritis among Turkish psoriatic patients**
IŞIK S., SILAN F., KILIÇ S., Hiz M. M. , Ogretmen Z., ÖZDEMİR Ö.
EGYPTIAN RHEUMATOLOGIST, vol.38, no.4, pp.313-317, 2016 (Journal Indexed in ESCI)
- XXIV. **Hypermethylated promoter profiles for tumoursupressor APC p53 MSH6 and MGMT genes in CRCtumours**
Şık E., ÖZDEMİR Ö., KÜÇÜK KURTULGAN H., URFALI M., SILAN F.
Pyrex Journal of Biomedical Research, 2016 (Refereed Journals of Other Institutions)
- XXV. **Comparison of the thrombophilic gene polymorphismsand recurrent pregnancy loss Results on combined geneeffect of FV Leiden FVR2 FXIII MTHFR A1298Cand C677T PAI 1 4G 5G and ACE I D genes in RPLWomen from Misk Belarus and Canakkale Sivas Turkey**
YILDIZ O., SILAN F., ÖZDEMİR Ö., Kuru B., Mosse I., Kilchevsky A., Gonchar A., Seldyar N.
Biomedical Genetics and Genomics, 2016 (Refereed Journals of Other Institutions)
- XXVI. **An infertile case of 47 XYY syndromewithout autistic spectrum Cost effectivewell define of extra Y chromosome byGTG C bandings QF PCR and FISHanalyses**
ÖZDEMİR Ö., PAKSOY B., Gürgen A., URFALI M., YILDIZ O., ULUDAĞ A., SILAN F.
Cumhuriyet Tıp Dergisi, 2016 (Refereed Journals of Other Institutions)
- XXVII. **GÜNGÖR ANC SILAN F KILINC N GENCER M ULUDAĞ A COŞAR E KOC E OZDEMIR O Is 9qh Positivity A Risk Factor For Molar Pregnancy 2016 14 15**
Güngörçakar A., SILAN F., GENCER M., ULUDAĞ A., COŞAR E., KOÇ E., ÖZDEMİR Ö.
Jinekoloji - Obstetrik ve Neonatoloji Tıp Dergisi, pp.14-15, 2016 (Other Refereed National Journals)
- XXVIII. **9qh liđi Molar Gebelik İçin Bir Risk Faktörü mü**
ÇAKIR GÜNGÖR A. N. , SILAN F., KILINÇ N., GENCER M., ULUDAĞ A., COŞAR E., KOÇ E., ÖZDEMİR Ö.
Jinekoloji - Obstetrik ve Neonatoloji Tıp Dergisi, 2016 (Other Refereed National Journals)
- XXIX. **Triploidy Diploidy Mosaisizm Diandry and Uniparental Isodisomy Fetus withOmphalocele and Contracted Finger**
SILAN F., ÇAKIR GÜNGÖR A. N. , URFALI M., ULUDAĞ A., ÖZDEMİR Ö.
Family Medicine & Medical ScienceResearch, 2015 (Refereed Journals of Other Institutions)
- XXX. **Multiple Inherited Thrombophilic Gene Polymorphisms in Spontaneous Abortions in Turkish Population**
Yalcintepe S., ÖZDEMİR Ö., HACIVELİOĞLU S. Ö. , Akurut C., Koc E., ULUDAĞ A., Cosar E., SILAN F.
INTERNATIONAL JOURNAL OF MOLECULAR AND CELLULAR MEDICINE, vol.4, no.2, pp.120-127, 2015 (Journal Indexed in ESCI)
- XXXI. **Tiroid Kanserinde Moleküler Etyolojik Faktörler Derleme**
ÖZDEMİR S., ÖZDEMİR Ö.
Cumhuriyet Journal Medical, vol.36, pp.128-146, 2014 (Other Refereed National Journals)
- XXXII. **Tiyopentalin rat karaciđer NADPH sitokrom P 450 redüktaz NADH Sitokrom b5 redüktaz ve N Nitrozodimetilamin demetilaz enzim aktiviteleri üzerindeki etkisi**
ÖZDEMİR S., SİLİÇ Y., ÖZDEMİR Ö., ÇETİNKAYA Ö.
Cumhuriyet Üniversitesi Tıp Fakültesi Dergisi, vol.22, no.4, pp.193-195, 2000 (Other Refereed National Journals)

Books & Book Chapters

- Biyoteknoloji esasları ve tıbbi genetikte yeni uygulama alanları**
ÖZDEMİR Ö., SILAN F.
in: Tıbbi Genetik ve Klinik Uygulamaları, Munis DÜNDAR, Editor, MG GRUP MATBAACILIK, KAYSERİ 2016;syf 935-958, Kayseri, pp.539-556, 2016

II. Kardiyovasküler sistem hastalıklarına genetik yaklaşım

SILAN F., ÖZDEMİR Ö.

in: Tıbbi Genetik ve Klinik Uygulamaları, Munis DÜNDAR, Editor, MG GRUP MATBAACILIK, Kayseri, pp.935-958, 2016

III. Kardiyovasküler Hastalıkların Genetiği

SILAN F., ÖZDEMİR Ö.

in: Tıbbi Genetik ve Klinik Uygulamaları, Munis DüNDAR, Editor, Erciyes Üniversitesi, 2016

Refereed Congress / Symposium Publications in Proceedings

- I. **Tekrarlayan Gebelik Kaybı olan olguların kopya sayısı değişikliklerinin profillemesi**
YILDIZ O., SILAN F., KARAKAYA T., ÖZDEMİR Ö.
Ulusal tıbbi genetik kongresi 2020, Online, Turkey, 20 November 2020
- II. **Multiple pterygia, joint contractures, hypoplastic lungs and hydrops: A rare case of lethal multiple pterygium syndrome**
BAKIOĞLU KAYA D., AKCAN M. B. , SÖNMEZ V., KABLAN A., ÖZDEMİR Ö., SILAN F.
Ulusal tıbbi genetik kongresi 2020, Online, Turkey, 20 November 2020
- III. **HbF veya HbA2 seviyelerinde farklılığı bulunan bireylerde Kruppel-like factor 1 (KLF1) ve Hemoglobin subunit delta (HBD) genlerinde saptanan mutasyonların genotip-fenotip ilişkisinin incelenmesi**
KARAKAYA T., SILAN F., ÖZDEMİR Ö.
Ulusal tıbbi genetik kongresi 2020, Online, Turkey, 20 November 2020
- IV. **ÇOMÜ Tıbbi Genetik Tanı Merkezi'nde Spinal Musküler Atrofi Taşıyıcılığı Açısından Genotiplendirilen Olguların SMN1/SMN2 Genlerinin Kopya Sayılarının Analizi**
ÖZTÜRK M., ÖZDEMİR Ö., SILAN F.
Ulusal tıbbi genetik kongresi 2020, Online, Turkey, 20 November 2020
- V. **Oculocutaneous albinism caused by a compound heterozygous mutations in TYR gene: a case report**
AKCAN M. B. , BAKIOĞLU KAYA D., SÖNMEZ V., KABLAN A., ÖZDEMİR Ö., SILAN F.
Ulusal tıbbi genetik kongresi 2020, Online, Turkey, 20 November 2020
- VI. **Investigation of autoinflammatory genes in patients with conventional treatment-resistant dermatological diseases Poster presentation,**
IŞIK S., ALBUZ B., AKBAŞ N. E. , SILAN F., ÖZDEMİR Ö.
European Biotechnology Congress 2020, Prag, Czech Republic, 24 September 2020
- VII. **A rare lethal multiple pterygium syndrome caused by a homozygous point mutation In CHRNG gene:a case report.**
AKCAN M. B. , ALBUZ B., KABLAN A., ÖZDEMİR Ö., SILAN F.
European Biotechnology Congress 2020, Czech Republic, 24 September 2020
- VIII. **The Genomics of Arthrogryposis, a Complex Trait: Candidate Genes and Further Evidence for Oligogenic Inheritance**
ÖZDEMİR S., SILAN F., AKGÜN M. Y. , KOÇ ÖZTÜRK F., ÖZDEMİR Ö.
European Biotechnology Congress 2020, Prag, Czech Republic, 24 September 2020
- IX. **A novel variant for SQSTM1 gene in a patient with frontotemporal dementia.**
OCAK Ö., KARAKAYA T., ALBUZ B., AKBAŞ N. E. , SÖNMEZ V., AKCAN M. B. , SILAN F., ÖZDEMİR Ö.
European Biotechnology Congress 2020, Prag, Czech Republic, 24 September 2020
- X. **Additional point mutations in ACAN and GATA4 genes in an atypical achondroplasia patient with platyspondyly and congenital heart defect.**
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European Biotechnology Congress 2020, Prag, Czech Republic, 24 September 2020
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ALBUZ B., AKBAŞ N. E. , KARAKAYA T., SÖNMEZ V., AKCAN M. B. , SILAN F., ÖZDEMİR Ö.

European Biotechnology Congress 2020, Prag, Czech Republic, 24 September 2020

XII. A Homozygous Nonsense Mutation In MTHFR Gene Causes A Severe Phenotype: Hypotonia And Hydrocephalus. Poster presentation

ALBUZ B., MALÇOK Ü. A. , SILAN F., ÖZDEMİR Ö.

European Biotechnology Congress 2020, Prag, Czech Republic, 24 September 2020

XIII. Chromatin remodelling dysfunction and CHD2 related epilepsy: Reports of two unrelated child.

KABLAN A., ALBUZ B., AKBAŞ N. E. , KARAKAYA T., SÖNMEZ V., AKCAN M. B. , SILAN F., ÖZDEMİR Ö.

European Biotechnology Congress 2020, Prag, Czech Republic, 24 September 2020

XIV. A Rare Chromosomal Aneuploidy That Diagnosed With Array CGH And MLPA Validation : Trisomy 4 in a fetus

SÖNMEZ V., ALBUZ B., KARAKAYA T., AKBAŞ N. E. , SILAN F., ÖZDEMİR Ö.

V. International Participated Erciyes Medical Genetics Days Congress, Nevşehir, Turkey, 20 February 2020

XV. HOMOZYGOUS FRAMESHİFT MUTATION İN DDB2 GENE İNDUCED SQUAMOUS CELL AND BASAL CELL CARCİNOMAS İN A CHİLD WİTH XERODERMA PİGMENTOSUM

AKBAŞ N. E. , ALBUZ B., SILAN F., ÖZDEMİR Ö.

V. International Participated Erciyes Medical Genetics Days Congress, Nevşehir, Turkey, 20 February 2020

XVI. A Novel Combined C.1630T>C AND C.1579G>A Point Mutations in ALOX12B Gene in A Rare Autosomal Recessive Congenital Ichthyosis : A Case Report

ALBUZ B., AKBAŞ N. E. , AYLANÇ H., SILAN F., ÖZDEMİR Ö.

1st Bursa International Genetics Days: Dermatogenetics Symposium, Bursa, Turkey, 09 January 2020

XVII. Novel Missense Mutation in Phospholipase C-Gamma-2 Gene (PLCG2) Causes Cold-Induced Psoriasis : A Case report

SILAN F., ALBUZ B., SÖNMEZ V., ÖZDEMİR Ö.

1st Bursa International Genetics Days: Dermatogenetics Symposium, Bursa, Turkey, 09 January 2020

XVIII. BCL11B gene may be a candidate gene for mastocytosis in a patient with partial trisomy of distal 14q

ALBUZ B., SILAN F., ÖZTÜRK M., YILDIZ O., ÖZDEMİR Ö.

13th Balkan Congress of Human Genetics, Edirne, Turkey, 17 - 20 April 2019, vol.22, pp.156-1156

XIX. A rare heterochromatin polymorphism of chromosome 6 associated with recurrent miscarriage: A case report

ALBUZ B., SILAN F., ÖZTÜRK M., KARAKAYA T., ÖZDEMİR Ö.

13. Balkan Congress of Human Genetics, Edirne, Turkey, 17 - 20 April 2019, vol.0

XX. Mosaic Trisomy 9 Presenting With Congenital Diaphragmatic Hernia And Facial Dysmorphism

Öztürk M., DEMİR B., Arayıcı S., Demir S., ÖZDEMİR Ö., SILAN F.

13th Balkan Human Genetic Congress, Edirne, Turkey, 17 - 20 April 2019

XXI. The c.1397CG and c.3209GA mutations in exon 10 of CFTR gene in an infertile men with oligoastenozoospermia

SILAN F., DİNÇSOY BİR F., ERSAY A. R. , KARAKAYA T., ÖZDEMİR Ö.

VII Baltic Genetics Congress, Riga, Latvia, 24 - 27 October 2018, vol.16, pp.255

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European Biotechnology Congress, Athens, Greece, 26 - 28 April 2018, vol.280
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SILAN F., YÖRÜK YAYAR Ö., DİNÇSOY BİR F., KARAKAYA T., PAKSOY B., ÖZDEMİR Ö.
Erciyes Tıp Genetik Günleri 2018, Kayseri, Turkey, 7 - 10 March 2018, vol.40, pp.74
- XXIX. **Çanakkale popülasyonunda HBB(Hemoglobin Subunit Beta) geni mutasyon profilleri**
SILAN F., KARAKAYA T., DİNÇSOY BİR F., ÖZDEMİR Ö.
3.Hematolojik Genetik Sempozyumu, İzmir, Turkey, 14 - 16 February 2018, pp.65
- XXX. **Kliniğimizde Takip Edilen Kadın Hastalarda Servikal Neoplaziyi Değerlendirmek İçin Kullanılan Pap Smear ve HPV DNA Test Sonuçlarının Retrospektif Analizi**
BEYAZİT F., SILAN F., GENCER M., AYDIN B., PAKSOY B., ÜNSAL M. A. , ÖZDEMİR Ö.
İstanbul Üniversitesi 7. Kadın Doğum Günleri, İstanbul, Turkey, 7 - 10 December 2017
- XXXI. **Whole exome sequencing reveals potential oligogenic inheritance and candidate novel genes in patients with arthrogyriposis**
BAYRAM Y., ULUDAĞ ALKAYA D., pehlivan d., Gezdirici A., SILAN F., ÖZDEMİR Ö., ELÇİOĞLU H. N. , YILDIZ O., yavuz ş., TÜYSÜZ B.
American Society of Human Genetics 67th Annual Meeting, 17 - 21 October 2017
- XXXII. **Two candidate genes for recurrent pregnancy loss and infertility: Could ZP3 and UPK3B give us new diagnostic and therapeutic approach?**
SILAN F., PAKSOY B., KARAKAYA T., YILDIZ O., URFALI M., ÖZDEMİR Ö.
World BioDiscovery Congress 2017, Sofia, Bulgaria, Sofya, Bulgaria, 17 - 19 July 2017, vol.20, pp.20113
- XXXIII. **Clinical and molecular characterization of SLC7A gene that located in 14q11.2 locus in a seconder infertile rare case with lysinuric protein intolerance**
SILAN F., Paksoy B., Urfali M., Karakaya T., ÖZDEMİR Ö.
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European Biotechnology Congress, Dubrovnik, Croatia, 25 - 27 May 2017, vol.256
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- XXXVIII. **A unique keratosis pattern in a case of epidermolytic hyperkeratosis: Report of a case in 46,XX,9qh karyotype"**
ÖĞRETMEN Z., SILAN F., ÖZDEMİR Ö.
2nd International Dermatology and Cosmetology Congress (INDERCOS 2017), İstanbul, Turkey, 15 - 18 March 2017
- XXXIX. **Association between severity of coronary artery disease with genetic polymorphisms MDR 1 and eNOS**
öztürk u., GAZİ E., ÖZDEMİR Ö., AKŞİT E., BEKLER A., TEMİZ A., BARUTÇU A.
32. Uluslararası Katılımlı Kardiyoloji Kongresi, 20 - 23 October 2016
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GAZİ E., öztürk u., BEKLER A., BARUTÇU A., TEMİZ A., AKŞİT E., ÖZDEMİR Ö.
32. Uluslararası Katılımlı Türk Kardiyoloji Kongresi, 20 - 23 October 2016
- XLI. **Is MDR polymorphism related to acute coronary syndrome live under antiaggregant treatment**
GAZİ E., BEKLER A., BARUTÇU A., ÖZDEMİR Ö.
Uluslararası katılımlı TürkKardiyoloji Kongresi, 20 - 23 October 2016
- XLII. **Mental ve motor geriliği olan dismorfik olguda heterozigot 1p36 delesyonu Sitogenetik moleküler sitogenetik ArrayCGH FISH ve MLPA tekniklerinin heterozigot delesyon saptama etkinliklerinin karşılaştırılması**
ÖZDEMİR Ö., URFALI M., YILDIZ O., SILAN F.
XII. Ulusal Tıbbi Genetik Kongresi. Official Journal of Turkish Society of medical Genetics, İzmir, Turkey, 5 - 09 October 2016
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YILDIRIM M. E. , KÜÇÜK KURTULGAN H., Kılıçgün H., ÖZDEMİR Ö., BETON O., TEKİN Y. K.
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SILAN F., ÖĞRETMEN Z., ÖZDEMİR Ö.
25TH EADV, 28 September - 02 October 2016
- XLVI. **The thrombophilic gene polymorphisms and recurrent pregnancy loss dilemma: From Minsk/Belarus and Canakkale - Sivas/Turkish populations**
SILAN F., Mosse I., Gonchar A., Sedlyar N., Kilchevsky A. V. , Kuru B., ÖZDEMİR Ö., Ozdemir O.
European Biotechnology Conference, Riga, Latvia, 5 - 07 May 2016, vol.231
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Urfali M., SILAN F., Tan Y. Z. , Celiker F., Guler Z., ÖZDEMİR Ö.
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European Biotechnology Conference, Latvia, 5 - 07 May 2016, vol.231

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BAĞCI B., bağci g., huzmeli c., SEZGİN İ., ÖZDEMİR Ö.
European Human Genetics Conference 2016, Barcelona, Spain, 21 - 24 May 2016
- LII. **The thrombophilic gene polymorphisms and recurrent pregnancy loss dilemma From Minsk Belarus and Canakkale Sivas Turkish populations 05 07 May 2016 Riga LATVIA Journal of Biotechnology 231 S20**
SILAN F., Mosse I., Gonchar A., Sedlyar N., Kilchevsky A., YILDIZ O., Kuru B., ÖZDEMİR Ö.
European Biotchnology Congress, Riga, Latvia, 5 - 07 May 2016
- LIII. **the thrombophilic gene polymorphisms and Recurrent Pregnancy Loss dilemma from Misk Belarus and Canakkale Sivas Turkish populations**
SILAN F., Mosse I., Kilchevsky A., Gonchar A., Sedlyar N., Kuru B., ÖZDEMİR Ö.
European Biotechnology Conference, Riga, Latvia, 5 - 07 May 2016
- LIIII. **Assessment of BMP 6 polymorphism and relationship with disease activity in Ankylosing Spondylitis patients 05 07 May 2016 Riga LATVIA Journal of Biotechnology 231 S23**
Oztopuz O., SILAN F., AKBAL A., Coskun O., ÖZDEMİR Ö.
European Biotchnology Congress, Riga, Latvia, 5 - 07 May 2016
- LIV. **A Case report of an infertile man with Isodicentric Y Chromosome mosaicism with duplicated SRY SHOX and deleted AZF regions**
SILAN F., URFALI M., ÖZDEMİR Ö.
Eurobiotech2016, 5 - 07 May 2016
- LIV. **Assesment of BMP-6 polymorphism and relationship with disease acitvity in Ankylosing Spondylitis patients**
ÖZTOPUZ R. Ö. , SILAN F., AKBAL A., COŞKUN Ö., ÖZDEMİR Ö.
European Biotechnology Congress, 5 - 07 May 2016
- LVI. **Mental Retardasyon Kısa Boy ve Dismorfik Yüz Bulguları ile Karakterize DiGeorge Sendromlu Olgu FISH Yöntemi ile Doğru Tanı Çanakkale Deneyimi 29 30 Nisan 2016 Çanakkale**
Kuru B., SILAN F., ULUDAĞ A., URFALI M., YILDIZ O., ÖZDEMİR Ö.
1. Trakya Üniversiteler Birliği Yüksek Lisans Öğrenci Kongresi, Çanakkale, Turkey, 29 - 30 April 2016
- LVII. **UGT1A1 GENE MUTATIONS MAY CAUSE MYCOPHENOLATE MOFETIL INDUCED LEUCOPENIA AFTER RENAL TRANSPLANTATION A CASE REPORT**
BAKIRDÖĞEN S., SILAN F., ÖZKUL F., ARIK M. K. , SILAN C., ALTINIŞIK H. B. , ÖZDEMİR Ö.
Gevher Nesibe Tıp Günleri 2016 ve Tıbbi Genetik ve Klinik Uygulamaları Kongresi, Kayseri, Turkey, 11 - 13 February 2016
- LVIII. **The microdeletion/microduplication profiles in spontaneously aborted fetal materials: Double blind results of QF-PCR and MLPA techniques**
SILAN F., Ari E., ULUDAĞ A., Yildiz O., Isin B., Paksoy B., Urfali M., ÖZDEMİR Ö.
European Biotechnology Congress, Bucharest, Romania, 7 - 09 May 2015, vol.208
- LIX. **The RFLP profiles at BRAF V600E mutations in thyroid FNAB nodules**
ÖZDEMİR S., Asik M., SILAN F., ÖZDEMİR Ö., Tan Y. Z. , ARI E., EROGLU M., Ukinc K.
European Biotechnology Congress, Bucharest, Romania, 7 - 09 May 2015, vol.208
- LX. **Microtia, micrognati, facial dysmorphism, short stature and mental retardation: A rare case with Meirer-Gorlin syndrome**
Paksoy B., SILAN F., Yildiz O., ÖZDEMİR Ö., Tas Z. T.
European Biotechnology Congress, Bucharest, Romania, 7 - 09 May 2015, vol.208
- LXI. **KRAS BRAF oncogene mutations and tissue spesific promoter hypermethylation oftumor supressor HIC 1 P16 DAPK1 SFRP2 and MGMT genes in colorectal cancer patients**
BAĞCI B., SARI M., KARADAYI K., ÖZDEMİR Ö.
EUROPEAN HUMAN GENETICS CONFERENCE, GLASKOW, England, 6 - 09 July 2015
- LXII. **The relationship between germ line MTHFR C677T and A1298C polymorphisms and psoriasis**

- KILIÇ S., ÖZDEMİR Ö., SILAN F., DAMLA K., SILAN C., ÖĞRETMEN Z.
International Biotechnology Congress, BÜKREŞ, Romania, 7 - 09 May 2015
- LXIII. **Familial X chromosome translocation Xqtriplcation and SHOX gene deletion with short stature**
SILAN F., PAKSOY B., YILDIZ O., ÖZDEMİR Ö.
International Biotechnology Congress, BÜKREŞ, Romania, 7 - 09 May 2015
- LXIV. **A case of 47 XYY syndrome without behavioral and emotional difficulties Cost effective well define of extra Y chromosome by GTG Cbandings and FISH analysis**
ÖZDEMİR Ö., SILAN F., PAKSOY B., CAMER G.
International Biotechnology Congress, BÜKREŞ, Romania, 7 - 09 May 2015
- LXV. **Short stature and other clinical findings of a case with karyotype 46 XYdel Y q11 2 12 mar**
ÖZDEMİR Ö., ÇOLAK A., EĞİLMEZ H.
. Abstracts of the 1th European Cytogenetics Conference, Turkey, 22 June 1997 - 25 March 2015
- LXVI. **TNF Alpha G308A polymorphism is a risk factor for diabet among psoriatic patients P 1650**
HIZ M. M. , ÖĞRETMEN Z., SILAN F., ÖZDEMİR Ö.
23rd EADV Congress Building Bridges, Amsterdam, Netherlands, 8 - 12 October 2014
- LXVII. **Association of eNOS Glu298Asp gene polymorphism with family history of psoriasis P 1109**
ÖĞRETMEN Z., HIZ M. M. , SILAN F., ÖZDEMİR Ö.
23rd EADV Congress Building Bridges, Amsterdam, Netherlands, 8 - 12 October 2014
- LXVIII. **The rates of eNOS Glu298Asp gene polymorphism among psoriatic patients in Çanakkale Turkey J Exp Clin Med 31 2 128 2014**
HIZ M. M. , ÖĞRETMEN Z., SILAN F., ÖZDEMİR Ö.
3 rd Scientific Writing and Stereology Workshop, Turkey, 8 - 12 April 2014, vol.31, pp.128
- LXIX. **ÇANAKKALE DE AİLEVİ AKDENİZ ATEŞİ OLAN ÇOCUKLARDA Mefv GEN MUTASYONLARI VE FENOTİP GENOTİP İLİŞKİSİ**
BATTAL F., SILAN F., TOPALOĞLU N., YILDIRIM Ş., AYLANÇ H., BİNNETOĞLU F. K. , TEKİN M., KAYMAZ N., ÖZDEMİR Ö.
10. ULUDAĞ PEDIATRİ KİŞ KONGRESİ, Turkey, 16 - 19 March 2014, pp.35-36
- LXX. **The relationship between psoriasis and TNF alpha G308A and G238T polymorphisms P 180**
ÖĞRETMEN Z., SILAN F., HIZ M. M. , ATİK YALÇINTEPE S., KIRILMAZ B., ÖZDEMİR Ö.
4th Congress of the Psoriasis International Network, 4 - 06 July 2013, vol.27, pp.1-77
- LXXI. **The relationship between psoriasis and tnf alpha g308a and g238a polymorphisms**
ÖĞRETMEN Z., SILAN F., merve meliha h., Sinem Atik Y., KIRILMAZ B., ÖZDEMİR Ö.
JOURNAL OF THE EUROPEAN ACADEMY OF DERMATOLOGY AND VENEREOLOGY, 1 - 03 July 2013
- LXXII. **P06 205 Evaluation of CYP2C9 and CYP2D6 gene polymorphisms in thyroid cancer**
ULUDAĞ A., ÖZDEMİR S., SILAN C., ATİK S., SILAN F., ÖZDEMİR Ö.
European Human Genetics Conference. Nürnberg-GERMANY 23-26 June 2012, 23 - 26 June 2012, vol.20, pp.198
- LXXIII. **P06 206 The possible role of the xenobiotic transporter P glycoprotein polymorphism that encoded by the MDR1 3435 C T gene in the susceptibility of differentiated thyroid cancer**
ÖZDEMİR S., ULUDAĞ A., SILAN F., SİNEM A., TURGUT B., ÖZDEMİR Ö.
European Human Genetics Conference.Nürnberg-GERMANY 23-26 June 2012., NÜRNBERG, Germany, 23 - 26 June 2012, vol.20, pp.199
- LXXIV. **6 P76 Increased T allele frequency in MTHFR C677T gene in thyroid carcinoma**
ÖZDEMİR S., SILAN F., ERSELCAN T., ULUDAĞ A., ÜSTÜN F., ÇOLAK A., ATİK S., ÖZDEMİR Ö.
8th European Cytogenetic Conference, 02-05 July 2011, Porto, Portugal, 2 - 05 July 2011, vol.19, pp.175
- LXXV. **Adelosan İdiyopatik Skolyozda MATN 1 ve LCT C13910T Gen Polimorfizmi**
YILMAZ H., ZATERİ C., ULUDAĞ A., BAKAR C., KOŞAR Ş., ÖZDEMİR Ö.
23.Ulusal Fiziksel Tıp ve Rehabilitasyon Kongresi. Türk Fiz Rehab Derg 2011;57 Özel Sayı;1-334,p-146, Turkey, 11 - 15 May 2011
- LXXVI. **Anjiotensin konverting enzim gen polimorfizmi I D ve esansiyel hipertansiyon arasındaki ilişki Populasyon çalışması**
ATİK S., DEMİREL Y., SILAN F., DOĞAN S., AŞGÜN H. F. , ÖZDEMİR Ö.

I. Uluslararası Katılımlı Kök Hücre Sempozyumu, Samsun, Turkey, 29 September - 01 October 2010

- LXXVII. **Akciğer kanseri olgularında K ras kodon 12 13 mutasyon sıklığı ve tümör dokularına özgü TS promotör bölge metilasyon profilleri**
ARSLAN S., KÖKSAL B., ÖZDEMİR Ö., DOĞAN Ö. T. , YILDIZ F., AKKURT İ.
11. Ulusal Tıbbi Biyoloji ve Genetik Kongresi, Muğla, Turkey, 28 - 30 October 2009
- LXXVIII. **Tumoral tissue specific promoter hypermethylation of tumor supressor genes SFRp2 p16 DAPK1 HIC and MGMT in a case with non small cell carcinoma A case report**
ARSLAN S., DOĞAN T., KOKSAL B., YILDIRIM M., GUMUS C., ELAGOZ C., AKKURT I., ÖZDEMİR Ö.
7th European Cytogenetics Conference, 4 - 07 July 2009
- LXXIX. **P 4 14 In Vivo Evaluation Of Micronuclei In Rat Reticulocytes For The Relative Efficacy Of Genomic Instability In Medical Radiation Exposure**
EĞİLMEZ H., GÜL E., ERSELCAN T., GÜMÜŞ C., ÖZDEMİR S., ÖZDEMİR Ö.
5.th European Cytogenetics Conference, 4-7 June 2005, Madrid, Spain, 4 - 07 June 2005, vol.13, pp.109-110
- LXXX. **PS 68 Acut effect of I 131 treatment on chromosomes**
ERSELCAN T., SÜNGÜ S., ÖZDEMİR S., ORBAY D., ÖZDEMİR Ö.
Annual Congress of the European Association of Nuclear Medicine, 25-29 Aug 2001, NAPOLİ, Italy, 25 - 29 August 2001, vol.28, pp.1092
- LXXXI. **Sister chromatide exchange frequency in women that exposed to biomass in a village of central anatolia**
SÜNGÜ S., ÇINAR Z., AKKURT İ., ÖZDEMİR Ö., SEYFİKLİ Z.
Third European Conference, 7 - 10 July 2001
- LXXXII. **In vivo DNA methylation of E coli DH 5 and Top10F strains by cytosine 5 methyltransferase M Msp1**
ÖZDEMİR Ö., HORNBY P.
XIII. Ulusal Biyokimya Kongresi, Turkey, 26 - 30 March 1996
- LXXXIII. **5 Azodeoksisitidinin Baz analogu insan kromozomları üzerindeki etkisinin incelenmesi**
ÖZDEMİR Ö., SEZGİN İ., ÇOLAK A.
II. Ulusal Tıbbi Biyoloji Kongresi, Turkey, 22 - 24 September 1990
- LXXXIV. **Popülasyonda sentromer polimorfizmi**
ÖZDEMİR Ö., SEZGİN İ., ÇOLAK A.
Atatürk Üniversitesi X. Ulusal Biyoloji kongresi, Turkey, 18 - 20 July 1990
- LXXXV. **Kangal köpeklerinde Canis familiaris kromozomal arařtırmalar**
ÖZDEMİR Ö., SEZGİN İ., COLAK A.
IX. Ulusal Biyoloji Kongresi, Turkey, 21 - 23 September 1988

Supported Projects

SILAN C., SILAN F., ÖZDEMİR Ö., URFALI M., Project Supported by Higher Education Institutions, Ailesel Akdeniz Ateři olan hastalarda kolşisin tedavi yanıtlarının Sitokrom P450 CYP2D6, CYP2A3 ve CYP2A4 polimorfizmleri ile arasındaki ilişkinin arařtırılması, 2015 - 2017

GAZİ E., ÖZTÜRK U., ÖZDEMİR Ö., Project Supported by Higher Education Institutions, Akut koroner sendrom tanısı ile başvuran , asetilsalisilik asit veya klopidogrel kullanan hastalarda MDR1 ve eNOS T786C gen polimorfiziminin arařtırılması, 2014 - 2015

SILAN F., ÖZDEMİR Ö., ÇAKIR GÜNGÖR A. N. , AKURUT Ç., Project Supported by Higher Education Institutions, Maternal Kandan Fetal DNA İzolasyonu ve Fetal RhD Analizi, 2014 - 2014

GAZİ E., SILAN F., ÖZDEMİR Ö., Project Supported by Higher Education Institutions, Koroner yavaş akımı olan hastalarda ateroskleroz ile ilgili gen polimorfizmi varlığı ve carotis intima-media kalınlığı, endotel fonksiyonlarının ile ilişkisi, 2013 - 2014

ÖĞRETMEN Z., SILAN F., KIRILMAZ B., ÖZDEMİR Ö., Project Supported by Higher Education Institutions, Psoriasisli hastalarda Kardiyolojik risk faktörleri ile ACE, eNOS, FVL, GJB2 polimorfizmlerinin ilişkilerinin arařtırılması, 2011 - 2012

GÜÇLÜ O., DEREKÖY F. S. , SILAN F., ÖZDEMİR Ö., Project Supported by Higher Education Institutions, Maksiller Sinüsün

Citations

Total Citations (WOS):361

h-index (WOS):10

Scholarships

Radiodiagnogical, histopathological and FISH analysis of different tumoral tissues (epithelial and bone marrow) from long term N'-ethyl-N'-nitrosourea (ENU) treated rats, NATO, 2000 - Continues

The molecular mechanism of action of azacytidine (Desitabine), NATO, 1994 - Continues

Awards

ÖZDEMİR Ö., The RFLP profiles at BRAF V600E mutations in thyroid FNAB nodules. International Biotechnology Congress. 7-9 May 2015, Bucharest/ROMANIA. Bu araştırma Poster 2. 'İğİ ile ödüllendirilmiştir., En iyi poster 2'lük ödülü. Uluslararası Biyoteknoloji Kongresi, 7-9 Mayıs 2015 Butabeşte/ROMANIA/2015, May 2015

SILAN F., ÖZDEMİR Ö., The microdeletion/microduplication profiles in spontaneously aborted fetal materials: Double blind results of QF-PCR and MLPatechniques, European Biotechnology Thematic Asociation, May 2015

SILAN F., ÖZDEMİR Ö., Variable R.Msp1 fragmentation in genomic DNA due to DNA hypomethylation in CRF patients with MTHFR C677Tgene polymorphism: from genetics to epigenetics., European Human Genetics Conference, May 2014

ÖZDEMİR Ö., SILAN F., Sitogenetik Sonuçları Olan Bir Tek Gen Defekti: Prematur Chromatide Separation, Erişkin Yaşta Görülen Genetik Hastalılar Sempozyumu, December 2013

ÖZDEMİR Ö., Analysis of Ki-ras Exon 2 Gene Mutations in 3-Methylcholanthreneand Butylated Hydroxytoluene-Induced Rat Lung Tissues, Poster Teşfik (Birincilik) ödülü, Çukurova Üniversitesi ADANA / 2003, April 2003

ÖZDEMİR Ö., Increased cell proliferation and RMsp1 fragmentation in testicular tissue of rat induced by 5-aza-2'-deoxycytidine., Prof Dr. Altan GÜNALP 1. lik Ödülü Osman Gazi Üniversitesi / 2001, January 1997