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Education Information

Doctorate, Hacettepe University, Sağlık Bilimleri Enstitüsü, Tıbbi Biyoloji, Turkey 1992 - 1997
Postgraduate, Dokuz Eylul University, Sağlık Bilimleri Enstitüsü, Tıbbi Biyoloji, Turkey 1987 - 1989

Foreign Languages

English, B2 Upper Intermediate

Dissertations

Doctorate, Spinal Müsküler Atrofi Aday Genlerinde Delesyon ve CA Tekrar Analizleri, Hacettepe University, Sosyal Bilimler Enstitüsü, 1997
Postgraduate, Ampisilin, Amoksisilin ve Beta-Laktamaz İnhibitörü İçeren Değişik konsantrasyonlarının İdrar İzolatı Escherichia coli'lere Etkisi, Dokuz Eylul University, Sağlık Bilimleri Enstitüsü, 1989

Research Areas

Medicine, Health Sciences, Fundamental Medical Sciences, Medical Biology, Internal Medicine Sciences, Medical Genetics, Life Sciences, Population Biology, Population Genetics, Natural Sciences

Academic Titles / Tasks

Professor, Istanbul University, Istanbul Medical Faculty, Basic Medical Sciences, 2013 - Continues
Associate Professor, Gaziantep University, School Of Medıcıne, Department Of Basıc Medıcal Scıences, 2006 - 2013
Associate Professor, Ege University, Faculty Of Medicine, Dahili Tıp Bilimleri Bölümü, 2005 - 2006
Assistant Professor, Ege University, Sağlık Bilimleri Enstitüsü, 1998 - 2005

Research Assistant PhD, Ege University, Faculty Of Science, Biyoloji Bölümü, 1997 - 1998
Research Assistant, Hacettepe University, Tıp Fakültesi (Türkçe), Temel Tıp Bilimleri Bölümü, 1992 - 1997
Expert, Cukurova University, Tıp Fakültesi, Dahili Tıp Bilimleri, 1989 - 1992
Research Assistant, Dokuz Eylul University, Tıp Fakültesi, Temel Tıp Bilimleri Bölümü, 1987 - 1989

Academic and Administrative Experience

Deputy Chief Physician, Istanbul University, Istanbul Medical Faculty, Basic Medical Sciences, 2019 - Continues
Fakülte Yönetim Kurulu Üyesi, Istanbul University, Istanbul Medical Faculty, Basic Medical Sciences, 2019 - Continues

Advising Theses

Pehlivan S., Madde Kullanım Bozukluğu Olan Hastalarda Epigenetik Farklılıkların Araştırılması ve Klinik Parametrelerle Karşılaştırılması, Postgraduate, Y.Oyacı(Student), 2020
Pehlivan S., Madde kullanım bozukluğunda rolü olabilecek genlerin DNA düzeyinde araştırılması ve bulguların klinik parametrelerle karşılaştırılması, Postgraduate, S.Kurnaz(Student), 2017
Pehlivan S., Sigara bağımlılığında rolü olan dopaminerjik ve serotenerjik genlerin DNA,RNA ile epigenetik düzeyde araştırılması, Postgraduate, Ü.Sever(Student), 2017
Pehlivan S., Kronik Myelositer Lösemide kromozom (10-22 ve cinsiyet) Düzensizliklerinin MLPA Yöntemi ile Araştırılması, Postgraduate, T.Sever(Student), 2009
Pehlivan S., Kronik Myelositer Lösemide kromozom (1-9 nolu kromozomlar) Düzensizliklerinin MLPA Yöntemi ile Araştırılması., Postgraduate, C.Kılınçarslan(Student), 2009
Pehlivan S., İnfertil Erkek Hastalarda AZF Bölgesini Kapsayan Y Kromozom Mikrodelesyon Taraması , Postgraduate, Ö.Okutman(Student), 2001
Pehlivan S., Böbreğin Renal Hücreli Kanserinde Dinükleotit Tekrar ve AP-PCR Metodları Uygulamaları , Postgraduate, A.Kunt(Student), 2001

Published journal articles indexed by SCI, SSCI, and AHCI

I. **The effect of hereditary thrombotic factors and comorbidities on the severity of COVID-19 disease**
Kose M., Senkal N., Konyaoglu H., Emet A., Oyaci Y., PEHLİVAN S., Sayin G.
European Review for Medical and Pharmacological Sciences, vol.27, no.1, pp.395-403, 2023 (SCI-Expanded)

II. **The effect of DNA repair gene variants on COVID-19 disease: susceptibility, severity, and clinical course**
Senkal N., Serin I., PEHLİVAN S., Pehlivan M., Medetalibeyoglu A., Cebeci T., Konyaoglu H., Oyacı Y., Sayın G. Y., Isoglu-Alkac U., et al.
Nucleosides, Nucleotides and Nucleic Acids, 2023 (SCI-Expanded)

III. **Association of Tumor Necrosis Factor-Alpha (TNF-α) and Suppressor of Cytokine Signaling-1 (SOCS-1) Gene Variants in Children with COVID-19**
Uysalol M., Serin I., Oyacl Y., Yldlz R., Uysalol E., PEHLİVAN S.
Journal of Pediatric Infectious Diseases, vol.18, no.1, pp.38-45, 2023 (SCI-Expanded)

IV. **PERIOD3 (PER3) VNTR Variant Associated with Seasonal Pattern and Family History in Bipolar Disorder**
Aytac H. M., Pehlivan M., Oyaci Y., PEHLİVAN S.
Psychiatria Danubina, vol.34, no.4, pp.695-699, 2022 (SCI-Expanded)

V. **Association of intron 4 VNTR polymorphism in the NOS3 gene with rapid cycling and treatment resistance in bipolar disorder: a case-control study**
Aytac H. M., PEHLİVAN M., Oyaci Y., Pehlivan S.

NEUROSCIENCES, vol.27, no.4, pp.229-236, 2022 (SCI-Expanded)

VI. **Role of cytokines in multiple myeloma: IL-1RN and IL-4 VNTR polymorphisms**
Serin I., Oyaci Y., Pehlivan M., PEHLİVAN S.
Cytokine, vol.153, 2022 (SCI-Expanded)

VII. **Association of mannose-binding lectin 2 (MBL2) and suppressor of cytokine signaling-1 (SOCS1) gene variants in children with febrile neutropenia.**
Uysalol E. P., Uysalol M., Pehlivan M., Oyaci Y., Pehlivan S., Serin I.
Journal of infection and chemotherapy : official journal of the Japan Society of Chemotherapy, vol.28, no.5, pp.657-662, 2022 (SCI-Expanded)

VIII. **Investigation of <u>MBL2</u> and <u>NOS3</u> functional gene variants in suspected COVID-19 PCR (-) patients.**
Pehlivan S., Köse M., Mese S., Serin I., Senkal N., Oyacı Y., Medetalibeyoglu A., Pehlivan M., Sayın G., Isoglu-Alkac Ü., et al.
Pathogens and global health, vol.116, no.3, pp.178-184, 2022 (SCI-Expanded)

IX. **Quantitative detection of methylated SOCS-1 in schizophrenia and bipolar disorder considering SOCS-1-1478CA/del polymorphism and clinical parameters**
Aytac H. M., Pehlivan S., Pehlivan M., Oyaci Y.
IRISH JOURNAL OF MEDICAL SCIENCE, 2022 (SCI-Expanded)

X. **Cytokine gene polymorphism frequencies in Turkish population living in Marmara region**
Özdilli K., Ögret Y., Oğuz S. R., Sel F. A., Şentürk Çiftçi H., Çınar Ç., Pehlivan S., Oğuz F.
TURKISH JOURNAL OF BIOCHEMISTRY, vol.47, 2022 (SCI-Expanded)

XI. **DNA methylation pattern of gene promoters of MB-COMT, DRD2, and NR3C1 in Turkish patients diagnosed with schizophrenia**
Aytac H. M., Oyaci Y., Pehlivan M., PEHLİVAN S.
EUROPEAN JOURNAL OF HUMAN GENETICS, vol.30, no.SUPPL 1, pp.285, 2022 (SCI-Expanded)

XII. **Importance of mannose-binding lectin2 polymorphism (rs1800450) in infections in children**
Uysalol M., Gumus S., Yildiz R., Pasli Uysalol E., Pehlivan S., Pehlivan M., Serin I.
BIOMARKERS, vol.27, no.1, pp.44-49, 2022 (SCI-Expanded)

XIII. **COMTVal158Met polymorphism is associated with ecstasy (MDMA)-induced psychotic symptoms in the Turkish population**
Aytac H. M., Oyaci Y., Aydin P. C., Pehlivan M., Pehlivan S.
NEUROSCIENCES, vol.27, no.1, pp.24-30, 2022 (SCI-Expanded)

XIV. **The suppressor of cytokine signaling-1 (SOCS1) gene polymorphism and promoter methylation correlate with the course of COVID-19**
Tukek T., PEHLİVAN S., Medetalibeyoglu A., Serin I., Oyacı Y., Arıcı H., Senkal N., Pehlivan M., Isoglu-Alkac U., Kose M.
Pathogens and Global Health, 2022 (SCI-Expanded)

XV. **Tumor Necrosis Factor-alpha (TNF-alpha)-238 G/A Polymorphism Is Associated with the Treatment Resistance and Attempted Suicide in Schizophrenia**
Aytac H. M., Ozdilli K., Tuncel F. C., PEHLİVAN M., Pehlivan S.
IMMUNOLOGICAL INVESTIGATIONS, vol.51, no.2, pp.368-380, 2022 (SCI-Expanded)

XVI. **What are the roles of global DNA and APC 2 gene promotor hypermethylation in multiple myeloma?**
Pehlivan S., Serin I., NURSAL A. F., Oyaci Y., GÜNDEŞ İ., Pehlivan M.
MOLECULAR BIOLOGY REPORTS, vol.48, no.12, pp.7875-7882, 2021 (SCI-Expanded)

XVII. **Investigation of inflammation related gene polymorphism of the mannose-binding lectin 2 in schizophrenia and bipolar disorder.**
Aytac H. M., Yazar M. S., Erol A., Pehlivan S.
Neurosciences (Riyadh, Saudi Arabia), vol.26, no.4, pp.346-356, 2021 (SCI-Expanded)

XVIII. **Association of the Uncoupling Protein 2-866 G/A Polymorphism with Family History and Duration of Tobacco Use Disorder in a Turkish Population**
Aytac H. M., Pehlivan S., Kurnaz S., PEHLİVAN M., Aydin P. C.
PSYCHIATRY AND CLINICAL PSYCHOPHARMACOLOGY, vol.31, no.3, pp.280-285, 2021 (SCI-Expanded)

XIX. **Medication-related osteonecrosis of the jaw (MRONJ) and eNOS Polymorphisms in multiple myeloma patients: a single center experience.**
Taş Ozyurtseven B., Serin I., Nursal A. F., Pehlivan S., Pehlivan M.
BMC oral health, vol.21, pp.272, 2021 (SCI-Expanded)

XX. **Mannose binding lectin gene 2 (rs1800450) missense variant may contribute to development and severity of COVID-19 infection.**
Medetalibeyoglu A., Bahat G., Senkal N., Kose M., Avcı K., Sayın G., Isoglu-Alkac Ü., Tukek T., Pehlivan S.
Infection, genetics and evolution : journal of molecular epidemiology and evolutionary genetics in infectious diseases, vol.89, pp.104717, 2021 (SCI-Expanded)

XXI. **DNA Repair Genes and Chronic Myeloid Leukemia: ERCC2 (751), XRCC1 (399), XRCC4-Intron 3, XRCC4 (-1394) Gene Polymorphisms**
Ozdilli K., Pehlivan M., Serin I., Savran F., Tomatir A. G., PEHLİVAN S.
MEDITERRANEAN JOURNAL OF HEMATOLOGY AND INFECTIOUS DISEASES, vol.13, 2021 (SCI-Expanded)

XXII. **A new parameter in multiple myeloma: CYP3A4*1B single nucleotide polymorphism**
Serin I., Pehlivan S., GÜNDEŞ İ., Fidan Oyaci Y., PEHLİVAN M.
ANNALS OF HEMATOLOGY, vol.100, no.2, pp.421-427, 2021 (SCI-Expanded)

XXIII. **Global and glucocorticoid receptor gene-specific (NR3C1) DNA methylation analysis in patients with cannabinoid or synthetic cannabinoid use disorder.**
Pehlivan S., Aytac H. M., Cetinay Aydin P., Nursal A. F., Pehlivan M.
Psychiatry research, vol.298, pp.113774, 2021 (SCI-Expanded)

XXIV. **Detection of altered methylation of MB-COMT promotor and DRD2 gene in cannabinoid or synthetic cannabinoid use disorder regarding gene variants and clinical parameters.**
Oyaci Y., Aytac H. M., Pasin O., Cetinay Aydin P., Pehlivan S.
Journal of addictive diseases, vol.39, no.4, pp.526-536, 2021 (SSCI)

XXV. **A relationship between endothelial nitric oxide synthetase gene variants and substance use disorder**
Pehlivan S., Aydin P. C., Nursal A. F., Pehlivan M., Oyaci Y., Yazici A. B.
Endocrine, Metabolic and Immune Disorders - Drug Targets, vol.21, no.9, pp.1679-1684, 2021 (SCI-Expanded)

XXVI. **UCP2 and CFH Gene Variants with Genetic Susceptibility to Schizophre-nia in Turkish Population**
NURSAL A. F., Aydin P. C., PEHLİVAN M., Sever U., Pehlivan S.
ENDOCRINE METABOLIC & IMMUNE DISORDERS-DRUG TARGETS, vol.21, no.11, pp.2084-2089, 2021 (SCI-Expanded)

XXVII. **Macrophage Migration Inhibitory Factor-173 G/C Polymorphism is Associated With The Age of Onset and Insight in Schizophrenia in the Turkish Population**
Aytac H. M., Oyaci Y., Yazar M. S., Pehlivan S.
NEUROLOGICAL RESEARCH, vol.43, no.12, pp.977-984, 2021 (SCI-Expanded)

XXVIII. **Role of MIF -173G/C and Mbl2 codon 54A/B variants in risk of multiple myeloma: An association study.**
Pehlivan M., Nursal A., Gündeş İ., Oyacı Y., Kıvanç D., Pehlivan S.
Endocrine, metabolic & immune disorders drug targets, vol.21, pp.925-931, 2021 (SCI-Expanded)

XXIX. **The role of endothelial nitric oxide synthase gene polymorphisms in patients with lung cancer**
Kocer C., Benlier N., Balci S. O., PEHLİVAN S., Sanli M., Nacak M.
CLINICAL RESPIRATORY JOURNAL, vol.14, no.10, pp.948-955, 2020 (SCI-Expanded)

XXX. **Association of MIF and MBL2 gene polymorphisms with attempted suicide in patients diagnosed with schizophrenia or bipolar disorder**
Aytac H. M., Oyaci Y., Yazar M. S., EROL A., PEHLİVAN S.
JOURNAL OF CLINICAL NEUROSCIENCE, vol.78, pp.264-268, 2020 (SCI-Expanded)

XXXI. **Evaluation of COMT (rs4680), CNR2 (rs2501432), CNR2 (rs2229579), UCP2 (rs659366), and IL-17 (rs763780) gene variants in synthetic cannabinoid use disorder patients**
PEHLİVAN S., Aytac H. M., Kurnaz S., Pehlivan M., Aydin P. C.
JOURNAL OF ADDICTIVE DISEASES, vol.38, no.4, pp.495-505, 2020 (SSCI)

XXXII. **PER3 VNTR variant and susceptibility to smoking status/substance use disorder in a Turkish**

population

Nursal A. F., Aydin P. C., Uysal M. A., Pehlivan M., Oyac Y., PEHLİVAN S.

ARCHIVES OF CLINICAL PSYCHIATRY, vol.47, no.3, pp.71-74, 2020 (SCI-Expanded)

XXXIII. **Investigation of catechol-O-methyltransferase and cannabinoid receptor 2 gene variants in tobacco use disorder or tobacco use disorder and schizophrenia comorbidity Tütün kullanım bozukluğu veya tütün kullanım bozukluğu ve şizofreni komorbiditesinde katekol-o-metiltransferaz ve kannabinoid reseptör 2 gen varyantlarının incelenmesi**

Pehlivan S., Çetinay Aydin P., Aytaç H. M., Uysal M. A., Sever Ü., Pehlivan M.

Anadolu Psikiyatri Dergisi, vol.21, no.6, pp.572-578, 2020 (SCI-Expanded)

XXXIV. **TNF-α-308 G/A variant may be associated with bipolar disorder in a Turkish population**

Nursal A. F., Aytac H. M., Ciftci H. S., Yazar M. S., Oyaci Y., Pehlivan M., Pehlivan S.

Revista de Psiquiatria Clinica, vol.47, no.6, pp.176-179, 2020 (SCI-Expanded)

XXXV. **Dopamine D4 Receptor Gene Exon III VNTR Variant Influences Smoking Status in Turkish Population**

Uysal M. A., Sever U., NURSAL A. F., Pehlivan S., Bahadir A., Yurt S., Chousein E. G. U., Onur S. T., Bagci B. A., Hattatoglu D. G., et al.

NOROPSIKIYATRI ARSIVI-ARCHIVES OF NEUROPSYCHIATRY, vol.56, no.4, pp.248-252, 2019 (SCI-Expanded)

XXXVI. **CNR2 rs2229579 and COMT Val158Met variants, but not CNR2 rs2501432, IL-17 rs763780 and UCP2 rs659366, contribute to susceptibility to substance use disorder in the Turkish population**

Kurnaz S., Yazici A. B., NURSAL A. F., Aydin P. C., Atar A. O., Aydin N., Kincir Z., Pehlivan S.

PSYCHIATRY AND CLINICAL PSYCHOPHARMACOLOGY, vol.29, no.4, pp.847-853, 2019 (SCI-Expanded)

XXXVII. **Genetic factors associated with the predisposition to late onset Alzheimer's disease**

DURMAZ A., Kumral E., Durmaz B., ONAY H., Aslan G. I., Ozkinay F., Pehlivan S., Orman M., Cogulu O.

GENE, vol.707, pp.212-215, 2019 (SCI-Expanded)

XXXVIII. **CYP2A6 gene variants may explain smoking status in a Turkish cohort**

Pehlivan S., Uysal M. A., Cagatay T., NURSAL A. F., Cinar Ç., Erkan F., Sever U., Bingol Z., PEHLİVAN M., Pence S.

PSYCHIATRY AND CLINICAL PSYCHOPHARMACOLOGY, vol.29, no.3, pp.340-345, 2019 (SCI-Expanded)

XXXIX. **Paraoxonase-1 Polymorphisms (L55M/Q192R) and Activities (PONase/AREase) in Patients with Idiopathic Recurrent Early Pregnancy Loss: A Preliminary Study**

Ozturk E., Pehlivan S., Ozcan C., UĞUR M. G., Balat O.

GENETIC TESTING AND MOLECULAR BIOMARKERS, vol.23, no.7, pp.501-505, 2019 (SCI-Expanded)

XL. **THE ASSOCIATION OF GVHD WITH HLA DR ALLELES, IFN-GAMMA, TGF-BETA, AND MBL2 GENE POLYMORPHISM**

Oguz R., Ciftci H., Gokce M., Ogret Y., Karadeniz S., Pehlivan S., Aydin F.

HLA, vol.93, no.5, pp.368, 2019 (SCI-Expanded)

XLI. **Association of XRCC1 and XPD functional gene variants with nicotine dependence and/or schizophrenia: a case-control study and in silico analysis**

Pehlivan S., Aydin N., Nursal A. F., Uysal M. A., Pehlivan M., Tekcan A., Yavuz F. K., Sever U., Yavuzlar H., Kurnaz S., et al.

PSYCHIATRY AND CLINICAL PSYCHOPHARMACOLOGY, vol.29, no.1, pp.21-27, 2019 (SCI-Expanded)

XLII. **Effect of monoamine oxidase B A644G variant on nicotine dependence and/or schizophrenia risk**

Pehlivan S., Aydin P. C., Uysal M. A., Ciftci H. S., Sever U., Yavuz F. K., Aydin N., Nursal A. F.

ARCHIVES OF CLINICAL PSYCHIATRY, vol.46, no.1, pp.21-24, 2019 (SCI-Expanded)

XLIII. **Endothelial Nitric Oxide Synthase Gene Variants and Susceptibility to Chronic Myeloid Leukemia (Ph plus)**

PEHLİVAN M., Pehlivan S., KAYNAR L., Sever T., YILMAZ M., Eser B., Okan V., ÇETİN M., TOMATIR A. G.

INTERNATIONAL JOURNAL OF HUMAN GENETICS, vol.18, no.3, pp.247-252, 2018 (SCI-Expanded)

XLIV. **XRCC4 rs6869366 polymorphism is associated with susceptibility to both nicotine dependence and/or schizophrenia**

Pehlivan S., Uysal M. A., Aydin N., NURSAL A. F., PEHLİVAN M., Yavuzlar H., Sever U., Kurnaz S., Yavuz F. K., Uysal S., et al.

ARCHIVES OF CLINICAL PSYCHIATRY, vol.45, no.3, pp.53-56, 2018 (SCI-Expanded)

XLV. **Uteroglobin gene polymorphism (G38A) may be a risk factor in childhood idiopathic nephrotic syndrome**
KILIÇ B. D., BÜYÜKÇELİK M., OĞUZKAN BALCI S., Pehlivan S., KUL S., Col N., Balat A.
PEDIATRIC NEPHROLOGY, vol.33, no.2, pp.295-303, 2018 (SCI-Expanded)

XLVI. **XRCC4 rs6869366 polymorphism is associated with suspectibility to nicotine dependence and/or schizophrenia**
PEHLİVAN S., Uysal M. A., Aydın N., NURSAL A., PEHLIVAN M., Yavuzlar H., Sever U., Kurnaz S., Yavuz F. K., Aydın P.
SAO PAULO MEDICAL JOURNAL, vol.45, no.3, pp.53-56, 2018 (SCI-Expanded)

XLVII. **Association of the TNF-alpha, IL-2, and IL-2RB gene variants with susceptibility to psoriasis in a Turkish cohort**
Gulel A., Inaloz H. S., NURSAL A. F., Sever T., Pehlivan S.
CENTRAL EUROPEAN JOURNAL OF IMMUNOLOGY, vol.43, no.1, pp.50-57, 2018 (SCI-Expanded)

XLVIII. **Possible association between DNA repair gene variants and cannabis dependence in a Turkish cohort: a pilot study**
Pehlivan S., Yazici A. B., Aydin N., NURSAL A. F., Kurnaz S., Atar A. O., Sever U., Kincir Z., PEHLİVAN M., Aydin P. C.
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XLIX. **Can VNTR variants in eNOS and XRCC4 genes contribute to formation of nicotine dependence and/or schizophrenia?**
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BRATİSLAVA MEDİCAL JOURNAL-BRATİSLAVSKE LEKARSKE LİSTY, vol.118, no.8, pp.467-471, 2017 (SCI-Expanded)

L. **The functional variants of endothelial nitric oxide synthase gene associated with rheumatoid arthritis in Turkish adults**
Pehlivan S., Aydeniz A., Sever T., Altindag O., Pehlivan M., Gursoy S., Oguzkan-Balci S.
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LI. **MBL2 and MIF gene polymorphisms in cardiovascular patients with atherosclerotic lesions undergoing heart valve replacement**
Eksi F., Pehlivan S., Erdogan M. B., Bayram A., Oguzkan-Balci S., YAMAK B., Pehlivan M.
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LII. **Effect of Cytokine Genes in the Pathogenesis and on the Clinical Parameters for the Treatment of Multiple Myeloma**
Haydaroglu H., Balci S. O., Pehlivan S., Ozdilli K., Gundogan E., Okan V., Nursal A. F., Pehlivan M.
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LIII. **eNOS and XRCC4 VNTR variants contribute to formation of nicotine dependence and/or schizophrenia**
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LIV. **The Endothelial Nitric Oxide Synthase Gene Variants as a Risk Factor for Chronic Lymphocytic Leukemia**
PEHLİVAN M., TOMATIR A. G., Nursal A. F., ŞAHİN H. H., Pehlivan S.
UHOD-ULUSLARARASI HEMATOLOJI-ONKOLOJI DERGISI, vol.27, no.2, pp.85-90, 2017 (SCI-Expanded)

LV. **Relationship between cigarette (nicotine) addiction and dopamine receptor D2 gene variants (TaqlA and-141C Ins/Del)**
Sever U., Uysal M. A., PEHLİVAN S.
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LVI. **The Associations of IL-6, IFN-gamma, TNF-alpha, IL-10, and TGF-beta 1 Functional Variants with Acute Myeloid Leukemia in Turkish Patients**
Nursal A. F., PEHLİVAN M., ŞAHİN H. H., Pehlivan S.
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Uysal M. A., Sever U., PEHLİVAN S.
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LVIII. **Analysis of CYP2A6 and CYP2A13 genes using new generation sequencing method in smokers and non-smokers**
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LIX. **Effects of TNFalpha, NOS3, MDR1 gene polymorphisms on clinical parameters, prognosis and survival of multiple myeloma cases**
Basmaci C., Pehlivan M., Tomatir A., Sever T., Okan V., Yilmaz M., Oguzkan-Balci S., Pehlivan S.
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LX. **Effects of TNF?, NOS3, MDR1 Gene Polymorphisms on Clinical Parameters, Prognosis and Survival of Multiple Myeloma Cases**
PEHLİVAN S., TOMATIR A. G., PEHLIVAN M., OGUZKAN-BALCI S.
ASIAN PACIFIC JOURNAL OF CANCER PREVENTION, vol.17, no.3, pp.1009-1014, 2016 (SCI-Expanded)

LXI. **Genetic contributing factors to substance abuse: an association study between eNOS gene polymorphisms and cannabis addiction in a Turkish population**
Isir A. B., Nacak M., Balci S. O., Aynacioglu A. S., Pehlivan S.
AUSTRALIAN JOURNAL OF FORENSIC SCIENCES, vol.48, no.6, pp.676-683, 2016 (SCI-Expanded)

LXII. **Male Individualization Based on Y-Chromosomal Short Tandem Repeats: A Comparative Information Theoretical Analysis of 16 Y-STR Loci in Central Anatolia and Iraqi Populations**
ISIR A. B., Ozkorkmaz A., Baransel C., GÖKALP ÖZKORKMAZ E., Pehlivan S.
INTERNATIONAL JOURNAL OF HUMAN GENETICS, vol.15, no.4, pp.157-171, 2015 (SCI-Expanded)

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Awards

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