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International Researcher IDs

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Education Information

Expertise In Medicine, Istanbul University, Istanbul Medical Faculty, İç Hastalıkları, Turkey 1987 - 1993

Postgraduate, Istanbul University, Istanbul Medical Faculty, Turkey 1980 - 1987

Foreign Languages

English, B2 Upper Intermediate

Certificates, Courses and Trainings

Health&Medicine, Cancer Genetics, European School of Medical Genetics, 1999

Dissertations

Doctorate, Parafin içinde saklanan malign melanom biyopsi örneklerinde p53 geninin DGGE ve dizi analizi; p16, retinoblastoma ve CDK4 genlerinin FISH yöntemi ile incelenmesi, Istanbul University, Health Sciences Institute, İç Hastalıkları Anabilim Dalı, 2002

Expertise In Medicine, PROPAFENON'UN VENTRİKÜLER ARİTMİLER VE SİNYAL ORTALAMALI EKG PARAMETRELERİ ÜZERİNDEKİ ETKİSİNİN ARAŞTIRILMASI, Istanbul University, Istanbul Medical Faculty, İç Hastalıkları, 1993

Research Areas

Medicine, Health Sciences, Internal Medicine Sciences, Medical Genetics

Academic Titles / Tasks

Professor, Istanbul University, Istanbul Medical Faculty, Division of Medical Sciences , 1996 - Continues

Advising Theses

ÇEFLE K., Kronik Lenfositik Lösemide Kromozomal Aberasyonlar ve p53 Yolağındaki Gen Ekspresyonları Arasındaki İlişki, Doctorate, G.ÖZTAN(Student), Continues

ÇEFLE K., AML' de APAF-1 Promotör Metilasyonu, Kardeş Kromatid Değişimi, Kromozomal Anomaliler ile Klinik ve Laboratuvar Parametreleri Arasındaki İlişki, Postgraduate, Ö.Özgen(Student), 2012

ÇEFLE K., Kronik Lenfositik Lösemili Hastalarda XRCC1 (X-Ray Cross Complementing Group 1) Geninde Arg399Gln ve Arg194Trp Polimorfizmlerinin ve Kardeş Kromatid Değişimi Sıklığı ile Korelasyonlarının Araştırılması, Doctorate, N.Duman(Student), 2008

Published journal articles indexed by SCI, SSCI, and AHCI

- I. **The effect of Anzer honey on X-ray induced genotoxicity in human lymphocytes: An in vitro study**
Bagatir G., Kaya M., Suer İ., Çefle K., Palanduz A., Palanduz Ş., Becerir H. B., Koçyiğit Avcı M., Öztürk Ş.
MICROSCOPY RESEARCH AND TECHNIQUE, vol.85, no.6, pp.2241-2250, 2022 (SCI-Expanded)
- II. **OCT-1 Expression in Patients with Chronic Myeloid Leukemia: A Comparative Analysis with Respect to Response to Imatinib Treatment**
Bozkurt Bulakçı B., Aday A., Gürtekin B., Yavuz A. S., Öztürk Ş., Çefle K., Palanduz A., Palanduz Ş.
INDIAN JOURNAL OF HEMATOLOGY AND BLOOD TRANSFUSION, vol.1, no.1, pp.1-7, 2022 (SCI-Expanded)
- III. **Clinical Characteristics and Mutation Spectrum of Neurofibromatosis Type 1 in 27 Turkish Families**
Sharifi S., Kalayci T., Palanduz S., Ozturk S., Cefle K.
BALKAN MEDICAL JOURNAL, vol.38, no.6, pp.365-373, 2021 (SCI-Expanded)
- IV. **Skeletal and molecular findings in 51 Cleidocranial dysplasia patients from Turkey**
Berkay E. G., Elkanova L., Kalayci T., ULUDAĞ ALKAYA D., Altunoglu U., Cefle K., Mihci E., NUR B., Tasdelen E., Bayramoglu Z., et al.
AMERICAN JOURNAL OF MEDICAL GENETICS PART A, vol.185, no.8, pp.2488-2495, 2021 (SCI-Expanded)
- V. **Dysregulation of MS4A3 and PRDX5 Gene Expression in Multiple Myeloma Patients**
Suer İ., Aday A., Sariman M., Ayer M., Hindilerden I. Y., Ekmekci S. S., Abacı N., Palanduz Ş., Çefle K., Öztürk Ş.
UHOD-ULUSLARARASI HEMATOLOJİ-ONKOLOJİ DERGİSİ, vol.31, no.4, pp.205-213, 2021 (SCI-Expanded)
- VI. **RELATIONSHIP BETWEEN CHROMOSOMAL ABERRATIONS AND GENE EXPRESSIONS IN THE p53 PATHWAY IN CHRONIC LYMPHOCYTIC LEUKEMIA**
ÖZTAN G., Aktan M., Palanduz Ş., İŞSEVER H., ÖZTÜRK Ş., Nikerel E., Ucur A., Bagatir G., BAYRAK A. G., ÇEFLE K.
BALKAN JOURNAL OF MEDICAL GENETICS, vol.23, no.1, pp.15-23, 2020 (SCI-Expanded)
- VII. **DNA damage effects of inhalation anesthetics in human bronchoalveolar cells**
ÇUKUROVA Z., Cetingok H., Ozturk S., Gedikbasi A., HERGÜNSEL O., Ozturk D., Don B., Cefle K., Palanduz S., Ertem D. H.
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- VIII. **Investigation of Gene Expressions of Myeloma Cells in the Bone Marrow of Multiple Myeloma Patients by Transcriptome Analysis**
Sariman M., Abacı N., Ekmekci S., Cakiris A., Pacal F., Ustek D., Ayer M., Yenerel M. N., Besisik S., Cefle K., et al.
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- IX. **Clinical features and molecular genetic analysis in a Turkish family with oral white sponge nevus.**
Kurklu E., Ozturk S., Cassidy A. J., Ak G., Koray M., Cefle K., Palanduz S., Gulluoglu M., Tanyeri H., McLean W.
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- X. **REST Final-Exon-Truncating Mutations Cause Hereditary Gingival Fibromatosis**
BAYRAM Y., WHITE J. J., Elcioglu N., CHO M. T., ZADEH N., Gedikbasi A., Palanduz S., Ozturk Ş., Cefle K., Kasapcopur O., et al.
AMERICAN JOURNAL OF HUMAN GENETICS, vol.101, no.1, pp.149-156, 2017 (SCI-Expanded)
- XI. **WRN Mutation Update: Mutation Spectrum, Patient Registries, and Translational Prospects**
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HUMAN MUTATION, vol.38, no.1, pp.7-15, 2017 (SCI-Expanded)

- XII. **The frequency of C609T polymorphism in the NQO1 gene and its relation to cytogenetic abnormalities in patients with myelodysplastic syndrome.**
Bagatır G., Sirma S. Ö., Palanduz S., Ozturk S., Cefle K., Ozbek U., Yenerel M. N., Nalcacı M.
Cellular and molecular biology (Noisy-le-Grand, France), vol.62, pp.61-5, 2016 (SCI-Expanded)
- XIII. **Mutations in RAD21 Disrupt Regulation of APOB in Patients With Chronic Intestinal Pseudo-Obstruction**
BONORA E., BIANCO F., Cordeddu L., Bamshad M., Francescatto L., Dowless D., STANGHELLINI V., COGLIANDRO R. F., Lindberg G., Mungan Z., et al.
GASTROENTEROLOGY, vol.148, no.4, pp.771-793, 2015 (SCI-Expanded)
- XIV. **Evaluation of micronuclear frequencies in both circulating lymphocytes and buccal epithelial cells of patients with oral lichen planus and oral lichenoid contact reactions**
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ORAL DISEASES, vol.20, no.5, pp.521-527, 2014 (SCI-Expanded)
- XV. **Genotoxicity of fixation devices analyzed by the frequencies of sister chromatid exchange**
Aydil B. A., Kocak Berberoglu H., Ozturk S., Cefle K., Palanduz S., Erkal H.
ULUSAL TRAVMA VE ACIL CERRAHI DERGISI-TURKISH JOURNAL OF TRAUMA & EMERGENCY SURGERY, vol.19, no.4, pp.299-304, 2013 (SCI-Expanded)
- XVI. **A Turkish trichothiodystrophy patient with homozygous XPD mutation and genotype-phenotype relationship**
Pehlivan D., Cefle K., Raams A., Ozturk Ş., Baykal C., Kleijer W. J., Palanduz S., Jaspers N. G. J.
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- XVII. **Prostaglandin transporter mutations cause pachydermoperiostosis with myelofibrosis**
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- XVIII. **A novel two bases deletion in the albumin gene causes analbuminaemia in a young Turkish man**
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- XX. **Investigation of Arg399Gln and Arg194Trp Polymorphisms of the XRCC1 (X-Ray Cross-Complementing Group 1) Gene and Its Correlation to Sister Chromatid Exchange Frequency in Patients with Chronic Lymphocytic Leukemia**
Duman N., Aktan M., Ozturk S., Palanduz S., Cakiris A., Ustek D., Ozbek U., Nalcaci M., Cefle K.
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- XXI. **A novel frameshift deletion in the albumin gene causes analbuminemia in a young Turkish woman**
Dagnino M., Caridi G., Aydin Z., Ozturk S., Karaali Z., Kazancioglu R., Cefle K., Gursu M., Campagnoli M., Galliano M., et al.
Clinica Chimica Acta, vol.411, pp.1711-1715, 2010 (SCI-Expanded)
- XXII. **WRN mutations in Werner syndrome patients: genomic rearrangements, unusual intronic mutations and ethnic-specific alterations**
Friedrich K., Lee L., Leistritz D. F., Nuernberg G., Saha B., Hisama F. M., Eyman D. K., Lessel D., Nuernberg P., Li C., et al.
HUMAN GENETICS, vol.128, no.1, pp.103-111, 2010 (SCI-Expanded)
- XXIII. **NILOTINIB EFFICACY IN 21 IMATINIB-RESISTANT OR-INTOLERANT T (9;22) POSITIVE CHRONIC MYELOID LEUKEMIA PATIENTS WITH AND WITHOUT ADDITIONAL CHROMOSOMAL CHANGES**
Yavuz A. S., Elcioglu O. C., Akpinar T. S., Cosan F., Ucur A., Bayrak A., Cefle K., Oeztuerk S., Palanduz S., Yenerel M. N., et al.

- NOBEL MEDICUS, vol.6, no.2, pp.57-62, 2010 (SCI-Expanded)
- XXIV. **A POSSIBLE DELETERIOUS EFFECT OF INCREASED SERUM COPPER ON MYOCARDIAL FUNCTION IN PATIENTS WITH DILATED CARDIOMYOPATHY AWAITING TRANSPLANTATION**
Cefle K., Ercag E., Gezertas S., Uzer A., Oeztuerk Ş., Cefle A., Palanduz S., Gueler K.
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- XXV. **TRANSPLANTASYON BEKLEYEN DİLATE KARDİYOMİYOPATİLİ HASTALARDA YÜKSEK SERUM BAKIR DÜZEYİNİN MİYOKARD İŞLEVİ ÜZERİNDEKİ MUHTEMEL KÖTÜ ETKİSİ**
PALANDUZ Ş., ÇEFLE K.
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- XXVI. **İMATİNİBE DİRENÇLİ VEYA ENTOLERANS GÖSTEREN, KROMOZOMAL DEĞİŞİKLİKLERİ OLAN VE OLMAYAN T(9;22) POZİTİF KRONİK MYELOİD LÖSEMİLİ 21 HASTADA NİLOTİNİB'İN ETKİNLİĞİ**
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- XXVII. **Cytogenetic Analysis and Examination of SOS1 Gene Mutation in a Turkish Family with Hereditary Gingival Fibromatosis**
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- XXVIII. **Micronuclear and sister chromatid exchange analyses in peripheral lymphocytes of patients with oral lichen planus - a pilot study**
Ergun S., Warnakulasuriya S., Duman N., Saruhanoglu A., Sevinc B., Öztürk S., Ozel S., Cefle K., Palanduz S., Tanyeri H.
ORAL DISEASES, vol.15, no.7, pp.499-504, 2009 (SCI-Expanded)
- XXIX. **Comparison of the Cytogenetic and Molecular Analyses in the Assessment of Imatinib Response in Chronic Myelocytic Leukemia**
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- XXX. **Left Ventricular Thickness Is Increased in Nonhypertensive Turner's Syndrome**
Sozen A. B., Cefle K., Kudat H., Ozturk Ş., Oflaz H., Akkaya V., Palanduz S., Demirel S., Özcan M., Goren T., et al.
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- XXXI. **ALTERATIONS IN LYMPHOCYTE MEMBRANE PROTEIN CONTENT AND INCREASED LYMPHOCYTE RIGIDITY IN CATS WITH DIABETES MELLITUS**
ÇEFLE K.
JOURNAL OF PHYSIOLOGICAL SCIENCES, no.59, pp.505, 2009 (SCI-Expanded)
- XXXII. **The effect of parental consanguinity on the clinical and laboratory findings of rheumatoid arthritis**
Cefle K., ÇEFLE A., YAZICI A., SELEK A.
INTERNATIONAL JOURNAL OF CLINICAL PRACTICE, vol.63, no.7, pp.1056-1060, 2009 (SCI-Expanded)
- XXXIII. **ALTERATIONS IN LYMPHOCYTE MEMBRANE PROTEIN CONTENT AND INCREASED LYMPHOCYTE RIGIDITY IN CATS WITH DIABETES MELLITUS**
Tamer S. A., Cefle K., Kaymaz A. A., Albeniz I.
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Koeseoglu B., Oeztuerk Ş., Kocak H., Palanduz S., Cefle K.
Yonsei medical journal, vol.49, no.5, pp.742-7, 2008 (SCI-Expanded)
- XXXV. **Effect of Cyclosporin A and Tacrolimus on sister chromatid exchange frequency in renal transplant patients**
Ozturk Ş., Ayna T. K., Cefle K., Palanduz S., Ciftci H. S., Kaya Ş., Diler A. S., Turkmen A., Gurtekin M., Sever M. S., et al.
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- XXXVI. **Atrial and ventricular arrhythmogenic potential in Turner syndrome**
Sozen A. B., Cefle K., Kudat H., Ozturk Ş., Oflaz H., Pamukcu B., Akkaya V., Isguven P., Palanduz S., Özcan M., et al.
PACE-PACING AND CLINICAL ELECTROPHYSIOLOGY, vol.31, no.9, pp.1140-1145, 2008 (SCI-Expanded)

- XXXVII. **A novel locus for syndromic chronic idiopathic intestinal pseudo-obstruction maps to chromosome 8q23-q24**
 Deglincerti A, De Giorgio R, Cefle K, Devoto M, Pippucci T, Castegnaro G, Panza E, Barbara G, Cogliandro R. F., Mungan Z., et al.
 EUROPEAN JOURNAL OF HUMAN GENETICS, vol.15, no.8, pp.889-897, 2007 (SCI-Expanded)
- XXXVIII. **The genotoxic effects in lymphocyte cultures of children treated with radiosynovectomy by using yttrium-90 citrate colloid**
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- XXXIX. **Definition of C282Y mutation in a hereditary hemochromatosis family from Turkey**
 Yoenal O., Hatirnaz O., Akyuez F., KOROGLU G., Ozbeik U., Cefle K., Mungan Z.
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- XL. **Comparison of rheological parameters in patients with post hepatic and alcoholic cirrhosis**
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- XLI. **Initial maternal meiotic I error leading to the formation of a maternal i(2q) and a paternal i(2p) in a healthy male**
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- XLII. **Monitoring the genotoxic effects of radiosynovectomy with Re-186 in paediatric age group undergoing therapy for haemophilic synovitis**
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- XLIV. **Increased sister chromatid exchange frequency in young women with breast cancer and in their first-degree relatives.**
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- XLV. **A case of progressive pseudorheumatoid arthropathy of 'childhood' with the diagnosis delayed to the fifth decade**
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- XLVI. **The genotoxic effects in lymphocyte cultures of infants treated with radiosynovectomy by using Yttrium-90 citrate colloid.**
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- XLVII. **A different approach to telomere analysis with ddPRINS in chronic lymphocytic leukemia**
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- XLVIII. **Two sibilings with distal pachydermodactyly**
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- XLIX. **A solitary calvarial lytic lesion with typical histopathological findings of juvenile hyaline fibromatosis.**
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- L. **Clinical and molecular characterization of two adults with autosomal recessive Robinow syndrome**
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- LI. Genotoxicity and sister chromatid exchange in patients with myelodysplastic disorders.**
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- LII. Alterations in rheological properties and erythrocyte membrane proteins in cats with diabetes mellitus.**
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- LIV. Peutz-Jeghers syndrome: report of 6 cases in a family and management of polyps with intraoperative endoscopy.**
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- LV. A case of mandibuloacral dysplasia presenting with features of scleroderma**
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- LIX. Relationship between insulin-like growth factor-I and bone mineral density in men aged over 65 years**
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- LXV. **A case of turner syndrome with a rare reciprocal translocation between an autosome and the X chromosome**
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- 11.Ulusal Tıbbi Genetik Kongresi /46, XY,t(4;6) (p15.3;q23) Kriptik Dengeli Resiprokal Translokasyonunu Taşıyan İnfertil Olgu, Attendee, Turkey, 2014
- 11.Ulusal Tıbbi Genetik Kongresi /İnhalasyon Anestezisi İle Oluşan Genotoksik Etkilerin Bronkoalveoler Lavaj Sıvısında Tek Hücre Jel Elektoroforezi, Komet Yöntemi İle İncelenmesi, Attendee, Turkey, 2014
- 11.Ulusal Tıbbi Genetik Kongresi /mos 46, XX/ 47, XXX/ 48,XXXX Karyotipli Cinsel Kimlik Bozukluğu Tanılı Olgu, Attendee, Turkey, 2014
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- 11.Ulusal Tıbbi Genetik Kongresi /Akut Lökoz Tanılı Bir Olguda i(11)(q10), i(11)(p10),+11 Bulgusu, Attendee, Turkey, 2014
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