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Personal Information

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Education Information

Doctorate, Gazi University, Sağlık Bilimleri Enstitüsü, Tıbbi Biyoloji Ve Genetik (Dr), Turkey 1994 - 1999
Under Graduate, Gazi University, Tıp Fakültesi, Turkey 1985 - 1992

Foreign Languages

English, C1 Advanced

Dissertations

Doctorate, Türkiye'xxde alkol bağımlılarında dopamin D2 reseptör (DRD2) gen lokusu polimorfizmlerinin Taq 1 enzimi kullanılarak PCR yöntemi ile belirlenmesi, Gazi University, Sağlık Bilimleri Enstitüsü, Tıbbi Biyoloji Ve Genetik (Dr), 1999

Research Areas

Health Sciences

Academic Titles / Tasks

Professor, Gazi University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2011 - Continues
Associate Professor, Gazi University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2006 - 2011
Assistant Professor, Gazi University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2003 - 2006
Assistant Professor, Gazi University, Tıp Fakültesi, Temel Tıp Bilimleri Bölümü, 2003 - 2003
Lecturer, Gazi University, Tıp Fakültesi, Temel Tıp Bilimleri Bölümü, 2002 - 2003
Expert, Gazi University, Tıp Fakültesi, Temel Tıp Bilimleri Bölümü, 1999 - 2002
Research Assistant, Gazi University, Tıp Fakültesi, Temel Tıp Bilimleri Bölümü, 1994 - 1999

Professional Experience

Head of Department, Gazi University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2016 - 2019
Fakülte Kurulu Üyesi, Gazi University, Tıp Fakültesi, 2014 - 2017
Fakülte Yönetim Kurulu Üyesi, Gazi University, Tıp Fakültesi, 2014 - 2017
Yıl Koordinatörü, Gazi University, Tıp Fakültesi, 2004 - 2007
Head of Department, Gazi University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2003 - 2004

Advising Theses

YİRMİBEŞ KARAOĞUZ M., Doğumsal kalp anomalili çocukluk yaş grubu hastalarının klinik ve genetik analiz sonuçlarının retrospektik olarak değerlendirilmesi, Expertise In Medicine, S.MERMER(Student), 2018
YİRMİBEŞ KARAOĞUZ M., ERGÜN M. A. , SHOX gene analysis with Fluoresant in situ Hybridization and sequence methods in idiopathic short stature patients, Expertise In Medicine, A.BAKIR(Student), 2013
YİRMİBEŞ KARAOĞUZ M., PERÇİN F. E. , Prader-Willi ve Angelman Sendromlu Hastaların Tanısında Konvansiyonel Sitogenetik, Moleküler Sitogenetik Ve Moleküler Genetik Yöntemlerin Kullanımı, Expertise In Medicine, A.ÖZTÜRK(Student), 2011
YİRMİBEŞ KARAOĞUZ M., Ailesel olan ve ailesel olmayan şizofreni hastalarında kromozomal yeni düzenlenimlerin konvansiyonel sitogenetik ve moleküler sitogenetik yöntemlerle incelenmesi, Expertise In Medicine, A.KOÇ(Student), 2008

Jury Memberships

Associate Professor Exam, Associate Professor Exam, Gazi Üniversitesi, October, 2020
Appointment Academic Staff, Appointment Academic Staff, İstanbul Üniversitesi, November, 2019
Associate Professor Exam, Associate Professor Exam, Gazi Üniversitesi, November, 2017
Associate Professor Exam, Associate Professor Exam, Gazi Üniversitesi, April, 2017

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

I. **Expression of the syncytin-1 and syncytin-2 genes in the trophoblastic tissue of the early pregnancy losses with normal and abnormal karyotypes**
Tuğ E., Yirmibeş Karaoğuz M., Nas T.
GENE, vol.741, 2020 (Journal Indexed in SCI)

II. **Microdeletion and mutation analysis of the SHOX gene in patients with idiopathic short stature with FISH and sequencing**
Bakir A., YİRMİBEŞ KARAOĞUZ M., Emriye Percin F., TUĞ E., CİNAZ P., ERGÜN M. A.
TURKISH JOURNAL OF MEDICAL SCIENCES, vol.48, no.2, pp.386-390, 2018 (Journal Indexed in SCI)

III. **Confirmation of the prenatal mosaic trisomy 2 via fetal USG and cytogenetic analyses**
TUĞ E., Karcaaltincaba D., YİRMİBEŞ KARAOĞUZ M., Saat H., Ozek A.
JOURNAL OF MATERNAL-FETAL & NEONATAL MEDICINE, vol.30, no.13, pp.1579-1583, 2017 (Journal Indexed in SCI)

IV. **AN INTERCHANGE TRISOMY RESULTED IN DOWN SYNDROME**
TUĞ E., Karaoguz M., Bakir A.
GENETIC COUNSELING, vol.27, no.3, pp.429-431, 2016 (Journal Indexed in SCI)

V. **Chromosomal-array analysis reveals partial 11q duplication and partial 12p deletion in a mildly affected case**
TUĞ E., YİRMİBEŞ KARAOĞUZ M., Kayhan G., ERGÜN M. A. , PERÇİN F. E.
AMERICAN JOURNAL OF MEDICAL GENETICS PART A, vol.164, no.7, pp.1770-1776, 2014 (Journal Indexed in SCI)

VI. **Molecular karyotyping of an isolated partial trisomy 11q patient with additional findings**
KAYHAN G., Cavdarli B., YİRMİBEŞ KARAOĞUZ M., PERÇİN F. E. , Kaymak A. O. , Biri A., ERGÜN M. A.
GENE, vol.524, no.2, pp.355-360, 2013 (Journal Indexed in SCI)

VII. **Prenatally detected tetrasomy 18p and trisomy 21q due to i(18p) and i(21q) by using cytogenetic and molecular techniques**
YİRMİBEŞ KARAOĞUZ M., Percin E. F. , Pala E., Kaymak A. O. , TUĞ E., Biri A. A.
CHROMOSOME RESEARCH, vol.21, 2013 (Journal Indexed in SCI)

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with Mild Phenotype
TUĞ E., YİRMİBEŞ KARAOĞUZ M., KAYHAN G., ERGÜN M. A. , PERÇİN F. E.
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IX. **A patient with 9q subtelomeric deletion syndrome with additional findings.**
TUĞ E., Cavdarli B., Karaoguz M., PERÇİN F. E.
Genetic counseling (Geneva, Switzerland), vol.23, no.4, pp.465-71, 2012 (Journal Indexed in SCI Expanded)

X. **Detection of Marker Chromosome in the Abortion Material; Does It Reflect the Karyotype of the Pregnancy Lost Tissue or the Maternal Decidual Tissue? Case Report**
Koc A., YİRMİBEŞ KARAOĞUZ M., Pala E., PERÇİN F. E. , ERDEM M., Karaer K., Kaymak A. O.
TURKIYE KLINIKLERI TIP BILIMLERI DERGISI, vol.31, no.5, pp.1293-1297, 2011 (Journal Indexed in SCI)

XI. **Monosomy 1p36 Syndrome: The First Case Report from Turkey**
Karaer K., YİRMİBEŞ KARAOĞUZ M., PERÇİN F. E.
TURKIYE KLINIKLERI TIP BILIMLERI DERGISI, vol.31, no.1, pp.280-284, 2011 (Journal Indexed in SCI)

XII. **Comparison of DRD2 rs1800497 (TaqIA) polymorphism between schizophrenic patients and healthy controls: Lack of association in a Turkish sample**
ASLAN S., YİRMİBEŞ KARAOĞUZ M., Eser H. Y. , Karaer D. K. , Taner E.
INTERNATIONAL JOURNAL OF PSYCHIATRY IN CLINICAL PRACTICE, vol.14, no.4, pp.257-261, 2010 (Journal Indexed in SCI)

XIII. **The importance of systematic genetic approach to familial schizophrenia cases and discussion of cryptic mosaic X chromosome aneuploidies in schizophrenia pathogenesis**
Koc A., YİRMİBEŞ KARAOĞUZ M., COŞAR B., PERÇİN F. E. , Sahin S., Baysak E., Acikyurek K.
INTERNATIONAL JOURNAL OF PSYCHIATRY IN CLINICAL PRACTICE, vol.14, no.3, pp.204-211, 2010 (Journal Indexed in SCI)

XIV. **The Apolipoprotein E Gene and Taq1A Polymorphisms in Childhood Obesity**
ERGÜN M. A. , YİRMİBEŞ KARAOĞUZ M., Koc A., Camurdan O., BİDECİ A., Yazici A. C. , CİNAZ P.
GENETIC TESTING AND MOLECULAR BIOMARKERS, vol.14, no.3, pp.343-345, 2010 (Journal Indexed in SCI)

XV. **SRY GENE AMPLIFICATIONS AND GENOTYPINGS REVEALED THE OCCURRENCE OF THE HIDDEN MATERNAL DECIDUAL CELLS IN 46,XX KARYOTYPED SPONTANEOUS ABORTIONS**
Karaoguz M., PERÇİN F. E. , Pala E., Biri A. A. , Korucuoglu U.
GENETIC COUNSELING, vol.21, no.1, pp.9-17, 2010 (Journal Indexed in SCI)

XVI. **Prenatal diagnosis of mosaic ring 22 duplication/deletion with terminal 22q13 deletion due to abnormal first trimester screening and choroid plexus cyst detected on ultrasound**
Koc A., Arisoy O., Pala E., ERDEM M., Kaymak A. O. , Erkal O., YİRMİBEŞ KARAOĞUZ M.
JOURNAL OF OBSTETRICS AND GYNAECOLOGY RESEARCH, vol.35, no.5, pp.978-982, 2009 (Journal Indexed in SCI)

XVII. **An unexpected finding in a child with neurological problems: mosaic ring chromosome 18**
Koc A., Kan D., Karaer K., ERGÜN M. A. , YİRMİBEŞ KARAOĞUZ M., GÜCÜYENER K., Hinreiner S., Liehr T., PERÇİN F. E.
EUROPEAN JOURNAL OF PEDIATRICS, vol.167, no.6, pp.655-659, 2008 (Journal Indexed in SCI)

XVIII. **A case with a ring chromosome 22**
Koc A., Karaer K., ERGÜN M. A. , YİRMİBEŞ KARAOĞUZ M., Kan D., Cansu A., PERÇİN F. E.
TURKISH JOURNAL OF PEDIATRICS, vol.50, no.2, pp.193-196, 2008 (Journal Indexed in SCI)

XIX. **Double aneuploidy in spontaneous miscarriages: Two case reports and review of the literature**
Korucuoglu U., ERDEM M., Pala E., YİRMİBEŞ KARAOĞUZ M., ERDEM A., Biri A.
FETAL DIAGNOSIS AND THERAPY, vol.24, no.2, pp.106-110, 2008 (Journal Indexed in SCI)

XX. **Apolipoprotein e gene polymorphism in nonalcoholic fatty liver disease**
Demirag M. D. , Onen H. İ. , Karaoguz M., Dogan İ., Karakan T., Ekmekci A., Guz G.
DIGESTIVE DISEASES AND SCIENCES, vol.52, no.12, pp.3399-3403, 2007 (Journal Indexed in SCI)

XXI. **A boy with small supernumerary marker chromosome X identified by FISH.**
Koc A., Karaoguz M., Pala E., Kan D., Karaer K., Guecueyener K., PERÇİN F. E.
Genetic counseling (Geneva, Switzerland), vol.18, no.4, pp.393-9, 2007 (Journal Indexed in SCI Expanded)

XXII. **Cytogenetic results of amniocentesis materials: Incidence of abnormal karyotypes in the Turkish**

collaborative study
Karaoguz M., Bal F., Yakut T., Ercelen N. O. , ERGÜN M. A. , Gokcen A. B. , Biri A. A. , Kimya Y., Urman B., Gultomruk M., et al.
GENETIC COUNSELING, vol.17, no.2, pp.219-230, 2006 (Journal Indexed in SCI)

XXIII. **Is cytogenetic diagnosis of 46,XX karyotype spontaneous abortion specimens erroneous? Fluorescence in situ hybridization as a confirmatory technique**
Karaoguz M., Nas T., Konac E., Ince D., Pala E., Menevse S.
JOURNAL OF OBSTETRICS AND GYNAECOLOGY RESEARCH, vol.31, no.6, pp.508-513, 2005 (Journal Indexed in SCI)

XXIV. **The in vitro effect of beta-carotene and mitomycin C on SCE frequency in Down's syndrome lymphocyte cultures.**
Bal F., Sahin F., Yirmibes M., Balci A., Menevse S.
The Tohoku journal of experimental medicine, vol.184, no.4, pp.295-300, 1998 (Journal Indexed in SCI Expanded)

Articles Published in Other Journals

I. **Prenatal and Postnatal Clinical Spectrum of a Mosaic Small Supernumerary Marker Chromosome 22**
TUĞ E., YİRMİBEŞ KARAOĞUZ M., ERGÜN M. A.
International Journal of Pediatrics and Child Health, vol.7, pp.36-39, 2019 (Refereed Journals of Other Institutions)

II. **A new family with 3q27.3q29 İnterstitial Deletion**
KAYHAN G., SAVAŞ A., YİRMİBEŞ KARAOĞUZ M., PERÇİN F. E.
Gazi Medical Journal, vol.30, no.1, pp.1-101, 2019 (Refereed Journals of Other Institutions)

III. **Diagnostic Yield of Molecular Karyotyping of Idiopathic Intellectual Disability Patients Ended with One Causative Anomaly; Duplication 9q34 Syndrome**
Cavdarli B., Percin E. F. , YİRMİBEŞ KARAOĞUZ M., ERGÜN M. A.
GAZI MEDICAL JOURNAL, vol.30, no.3, pp.252-257, 2019 (Journal Indexed in ESCI)

IV. **A new family with 3q27.3.3q29 interstitial deletion**
KAYHAN G., SAVAŞ A., YİRMİBEŞ KARAOĞUZ M., PERÇİN F. E.
Gazi Medical Journal, vol.30, no.1, pp.32, 2019 (Refereed Journals of Other Institutions)

V. **The algorithm of mosaicism during prenatal diagnosis**
YİRMİBEŞ KARAOĞUZ M.
Gazi Medical Journal, vol.30, no.1, pp.7, 2019 (Refereed Journals of Other Institutions)

VI. **Elit Türk Sporcularında Anjotensin Dönüştürücü Enziim (I/D) Polimorfizmi ile Sportif Performans Arasındaki İlişkinin İncelenmesi**
KURTULUŞ M., GÜNAY M., ÇETİN E., ÇELENK Ç., CİCİOĞLU H. İ. , ALP E., YİRMİBEŞ KARAOĞUZ M., Kesici T.
Gaziantep Üniversitesi Spor Bilimleri Dergisi, vol.3, no.4, pp.122-137, 2018 (Refereed Journals of Other Institutions)

VII. **Lethal Multıple Pterygıum Syndrome related wıth RYR1 gene gene mutatıon**
KAYHAN G., SEZER A., Ozdemır H., ERGÜN M. A. , BAYRAM M., YİRMİBEŞ KARAOĞUZ M., PERÇİN F. E.
ERCIYES MEDICAL JOURNAL, vol.40, no.2, 2018 (Refereed Journals of Other Institutions)

VIII. **Clinical findings of the two fetuses with the pericentric inversion of chromosome Y relevant or coincidental**
SAVAŞ A., SEZER A., KAYHAN G., YİRMİBEŞ KARAOĞUZ M.
ERCIYES MEDICAL JOURNAL, vol.40, no.2, 2018 (Refereed Journals of Other Institutions)

IX. **Background of a carrier family with along inversion of chromosome 2 detected via karyotyping and aCGH analysis**
YİRMİBEŞ KARAOĞUZ M., SEZER A., KAYHAN G.
ERCIYES MEDICAL JOURNAL, vol.40, no.2, 2018 (Refereed Journals of Other Institutions)

X. **Lethal multiple pterygıum syndrome related with RYR1 gene mutation**
KAYHAN G., SEZER A., ÖZDEMİR H., ERGÜN M. A. , BAYRAM M., YİRMİBEŞ KARAOĞUZ M., PERÇİN F. E.
ERCIYES MEDICAL JOURNAL, vol.40, no.2, 2018 (Refereed Journals of Other Institutions)

XI. **Holoprosencephaly: A Rare Finding in Mosaic Trisomy 9 Syndrome**
Mermer S., Ozek M., Percin F. E. , YİRMİBEŞ KARAOĞUZ M., BAYRAM M.
ERCIYES MEDICAL JOURNAL, vol.40, no.1, pp.54-56, 2018 (Journal Indexed in ESCI)

XII. **Clinical and submicroscopic findings of two prenatal cases with inv dup del (8p) syndrome**
Sezer A., Bayram M., Kayhan G., Unal A., Ozdemir H., Karcaaltincaba D., Karaoguz M.
GENE REPORTS, vol.10, pp.75-78, 2018 (Journal Indexed in ESCI)

XIII. **Fetal karyotyping via invasive tests: When, to whom, and by which procedure?:Ne zaman, kimlere ve hangi yöntemle?**
YİRMİBEŞ KARAOĞUZ M.
Türkiye Klinikleri Tıbbi Genetik, vol.3, no.1, pp.44-47, 2018 (Other Refereed National Journals)

XIV. **Thrombophilic Status of Extracted Fetal Tissues of Spontaneously Aborted Embroys**
Pala E., YİRMİBEŞ KARAOĞUZ M., Balabanli B. K. , PERÇİN F. E. , ERDEM O. A.
GAZI MEDICAL JOURNAL, vol.28, no.1, pp.35-38, 2017 (Journal Indexed in ESCI)

XV. **Diagnosis of the Genomic Imprinting Diseases by the Usage of Conventional and Molecular Analyses**
Kaymak A. O. , YİRMİBEŞ KARAOĞUZ M., GÜCÜYENER K., PERÇİN F. E.
GAZI MEDICAL JOURNAL, vol.28, no.3, pp.200-203, 2017 (Journal Indexed in ESCI)

XVI. **A case with 22q11 deletion syndrome and anal anomalies**
ÖZYAVUZ ÇUBUK P., YİRMİBEŞ KARAOĞUZ M., PERÇİN F. E.
Erciyes Medical Journal, vol.38, no.1, pp.24, 2016 (Refereed Journals of Other Institutions)

XVII. **Holoprocencephaly noted in a case of mosaic trisomy 9 syndrome**
MERMER S., PERÇİN F. E. , Özek A. M. , YİRMİBEŞ KARAOĞUZ M.
Erciyes Medical Journal, vol.38, no.1, pp.46, 2016 (Refereed Journals of Other Institutions)

XVIII. **A new case with mosaic trisomy 19q**
ÜNAL A., KAYHAN G., YİRMİBEŞ KARAOĞUZ M., PERÇİN F. E.
Erciyes Medical Journal, vol.38, no.1, pp.32, 2016 (Refereed Journals of Other Institutions)

XIX. **Genetic aspects of recurrent pregnancy loss Genetik açıdan tekrarlayan gebelik kaybı**
YİRMİBEŞ KARAOĞUZ M.
Turkiye Klinikleri J Med Genet-Special Topics, vol.1, no.1, pp.23-27, 2016 (Other Refereed National Journals)

XX. **Farklı bulguları olan spina bifida ve renal anomalili iki olgu sunumu**
Aydın H., Yoldaş M., Yeşiller E., Geçkinli B., Karaman A., TUĞ E., YİRMİBEŞ KARAOĞUZ M., ATASOY H. İ.
J Kartal TR, vol.26, no.3, pp.272-276, 2015 (Other Refereed National Journals)

XXI. **Does ovulation induction increase the risk of aneuploid conception Comparison of first trimester miscarriages after FSH stimulated cycles and naturallly conceived cylcles**
Telli C., ERDEM M., BOZKURT N., YİRMİBEŞ KARAOĞUZ M., ÖKTEM M., ERDEM A., KARABACAK R. O. , Celtemen M., KAYHAN G.
International Jornal of Women's Health and Reproduction Sciences, vol.2, no.4, pp.225-228, 2014 (Refereed Journals of Other Institutions)

XXII. **A prenatal tertiary trisomy resulting from balanced maternal 8 9 translocation**
KAYHAN G., ERGÜN M. A. , BİRİ A., YİRMİBEŞ KARAOĞUZ M.
JOURNAL OF THE TURKISH-GERMAN GYNECOLOGICAL ASSOCIATION, 2011 (Refereed Journals of Other Institutions)

XXIII. **Karakteristik EEG bulgularına sahip Wolf Hirschhorn sendromlu bir olgu klinik ve moleküler sitogenetik tanı**
Karaer K., Koç A., YİRMİBEŞ KARAOĞUZ M., Cansu A., PERÇİN F. E.
Türkiye Klinikleri J Pediatr, vol.19, no.2, pp.171-175, 2010 (Other Refereed National Journals)

XXIV. **Estimates of sperm sex chromosome aneuploidy rates by fluorescence in situ hybridization in low level 47, XXY mosaicism**
YİRMİBEŞ KARAOĞUZ M., KONAÇ E., ERDEM A., ERDEM M.
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XXV. **Cytogenetic evaluation of cordocentesis materials in prenatal diagnosis and application of FISH as an additional technique**

Konaç E., Yirmibeş Karaoğuz M., Ergün M. A. , Nas T., Menevşe E. S.
Gazi Medical Journal, vol.15, no.3, pp.97-100, 2004 (National Refreed University Journal)

XXVI. **Cytogenetic Evaluation of Cordocentesis Materials in Prenatal Diagnosis and Application of the FISH Technique to Detect the Maternal Cell Contamination**
KONAÇ E., YİRMİBEŞ KARAOĞUZ M., ERGÜN M. A.
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XXVII. **Chromosomal Abnormalities in Habitual Abortions A Study of 192 Couples**
YİRMİBEŞ KARAOĞUZ M., ince g., ERGÜN M. A. , NAS T.
Gülhane Tıp Dergisi, vol.44, pp.40-42, 2002 (Other Refereed National Journals)

Refereed Congress / Symposium Publications in Proceedings

I. **Prenatal tanıda mozaisizm algoritmaları**
YİRMİBEŞ KARAOĞUZ M.
Uluslararsı katılımlı 13. Ulusal Tıbbi Genetik Kongresi, Turkey, 7 - 11 November 2018

II. **A mosaic double aneuploidy: mos 45,X/47,XX,18 with mild phenotype**
DEMİRBAŞ M. H. , HABİLOĞLU E., ERGÜN M. A. , PERÇİN F. E. , YİRMİBEŞ KARAOĞUZ M.
European Human Genetics Conference, Milano, İtalya, 16 - 19 June 2018

III. **Prenatal tanıya giriş ve invasiv testler**
YİRMİBEŞ KARAOĞUZ M.
Erciyes Tıp Genetik Günleri 2017, 11-13 Mayıs 2017, Kayseri, Turkey, 11 - 13 May 2017

IV. **Is there any relationship between NRG1 gene duplication and cardiac findings in two prenatal cases with invdupdel(8p) syndrome?**
SEZER A., BAYRAM M., KAYHAN G., ÜNAL A., ÖZDEMİR H., KARÇAALTINCABA D., YİRMİBEŞ KARAOĞUZ M.
European Human Genetics Conference, Kopenhag, Danimarka, Kopenhag, Denmark, 27 - 30 May 2017

V. **A Case with 22q11 deletion Syndrome and Anal Anomalies**
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VI. **Holoprocencephaly noted in case of Mosaic Trisomy 9 Syndrome**
MERMER S., PERÇİN F. E. , Özek A. M. , BAYRAM M., YİRMİBEŞ KARAOĞUZ M.
Medical Genetics and Clinical Applications, Erciyes Medical Journal, Turkey, 11 - 13 February 2016, vol.38, pp.46

VII. **A New Case with Mosaic Trisomy 19Q**
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VIII. **Primary amenorrhea visual impairment and intellectual disability in a girl with a complex rearrangement involving 5q33 3 and 9q21 2 microdeletions**
KAYHAN G., ERGÜN M. A. , Thomas L., YİRMİBEŞ KARAOĞUZ M., PERÇİN F. E.
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IX. **Fraser Syndrome a case report diagnosed prenatally at 17 weeks old and postpartum examinations**
BİRİ A., GÜLER İ., HİMMETOĞLU Ö., YİRMİBEŞ KARAOĞUZ M., BALCI S.
Altıncı Türk-Alman Jinekoloji Derneği Kongresi, Antalya, Turkey, 18 - 22 May 2005

X. **Chromosomal Array AnalysisReveals Partial 11q Duplication and Partial 12p Deletion in A Mildly Affected Case**
TUĞ E., YİRMİBEŞ KARAOĞUZ M., KAYHAN G., ERGÜN M. A. , PERÇİN F. E.
10. Ulusal tıbbiGenetik Kongresi, Bursa, 2012., Bursa, Turkey, 18 - 22 December 2012

XI. **Kromozomal yeniden düzenlenmelerde moleküler karyotipleme ile genotip fenotip ilişkisinin belirlenmesi**
KAYHAN G., ERGÜN M. A. , YİRMİBEŞ KARAOĞUZ M., PERÇİN F. E.
11. Ulusal Tıbbi genetil Kongresi, İstanbul, Turkey, 24 - 27 September 2014

XII. **Chromosomal abnormalities identifiedin 836 abortions nine years experience**

YİRMİBEŞ KARAOĞUZ M., KAYHAN G., öztürk kaymak a., PALA E.
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XIII. **Meningomyelocele and Renal Hypoplasia: A Rare Case Report**
Aydin H., Tug E., Duzenli S., Erkal O., Yoldas A., Karaoguz M.
7th European Cytogenetics Conference, Stockholm, Sweden, 4 - 07 July 2009, vol.17, pp.57

XIV. **2q37 delesyonlu bir olgu**
KAYHAN G., erkal ö., YİRMİBEŞ KARAOĞUZ M., PERÇİN F. E.
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