Asst. Prof. SİNEM KOCAGİL

Personal Information

Email: skocagil@ogu.edu.tr
Web: https://avesis.ogu.edu.tr/2305

International Researcher IDs
ORCID: 0000-0003-2595-3919
Yoksis Researcher ID: 246530

Education Information

Undergraduate, Anadolu University, Department Of Sogıology (External), Turkey 2017 - 2022
Expertise In Medicine, Eskisehir Osmangazi University, TIP FAKÜLTESİ, DAHİLİ TIP BİLİMLERİ BÖLÜMÜ, Turkey 2016 - 2020
Undergraduate, Akdeniz University, Faculty Of Medicine, Turkey 2009 - 2015

Foreign Languages

English, C1 Advanced

Certificates, Courses and Trainings

Health&Medicine, European Diploma in Medical Genetics and Genomics, European Union of Medical Specialists, 2021
Health&Medicine, 12th Goldrain Course in Clinical Cytogenetics, ECA - European Cytogeneticists Association, 2017

Dissertations

Expertise In Medicine, NON-sendromik konjenital kalp hastalıklarında aday genlerin değerlendirilmesi, Eskisehir Osmangazi University, TIP FAKÜLTESİ, DAHİLİ TIP BİLİMLERİ BÖLÜMÜ, 2020

Research Areas

Medical Genetics

Academic Titles / Tasks

Assistant Professor, Eskisehir Osmangazi University, TIP FAKÜLTESİ, DAHİLİ TIP BİLİMLERİ BÖLÜMÜ, 2022 - Continues
Lecturer, Eskisehir Osmangazi University, TIP FAKÜLTESİ, DAHİLİ TIP BİLİMLERİ BÖLÜMÜ, 2021 - Continues
Research Assistant, Eskisehir Osmangazi University, TIP FAKÜLTESİ, DAHİLİ TIP BİLİMLERİ BÖLÜMÜ, 2016 - 2021

Courses

Klinik Genetiğe Giriş, Postgraduate, 2023 - 2024
Preimplantasyon Genetik Tanı, Undergraduate, 2023 - 2024, 2022 - 2023
Klinik Genetiğe Giriş, Undergraduate, 2023 - 2024
Genetik Danışmanlık , Undergraduate, 2023 - 2024
Prenatal Tanı Yöntemleri, Undergraduate, 2023 - 2024, 2022 - 2023
Kan Hastalıklarının Genetik Yönü: Hemofili, Undergraduate, 2023 - 2024
Genomik Varyasyonların Moleküler Tanısı, Undergraduate, 2023 - 2024, 2022 - 2023
Kan Hastalıklarının Genetik Yönü: Talassemi, Undergraduate, 2023 - 2024
Santral Sinir Sistemi Tümör Genetiği, Undergraduate, 2023 - 2024, 2022 - 2023
Genetik Hastalıkların Tanısında Kullanılan Moleküler Yöntemler, Undergraduate, 2023 - 2024, 2022 - 2023
Teratojenler ve Mutajenler, Undergraduate, 2023 - 2024, 2022 - 2023, 2021 - 2022
Beyin ve İskelet Sistemi Gelişimi Anomalilerinin Genetik Yönü, Undergraduate, 2023 - 2024, 2022 - 2023

Published journal articles indexed by SCI, SSCI, and AHCI

I. **Late Diagnosis of Congenital Chloride Diarrhea Mimicking Hirschsprung's Disease**
Çelik A. T., Barış Z., Aydemir Y., Kocagil S.
CLINICAL PEDIATRICS, vol.63, no.11, pp.1494-1497, 2024 (SCI-Expanded)

II. **Interstitial 3p25.3 deletion syndrome: 13 years'-long follow-up of an affected individual.**
Kocagil S., Susam E., Yimenicioğlu S., Aynaci S., Gökalp E. E., Artan S.
Clinical dysmorphology, 2024 (SCI-Expanded)

III. **Screening of Mutations in Maturity-onset Diabetes of the Young-related Genes and RFX6 in Children with Autoantibody-negative Type 1 Diabetes Mellitus.**
Şimşek E., Cilingir O., Simsek T., Kocagil S., Erzurumluoglu Gokalp E., Demiral M., Binay C.
Journal of clinical research in pediatric endocrinology, vol.16, pp.137-145, 2024 (SCI-Expanded)

IV. **Further Evidence for RFWD3 Gene Causing Fanconi Anemia Complementation Group W: Detailed Clinical Report of the Second Case in the Literature**
Kocagil S., Şafak İ. N., Saraç E., Aydın C., Artan S., Kırel B.
MOLECULAR SYNDROMOLOGY, vol.14, no.4, pp.1, 2023 (SCI-Expanded)

V. **Clinical and molecular evaluation of MEFV gene variants in the Turkish population: a study by the National Genetics Consortium**
Dundar M., Fahrioglu U., Yildiz S. H., Bakir-Gungor B., Temel S. G., Akin H., Artan S., Cora T., Sahin F. I., Dursun A., et al.
FUNCTIONAL & INTEGRATIVE GENOMICS, vol.22, no.3, pp.291-315, 2022 (SCI-Expanded)

VI. **A case of familial recurrent 17q12 microdeletion syndrome presenting with severe diabetic ketoacidosis**
Aydın C., Kıral E., Susam E., Tufan A., YARAR C., Çetin N., Kocagil S., Kırel B.
TURKISH JOURNAL OF PEDIATRICS, vol.64, no.3, pp.558-565, 2022 (SCI-Expanded)

VII. **Spectrum of PAH gene mutations and genotype-phenotype correlation in patients with phenylalanine hydroxylase deficiency from Turkey**
Cinar M., Yildirim G. K., KOCAGİL S., ÇİLİNGİR O.
JOURNAL OF PEDIATRIC ENDOCRINOLOGY & METABOLISM, vol.35, no.5, pp.639-647, 2022 (SCI-Expanded)

VIII. **Assessment of clinical characteristics of cardiac amyloidosis as a potential underlying etiology in patients diagnosed with heart failure with preserved ejection fraction**
Murat S., Cavusoglu Y., Yalvac H. E., Sivrikoz I. A., Kocagil S.
KARDIOLOGIA POLSKA, vol.80, no.6, pp.672-678, 2022 (SCI-Expanded)

IX. **NDE1-related disorders: A recurrent NDE1 pathogenic variant causing Lissencephaly 4 can also be associated with microhydranencephaly**
BAŞ H., ŞAYLISOY S., ÇİLİNGİR O., Gokalp E., KOCAGİL S., YARAR C., Aras B., ARTAN S.
AMERICAN JOURNAL OF MEDICAL GENETICS PART A, vol.188, no.1, pp.326-331, 2022 (SCI-Expanded)

X. **An Anomaly with Potential as a New Prognostic Marker in CLL with del(13q): Gain of 16p13.3**
IŞIK S., Gunden G., GÜNDÜZ E., Akay O. M., Aslan A., ÖZEN H., ÇİLİNGİR O., ERZURUMLUOĞLU GÖKALP E., KOCAGİL S., ARTAN S., et al.
CYTOGENETIC AND GENOME RESEARCH, vol.161, no.10-11, pp.479-487, 2021 (SCI-Expanded)

XI. **A pediatric BAL case with double Ph chromosomes and trisomy 5**
Gunden G., Işık S., Özdemir C., Çilingir O., Bör Ö., Gokalp E. E., Kocagil S., Artan S., Aras B. D.
Cancer Genetics, vol.258, pp.7-9, 2021 (SCI-Expanded)

XII. **Which prognostic marker is responsible for the clinical heterogeneity in CLL with 13q deletion?**
Durak Aras B., Isik S., Uskudar Teke H., Aslan A., Yavasoglu F., Gulbas Z., Demirkan F., Ozen H., Cilingir O., Inci N. S., et al.
Molecular cytogenetics, vol.14, no.1, pp.2, 2021 (SCI-Expanded)

XIII. **The association between repeat number in C9orf72 and phenotypic variability in Turkish patients with frontotemporal lobar degeneration**
ERZURUMLUOĞLU GÖKALP E., ÇİLİNGİR O., Adapinar B. D. O., Bilgic B., Kocagil S., ÖZEN H., DURAK ARAS B., YENİLMEZ Ç., ARTAN S.
Neurobiology of Aging, vol.76, 2019 (SCI-Expanded)

Articles Published in Other Journals

I. **ERCC8 related Cockayne syndrome type-1: A rare entity diagnosed in a Turkish boy**
Kocagil S., Keklikci A. R., Aydemir Y., Çilingir O., Aynacı S., Erzurumluoğlu Gökalp E., Durak Aras B., Artan S.
Journal of Surgery and Medicine, vol.7, no.10, pp.719-721, 2023 (Peer-Reviewed Journal)

II. **Amiloid Olgu Serisi /Transtiretin tip kardiyak amiloidoz/AV blok**
Murat S., Çavuşoğlu Y., Ak Sivrikoz İ., Kocagil S., Yalvaç H. E., Gündüz E., Üsküdar Teke H.
updates in cardiology, vol.4, no.1, pp.3-5, 2022 (Non Peer-Reviewed Journal)

III. **Wiedemann-Steiner Syndrome: A Rare Differential Diagnosis of Neurodevelopmental Delay and Dysmorphic Features**
ÇARMAN K. B., Kaplan E., Aslan C. N., Kocagil S., Cilinigr O., YARAR C.
JOURNAL OF PEDIATRIC GENETICS, vol.11, no.02, pp.162-164, 2022 (ESCI)

IV. **Evaluation of the Effect of Circulating IncRNAs in Colorectal Cancers: As a Potential Biomarker**
ARSLAN S., DİNCER M., BAYIR D., EKER SARIBOYACI A., ERZURUMLUOĞLU GÖKALP E., KOCAGİL S., ARIK D., ÖZEN H., DURAK ARAS B., ARTAN S., et al.
TURK ONKOLOJI DERGISI, vol.37, pp.305-313, 2022 (ESCI)

V. **Analyzing The Mutations Of Notch1 And Sf3b1 Genes In Cases With CLL Detected Isolated 13q Deletion**
Günden G., Işık S., Üsküdar Teke H., Çilingir O., Oğuz Davutoğlu N., Erzurumluoğlu Gökalp E., Kocagil S., Durak Aras B.
Osmangazi Tıp Dergisi, vol.45, no.2, pp.480-484, 2021 (Peer-Reviewed Journal)

VI. **Türk Popülasyonunda APOE Polimorfizmleri ve Alzheimer Hastalığı Arasındaki İlişki**
ÇİLİNGİR O., ÖZBABALIK ADAPINAR B. D., DURAK ARAS B., ERZURUMLUOĞLU GÖKALP E., ÖZKAN S., ARSLAN S., HAZİYEVA K., KOCAGİL S., BİLGİN M., ARTAN S.
OSMANGAZİ JOURNAL OF MEDICINE, 2020 (Peer-Reviewed Journal)

VII. **Ailesel Akdeniz Ateşi Tanısı Alan Olgularda MEFV Geni Mutasyonlarının ve Allel Frekanslarının Dağılımı - Tek Merkez Deneyimi**
ÇİLİNGİR O., DURAK ARAS B., ARSLAN S., KUTLAY Ö., ERZURUMLUOĞLU E., KOCAGİL S., ARTAN S.
Osmangazi Tıp Dergisi, vol.40, no.2, pp.39-46, 2018 (Peer-Reviewed Journal)

Books & Book Chapters

I. **Spinobulbar Musküler Atrofiye Genetik Yaklaşım**
ÇİLİNGİR O., KOCAGİL S.
in: Nükleotid Tekrar Artışı Hastalıkları - 2023, Prof. Dr. Ayşe Gül ZAMANİ, Editor, Türkiye klinikleri, ONLINE, pp.55-61, 2023

Refereed Congress / Symposium Publications in Proceedings

I. **CHD8 missense variants cause a variable neurodevelopmental disorder with incomplete penetrance**
Godfrey M., Levy M., Leonardi E., Campbell C., Demain L., Jenkinson S., Hilton S., Almoguera Castillo B., Balasubramanian M., Bijlsma E. K., et al.
56th Annual Conference of the European-Society-of-Human-Genetics (ESHG), Glasgow, England, 10 - 13 June 2023, pp.721

II. **Report of first Endocrine-cerebro-osteodysplasia patient to reach childhood age**
KOCAGİL S., ÖZBAKIR D. H., Kabaoglu U., ERZURUMLUOĞLU GÖKALP E., ÇİLİNGİR O.
56th Annual Conference of the European-Society-of-Human-Genetics (ESHG), Glasgow, England, 10 - 13 June 2023, pp.145

III. **REPORT OF THE FIRST ENDOCRINE-CEREBRO-OSTEODYSPLASIA PATIENT TO REACH CHILDHOOD AGE**
Özbakır D. H., Kocagil S., Erzurumluoğlu Gökalp E., Çilingir O.
EURODYSMORPHO 2023, Lisbon, Portugal, 13 - 16 September 2023, pp.28

IV. **EXPANDING THE PHENOTYPIC SPECTRUM OF INTELLECTUAL DEVELOPMENTAL DISORDER-70**
Kocagil S., Keklikci A. R., Aynacı S., Susam E.
EuroDysmorpho 2023, Lisbon, Portugal, 13 - 16 September 2023, pp.89-90

V. **Complete paternal isodisomy of chromosome 15 in a patient with atypical presentation of Angelman syndrome**
Çilingir O., Saraç E., Aynacı S., Kocagil S.
14th European Cytogenomics Conference, Montpellier, France, 1 - 04 July 2023, pp.48

VI. Yalvaç H. E., Murat S., Çavuşoğlu Y., Kocagil S., Ak Sivrikoz İ., Üsküdar Teke H.
2. Ulusal KY Toplantısı, İstanbul, Turkey, 2 - 03 June 2023, pp.13-14

VII. **Fanconi Aplastik Anemili 5 Olgu: FANCA, ERCC4, RFWD3 Genlerinde 3 Novel/Nadir Varyant**
Özbakır D. H., Saraç E., Susam E., Kocagil S., Aynacı S., Kırel B., Çilingir O.
2. Ulusal HematoOnkoGenetik Kongresi, İskele, Cyprus (Kktc), 4 - 07 May 2023, pp.52

VIII. **Farklı Yapısal Y Kromozom Anomalisi Saptanan Olgularda Klinik ve Genetik Sonuçların İncelenmesi**
Tosumoğlu E., Saraç E., Üre İ., Kocagil S.
15. Ulusal Tıbbi Genetik Kongres, Muğla, Turkey, 9 - 13 November 2022, pp.240

IX. **Oldukça Nadir Bir Otozomal Resesif Ehlers-Danlos Sendromu Alt tipi: TNXB Geni İlişkili Klasik Bulgularla Seyreden Bir Aile**
Kocagil S., Aynacı S.
15. Ulusal Tıbbi Genetik Kongresi , Muğla, Turkey, 9 - 13 November 2022, pp.99

X. **NANCE-HORAN SYNDROME: AN ULTRA RARE PHENOTYPE DIGNOSED IN A FAMILY WITH FEMALE SIBLING AFFECTED AS SEVERE AS THE MALE**
Kocagil S., Kaplan E., Erzurumluoğlu Gökalp E.
EuroDysmorpho 2022, Barcelona, Spain, 14 - 17 September 2022, pp.95-96

XI. **CRPPA gene-associated Walker-Warburg syndrome: A rare prenatal case with intragenic homozygous deletion**
Özbakır D. H., Bütün Z., Kocagil S.
7. Uluslararası Erciyes Tıp Tıbbi Genetik Kongresi, Kayseri, Turkey, 26 - 28 May 2022, pp.102-103

XII. **A Brittle cornea syndrome with a novel pathogenic variant of PRDM5 gene**
Susam E., Yıldırım N., Kocagil S., Çilingir O.

54th European Society of Human Genetics (ESHG) Conference:Virtual Conference, 28 - 31 August 2021, vol.30, pp.119-120

XIII. **Peutz-Jeghers Sendromu:İki Nadir Varyant, İki Aile**
Kocagil S., Çilingir O.
1. Ulusal HematoOnkoGenetik Kongresi, Antalya, Turkey, 25 - 28 November 2021, pp.166

XIV. **Hiperfenilalaninemi ve Fenilketonüri Tanılı Hastalarda PAH GeniMutasyon Spektrumu ve Genotip-Fenotip Korelasyonu**
Akyüz M., Kılıç Yıldırım G., Kocagil S., Çilingir O.
65. Türkiye Milli Pediatri Kongresi, Antalya, Turkey, 3 - 07 November 2021

XV. **A NOVEL FRAMESHIFT VARIANT IN A PATIENT WITH CHD8-RELATED OVERGROWTH SYNDROME**
KOCAGİL S., KEKLİKCİ A. R., KILIÇ YILDIRIM G., ÇİLİNGİR O., ERZURUMLUOĞLU GÖKALP E., DURAK ARAS B., ARTAN S.
31TH EUROPEAN MEETING ON DYSMORPHOLOGY ONLINE, Germany, 23 - 25 September 2021, vol.1, pp.34-35

XVI. **A complex chromosomal rearrangement in a patient with developmental delay and dysmorphic features**
AYNACI S., TOSUMOĞLU E., KEKLİKÇİ A. R., KOCAGİL S., ERZURUMLUOĞLU GÖKALP E., ÇİLİNGİR O., DURAK ARAS B., ARTAN S.
14. ULUSAL TIBBİ GENETİK KONGRESİ'xx'xxULUSLARAARSI KATILIMLI'xx'xx, Turkey, 20 - 22 November 2020, vol.31, pp.94

XVII. **A Rare Form Of Constitutional Chromoanasynthesis: Ring Chromosome 18**
Susam E., Erzurumluoğlu Gökalp E., Tosumoğlu E., Kocagil S., Çilingir O., Durak Aras B., Artan S.
14. ULUSAL TIBBİ GENETİK KONGRESİ'xx'xxULUSLARAARSI KATILIMLI'xx'xx, Ankara, Turkey, 20 - 22 November 2020, vol.31, pp.15

XVIII. **Atipik bulguları olan Smith-Magenis sendromlu bir olgu sunumu**
Saraç E., Özbakır D. H., Kocagil S., Erzurumluoğlu Gökalp E., Çilingir O., Durak Aras B., Artan S.
14. ULUSAL TIBBİ GENETİK KONGRESİ'xx'xxULUSLARAARSI KATILIMLI'xx'xx, Eskişehir, Turkey, 20 - 22 November 2020, vol.31, pp.92

XIX. **A rare homozygous variant in CUL7 gene in two syblings with variable features of 3M syndrome**
KOCAGİL S., KILIÇ YILDIRIM G., ARTAN S.
14. ULUSAL TIBBİ GENETİK KONGRESİ"Uluslararası Katılımlı", Turkey, 20 - 22 November 2020

XX. **A Novel Mutation of DYSF Gene in A Patient with Limb Girdle Muscular Dystrophy Type 2b**
KOCAGİL S., KAPLAN E., SUSAM E., DURAK ARAS B., ARTAN S., ERZURUMLUOĞLU GÖKALP E., ÇİLİNGİR O.
European Human Genetics Virtual Conference, 06 June 2020

XXI. **An interstitial 6q25.1 microdeletion syndrome in a patient with dysmorphic features, intellectual dysability and stereotypical movements**
KOCAGİL S., DURAK ARAS B., ERZURUMLUOĞLU GÖKALP E., ÇİLİNGİR O., AYNACI S., ARTAN S.
13th Balkan Congress of Human Genetics, Edirne, Turkey, 17 - 20 April 2019, vol.22, pp.85

XXII. **A rare case of severe microcephaly caused by pathogenic variant of NDE1**
BAŞ H., ÇİLİNGİR O., DURAK ARAS B., KOCAGİL S., ARTAN S.
13th Balkan Congress of Human Genetics, Edirne, Turkey, 17 - 20 April 2019, vol.22, pp.67

XXIII. **The 15q11.2 BP1–BP2 Microdeletion Syndrome with Variable Expressivity**
ERZURUMLUOĞLU GÖKALP E., ÇİLİNGİR O., DURAK ARAS B., KOCAGİL S., SUSAM E., ARTAN S.
13th Balkan Congress of Human Genetics, Edirne, Turkey, 17 - 20 April 2019, vol.22, pp.95

XXIV. **DO INTERMEDIATE REPEAT EXPANSION LENGTH IN C9ORF72 HAVE AN EFFECT ON CLINICS IN CASES WITH FRONTOTEMPORAL LOBAR DEGENERATION?**
ERZURUMLUOĞLU GÖKALP E., ÖZBABALIK ADAPINAR B. D., TEPGEÇ F., ÇİLİNGİR O., BİLGİÇ B., KOCAGİL S., ÖZEN H., DURAK ARAS B., YENİLMEZ Ç., ARTAN S.
FENS Regional Meeting 2019, Belgrade, Serbia, 10 - 13 July 2019

XXV. **Clinical use of chromosomal microarray analysis in detection of fetal chromosomal abnormalities**
ARTAN S., BAŞ H., ERZURUMLUOĞLU GÖKALP E., VELİPAŞAOĞLU M., KOCAGİL S., DURAK ARAS B., TEMENA M. A., PANAL G., ANSARİ S. K., ÇİLİNGİR O.

12th EuropeanCytogenomics Conference 2019, Salzburg, Austria, 6 - 09 July 2019, vol.12

XXVI. **Macroorchidism as a Uniq Sign in 3q13.31 Deletion Syndrome.**
SUSAM E., ÇİLİNGİR O., DURAK ARAS B., ERZURUMLUOĞLU GÖKALP E., KOCAGİL S., ARTAN S., ŞİMŞEK E.
European Human Genetics Conference, Gothenburg, Sweden, 15 June 2019, vol.27, pp.1843

XXVII. **Nöronal Seroid Lipofusinozis Tip 11 Tanısı Alan 3 Kardeş**
Yarar C., Kaplan E., Kocagil S., Çarman K. B., Çilingir O., Kıral E., Bozan G., Laçinel Gürlevik S., Kılıç Yıldırım G., Dinleyici E. Ç.
21. Uluslararası Katılımlı Ulusal Çocuk Nörolojisi Kongresi, Muğla, Turkey, 1 - 05 May 2019, pp.112-113

XXVIII. **De Novo t(X;5) in a Patient with Premature Ovarian Failure and Reccurrent Vertebrae Fractures.**
AYNACI S., KOCAGİL S., TOSUMOĞLU E., PANAL G., ÇİLİNGİR O., DURAK ARAS B., ARTAN S.
Erciyes Medical Genetics Days, Kayseri, Turkey, 21 February 2019, vol.22, pp.23

XXIX. **NPHP1 homozigot gen delesyonu saptanan Joubert Sendromu tip 4 tanılı iki olgu sunumu**
KOCAGİL S., EREN M. C., ELMAS M., YARAR C., ERZURUMLUOĞLU E., ÇİLİNGİR O., ARTAN S.
Uluslararası Katılımlı XIII. Ulusal Tıbbi Genetik Kongresi, Antalya, Turkey, 7 - 11 November 2018

XXX. **CYP21A2 mutations in congenital adrenal hyperplasia due to 21 hydroxylase deficiency in Turkish population**
ÇİLİNGİR O., ŞİMŞEK E., DURAK ARAS B., ERZURUMLUOĞLU E., TEMENA A., KOCAGİL S., ARTAN S.
European Human Genetics Conference 2018, Milan, Italy, 16 - 19 June 2018

XXXI. **Comparison of phenotypic variability with C9orf72 gene GGGGCC hexanucleotide repeat expansion in frontotemporal lobar degeneration spectrum**
ERZURUMLUOĞLU E., ÇİLİNGİR O., ÖZBABALIK ADAPINAR B. D., BİLGİÇ B., KOCAGİL S., DURAK ARAS B., YENİLMEZ Ç., ARTAN S.
European Human Genetics Conference 2018, Milan, Italy, 16 - 19 June 2018

XXXII. **SEVERE PELIZAEUS - MERZBACHER DISEASE ON A CASE WITH DEVELOPMENTAL DELAY AND ABNORMAL MYELINATION**
BAŞ H., YARAR C., ERZURUMLUOĞLU E., DURAK ARAS B., KOCAGİL S., ARTAN S.
Erciyes Medical Genetics Days 2018, Turkey, 7 - 10 March 2018

XXXIII. **Parsiyel Trizomi 15 tanımlanan 2 kardeş olgu**
KOCAGİL S., TEKİN A. N., ERZURUMLUOĞLU E., ÇAKIL SAĞLIK A., Tosumoğlu E., ARTAN S.
Ulusal Neonatoloji Kongresi (UNEKO-26), Girne, Cyprus (Kktc), 14 - 18 April 2018, pp.221

XXXIV. **De novo 7q31 deletion involving FOXP2 gene associated with speech disability**
ERZURUMLUOĞLU E., ÖZDEMİR M., KOCAGİL S., ÇİLİNGİR O., TOSUMOĞLU E., DURAK ARAS B., ARTAN S.
11th European Cytogenetic Conference 2017, 1 - 04 July 2017, vol.10

XXXV. **A New Mutation Associated With Bannayan Riley Ruvalcaba Syndrome**
ÇİLİNGİR O., ÖZDEMİR M., YARAR C., ERZURUMLUOĞLU E., DURAK ARAS B., KOCAGİL S., KHADEM ANSARİ S., BAŞ H., ARTAN S.
Erciyes Tıp Genetik Günleri, Turkey, 11 - 13 May 2017, vol.39

XXXVI. **A NOVEL INDEL MUTATION IN THE TCOF1 GENE FOUND IN ANEWBORN WITH TREACHER COLLINS SYNDROME**
KOCAGİL S., ÇİLİNGİR O., ÇARMAN K. B., AYNACI S., DURAK ARAS B., BAŞ H., ÖZDEMİR M., ARTAN S.
Erciyes Medical Genetics Days, Turkey, 11 - 13 May 2017

XXXVII. **The frequencies of CFTR M470V, intron 8 poly-T and pathogenic mutations in cystic fibrosis patients**
ARTAN S., ÇİLİNGİR O., DURMUŞ AYDOĞDU S., ERZURUMLUOĞLU E., KOCAGİL S., DURAK ARAS B., ÖZDEMİR M.
European Human Genetics Conference, 27 - 30 May 2017

XXXVIII. **Beckwith Wiedemann Sendromu: Bir Olgu Sunumu.**
ASLAN H., ÖZDEMİR M., ÇİLİNGİR O., KOCAGİL S., KÜÇÜK H., DURAK ARAS B., ARTAN S.
XII. Ulusal Tıbbi Genetik Kongresi, İzmir, Turkey, 05 October 2016, vol.2, pp.330

Supported Projects

KOCAGİL S., DURAK ARAS B., ÇİLİNGİR O., Project Supported by Higher Education Institutions, NON-SENDROMİK KONJENİTAL KALP HASTALIKLARINDA ADAY GENLERİN DEĞERLENDİRİLMESİ, 2019 - Continues
DURMAZ ÇELİK F. N., KOCAGİL S., DURAK ARAS B., ERZURUMLUOĞLU GÖKALP E., ÖZKAN S., Project Supported by Higher Education Institutions, İdiyopatik Parkinson olgularında C9orf72 geni G4C2 tekrar sayılarının değerlendirilmesi, 2021 - 2023
Arslantaş A., Artan S., Durak Aras B., Özbek Z., Özkara E., Kocagil S., Erzurumluoğlu Gökalp E., Aykaç Ö., Çilingir O., Project Supported by Higher Education Institutions, Intrakraniyal Sakküler Anevrizmalarda Aday Genlerin Değerlendirilmesi, 2018 - 2019

Memberships / Tasks in Scientific Organizations

European Society of Human Genetics, Principal Member, 2023 - Continues, Austria
Tıbbi Genetik Derneği , Member, 2017 - Continues, Turkey

Scientific Refereeing

R&D Project of Group B, Eskisehir Osmangazi University, Turkey, June 2024
R&D Project of Group B, Eskisehir Osmangazi University, Turkey, June 2024
R&D Project of Group B, Eskisehir Osmangazi University, Turkey, June 2024
R&D Project of Group B, Eskisehir Osmangazi University, Turkey, June 2024
Osmangazi Tıp Dergisi, National Scientific Refreed Journal, January 2023

Tasks In Event Organizations

Durak Aras B., Erzurumluoğlu Gökalp E., Artan S., Kocagil S., Çilingir O., 7. Uluslararası Erciyes Tıp Tıbbi Genetik Kongresi, Scientific Congress, Turkey, Mayıs 2022

Metrics

Publication: 62
Citation (WoS): 14
Citation (Scopus): 27
H-Index (WoS): 2
H-Index (Scopus): 3

Congress and Symposium Activities

Tıbbi Genetik Derneği Bahar Okulu, Invited Speaker, Ankara, Turkey, 2024
7. Uluslararası Erciyes Tıp Tıbbi Genetik Kongresi, Invited Speaker, Kayseri, Turkey, 2022

Non Academic Experience

T.C SAĞLIK BAKANLIĞI BODRUM DEVLET HASTANESİ