

Asst. Prof. SİNEM KOCAGİL

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International Researcher IDs

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Education Information

Undergraduate, Anadolu University, Department Of Sociology (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. **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)
- II. **Late Diagnosis of Congenital Chloride Diarrhea Mimicking Hirschsprung's Disease**
Çelik A. T., Barış Z., Aydemir Y., Kocagil S.
CLINICAL PEDIATRICS, 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., Kirel 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., Kiral E., Susam E., Tufan A., YARAR C., Çetin N., Kocagil S., Kirel 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., Sivriköz 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 Sivriköz 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 lncRNAs 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 ONKOLOJİ DERGİSİ, vol.37, pp.305-313, 2022 (ESCI)
- V. **Analyzing The Mutations Of Notch1 And Sf3b1 Genes In Cases With CLL Detected Isolated 13q Deletion**
Gunden 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 Gökalg 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, RFW3 Genlerinde 3 Novel/Nadir Varyant**
Özbakır D. H., Saraç E., Susam E., Kocagil S., Aynacı S., Kirel 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 Gökalg 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., KEKLİKÇİ 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 Chromoanasythesis: 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 disability 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., VELİPAŞAĞLU M., KOCAGİL S., DURAK ARAS B., TEMENA M. A., PANAL G., ANSARI 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., Kiral 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 Recurrent 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 GGGCC 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 ANSARI 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

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Tasks In Event Organizations

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Metrics

Publication: 62

Citation (WoS): 14

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Non Academic Experience

T.C SAĞLIK BAKANLIĞI BODRUM DEVLET HASTANESİ