

## Doç. Dr. GÜLAY CEYLANER

### Kişisel Bilgiler

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### Uluslararası Araştırmacı ID'leri

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### SCI, SSCI ve AHCI İndekslerine Giren Dergilerde Yayımlanan Makaleler

- I. **The outcome of 41 Late-Diagnosed Turkish GA-1 Patients: A Candidate for the Turkish NBS**  
Klavuz S., Bulut D., KOR D., Şeker-Yılmaz B., ÖZCAN N., İNCECİK F., Onan B., CEYLANER G., Önenli-Mungan N.  
Neuropediatrics, cilt.52, sa.5, ss.358-369, 2021 (SCI-Expanded)
- II. **Isolated congenital diaphragm hernia associated with homozygous SLIT3 gene variant in dizygous twins**  
Kaya T. B., AYDEMİR Ö., CEYLANER S., Ceylaner G., TEKİN A. N.  
European Journal of Medical Genetics, cilt.64, sa.7, 2021 (SCI-Expanded)
- III. **Fabry disease screening in patients with kidney transplant: A single-center study in turkey**  
Erdogmus S., KUTLAY S., Kumru G., Sendogan D. O., ERTÜRK Ş., KEVEN K., CEYLANER G., ŞENGÜL Ş.  
Experimental and Clinical Transplantation, cilt.18, sa.4, ss.444-449, 2020 (SCI-Expanded)
- IV. **Investigation of MKRN3 mutation in patients with familial central precocious puberty**  
AYCAN Z., Savaş-Erdeve Ş., Çetinkaya S., Kurnaz E., Keskin M., Şahin N. M., Bayramoğlu E., CEYLANER G.  
JCRPE Journal of Clinical Research in Pediatric Endocrinology, cilt.10, sa.3, ss.223-229, 2018 (SCI-Expanded)
- V. **Congenital Glucose-Galactose Malabsorption in a Turkish Newborn: A Novel Mutation of Na<sup>+</sup>/Glucose Cotransporter Gene**  
Atay F. Y., Derme T., Uras N., CEYLANER G., CEYLANER S., Sari F. N., Oguz S. S.  
Digestive Diseases and Sciences, cilt.62, sa.1, ss.280-281, 2017 (SCI-Expanded)
- VI. **Impaired glucose tolerance in fanconi-bickel syndrome: Eight patients with two novel mutations**  
Şeker-Yılmaz B., Kör D., BULUT F. D., YÜKSEL B., Karabay-Bayazıt A., TOPALOĞLU A. K., CEYLANER G., Önenli-Mungan N.  
Turkish Journal of Pediatrics, cilt.59, sa.4, ss.434-441, 2017 (SCI-Expanded)
- VII. **A novel missense mutation, p.(R102W) in WNT7A causes Al-Awadi Raas-Rothschild syndrome in a fetus**  
Mutlu M. B., Cetinkaya A., Koc N., CEYLANER G., Erguner B., Aydın H., Karaman S., Demirci O., Goksu K., Karaman A.  
European Journal of Medical Genetics, cilt.59, sa.11, ss.604-606, 2016 (SCI-Expanded)
- VIII. **Genotypic and phenotypic features of the cystinosis patients from the South Eastern part of Turkey**  
Önenli-Mungan N., KOR D., Karabay-Bayazıt A., Cengiz N., Yavuz S., Noyan A., CEYLANER G., Şeker-Yılmaz B., TOPALOĞLU A. K., YÜKSEL B., et al.  
Turkish Journal of Pediatrics, cilt.58, sa.4, ss.362-370, 2016 (SCI-Expanded)
- IX. **Ring chromosome 9 in a newborn**  
Aldemir O., Celik I., Karaer K., CEYLANER G.  
Genetic Counseling, cilt.24, sa.4, ss.357-360, 2013 (SCI-Expanded)
- X. **Prenatal diagnosis of goldenhar syndrome with unusual features by 3D ultrasonography**  
Guzelmansur I., CEYLANER G., CEYLANER S., CEYLAN N., Daplan T.  
Genetic Counseling, cilt.24, sa.3, ss.319-325, 2013 (SCI-Expanded)

- XI. **Nephron-sparing surgery for renal cell carcinoma of the allograft after renal transplantation: Report of two cases**  
TÜZÜNER A., Çakir F., AKYOL C., Çelebi Z., CEYLANER S., CEYLANER G., ŞENGÜL Ş., KEVEN K.  
Transplantation Proceedings, cilt.45, sa.3, ss.958-960, 2013 (SCI-Expanded)
- XII. **A new family with autosomal dominant porencephaly with a novel COL4A1 mutation. Are arachnoid cysts related to COL4A1 mutations?**  
Değerliyurt A., CEYLANER G., Koçak H., Bilginer Gürbüz B., Cihan B., Rizzu P., CEYLANER S.  
Genetic Counseling, cilt.23, sa.2, ss.185-193, 2012 (SCI-Expanded)
- XIII. **High frequency of p.Thr93Met in Smith-Lemli-Opitz syndrome patients in Turkey**  
Kalb S., ÇAĞLAYAN A. O., Degerliyurt A., Schmid S., CEYLANER S., HATİPOĞLU N., Hinderhofer K., Rehder H., Kurtoglu S., CEYLANER G., et al.  
Clinical Genetics, cilt.81, sa.6, ss.598-601, 2012 (SCI-Expanded)
- XIV. **Ovulation induction with gonadotropins causes increased sister chromatid exchanges**  
Tonguç E., CEYLANER G., Var T., Zülfikaroğlu E., CEYLANER S.  
Genetic Counseling, cilt.22, sa.2, ss.193-198, 2011 (SCI-Expanded)
- XV. **A novel mutation in the RPS6KA3 gene in a patient with Coffin-Lowry syndrome**  
Senel S., CEYLANER S., CEYLANER G., Hanli Sahin A., Andrieux J., Delaunoy J.  
Genetic Counseling, cilt.22, sa.1, ss.21-24, 2011 (SCI-Expanded)
- XVI. **Vascular endothelial growth factor +405 C/G polymorphism is highly associated with an increased risk of endometriosis in Turkish women**  
Altinkaya S. O., Ugur M., CEYLANER G., Ozat M., Gungor T., CEYLANER S.  
Archives of Gynecology and Obstetrics, cilt.283, sa.2, ss.267-272, 2011 (SCI-Expanded)
- XVII. **Partial trisomy due to a de novo duplication 22q11.1-22q13.1: A cat-eye syndrome variant with brain anomalies**  
Karcaaltincaba D., CEYLANER S., CEYLANER G., Dalkilic S., Karli-Oguz K., Kandemir O.  
Genetic Counseling, cilt.21, sa.1, ss.19-24, 2010 (SCI-Expanded)
- XVIII. **Genetic abnormalities in Turkish women with premature ovarian failure**  
CEYLANER G., Altinkaya S. O., Mollamahmutoglu L., CEYLANER S.  
International Journal of Gynecology and Obstetrics, cilt.110, sa.2, ss.122-124, 2010 (SCI-Expanded)
- XIX. **Familial primary carpal tunnel syndrome with possible skipped generation.**  
Senel S., CEYLANER G., Yuksel D., Erkek N., KARACAN C. D.  
European journal of pediatrics, cilt.169, sa.4, ss.453-455, 2010 (SCI-Expanded)
- XX. **Mitral valve prolapse in two siblings with pyknodysostosis**  
Senel S., KARACAN C. D., Orun U., Erkek N., CEYLANER G.  
Genetic Counseling, cilt.20, sa.4, ss.397-401, 2009 (SCI-Expanded)
- XXI. **Prenatal diagnosis of a case with anencephaly-omphalocele-unilateral absent radial ray**  
CEYLANER S., CEYLANER G., Altun M., Coşkun A., Danişman N.  
Genetic Counseling, cilt.20, sa.2, ss.189-193, 2009 (SCI-Expanded)
- XXII. **Prenatal diagnosis of a case probably with oral-facial-digital syndrome - Gabrielli type**  
Güven M., CEYLANER G., CEYLANER S., ÜZEL M., Coskun A.  
Genetic Counseling, cilt.20, sa.2, ss.167-172, 2009 (SCI-Expanded)
- XXIII. **A boy with trisomy 13 presenting with a subtle clinical picture and metopic synostosis**  
ÜNAL S., Celik F. C., Soy D., CEYLANER S., Ceylaner G.  
American Journal of Medical Genetics, Part A, cilt.149, sa.7, ss.1608-1609, 2009 (SCI-Expanded)
- XXIV. **Frontonasal dysplasia: A family presenting autosomal dominant inheritance pattern**  
Koçak H., CEYLANER G.  
Genetic Counseling, cilt.20, sa.1, ss.63-68, 2009 (SCI-Expanded)
- XXV. **A prenatally diagnosed case of sirenomelia with polydactyly and vestigial tail**  
Güven M., ÜZEL M., CEYLANER S., Coskun A., CEYLANER G., Gungoren A.  
Genetic Counseling, cilt.19, sa.4, ss.419-424, 2008 (SCI-Expanded)
- XXVI. **Azoospermia and cryptorchidism in a male with a de novo reciprocal t(Y;16) translocation**

- Gunel M., Cavkaytar S., CEYLANER G., Batioglu S.  
Genetic Counseling, cilt.19, sa.3, ss.277-280, 2008 (SCI-Expanded)
- XXVII. **Neu-Laxova syndrome in an appropriate for gestational age newborn**  
Dilli D., Yaşar H., Dilmen U., CEYLANER G.  
Indian Journal of Dermatology, Venereology and Leprology, cilt.74, sa.5, ss.487-489, 2008 (SCI-Expanded)
- XXVIII. **GJB2 and mitochondrial A1555G gene mutations in nonsyndromic profound hearing loss and carrier frequencies in healthy individuals**  
Baysal E., Bayazit Y., CEYLANER S., Alatas N., Donmez B., CEYLANER G., San I., Korkmaz B., Yilmaz A., Menevse A., et al.  
Journal of Genetics, cilt.87, sa.1, ss.53-57, 2008 (SCI-Expanded)
- XXIX. **Autosomal dominant inheritance of congenital dislocation of the hip in 16 members of a family Bir ailenin 16 üyesinde otozomal dominant kalıtım gösteren gelişimsel kalça çıkığı**  
CEYLANER G., CEYLANER S., Ustunkan F., Inan M.  
Acta Orthopaedica et Traumatologica Turcica, cilt.42, sa.4, ss.289-291, 2008 (SCI-Expanded)
- XXX. **Chromosomal heteromorphisms may help for the diagnosis of uniparental disomy (UPD): A case report**  
CEYLANER G., CEYLANER S., Danişman N., Ergün A., Ekici E., Schinzel A., Baumer A.  
Prenatal Diagnosis, cilt.27, sa.11, ss.1072-1074, 2007 (SCI-Expanded)
- XXXI. **Evaluation of 2407 fetuses in a Turkish population**  
CEYLANER G., CEYLANER S., Günyeli I., Ekici E., Celasun B., Danişman N.  
Prenatal Diagnosis, cilt.27, sa.9, ss.800-807, 2007 (SCI-Expanded)
- XXXII. **Prenatal diagnosis of a Turkish Bartsocas-Papas syndrome case with upper limb pterigia**  
CEYLANER G., Güven M. A., CEYLANER S., ÜZEL M., Müftüoğlu K.  
Prenatal Diagnosis, cilt.27, sa.6, ss.563-565, 2007 (SCI-Expanded)
- XXXIII. **Prenatal diagnosis of Meckel Gruber syndrome presenting with renal agenesis: report of a case.**  
Güven M., CEYLANER S., CEYLANER G., Gul D., Ertas L.  
Genetic counseling (Geneva, Switzerland), cilt.17, sa.1, ss.65-68, 2006 (SCI-Expanded)
- XXXIV. **Extremely skewed X-chromosome inactivation patterns in women with recurrent spontaneous abortion**  
Bagıslar S., Ustuner I., Cengiz B., Soylemez F., Akyerli C. B., CEYLANER S., CEYLANER G., Acar A., Ozcelik T.  
Australian and New Zealand Journal of Obstetrics and Gynaecology, cilt.46, sa.5, ss.384-387, 2006 (SCI-Expanded)
- XXXV. **An infant with diaphragmatic hernia, anophthalmia and cardiac defect: Evaluation by magnetic resonance imaging autopsy**  
CEYLANER S., Gozer H., CEYLANER G., Ertas I., Kizilates S., Edguer T.  
Genetic Counseling, cilt.17, sa.2, ss.231-236, 2006 (SCI-Expanded)
- XXXVI. **Postmortem evaluation of 220 prenatally diagnosed fetuses with neural tube defects: Detection of associated anomalies in a Turkish population**  
CEYLANER S., CEYLANER G., Günyeli I., Ekici E., Tuğ M., Taner D., Ekerbicer H., Mollamahmutoğlu L., Danişman N.  
Prenatal Diagnosis, cilt.26, sa.2, ss.147-153, 2006 (SCI-Expanded)
- XXXVII. **A case of fetal anticonvulsant syndrome with severe bilateral upper limb defect**  
Güven M. A., Batukan C., CEYLANER S., CEYLANER G., ÜZEL M.  
Journal of Maternal-Fetal and Neonatal Medicine, cilt.19, sa.2, ss.115-117, 2006 (SCI-Expanded)
- XXXVIII. **Volume of sampled amniotic fluid and prenatal cytogenetic diagnosis**  
Güven M., CEYLANER G., Coskun A.  
International Journal of Gynecology and Obstetrics, cilt.95, sa.2, ss.157-158, 2006 (SCI-Expanded)
- XXXIX. **Craniosynostosis and ectopia lentis in a propositus whose parents are cousins [4]**  
Güven D., Kalayci D., Hasiripi H., CEYLANER S., CEYLANER G.  
American Journal of Medical Genetics, cilt.134 A, sa.2, ss.231, 2005 (SCI-Expanded)
- XL. **A novel mutation in the lysyl hydroxylase 1 gene causes decreased lysyl hydroxylase activity in an Ehlers-danlos VIA patient**  
Walker L. C., Overstreet M. A., Siddiqui A., De Paepe A., CEYLANER G., Malfait F., Symoens S., Atsawasuwan P.,

## **Diğer Dergilerde Yayınlanan Makaleler**

- I. **New mutation identified in the SRY gene high mobility group (HMG) Sry geni high mobility group (HMG) bölgesinde tanımlanan yeni mutasyon**  
Şahin F. i., Esi S., CEYLANER G., Emin E. K., Enver O. Ö., Terzi Y. K., Çelik Z. Y.  
Turk Jinekoloji ve Obstetrik Dernegi Dergisi, cilt.10, sa.2, ss.118-121, 2013 (ESCI)
- II. **A child with XYY karyotype and epilepsy**  
Değerliyurt A., CEYLANER G., CEYLANER S.  
Journal of Pediatric Neurology, cilt.9, sa.2, ss.255-258, 2011 (ESCI)
- III. **Prenatally diagnosed partial trisomy 3q case with an omphalocele and less severe phenotype Prenatal tanısı konulmuş, omfalosel ve hafif fenotipik anormalliklere sahip kısmi trizomi 3q olgusu**  
Arikan D. C., Coşkun A., Arikan I., Kiran G., CEYLANER G.  
Journal of the Turkish German Gynecology Association, cilt.11, sa.4, ss.228-232, 2010 (ESCI)
- IV. **A prenatally diagnosed pentology of cantrell case with encephalocele: A rare variant Prenatal tanısı konmuş Cantrell Pentalojisi olgusu: Ensefaloselin eşlik ettiği nadir bir varyant**  
Güven M. A., CEYLANER G., CEYLANER S., Coşkun A., Bayazit H.  
Turk Jinekoloji ve Obstetrik Dernegi Dergisi, cilt.6, sa.2, ss.123-127, 2009 (Scopus)
- V. **A case with dextrocardia and multiple cardiac anomalies in a family with congenital heart malformations**  
CEYLANER S., Güven M. A., CEYLANER G., Çiragil G., Tuğ M., Ertaş I. E.  
Journal of the Turkish German Gynecology Association, cilt.6, sa.2, ss.158-160, 2005 (ESCI)

## **Metrikler**

Yayın: 45

Atf (WoS): 237

Atf (Scopus): 268

H-İndeks (WoS): 9

H-İndeks (Scopus): 10