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SCI, SSCI ve AHCI İndekslerine Giren Dergilerde Yayınlanan Makaleler

I. **The outcome of 41 Late-Diagnosed Turkish GA-1 Patients: A Candidate for the Turkish NBS**
Kllavuz S., Bulut D., KOR D., Şeker-Yllmaz 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.
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III. **Fabry disease screening in patients with kidney transplant: A single-center study in turkey**
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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+/Glucose Cotransporter Gene**
Atay F. Y., Derme T., Uras N., CEYLANER G., CEYLANER S., Sari F. N., Oguz S. S.
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VI. **Impaired glucose tolerance in fanconi-bickel syndrome: Eight patients with two novel mutations**
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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.
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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.
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X. **Prenatal diagnosis of goldenhar syndrome with unusual features by 3D ultrasonography**
Guzelmansur I., CEYLANER G., CEYLANER S., CEYLAN N., Daplan T.

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XII. **A new family with autosomal dominant porencephaly with a novel COL4A1 mutation. Are arachnoid cysts related to COL4A1 mutations?**
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XIV. **Ovulation induction with gonadotropins causes increased sister chromatid exchanges**
Tonguç E., CEYLANER G., Var T., Zülfikaroğlu E., CEYLANER S.
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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.
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XVI. **Vascular endothelial growth factor +405 C/G polymorphism is highly associated with an increased risk of endometriosis in Turkish women**
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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., Kandemr O.
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XVIII. **Genetic abnormalities in Turkish women with premature ovarian failure**
CEYLANER G., Altinkaya S. O., Mollamahmutoglu L., CEYLANER S.
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XIX. **Familial primary carpal tunnel syndrome with possible skipped generation.**
Senel S., CEYLANER G., Yuksel D., Erkek N., KARACAN C. D.
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XX. **Mitral valve prolapse in two siblings with pyknodysostosis**
Senel S., KARACAN C. D., Orun U., Erkek N., CEYLANER G.
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XXI. **Prenatal diagnosis of a case with anencephaly-omphalocele-unilateral absent radial ray**
CEYLANER S., CEYLANER G., Altun M., Coşkun A., Danişman N.
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XXII. **Prenatal diagnosis of a case probably with oral-facial-digital syndrome - Gabrielli type**
Güven M., CEYLANER G., CEYLANER S., ÜZEL M., Coskun A.
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XXIII. **A boy with trisomy 13 presenting with a subtle clinical picture and metopic synostosis**
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XXIV. **Frontonasal dysplasia: A family presenting autosomal dominant inheritance pattern**
Koçak H., CEYLANER G.
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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.
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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.
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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.
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XXX. **Chromosomal heteromorphisms may help for the diagnosis of uniparental disomy (UPD): A case report**
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XXXI. **Evaluation of 2407 fetuses in a Turkish population**
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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.
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XXXIII. **Prenatal diagnosis of Meckel Gruber syndrome presenting with renal agenesis: report of a case.**
Guven M., CEYLANER S., CEYLANER G., Gul D., Ertas L.
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XXXIV. **Extremely skewed X-chromosome inactivation patterns in women with recurrent spontaneous abortion**
Bagislar 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.
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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.
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XXXVII. **A case of fetal anticonvulsant syndrome with severe bilateral upper limb defect**
Guven 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**
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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., Yamauchi M., CEYLANER S., et al.
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Diğer Dergilerde Yayınlanan Makaleler

I. **New mutation identified in the SRY gene high mobility group (HMG) Sry genı high mobility group (HMG) bölges ınde tanimlanan yenı mutasyon**
Şahin F. i., Esi S., CEYLANER G., Emin E. K., Enver O. Ö., Terzi Y. K., ÇeliK Z. Y.
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II. **A child with XYY karyotype and epilepsy**
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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.
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IV. **A prenatally diagnosed pentalogy of cantrell case with encephaloce: A rare variant Prenatal tanisi konmuş Cantrell Pentalojisiolgusu: Ensefaloselin eşlik ettiğinadir bir varyant**
Güven M. A., CEYLANER G., CEYLANER S., Coşkun A., Bayazit H.
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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.
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Metrikler

Yayın: 45
Atıf (WoS): 237
Atıf (Scopus): 268
H-İndeks (WoS): 9
H-İndeks (Scopus): 10