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Published journal articles indexed by SCI, SSCI, and AHCI

- 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, vol.52, no.5, pp.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, vol.64, no.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, vol.18, no.4, pp.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.
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- 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**
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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-Bayazit A., Cengiz N., Yavuz S., Noyan A., CEYLANER G., Şeker-Yılmaz B., TOPALOĞLU A. K., YÜKSEL B., et al.
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- IX. **Ring chromosome 9 in a newborn**
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Genetic Counseling, vol.24, no.4, pp.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.
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- XI. **Nephron-sparing surgery for renal cell carcinoma of the allograft after renal transplantation: Report of two cases**
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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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- XIII. **High frequency of p.Thr93Met in Smith-Lemli-Opitz syndrome patients in Turkey**
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- XIV. **Ovulation induction with gonadotropins causes increased sister chromatid exchanges**
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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**
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- XVIII. **Genetic abnormalities in Turkish women with premature ovarian failure**
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- XIX. **Familial primary carpal tunnel syndrome with possible skipped generation.**
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- XX. **Mitral valve prolapse in two siblings with pyknodysostosis**
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- XXI. **Prenatal diagnosis of a case with anencephaly-omphalocele-unilateral absent radial ray**
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- XXII. **Prenatal diagnosis of a case probably with oral-facial-digital syndrome - Gabrielli type**
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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**
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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.
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- XXVII. **Neu-Laxova syndrome in an appropriate for gestational age newborn**
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- XXVIII. **GJB2 and mitochondrial A1555G gene mutations in nonsyndromic profound hearing loss and carrier frequencies in healthy individuals**
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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**
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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**
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- XXXIII. **Prenatal diagnosis of Meckel Gruber syndrome presenting with renal agenesis: report of a case.**
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- XXXIV. **Extremely skewed X-chromosome inactivation patterns in women with recurrent spontaneous abortion**
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- XXXV. **An infant with diaphragmatic hernia, anophthalmia and cardiac defect: Evaluation by magnetic resonance imaging autopsy**
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- XXXVII. **A case of fetal anticonvulsant syndrome with severe bilateral upper limb defect**
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- 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]**
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- XL. **A novel mutation in the lysyl hydroxylase 1 gene causes decreased lysyl hydroxylase activity in an Ehlers-danlos VIA patient**
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Articles Published in Other Journals

- I. **New mutation identified in the SRY gene high mobility group (HMG) Sry geni high mobility group (HMG) bölgesinde tanımlanan yeni mutasyon**
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Journal of Pediatric Neurology, vol.9, no.2, pp.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 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.
Türk Jinekoloji ve Obstetrik Derneği Dergisi, vol.6, no.2, pp.123-127, 2009 (Scopus)
- V. **A case with dextrocardia and multiple cardiac anomalies in a family with congenital heart malformations**
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