

# Prof. DİLEK YALNIZOĞLU

## Personal Information

Office Phone: [+90 312 305 1185](tel:+903123051185)

Office Phone: [+90 312 305 1165](tel:+903123051165)

Email: [dileky@hacettepe.edu.tr](mailto:dileky@hacettepe.edu.tr)

Web: <https://avesis.hacettepe.edu.tr/dileky>

## International Researcher IDs

ORCID: 0000-0001-9294-4305

Yoksis Researcher ID: 169372

## Education Information

Expertise In Medicine, Harvard University, Hms/Children's Hospital Boston, Klinik Nörofizyoloji Ve Epilepsi/Nöroloji, United States Of America 1997 - 1999

Expertise In Medicine, Hacettepe University, Tıp Fakültesi/Çocuk Sağlığı Ve Hastalıkları, Çocuk Nörolojisi, Turkey 1994 - 1997

Expertise In Medicine, Hacettepe University, Tıpta Uzmanlık, Çocuk Sağlığı Ve Hastalıkları, Turkey 1990 - 1994

Undergraduate, Ankara University, Tıp Fakültesi, Turkey 1983 - 1989

## Foreign Languages

English, C1 Advanced

## Research Areas

Medicine, Health Sciences, Internal Medicine Sciences, Child Health and Diseases, Pediatric Neurology

## Academic Titles / Tasks

Professor, Hacettepe University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2011 - Continues

Associate Professor, Hacettepe University, Tıp Fakültesi (İngilizce), Dahili Tıp Bilimleri Bölümü, 2005 - 2011

Assistant Professor, Hacettepe University, Tıp Fakültesi (İngilizce), Dahili Tıp Bilimleri Bölümü, 1999 - 2005

## Published journal articles indexed by SCI, SSCI, and AHCI

- Exploring metabolic alterations in PYCR2 deficiency: Unveiling pathways and clinical presentations of hypomyelinating leukodystrophy 10**  
Gurbuz B. B., GÜLBAKAN B., ÖZGÜL R. K., YALNIZOĞLU D., Yılmaz D. Y., GÖÇMEN R., KOŞUKCU C., KANDEMİR N., Acar N. V., SALİH B., et al.  
AMERICAN JOURNAL OF MEDICAL GENETICS PART A, no.9, 2024 (SCI-Expanded)
- Frequency and Characteristics of Sleep Apnea Syndrome in Pediatric Patients with Duchenne Muscular Dystrophy**

Yavuz B. C., Yildiz S. O., Buyuksahin H. N., ÖZEL E., EMİRALİOĞLU ORDUKAYA N., AYKAN H. H., Yalcin E., DOĞRU ERSÖZ D., KİPER E. N., ÖZTOPRAK Ü., et al.

EUROPEAN RESPIRATORY JOURNAL, 2024 (SCI-Expanded)

- III. **Predictors of Clinically Important Neuroimaging Findings in Children Presenting Pediatric Emergency Department**  
GÜNGÖR E., Haliloglu G., YALNIZOĞLU D., Oguz K. K., TEKŞAM Ö.  
PEDIATRIC EMERGENCY CARE, no.6, pp.474-479, 2024 (SCI-Expanded)
- IV. **Delineation of ADPRHL2 Variants: Report of Two New Patients with Review of the Literature**  
Yildiz S. O., Yalnizoglu D., ŞİMŞEK KİPER P. Ö., GÖÇMEN R., Sogukpinar M., Utine G. E., Haliloglu G.  
NEUROPEDIATRICS, no.02, pp.117-123, 2024 (SCI-Expanded)
- V. **Electro-clinical features and long-term outcomes in guanidinoacetate methyltransferase (GAMT) deficiency**  
YILDIZ Y., Ardiçlı D., GÖÇMEN R., YALNIZOĞLU D., Topçu M., Coşkun T., TOKATLI A., Haliloğlu G.  
European Journal of Paediatric Neurology, vol.49, pp.66-72, 2024 (SCI-Expanded)
- VI. **Non-epileptic paroxysmal events at pediatric video-electroencephalography monitoring unit over a 15-year period**  
Yavuz P., GÜNBEY C., KARAHAN S., TOPÇU M., Turanlı G., YALNIZOĞLU D.  
Seizure, vol.108, pp.89-95, 2023 (SCI-Expanded)
- VII. **Claude Syndrome in Childhood Associated with Probable Neuro-Behcet Disease**  
Yavuz P., SOLMAZ İ., Kaya U. A., Akgoz A., Oguz K. K., Aytac S., ÖZEN S., YALNIZOĞLU D.  
Neuropediatrics, vol.54, no.1, pp.82-87, 2023 (SCI-Expanded)
- VIII. **Clinically important intracranial abnormalities in children presenting with first focal seizure**  
Kasap T., TEKŞAM Ö., Turanlı G., KONUŞKAN B., Oğuz K. K., Haliloğlu G., YALNIZOĞLU D.  
Turkish Journal of Pediatrics, vol.65, no.1, pp.96-108, 2023 (SCI-Expanded)
- IX. **Practices of pediatric emergency physicians on the first febrile and afebrile seizures: a research in European Pediatric Emergency Medicine Survey Study**  
TEKŞAM Ö., Serdaroglu E., Haliloglu G., KONUŞKAN B., YALNIZOĞLU D.  
EUROPEAN JOURNAL OF EMERGENCY MEDICINE, vol.29, no.6, pp.455-457, 2022 (SCI-Expanded)
- X. **Neurologic manifestations in children with COVID-19**  
Gürlevik S. L., GÜNBEY C., ÖZSÜREKÇİ Y., KESİCİ S., GÖÇMEN R., TEMUÇİN Ç. M., ÖZEN S., CENGİZ A. B., YALNIZOĞLU D.  
European Journal of Paediatric Neurology, vol.39, pp.118-119, 2022 (SCI-Expanded)
- XI. **Synthetic MRI in children with tuberous sclerosis complex**  
ÇOBAN ÇİFÇİ G., GÜMELER E., PARLAK SAĞOL Ş., KONUŞKAN B., KARAKAYA KARABULUT J., YALNIZOĞLU D., Anlar B., Oguz K. K.  
INSIGHTS INTO IMAGING, vol.13, no.1, 2022 (SCI-Expanded)
- XII. **Evaluation of changes in physician behavior after introduction of pediatric syncope approach protocol in the emergency department**  
Yildiz L. A., Haliloglu G., YALNIZOĞLU D., ERTUĞRUL İ., ALEHAN D., TEKŞAM Ö.  
AMERICAN JOURNAL OF EMERGENCY MEDICINE, vol.55, pp.57-63, 2022 (SCI-Expanded)
- XIII. **Lesional resective epilepsy surgery in childhood: Comparison of two decades and long-term seizure outcome from a single center**  
GÜNBEY C., BİLGİNER B., Oguz K. K., Soylemezoglu F., Ergun E. L., AKALAN N., TOPÇU M., Turanlı G., YALNIZOĞLU D.  
EPILEPSY RESEARCH, vol.181, 2022 (SCI-Expanded)
- XIV. **Comparison of physical fitness, activity, and quality of life of the children with epilepsy and their healthy peers**  
SIRTBAŞ G., YALNIZOĞLU D., LİVANELİOĞLU A.  
Epilepsy Research, vol.178, 2021 (SCI-Expanded)
- XV. **Biallelic mutations in ELFN1 gene associated with developmental and epileptic encephalopathy and joint laxity**

DURŞUN A., YALNIZOĐLU D., Yılmaz D. Y., Oğuz K. K., GÜLBAKAN B., KOŞUKCU C., AKAR H. T., Kahraman A. B., Acar N. V., GÜNBEY C., et al.

EUROPEAN JOURNAL OF MEDICAL GENETICS, vol.64, no.11, 2021 (SCI-Expanded)

- XVI. **Long-term effects of vagus nerve stimulation in refractory pediatric epilepsy: A single-center experience**  
YALNIZOĐLU D., ARDIĐLI D., BİLGİNER B., KONUŞKAN B., Oğuz K. K., AKALAN N., Turanlı G., SAYGI S., TOPÇU M.  
EPILEPSY & BEHAVIOR, vol.110, 2020 (SCI-Expanded)
- XVII. **The training and organization of Paediatric Neurology in Europe: Special report of the European Paediatric Neurology Society & Committee of National Advisors**  
Craiu D., Haataja L., Hollody K., Krsek P., Lagae L., Mall V., Parker A. P. J., Steinlin M., YALNIZOĐLU D., Catsman-Berrevoets C.  
EUROPEAN JOURNAL OF PAEDIATRIC NEUROLOGY, vol.28, pp.6-15, 2020 (SCI-Expanded)
- XVIII. **Acute Cerebellitis or Postinfectious Cerebellar Ataxia? Clinical and Imaging Features in Acute Cerebellitis**  
Yıldırım M., GÖÇMEN R., KONUŞKAN B., PARLAK Ş., YALNIZOĐLU D., Anlar B.  
JOURNAL OF CHILD NEUROLOGY, vol.35, no.6, pp.380-388, 2020 (SCI-Expanded)
- XIX. **Risk factors for seizure recurrence in a pediatric observation unit**  
GÜLTEKİNGİL KESER A., TEKŞAM Ö., Halilolu G., YALNIZOĐLU D.  
AMERICAN JOURNAL OF EMERGENCY MEDICINE, vol.37, no.12, pp.2151-2154, 2019 (SCI-Expanded)
- XX. **Clinical highlights of a very rare phospholipid remodeling disease due to MBOAT7 gene defect**  
DURŞUN A., YALNIZOĐLU D., ÖZGÜL R. K., Oğuz K. K., Yucel-Yılmaz D.  
AMERICAN JOURNAL OF MEDICAL GENETICS PART B-NEUROPSYCHIATRIC GENETICS, 2019 (SCI-Expanded)
- XXI. **Epilepsy in neurofibromatosis type 1: Diffuse cerebral dysfunction?**  
Serdarolu E., KONUŞKAN B., Oğuz K. K., Gurler G., YALNIZOĐLU D., Anlar B.  
EPILEPSY & BEHAVIOR, vol.98, pp.6-9, 2019 (SCI-Expanded)
- XXII. **Expanding the phenotype of phospholipid remodelling disease due to MBOAT7 gene defect**  
YALNIZOĐLU D., ÖZGÜL R. K., Oğuz K. K., Ozer B., Yucel-Yılmaz D., Gurbuz B., Serdarolu E., Erol I., Topcu M., DURŞUN A.  
JOURNAL OF INHERITED METABOLIC DISEASE, vol.42, no.2, pp.381-388, 2019 (SCI-Expanded)
- XXIII. **Behavioral problems of preschool children with new-onset epilepsy and one-year follow-up - A prospective study**  
Yoldas T. C., GÜNBEY C., Degerliyurt A., Erol N., Ozmert E., YALNIZOĐLU D.  
EPILEPSY & BEHAVIOR, vol.92, pp.171-175, 2019 (SCI-Expanded)
- XXIV. **Response to Early Coenzyme Q10 Supplementation Is not Sustained in CoQ10 Deficiency Caused by CoQ2 Mutation**  
Eroglu F. K., ÖZALTIN F., Gonc N., Nalcaciolu H., Ozcakar Z. B., YALNIZOĐLU D., Gucer S., ORHAN D., Eminolu F. T., GÖÇMEN R., et al.  
PEDIATRIC NEUROLOGY, vol.88, pp.71-74, 2018 (SCI-Expanded)
- XXV. **Clinical phenotype of hereditary spastic paraplegia due to KIF1C gene mutations across life span**  
Yucel-Yılmaz D., Yucesan E., YALNIZOĐLU D., Oğuz K. K., Sagiroglu M. S., Ozbek U., Serdarolu E., Bilgic B., Erdem S., Iseri S. A. U., et al.  
BRAIN & DEVELOPMENT, vol.40, no.6, pp.458-464, 2018 (SCI-Expanded)
- XXVI. **Electroencephalographic findings in anti-N-methyl-D-aspartate receptor encephalitis in children: A series of 12 patients**  
YILDIRIM M., KONUŞKAN B., YALNIZOĐLU D., Topalolu H., Erol I., Anlar B.  
EPILEPSY & BEHAVIOR, vol.78, pp.118-123, 2018 (SCI-Expanded)
- XXVII. **A probable new syndrome with the storage disease phenotype caused by the VPS33A gene mutation**  
DURŞUN A., YALNIZOĐLU D., Gerdan O. F., Yucel-Yılmaz D., Sagiroglu M. S., YUKSEL B., GUCER S., Sivri S., ÖZGÜL R. K.  
CLINICAL DYSMORPHOLOGY, vol.26, no.1, pp.1-12, 2017 (SCI-Expanded)
- XXVIII. **The seizure semiology consistent with frontal lobe symptomatogenic zone in children**

- Oztoprak Ü., Yalnizoğlu D., Oguz K. K., Ergün E., Soylemezoglu F., Bilginer B., Akalan N., Topçu M., Turanli G.  
TURKISH JOURNAL OF PEDIATRICS, vol.58, no.6, pp.583-591, 2016 (SCI-Expanded)
- XXIX. **Cerebral Hyperperfusion in a Child with Stroke-Like Migraine Attacks after Radiation Therapy Syndrome**  
Ardicli D., GÖÇMEN R., Oguz K. K., VARAN A., YALNIZOĞLU D.  
NEUROPEDIATRICS, vol.47, no.4, pp.259-262, 2016 (SCI-Expanded)
- XXX. **SPECT-PET in Epilepsy and Clinical Approach in Evaluation**  
ERGUN E. L., SAYGI S., YALNIZOĞLU D., OGUZ K. K., Erbas B.  
SEMINARS IN NUCLEAR MEDICINE, vol.46, no.4, pp.294-307, 2016 (SCI-Expanded)
- XXXI. **Acute Abducens Nerve Paralysis in the Pediatric Emergency Department Analysis of 14 Patients**  
TEKŞAM Ö., GÜLTEKİNGİL KESER A., KONUŞKAN B., Haliloglu G., Oguz K. K., YALNIZOĞLU D.  
PEDIATRIC EMERGENCY CARE, vol.32, no.5, pp.307-311, 2016 (SCI-Expanded)
- XXXII. **Evaluation of central nervous system in patients with glycogen storage disease type 1a**  
Aydemir Y., Gurakan F., Temizel İ. N., DEMİR H., Oguz K. K., YALNIZOĞLU D., Topcu M., ÖZEN H., YÜCE A.  
TURKISH JOURNAL OF PEDIATRICS, vol.58, no.1, pp.12-18, 2016 (SCI-Expanded)
- XXXIII. **Mesenchymal stem cell application in children with subacute sclerosing panencephalitis**  
KUSKONMAZ B. B., UCKAN D., YALNIZOĞLU D., GUENEL M., OGUZ K. K., Konuskan B., ANLAR B.  
DEVELOPMENTAL MEDICINE AND CHILD NEUROLOGY, vol.57, no.9, pp.880-883, 2015 (SCI-Expanded)
- XXXIV. **Childhood epilepsy with occipital paroxysm: classification, atypical evolution and long-term prognosis in 35 patients**  
Aksoy A., Haliloglu G., YALNIZOĞLU D., Turanli G.  
TURKISH JOURNAL OF PEDIATRICS, vol.57, no.5, pp.439-452, 2015 (SCI-Expanded)
- XXXV. **Intelligence quotient improves after antiepileptic drug withdrawal following pediatric epilepsy surgery**  
Boshuisen K., van Schooneveld M. M. J., Uiterwaal C. S. P. M., Cross J. H., Harrison S., Polster T., Daehn M., Djimjadi S., YALNIZOĞLU D., Turanli G., et al.  
ANNALS OF NEUROLOGY, vol.78, no.1, pp.104-114, 2015 (SCI-Expanded)
- XXXVI. **Electrical status epilepticus during sleep: A study of 22 patients**  
Degerliyurt A., YALNIZOĞLU D., Bakar E. E., TOPÇU M., Turanli G.  
BRAIN & DEVELOPMENT, vol.37, no.2, pp.250-264, 2015 (SCI-Expanded)
- XXXVII. **Semiological seizure classification of epileptic seizures in children admitted to video-EEG monitoring unit**  
Alan S., YALNIZOĞLU D., Turanli G., Karli-Oguz K., Lay-Ergun E., Soylemezoglu F., AKALAN N., TOPÇU M.  
TURKISH JOURNAL OF PEDIATRICS, vol.57, no.4, pp.317-323, 2015 (SCI-Expanded)
- XXXVIII. **Human intracellular ISG15 prevents interferon-alpha/beta over-amplification and auto-inflammation**  
Zhang X., Bogunovic D., Payelle-Brogard B., Francois-Newton V., Speer S. D., Yuan C., Volpi S., Li Z., Sanal O., Mansouri D., et al.  
NATURE, vol.517, no.7532, pp.89-103, 2015 (SCI-Expanded)
- XXXIX. **Two Turkish siblings with MEGDEL syndrome due to novel SERAC1 genemutation.**  
ÜNAL Ö., ÖZGÜL R. K., YÜCEL YILMAZ D., YALNIZOĞLU D., TOKATLI A., SİVRİ H. S., HİŞMİ B., COŞKUN T., DURSUN A.  
Turkish Journal Of Pediatrics, vol.57, pp.388-393, 2015 (SCI-Expanded)
- XL. **Successful Treatment of Severe Myasthenia Gravis Developed After Allogeneic Hematopoietic Stem Cell Transplantation With Plasma Exchange and Rituximab**  
ÜNAL Ş., SAĞ E., KUSKONMAZ B. B., Kesici S., BAYRAKÇI B., AYVAZ D. N., TEZCAN I., YALNIZOĞLU D., UCKAN D.  
PEDIATRIC BLOOD & CANCER, vol.61, no.5, pp.928-930, 2014 (SCI-Expanded)
- XLI. **Usefulness of long-term video-EEG monitoring in children at a tertiary care center**  
Onay S., YALNIZOĞLU D., TOPÇU M., Turanli G.  
TURKISH JOURNAL OF PEDIATRICS, vol.55, no.6, pp.591-597, 2013 (SCI-Expanded)
- XLII. **Developmental abnormalities and mental retardation: diagnostic strategy**  
TOPÇU M., YALNIZOĞLU D.  
PEDIATRIC NEUROLOGY, PT I, vol.111, pp.211-217, 2013 (SCI-Expanded)

- XLIII. **Identification of a Novel Twinkle Mutation in a Family With Infantile Onset Spinocerebellar Ataxia by Whole Exome Sequencing**  
DUNDAR H., ÖZGÜL R. K., YALNIZOĞLU D., Erdem S., Oguz K. K., Tuncel D., TEMUÇİN Ç. M., DURSUN A.  
PEDIATRIC NEUROLOGY, vol.46, no.3, pp.172-177, 2012 (SCI-Expanded)
- XLIV. **The classification and differential diagnosis of absence seizures with short-term video-EEG monitoring during childhood**  
UYSAL SOYER Ö., YALNIZOĞLU D., Turanli G.  
TURKISH JOURNAL OF PEDIATRICS, vol.54, no.1, pp.7-14, 2012 (SCI-Expanded)
- XLV. **Whole-exome sequencing identifies recessive WDR62 mutations in severe brain malformations**  
Bilguvar K., Ozturk A. K., Louvi A., Kwan K. Y., Choi M., Tatli B., YALNIZOĞLU D., Tuysuz B., Caglayan A. O., GÖKBEN S., et al.  
NATURE, vol.467, no.7312, pp.207-211, 2010 (SCI-Expanded)
- XLVI. **ASSOCIATION OF POLYNEUROPATHY, MENTAL RETARDATION, SENSORINEURAL HEARING LOSS, 6th NERVE PALSY, CONVULSIONS, AND ORAL DYSKINESIA; A PROPABLE NEW NEUROMETABOLIC DISORDER**  
DURSUN A., YALNIZOĞLU D., DÜNDAR H., ERDEM S., AKARSU A. N., ÖZGÜL R. K.  
JOURNAL OF INHERITED METABOLIC DISEASE, vol.33, 2010 (SCI-Expanded)
- XLVII. **Surgery for epilepsy in children with dysembryoplastic neuroepithelial tumor: clinical spectrum, seizure outcome, neuroradiology, and pathology**  
BİLGİNER B., YALNIZOĞLU D., Soylemezoglu F., TURANLI G., CİLA A., TOPÇU M., Akalan N.  
CHILDS NERVOUS SYSTEM, vol.25, no.4, pp.485-491, 2009 (SCI-Expanded)
- XLVIII. **Lamotrigine in children with refractory epilepsy**  
Celebi A., YALNIZOĞLU D., Turanli G., Topaloglu H., AYSÜN S., TOPÇU M.  
TURKISH JOURNAL OF PEDIATRICS, vol.50, no.5, pp.426-431, 2008 (SCI-Expanded)
- XLIX. **Neuropsychiatric involvement in juvenile systemic lupus erythematosus**  
Demirkaya E., BİLGİNER Y., AKTAY AYAZ N., YALNIZOĞLU D., Karli-Oguz K., İŞIKHAN V., Turker T., TOPALOĞLU R., BEŞBAŞ N., BAKKALOĞLU A., et al.  
TURKISH JOURNAL OF PEDIATRICS, vol.50, no.2, pp.126-131, 2008 (SCI-Expanded)
- L. **Interleukin-6 (IL-6), tumor necrosis factor- $\alpha$  (TNF- $\alpha$ ) levels and IL-6, TNF-polymorphisms in children with thrombosis**  
Una S., GÜMRÜK F., Aytac Ş. S., YALNIZOĞLU D., GÜRGEY A.  
Journal of Pediatric Hematology/Oncology, vol.30, no.1, pp.26-31, 2008 (SCI-Expanded)
- LI. **Familial isolated non-compaction of myocardium presenting as restrictive cardiomyopathy**  
Ozkutlu S., Hascelik S., Yalnizoglu D., Altinok G.  
PEDIATRICS INTERNATIONAL, vol.49, no.4, pp.536-539, 2007 (SCI-Expanded)
- LII. **Neurologic outcome in patients with MRI pattern of damage typical for neonatal hypoglycemia**  
Yalnizoglu D., Halilo G., Turanli G., Cila A., Topcu M.  
BRAIN & DEVELOPMENT, vol.29, no.5, pp.285-292, 2007 (SCI-Expanded)
- LIII. **Evaluation of central nervous system in patients with glycogen storage disease type IA**  
Gurakan F., Aydemir Y., YÜCE A., Saltik-Temizel İ. N., DEMİR H., YALNIZOĞLU D., TOPÇU M., Usta Y., ÖZEN H.  
JOURNAL OF PEDIATRIC GASTROENTEROLOGY AND NUTRITION, vol.44, pp.163, 2007 (SCI-Expanded)
- LIV. **Malformations of cortical development and epilepsy: evaluation of 101 cases (Part II)**  
Guengoer S., Yalnizoglu D., Turanli G., Saatci I., Erdogan-Bakar E., Topcu M.  
TURKISH JOURNAL OF PEDIATRICS, vol.49, no.2, pp.131-140, 2007 (SCI-Expanded)
- LV. **Malformations of cortical development: clinical spectrum in a series of 101 patients and review of the literature (Part I)**  
Guengoer S., Yalnizoglu D., Turanli G., Saatci I., Erdogan-Bakar E., Topcu M.  
TURKISH JOURNAL OF PEDIATRICS, vol.49, no.2, pp.120-130, 2007 (SCI-Expanded)
- LVI. **Outcome and long term follow-up after corpus callosotomy in childhood onset intractable epilepsy**  
Turanli G., Yalnizoglu D., Genc-Acikgoz D., Akalan N., Topcu M.  
CHILDS NERVOUS SYSTEM, vol.22, no.10, pp.1322-1327, 2006 (SCI-Expanded)

## Articles Published in Other Journals

- I. **Pediatric cardiology consultation at long-term video EEG monitoring**  
GÜNBEY C., AYKAN H. H., KARAGÖZ T., TURANLI G., TOPÇU M., YALNIZOĞLU D.  
Annals of Medical Research, vol.30, no.1, pp.116-120, 2023 (Peer-Reviewed Journal)
- II. **Glucose Transporter Type 1 Deficiency Syndrome: A Single-Center Case Series**  
Yildirim M., Babayigit O., ILGAZ F., YALNIZOĞLU D., TOPÇU M.  
TURKISH JOURNAL OF NEUROLOGY, vol.27, no.3, pp.343-346, 2021 (ESCI)
- III. **Intertechnique Agreement in Epilepsy Imaging**  
GÜNEŞ A., YALNIZOĞLU D., GÜNBEY C., VOLKAN SALANCI B., SÖYLEMEZOĞLU A. F., BİLGİNER B., ERGÜN E.,  
TURANLI G., ERBAŞ B., TOPÇU M., et al.  
Türkiye Çocuk Hastalıkları Dergisi, vol.13, no.4, pp.292-301, 2019 (Peer-Reviewed Journal)
- IV. **HOMOZYGOUS GNAL MUTATION ASSOCIATED WITH FAMILIAL CHILDHOOD-ONSET GENERALIZED DYSTONIA**  
MASUHO I., FANG M., GENG C., ZHANG J., JIANG H., ÖZGÜL R. K., Yılmaz D. Y., YALNIZOĞLU D., YÜKSEL D., YARROW  
A., et al.  
NEUROLOGY-GENETICS, vol.2, no.3, 2016 (ESCI)
- V. **A t(5;16) translocation is the likely driver of a syndrome with ambiguous genitalia, facial dysmorphism, intellectual disability, and speech delay**  
OZANTÜRK A., Davis E. E., Sabo A., Weiss M. M., Muzny D., Dugan-Perez S., Stermans E. A., Gibbs R. A., Ozgul K. R.,  
YALNIZOĞLU D., et al.  
COLD SPRING HARBOR MOLECULAR CASE STUDIES, vol.2, no.2, 2016 (ESCI)
- VI. **Intractable Epilepsy in Childhood: Presurgical Evaluation and Treatment**  
YALNIZOĞLU D., HIRFANOĞLU T., SERDAROĞLU A., Turanli G., TOPÇU M.  
EPILEPSI, vol.18, pp.7-14, 2012 (ESCI)

## Refereed Congress / Symposium Publications in Proceedings

- I. **Central Nervous System Vasculitis in Primary Immune Deficiency**  
Uzunyayla-Sayici B., Oz S., GÖÇMEN R., TEZCAN F. İ., ÇAĞDAŞ AYVAZ D. N., ESENBOĞA S., ŞENER S., BAŞARAN H. Ö.,  
KARA A., YALNIZOĞLU D.  
Pediatric Neurology, Ankara, Turkey, 21 July 2022
- II. **CMV Miyokarditi ve Ağır Hipotonisi Olan Bir Hastada ORAI1 Defekti**  
ESENBOĞA S., DEVECİ K., AKARSU A., OCAK M., BİLDİK H. N., YAZ İ., HIZARCIOĞLU GÜLŞEN H., ERTUĞRUL İ.,  
ÇAĞDAŞ AYVAZ D. N., YALNIZOĞLU D., et al.  
7. Klinik İmmünoloji Kongresi, Turkey, 6 - 09 October 2021, pp.64-65
- III. **Comparison of physical fitness, activity and quality of life of the children with epilepsy and their healthy peers**  
SIRTBAŞ G., YALNIZOĞLU D., LİVANELİOĞLU A.  
32nd European Academy Of Childhood Disability Annual Meeting 2020, Poznan, Poland, 25 November 2020,  
vol.62, pp.40
- IV. **Severe Motor Mental Retardation with Microcephaly and Hypomyelination due to PYCR2 Gene variant in a Large Family**  
BİLGİNER GÜRBÜZ B., EROĞLU ERTUĞRUL N. G., KOŞUKCU C., YÜCEL YILMAZ D., ÖZGÜL R. K., YALNIZOĞLU D.,  
DURSUN A.  
Annual Symposium of the Society for the Study of Inborn Errors of Metabolism 2019 (SSIEM-2019), Rotterdam,  
Netherlands, 3 - 06 September 2019, vol.42, pp.1-479
- V. **Epilepsili Çocukların ve Sağlıklı Yaşıtlarının Fiziksel Aktivite, Uygunluk, Performans ve Yaşam Kalitelerinin Karşılaştırılması**  
SIRTBAŞ G., LİVANELİOĞLU A., YALNIZOĞLU D.

XVII. Fizyoterapi ve Rehabilitasyonda Gelişmeler Kongresi 2018, Antalya, Turkey, 25 - 28 April 2018

- VI. **Acute cerebellitis in children: A series of eight cases**  
YILDIRIM M., GÖÇMEN R., KONUŞKAN B., YALNIZOĞLU D., ANLAR F. B.  
12th The European Paediatric Neurology Society (EPNS) Congress, Lyon, France, 20 - 24 June 2017, vol.21, pp.121
- VII. **Ethylmalonic encephalopathy without ethylmalonic aciduria**  
YÜCEL YILMAZ D., ÖZGÜL R. K., PEKTAŞ E., serdaroğlu e., YALNIZOĞLU D., DURSUN A.  
Annual Symposium of the Society for the Study of Inborn Errors of Metabolism (SSIEM), 6 - 09 September 2016
- VIII. **ethymelanonic encephalopathy without ethylmalonic aciduria**  
YÜCEL YILMAZ D., ÖZGÜL R. K., PEKTAŞ E., SERDAROĞLU E., YALNIZOĞLU D., DURSUN A.  
SSIEM ROMA, 6 - 09 September 2016
- IX. **Pontocerebellar hypoplasia type 6 a case with neonatal seizures hypotonia and microcephaly diagnosed by exome sequencing**  
SERDAROĞLU E., ÖZGÜL R. K., YÜCEL YILMAZ D., YALNIZOĞLU D., DURSUN A.  
Annual Symposium of the Society for the Study of Inborn Errors of Metabolism (SSIEM), 6 - 09 September 2016
- X. **A case with psychomotor regression and leukoencephalopathy due to RNASEH2B gene defect**  
ÖZGÜL R. K., YÜCEL YILMAZ D., SERDAROĞLU E., YALNIZOĞLU D., TOPÇU M., DURSUN A.  
Annual Symposium of the Society for the Study of Inborn Errors of Metabolism (SSIEM), 6 - 09 September 2016
- XI. **pontocerebellar hypoplasia type 6 a case with neonatal seizures hypotonia and microcephaly diagnosed by exome sequencing**  
DURSUN A., serdaroğlu e., ÖZGÜL R. K., YÜCEL YILMAZ D., YALNIZOĞLU D.  
SSIEM roma, 6 - 09 September 2016
- XII. **Thalamic Lesions Associated with Epilepsy and ESES**  
YILDIRIM M., GÖÇMEN R., TOPÇU M., YALNIZOĞLU D.  
14th International Child Neurology Congress, Amsterdam, Netherlands, 1 - 05 May 2016
- XIII. **Ekzom Dizi Analizi ile ATP8A2 Aminofosfolipid Transporter Protein Geninde Saptanan Yeni Bir Splaying Mutasyonu**  
YÜCEL YILMAZ D., DURSUN A., YALNIZOĞLU D., serdaroğlu e.  
3. Nörometabolik Dismorfoloji Sempozyumu, Turkey, 10 - 12 March 2016
- XIV. **ANALYSIS of MRI and 18F FDG PET Scan IN NON TRAUMATIC NON TUMORAL TEMPORAL LOBE EPILEPSY experience from an epilepsy center**  
GÜNEŞ A., VOLKAN SALANCI B., TEZER FİLİK F. İ., YALNIZOĞLU D., GÖÇMEN R., SAYGI S., SÖYLEMEZOĞLU A. F., ERGÜN E., KARLI OĞUZ H. K.  
The 2016 European Congress of Radiology, Viyana, Austria, 2 - 06 March 2016
- XV. **Splicing mutation in aminophospholipid transporter protein ATP8A2 in a Turkish family**  
DURSUN A., YALNIZOĞLU D., YÜCEL YILMAZ D., serdaroğlu E., Unal Ö., görmez Z., Demirci H., Sağıroğlu M., ÖZGÜL R. K.  
SSIEM, 4 - 06 September 2015
- XVI. **SNX14 Sorting Nexin 14 gene mutation causes a new syndromic form of cerebellar atrophy in a Turkish family**  
ÖZGÜL R. K., YÜCEL YILMAZ D., Ömer g., YALNIZOĞLU D., Mahmut s., DURSUN A.  
SSIEM, 4 - 07 September 2015
- XVII. **Vacuolar storage material in a family with juvenile parkinsonism and mutations in FBX07**  
ESRA S., ÖZGÜL R. K., YALNIZOĞLU D., MADEO M., MALANDRİNİ A., KLEE E., Lİ Y., TN J., KARLI OĞUZ H. K., YÜCEL YILMAZ D., et al.  
Annual Symposium of the Society for the Study of Inborn Errors of Metabolism (SSIEM), 1 - 04 September 2015
- XVIII. **SMART syndrome A rare complication of cranial radiotherapy**  
ARDIÇLI D., GÖÇMEN R., VARAN A., YALNIZOĞLU D.  
The European Paediatric Neurology Society (EPNS), 27 - 30 May 2015, vol.19, pp.42
- XIX. **Magnetic resonance imaging findings in pediatric tuberous sclerosis patients**  
ANLAR F. B., GÖÇMEN R., SERDAROĞLU E., KONUŞKAN B., YALNIZOĞLU D.  
The European Paediatric Neurology Society (EPNS), 27 - 30 May 2015, vol.19, pp.131

- XX. **Dirençli Hipoglisemi Hipertrofik Kardiyomiyopati ve Ensefalopatili Bir Hastada Ekzom Dizileme ile Mitokondriyal TSFM Gen Defekti**  
DURSUN A., ÖMER FARUK G., MELİS P., YÜCEL YILMAZ D., TOPÇU M., YALNIZOĞLU D., YİĞİT Ş., ORHAN D., MAHMUT S., ÖZGÜL R. K.  
Uluslararası Katılımlı XIII.Ulusal Metabolik Hastalıklar Ve Beslenme Kongresi, Turkey, 14 - 18 April 2015
- XXI. **Ekzom Dizi Analizi ile MTO1 Mitochondriyal tRNA Modifier Geninde Saptanan Yeni Bir Mutasyon**  
YÜCEL YILMAZ D., ÖZGÜL R. K., ÖMER FARUK G., ESRA S., BETÜL Y., SAĞIROĞLU M., YALNIZOĞLU D., DURSUN A.  
Uluslararası Katılımlı XIII.Ulusal Metabolik Hastalıklar Ve Beslenme Kongresi, Turkey, 14 - 18 April 2015
- XXII. **Lizozom Otofagozom Defekti Sonucu Serebellar Atrofiye Neden Olan Yeni Bir Gen SNX14 Tanımlanması**  
ÖZGÜL R. K., YÜCEL YILMAZ D., YALNIZOĞLU D., MAHMUT S., DURSUN A.  
Uluslararası Katılımlı XIII.Ulusal Metabolik Hastalıklar Ve Beslenme Kongresi, 1, Turkey, 14 - 18 April 2015
- XXIII. **Klasik Glutarik Asidüri Tip I GA I Bulguları Göstermeyen Bir Ailede GCDH Gen Defekti**  
ÖZGÜL R. K., YÜCEL YILMAZ D., SANIYE Ö., YALNIZOĞLU D., TURANLI G., ESRA S., SİVRİ H. S., TOKATLI A., COŞKUN T., DURSUN A.  
Uluslararası Katılımlı XIII.Ulusal Metabolik Hastalıklar Ve Beslenme Kongresi, Turkey, 14 - 18 April 2015
- XXIV. **Hereditary spastic paraplegia with predominant cerebellar signs due to KIF1C mutation in two brothers**  
ÖZGÜL R. K., Esra s., YALNIZOĞLU D., YÜCEL YILMAZ D., TOPÇU M., ZZ G., Sağiroğlu M., DURSUN A.  
SSIEM, 4 - 07 September 2015
- XXV. **VAGUS NERVE STIMULATION IN CHILDREN WITH RETT SYNDROME**  
TOPÇU M., YALNIZOĞLU D., Turanlı G., Bilginer B., Akalan N.  
62nd Annual Meeting of the American-Epilepsy-Society, Washington, United States Of America, 5 - 09 December 2008, vol.49, pp.297
- XXVI. **Classification and follow-up of pediatric patients with absence epilepsy**  
Soyuer O. U., Turanlı G., Yalnizoglu D., Bakar E. E., Topcu M.  
60th Annual Meeting of the American-Epilepsy-Society, California, United States Of America, 1 - 05 December 2006, vol.47, pp.152

## Supported Projects

- TEZER FİLİK F. İ., KARLI OĞUZ H. K., YALNIZOĞLU D., Project Supported by Higher Education Institutions, İlaça Dirençli Epilepsi Hastalarında StereoEEG ile Epileptojenik Alanın Belirlenmesi, 2017 - 2021
- DURSUN A., YALNIZOĞLU D., YÜCEL YILMAZ D., ÖZGÜL R. K., Project Supported by Higher Education Institutions, TANI KONULAMAYAN METABOLİK/NÖROMETABOLİK HASTALIKLARDA VEZİKÜLER TRAFİK BOZUKLUKLARININ ARAŞTIRILMASI, 2017 - 2021
- YALNIZOĞLU D., Project Supported by Higher Education Institutions, Pediatrik Epilepsi Cerrahisinde Stereoelektroensefalogram ile Epileptojenik Foküsün Belirlenmesi, 2019 - 2020
- ANLAR F. B., Tağıyev A., KONUŞKAN B., YALNIZOĞLU D., Project Supported by Higher Education Institutions, Temporal lob epilepsi hastalarında inflamatuvar immün belirteçler, 2018 - 2020
- YALNIZOĞLU D., Project Supported by Higher Education Institutions, American Epilepsy Society Annual Meeting adlı kongreye poster sunumu ile katılma, 2017 - 2019
- YALNIZOĞLU D., Project Supported by Higher Education Institutions, Video-EEG Monitorizasyon Ünitesine Yatırılan Hastalarda Kardiyolojik Problemler, 2016 - 2017
- YETİM AKMAN A. F., BAYRAKCI B., YALNIZOĞLU D., KESİCİ S., TANYILDIZ M., Project Supported by Higher Education Institutions, ÜÇ PEDİYATRİK HASTADA BEYİN ÖLÜMÜ TANISINDA DEKOMPRESİF KRANİEKTOMİ VEYA KRANİOTOMİYE BAĞLI ZORLUK, 2015 - 2015



## **Metrics**

Publication: 88

Citation (WoS): 917

Citation (Scopus): 235

H-Index (WoS): 12

H-Index (Scopus): 10

## **Non Academic Experience**

Harvard Medical School Boston Children's Hospital, Klinik Nörofizyoloji ve Epilepsi Departmanı

Hacettepe Üniversitesi Tıp Fakültesi

Hacettepe Üniversitesi Tıp fakültesi