# Prof. DİLEK YALNIZOĞLU

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## Education Information

Expertise In Medicine, Harvard University, Hms/Chıldren's Hospıtal 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

## Articles Published in Journals That Entered SCI, SSCI and AHCI Indexes

I. ** 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., Turanli G., YALNIZOĞLU D.
EPILEPSY RESEARCH, vol.181, 2022 (Journal Indexed in SCI)

II. **Biallelic mutations in ELFN1 gene associated with developmental and epileptic encephalopathy and joint laxity**
DURSUN A., YALNIZOĞLU D., Yilmaz D. Y. , Oguz 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 (Journal Indexed in SCI)

III. **Long-term effects of vagus nerve stimulation in refractory pediatric epilepsy: A single-center**

**experience**

YALNIZOĞLU D., ARDIÇLI D., BİLGİNER B., KONUŞKAN B., Oguz K. K. , AKALAN N., Turanli G., SAYGI S., TOPÇU M.

EPILEPSY & BEHAVIOR, vol.110, 2020 (Journal Indexed in SCI)

IV. **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., YALNIZOĞLU D., Catsman-Berrevoets C.

EUROPEAN JOURNAL OF PAEDIATRIC NEUROLOGY, vol.28, pp.6-15, 2020 (Journal Indexed in SCI)

V. **Acute Cerebellitis or Postinfectious Cerebellar Ataxia? Clinical and Imaging Features in Acute Cerebellitis**

Yildirim M., GÖÇMEN R., KONUŞKAN B., PARLAK Ş., YALNIZOĞLU D., Anlar B.

JOURNAL OF CHILD NEUROLOGY, vol.35, no.6, pp.380-388, 2020 (Journal Indexed in SCI)

VI. **Risk factors for seizure recurrence in a pediatric observation unit**

GÜLTEKİNGİL KESER A., TEKŞAM Ö., Haliloglu G., YALNIZOĞLU D.

AMERICAN JOURNAL OF EMERGENCY MEDICINE, vol.37, no.12, pp.2151-2154, 2019 (Journal Indexed in SCI)

VII. **Clinical highlights of a very rare phospolipid remodeling disease due to MBOAT7 gene defect**

DURSUN A., YALNIZOĞLU D., ÖZGÜL R. K. , Oguz K. K. , Yucel-Yilmaz D.

AMERICAN JOURNAL OF MEDICAL GENETICS PART B-NEUROPSYCHIATRIC GENETICS, 2019 (Journal Indexed in SCI)

VIII. **Epilepsy in neurofibromatosis type 1: Diffuse cerebral dysfunction?**

Serdaroglu E., KONUŞKAN B., Oguz K. K. , Gurler G., YALNIZOĞLU D., Anlar B.

EPILEPSY & BEHAVIOR, vol.98, pp.6-9, 2019 (Journal Indexed in SCI)

IX. **Expanding the phenotype of phospholipid remodelling disease due to MBOAT7 gene defect**

YALNIZOĞLU D., ÖZGÜL R. K. , Oguz K. K. , Ozer B., Yucel-Yilmaz D., Gurbuz B., Serdaroglu E., Erol I., Topcu M., DURSUN A.

JOURNAL OF INHERITED METABOLIC DISEASE, vol.42, no.2, pp.381-388, 2019 (Journal Indexed in SCI)

X. **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., YALNIZOĞLU D.

EPILEPSY & BEHAVIOR, vol.92, pp.171-175, 2019 (Journal Indexed in SCI)

XI. **Response to Early Coenzyme Q10 Supplementation Is not Sustained in CoQ10 Deficiency Caused by CoQ2 Mutation**

Eroglu F. K. , ÖZALTIN F., Gonc N., Nalcacioglu H., Ozcakar Z. B. , YALNIZOĞLU D., Gucer S., ORHAN D., Eminoglu F. T. , GÖÇMEN R., et al.

PEDIATRIC NEUROLOGY, vol.88, pp.71-74, 2018 (Journal Indexed in SCI)

XII. **Clinical phenotype of hereditary spastic paraplegia due to KIF1C gene mutations across life span**

Yucel-Yilmaz D., Yucesan E., YALNIZOĞLU D., Oguz K. K. , Sagiroglu M. S. , Ozbek U., Serdaroglu E., Bilgic B., Erdem S., Iseri S. A. U. , et al.

BRAIN & DEVELOPMENT, vol.40, no.6, pp.458-464, 2018 (Journal Indexed in SCI)

XIII. **Electroencephalographic findings in anti-N-methyl-D-aspartate receptor encephalitis in children: A series of 12 patients**

YILDIRIM M., KONUŞKAN B., YALNIZOĞLU D., Topaloglu H., Erol I., Anlar B.

EPILEPSY & BEHAVIOR, vol.78, pp.118-123, 2018 (Journal Indexed in SCI)

XIV. **A probable new syndrome with the storage disease phenotype caused by the VPS33A gene mutation**

DURSUN A., YALNIZOĞLU D., Gerdan O. F. , Yucel-Yilmaz D., Sagiroglu M. S. , YUKSEL B., GUCER S., Sivri S., ÖZGÜL R. K.

CLINICAL DYSMORPHOLOGY, vol.26, no.1, pp.1-12, 2017 (Journal Indexed in SCI)

XV. **The seizure semiology consistent with frontal lobe symptomatogenic zone in children**

Oztoprak U., YALNIZOĞLU D., Oguz K. K. , ERGÜN E., Soylemezoglu F., BİLGİNER B., AKALAN N., TOPÇU M., Turanli G.

TURKISH JOURNAL OF PEDIATRICS, vol.58, no.6, pp.583-591, 2016 (Journal Indexed in SCI)

XVI. **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 (Journal Indexed in SCI)

XVII. **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 (Journal Indexed in SCI)

XVIII. **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 (Journal Indexed in SCI)

XIX. **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 (Journal Indexed in SCI)

XX. **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 (Journal Indexed in SCI)

XXI. **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 (Journal Indexed in SCI)

XXII. **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 (Journal Indexed in SCI)

XXIII. **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 (Journal Indexed in SCI)

XXIV. **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 (Journal Indexed in SCI)

XXV. **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 (Journal Indexed in SCI)

XXVI. **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 (Journal Indexed in SCI Expanded)

XXVII. **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., BAYRAKCİ B., AYVAZ D. N. , TEZCAN I., YALNIZOĞLU D., UCKAN D.
PEDIATRIC BLOOD & CANCER, vol.61, no.5, pp.928-930, 2014 (Journal Indexed in SCI)

XXVIII. **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 (Journal Indexed in SCI)

XXIX. **Developmental abnormalities and mental retardation: diagnostic strategy**
TOPÇU M., YALNIZOĞLU D.
PEDIATRIC NEUROLOGY, PT I, vol.111, pp.211-217, 2013 (Journal Indexed in SCI)

XXX. **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 (Journal Indexed in SCI)

XXXI. **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 (Journal Indexed in SCI)

XXXII. **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 (Journal Indexed in SCI)

XXXIII. **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 (Journal Indexed in SCI)

XXXIV. **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 (Journal Indexed in SCI)

XXXV. **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 (Journal Indexed in SCI)

XXXVI. **Neuropsychiatric involvement in juvenile systemic lupus erythematosus**
Demirkaya E., BİLGİNER Y., AKTAY AYAZ N., YALNIZOĞLU D., Karli-Oguz K., IŞIKHAN V., Turker T., TOPALOĞLU R., BEŞBAŞ N., BAKKALOĞLU A., et al.
TURKISH JOURNAL OF PEDIATRICS, vol.50, no.2, pp.126-131, 2008 (Journal Indexed in SCI)

XXXVII. **Interleukin-6 (IL-6), tumor necrosis factor-α (TNF-α) 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 (Journal Indexed in SCI Expanded)

XXXVIII. **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 (Journal Indexed in SCI)

XXXIX. **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 (Journal Indexed in SCI)

XL. **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 (Journal Indexed in SCI)

XLI. **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 (Journal Indexed in SCI)

XLII. **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 (Journal Indexed in SCI)

XLIII. **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 (Journal Indexed in SCI)

## Articles Published in Other Journals

I. **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 (Journal Indexed in ESCI)

II. **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 (Other Refereed National Journals)

III. **HOMOZYGOUS GNAL MUTATION ASSOCIATED WITH FAMILIAL CHILDHOOD-ONSET GENERALIZED DYSTONIA**
MASUHO I., FANG M., GENG C., ZHANG J., JİANG H., ÖZGÜL R. K. , Yilmaz D. Y. , YALNIZOĞLU D., YÜKSEL D., YARROW A., et al.
NEUROLOGY-GENETICS, vol.2, no.3, 2016 (Journal Indexed in ESCI)

IV. **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., Sistermans E. A. , Gibbs R. A. , Ozgul K. R. , YALNIZOĞLU D., et al.
COLD SPRING HARBOR MOLECULAR CASE STUDIES, vol.2, no.2, 2016 (Journal Indexed in ESCI)

V. **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 (Journal Indexed in ESCI)

## Refereed Congress / Symposium Publications in Proceedings

I. **Severe Motor Mental Retardation with Microcepahlyand Hypomyelination due to PYCR2 Gene variant in aLarge 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

II. **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

III. **pontocerebellar hypoplasia type 6 a case with meonatal seizures hypotonia and microcephaly diagnosed by exome sequencing**
DURSUN A., sedaroğlu e., ÖZGÜL R. K. , YÜCEL YILMAZ D., YALNIZOĞLU D.
SSIEM roma, 6 - 09 September 2016

IV. **ethymelanonic encephaopathy without etilmelanoc acitüria**
YÜCEL YILMAZ D., ÖZGÜL R. K. , PEKTAŞ E., SERDARDĞLu e., YALNIZOĞLU D., DURSUN A.
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V. **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

VI. **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

VII. **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

VIII. **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

IX. **Vacuolar storage material in a family with juvenile parkinsonism andmutations in FBXO7**
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

X. **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

XI. **Magnetic resonance imaging findings in pediatric tuberoussclerosis 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

XII. **Hereditary spastic paraplegia with predominant cerebel lar signs due to KIF1C mutation in two brothers**
ÖZGÜL R. K. , Esra s., YALNIZOĞLU D., YÜCEL YILMAZ D., TOPÇU M., ZZ G., Sağıroğlu M., DURSUN A.
SSIEM, 4 - 07 September 2015

XIII. **VAGUS NERVE STIMULATION IN CHILDREN WITH RETT SYNDROME**
TOPÇU M., YALNIZOĞLU D., Turanli 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

XIV. **Classification and follow-up of pediatric patients with absence epilepsy**
Soyuer O. U. , Turanli 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

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
YETİMAKMAN A. F. , BAYRAKCİ 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

## Citations

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