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Education Information

Post Doctorate of Medicine, Hacettepe University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, Turkey 1997 - 2000
Expertise In Medicine, Hacettepe University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, Turkey 1991 - 1996
Undergraduate, Hacettepe University, Tıp Fakültesi, Tıp Pr., Turkey 1984 - 1991

Research Areas

Health Sciences

Academic Titles / Tasks

Associate Professor, Gazi University, Tıp Fakültesi, Dahili Tıp Bilimleri, 2006 - 2010
Associate Professor, Karadeniz Technical University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2005 - 2006
Assistant Professor, Karadeniz Technical University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2003 - 2005

Courses

PARALİTİK HASTALIKLAR (Dönem4), Undergraduate, 2017 - 2018
Hereditary and acquired motor neuron diseases (Phase3), Undergraduate, 2017 - 2018
PARALYTIC DISORDERS (Phase4), Undergraduate, 2017 - 2018
APPROACH TO HEADACHE (Phase4), Undergraduate, 2017 - 2018
BAŞAĞRISINA YAKLAŞIM (Dönem4), Undergraduate, 2017 - 2018
Primary muscle disorders (Phase3), Undergraduate, 2017 - 2018
Herediter ve akkiz motor nöron hastalıkları (Dönem3), Undergraduate, 2017 - 2018
Primer kas hastalıkları (Dönem3), Undergraduate, 2017 - 2018

Advising Theses

DEMİR E., Investigation of primary headache, comorbid diseases and their effects on the quality of life in childhood,

Expertise In Medicine, C.TOSUN(Student), 2014
DEMİR E., Investigation of the relation between oxidative stress and carnitine levels in epileptic children treated with valproic acid or carbamazepine, Expertise In Medicine, A.NEŞE(Student), 2012
DEMİR E., Investigation of the etiological aspects of pediatric patients with acute impairment of consciousness, Expertise In Medicine, S.ELMAOĞULLARI(Student), 2012

Published journal articles indexed by SCI, SSCI, and AHCI

I. **An ultra-rare cause of severe hypotonia mimicking Pompe disease in an infant: RRM2B related mitochon- drial DNA depletion syndrome with a novel mutation**
İNCİ A., OKUR İ., DEMİR E., BİBEROĞLU G., TÜMER L., SERDAROĞLU A., EZGÜ F. S.
NEUROLOGY ASIA, vol.27, no.1, pp.199-202, 2022 (SCI-Expanded)

II. **Hypophosphatasia: is it an underdiagnosed disease even by expert physicians?**
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III. **Milk of calcium: A rare manifestation of juvenile dermatomyositis**
Yildirim D., AKDULUM İ., TALİM B., DEMİR E., BUYAN N., BAKKALOĞLU EZGÜ S. A.
ARCHIVES OF RHEUMATOLOGY, vol.36, no.2, pp.302-304, 2021 (SCI-Expanded)

IV. **Evaluation of state and trait anxiety levels of parents and children before electroencephalography procedures: A prospective study from a tertiary epilepsy center**
Tekin L. O., Cebeci D., Unver E., Acar A. Ş., Demir E., Gücüyener K., Uğraş Dikmen A., Serdaroğlu A., Arhan E.
EPILEPSY & BEHAVIOR, vol.112, 2020 (SCI-Expanded)

V. **Temporary Seizure in an Infant Who Had Been Exposed to G-Amino Butyric Acid Receptor Antagonist Thiocolchicoside**
Havah C., GÜCÜYENER K., Gurkas E., DEMİR E.
PEDIATRIC EMERGENCY CARE, vol.35, no.5, 2019 (SCI-Expanded)

VI. **A child with atypically subtle clinical presentation of acute arterial ischaemic stroke in the middle cerebral artery**
Orgun L. T., Derinkuyu B. E., Havali C., Boyunaga Ö. L., Yenicesu I., DEMİR E., GÜCÜYENER K.
INTERVENTIONAL NEURORADIOLOGY, vol.24, no.6, pp.684-687, 2018 (SCI-Expanded)

VII. **Internal carotid artery dissection without intracranial infarct following a minor shoulder trauma: The second pediatric case and review of the literature**
Cebeci D., ARHAN E., DEMİR E., UÇAR M., Ucar H. K., Serdaroglu A., Ozturk Z.
JOURNAL OF CLINICAL NEUROSCIENCE, vol.56, pp.172-175, 2018 (SCI-Expanded)

VIII. **Homozygous Mutations in TBC1D23 Lead to a Non-degenerative Form of Pontocerebellar Hypoplasia**
Marin-Valencia I., Gerondopoulos A., Zaki M. S., Ben-Omran T., Almureikhi M., Demir E., Guemez-Gamboa A., Gregor A., Issa M. Y., Appelhof B., et al.
AMERICAN JOURNAL OF HUMAN GENETICS, vol.101, no.3, pp.441-450, 2017 (SCI-Expanded)

IX. **Regional brain volume reduction and cognitive outcomes in preterm children at low risk at 9 years of age**
ARHAN E., GÜCÜYENER K., Soysal S., Salvarli S., Gurses M. A., SERDAROĞLU A., DEMİR E., Ergenekon E., TÜRKYILMAZ C., Onal E. E., et al.
CHILDS NERVOUS SYSTEM, vol.33, no.8, pp.1317-1326, 2017 (SCI-Expanded)

X. **Sacral aneurysmal bone cyst in a child presenting with radiculopathy.**
Derinkuyu B. E., BOYUNAGA Ö. L., Tekin-Orgun L., Ozdemir-Gokce A., Firat H., DEMİR E.
The spine journal : official journal of the North American Spine Society, vol.16, no.6, 2016 (SCI-Expanded)

XI. **Opsoclonus-myoclonus syndrome following rotavirus gastroenteritis**
Gurkas E., GÜCÜYENER K., Yilmaz U., Havali C., DEMİR E.
PEDIATRICS INTERNATIONAL, vol.56, no.6, 2014 (SCI-Expanded)

XII. **The Effect of Topiramate on Body Weight and Ghrelin, Leptin, and Neuropeptide-Y Levels of Prepubertal Children With Epilepsy**
Ozcelik A. A., SERDAROĞLU A., Bided A., ARHAN E., Soysal S., DEMİR E., GÜCÜYENER K.
PEDIATRIC NEUROLOGY, vol.51, no.2, pp.220-224, 2014 (SCI-Expanded)

XIII. **Matrix Metalloproteinase-7, Matrix Metalloproteinase-9, and Disease Activity in Pediatric Multiple Sclerosis Response**
Yilmaz U., GÜCÜYENER K., Gurkas E., DEMİR E., SERDAROĞLU A., Atak A., Aral A., Oner Y. A.
PEDIATRIC NEUROLOGY, vol.48, no.3, pp.255-256, 2013 (SCI-Expanded)

XIV. **Retinal Thickness in Pediatric Multiple Sclerosis**
Yilmaz U., GÜCÜYENER K., Erin D. M., Yazar Z., Gurkas E., SERDAROĞLU A., Tepe N., DEMİR E.
JOURNAL OF CHILD NEUROLOGY, vol.27, no.12, pp.1621-1622, 2012 (SCI-Expanded)

XV. **Reduced Retinal Nerve Fiber Layer Thickness and Macular Volume in Pediatric Multiple Sclerosis**
Yilmaz U., GÜCÜYENER K., Erin D. M., Yazar Z., Gurkas E., SERDAROĞLU A., Tepe N., DEMİR E.
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XVI. **Matrix Metalloproteinase-7 and Matrix Metalloproteinase-9 in Pediatric Multiple Sclerosis**
Unsal Y., Kivilcim G., Aysegul A., Arzu A., Esra G., Ercan D., Ayse S.
PEDIATRIC NEUROLOGY, vol.47, no.3, pp.171-176, 2012 (SCI-Expanded)

XVII. **Molecular Genetic Analysis of the PLP1 Gene in 38 Families with PLP1-related disorders: Identification and Functional Characterization of 11 Novel PLP1 Mutations**
Grossi S., Regis S., Biancheri R., Mort M., Lualdi S., Bertini E., Uziel G., Boespflug-Tanguy O., Simonati A., Corsolini F., et al.
ORPHANET JOURNAL OF RARE DISEASES, vol.6, 2011 (SCI-Expanded)

XVIII. **Clinical, neuroradiological and genetic findings in pontocerebellar hypoplasia**
Yasmin Namavar Y., Barth P. G., Kasher P. R., Van Ruissen F., Brockmann K., Bernert G., Writzl K., Ventura K., Cheng E. Y., Demir E.
BRAIN : A JOURNAL OF NEUROLOGY, vol.134, no.1, pp.143-156, 2011 (SCI-Expanded)

XIX. **Are serum nitric oxide and vascular endothelial growth factor levels affected by packed red blood cell transfusions?**
Ergenekon E., Bozkaya D., Goktas T., Erbas D., ATAK YÜCEL A., Turan O., Hirfanoglu I., Onal E. E., TÜRKYILMAZ C., KOÇ E., et al.
HEMATOLOGY, vol.15, no.3, pp.170-173, 2010 (SCI-Expanded)

XX. **An unusual presentation of Muscle-Eye-Brain disease: Severe eye abnormalities with mild muscle and brain involvement**
DEMİR E., GÜCÜYENER K., Akturk A., TALİM B., Konus Ö. L., Del Bo R., Ghezzi S., Comi G. P.
NEUROMUSCULAR DISORDERS, vol.19, no.10, pp.692-695, 2009 (SCI-Expanded)

XXI. **Assessment of mothers' knowledge and perceptions of electroencephalography and determination of the short-term effect of an informational leaflet**
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EPILEPSY & BEHAVIOR, vol.15, no.4, pp.491-495, 2009 (SCI-Expanded)

XXII. **Atypical Presentations of Subacute Sclerosing Panencephalitis in Two Neurologically Handicapped Cases**
Demir E., Ozcelik A., Arhan E., Serdaroglu A., Gucuyener K.
NEUROPEDIATRICS, vol.40, no.4, pp.195-198, 2009 (SCI-Expanded)

XXIII. **3-Methylcrotonyl-CoA Carboxylase Deficiency: Phenotypic Variability in a Family**
Eminoglu F. T., Ozcelik A. A., OKUR İ., TÜMER L., BİBEROĞLU G., DEMİR E., Hasanoglu A., Baumgartner M. R.
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XXIV. **Atypical presentations of SSPE: a clinical study in four cases**
Demir E., Aksoy A., Anlar B., Soenmez F. M.
TURKISH JOURNAL OF PEDIATRICS, vol.49, no.3, pp.295-300, 2007 (SCI-Expanded)

XXV. **Novel truncating and missense mutations of the KCC3 gene associated with Andermann syndrome**
Uyanik G., Elcioglu N., Penzien J., Gross C., Yilmaz Y., Olmez A., Demir E., Wahl D., Scheglmann K., Winner B., et al.

NEUROLOGY, vol.66, no.7, pp.1044-1048, 2006 (SCI-Expanded)

XXVI. **Ultrastructural defects of collagen VI filaments in an Ullrich syndrome patient with loss of the alpha 3(VI) N10-N7 domains**
Squarzoni S., Sabatelli P., Bergamin N., Guicheney P., Demir E., Merlini L., Lattanzi G., Ognibene A., Capanni C., Mattioli E., et al.
JOURNAL OF CELLULAR PHYSIOLOGY, vol.206, no.1, pp.160-166, 2006 (SCI-Expanded)

XXVII. **Effect of antiepileptic drugs on plasma lipids, lipoprotein (a), and liver enzymes**
Sonmez F., Demir E., ÖREM A., YILDIRMIŞ S., Orhan F., Aslan A., TOPBAŞ M.
JOURNAL OF CHILD NEUROLOGY, vol.21, no.1, pp.70-74, 2006 (SCI-Expanded)

XXVIII. **Benign familial infantile convulsions: phenotypic variability in a family.**
Demir E., Turanii G., Yalntzoglu D., Topcu M.
Journal of child neurology, vol.20, no.6, pp.535-8, 2005 (SCI-Expanded)

XXIX. **Giant axonal neuropathy: clinical and genetic study in six cases**
Demir E., Bomont P., Erdem S., Cavalier L., Demirci M., Kose G., Muftuoglu S., Cakar A., Tan E., Aysun S., et al.
JOURNAL OF NEUROLOGY NEUROSURGERY AND PSYCHIATRY, vol.76, no.6, pp.825-832, 2005 (SCI-Expanded)

XXX. **Infantile convulsions and paroxysmal choreoathetosis in a consanguineous family**
Demir E., Prud'homme J., Topcu M.
PEDIATRIC NEUROLOGY, vol.30, no.5, pp.349-353, 2004 (SCI-Expanded)

XXXI. **Collagen VI status and clinical severity in Ullrich congenital muscular dystrophy: Phenotype analysis of 11 families linked to the COL6 loci**
Demir E., Ferreiro A., Sabatelli P., Allamand V., Makri S., Echenne B., Maraldi M., Merlini L., Topaloglu H., Guicheney P.
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XXXII. **Andermann syndrome in a Turkish patient**
Demir E., Irobi J., Erdem S., Demirci M., Tan E., Timmerman V., De Jonghe P., Topaloglu H.
JOURNAL OF CHILD NEUROLOGY, vol.18, no.1, pp.76-79, 2003 (SCI-Expanded)

XXXIII. **Exon deletions in the GCHI gene in two of four Turkish families with dopa-responsive dystonia**
Klein C., Hedrich K., Kabakci K., Mohrmann K., Wiegers K., Landt O., Hagenah J., Schwinger E., Pramstaller P., Ozelius L., et al.
NEUROLOGY, vol.59, no.11, pp.1783-1786, 2002 (SCI-Expanded)

XXXIV. **Mutations in COL6A3 cause severe and mild phenotypes of Ullrich congenital muscular dystrophy**
Demir E., Sabatelli P., Allamand V., Ferreiro A., Moghadaszadeh B., Makrelouf M., Topaloglu H., Echenne B., Merlini L., Guicheney P.
AMERICAN JOURNAL OF HUMAN GENETICS, vol.70, no.6, pp.1446-1458, 2002 (SCI-Expanded)

XXXV. **A case of Sandifer's syndrome with hand tremor.**
Demir E., Saka E., Aysun S.
The Turkish journal of pediatrics, vol.43, no.4, pp.348-50, 2001 (SCI-Expanded)

XXXVI. **The gene encoding gigaxonin, a new member of the cytoskeletal BTB/kelch repeat family, is mutated in giant axonal neuropathy**
Bomont P., Cavalier L., Blondeau F., Hamida C., Belal S., Tazir M., Demir E., Topaloglu H., Korinthenberg R., Tuysuz B., et al.
NATURE GENETICS, vol.26, no.3, pp.370-374, 2000 (SCI-Expanded)

XXXVII. **Giant axonal neuropathy locus refinement to a < 590 kb critical interval**
Cavalier L., BenHamida C., Amouri R., Belal S., Bomont P., Lagarde N., Gressin L., Callen D., Demir E., Topaloglu H., et al.
EUROPEAN JOURNAL OF HUMAN GENETICS, vol.8, no.7, pp.527-534, 2000 (SCI-Expanded)

XXXVIII. **A cross section of autosomal recessive limb-girdle muscular dystrophies in 38 families**
DİNÇER P. R., Akcoren Z., Demir E., Richard I., Sancak O., Kale G., Ozme S., KARADUMAN A. A., Tan E., Urtizberea J., et al.
JOURNAL OF MEDICAL GENETICS, vol.37, no.5, pp.361-367, 2000 (SCI-Expanded)

XXXIX. **Weight gain associated with valproate in childhood**

Demir E., Aysun S.
PEDIATRIC NEUROLOGY, vol.22, no.5, pp.361-364, 2000 (SCI-Expanded)

XL. **Effect of alendronate treatment on the clinical picture and bone turnover markers in chronic idiopathic hyperphosphatasia**
DEMİR E., Bereket A., Özkan B., Topçu M.
Journal of Pediatric Endocrinology and Metabolism, vol.13, no.2, pp.217-221, 2000 (SCI-Expanded)

XLI. **Clinical and genetic correlate in childhood onset Friedreich ataxia**
Alikasifoglu M., Topaloglu H., Tuncbilek E., Ceviz N., Anar B., Demir E., Ozme S.
NEUROPEDIATRICS, vol.30, no.2, pp.72-76, 1999 (SCI-Expanded)

XLII. **A new case of oculocerebral hypopigmentation syndrome (Cross syndrome) with additional findings**
Tezcan I., Demir E., Asan E., Kale G., Muftuoglu S., Kotiloglu E.
CLINICAL GENETICS, vol.51, no.2, pp.118-121, 1997 (SCI-Expanded)

XLIII. **Infantile spasms as the initial symptom of biotinidase deficiency**
Kalayci Ö., Coskun T., Tokatli A., Demir E., Erdem G., Güngör C., Yükselen A., Özalp I.
Journal of Pediatrics, vol.124, no.1, pp.103-104, 1994 (SCI-Expanded)

Articles Published in Other Journals

I. **The Relation of Epilepsy Type with Depression and Anxiety in Children**
Ozturk Z., Soysal S., GÜCÜYENER K., DEMİR E., ARHAN E., SERDAROĞLU A.
GAZI MEDICAL JOURNAL, vol.32, pp.273-275, 2021 (ESCI)

II. **SMN1 gen delesyonu dışlanmış Spinal Musküler Atrofi ön tanılı çocuklarda etiyolojinin tüm ekzom dizi analizi verilerine dayanarak retrospektif olarak araştırılması**
SEZER A., DEMİR E., KAYHAN G., ERGÜN M. A., TUĞ E.
Gazi Medical Journal, 2020 (ESCI)

III. **Dikkat Eksikliği Hiperaktivite Bozukluğu olan Hastaların Nöropsikolojik Test Performansının Değerlendirilmesi**
GÜCÜYENER K., SOYSAL ACAR A. Ş., öztürk z., DEMİR E., ARHAN E., SERDAROĞLU A.
Gazi Medical Journal, vol.30, no.2, 2019 (Scopus)

IV. **Evaluation of Neuropsychological Test Performance of Patients with Attention Deficit Hyperactivity Disorder**
SOYSAL ACAR A. Ş., Ozturk Z., GÜCÜYENER K., DEMİR E., ARHAN E., SERDAROĞLU A.
GAZI MEDICAL JOURNAL, vol.30, no.2, pp.114-118, 2019 (ESCI)

V. **Sepsis-Associated Encephalopathy in a Child with the Torsion of Meckel's Diverticulum**
Gurkas E., Yilmaz U., DEMİR E., GÜCÜYENER K.
JOURNAL OF CLINICAL AND ANALYTICAL MEDICINE, vol.7, pp.158-160, 2016 (ESCI)

VI. **Case: Valproat-Induced Hyperammonemic Encephalopathy**
Aksu A. U., ARHAN E., DEMİR E., Ozcelik A. A., Serdaroglu A., GÜCÜYENER K.
GAZI MEDICAL JOURNAL, vol.27, no.2, pp.85-86, 2016 (ESCI)

VII. **Spastic Paraplegia with SPG11 Gene delE39 in a Turkish Patient**
Ozturk Z., Havali C., DEMİR E., GÜCÜYENER K.
GAZI MEDICAL JOURNAL, vol.27, no.4, pp.205-206, 2016 (ESCI)

VIII. **Autism Spectrum Disorder Management Practices and Level of Knowledge Among General Pediatricians**
Ozcelik A. A., Soysal S., ARHAN E., DEMİR E., GÜCÜYENER K., SERDAROĞLU A.
GAZI MEDICAL JOURNAL, vol.26, no.4, pp.158-162, 2015 (ESCI)

IX. **Reactive haemophagocytic syndrome in a child with acalculous cholecystitis**
ERDURAN E., DEMİR E., SARI A., girişken i.
Haema, vol.8, no.1, pp.113-117, 2005 (Peer-Reviewed Journal)

Books & Book Chapters

I. **ARTROGRİPOZİS TANI VE TEDAVİ ALGORİTMASI**
ÖZBUDAK P., DEMİR E.
in: Pediatrik Nöroloji: Algoritmalar ve İlaç Rehberi, Kumandaş Sefer, Cankurtaran Mehmet, Editor, AKADEMİSYEN KİTABEVİ, Ankara, pp.501-502, 2022

II. **SMA TANI VE TEDAVİ ALGORİTMASI**
DEMİR E., ÖZBUDAK P.
in: Pediatrik Nöroloji: Algoritmalar ve İlaç Rehberi, Kumandaş Sefer, Cankurtaran Mehmet, Editor, AKADEMİSYEN KİTABEVİ, Ankara, pp.509-511, 2022

III. **Artrogripozis**
ÖZBUDAK P., DEMİR E.
in: Temel Pediatrik Nöroloji: Tanı ve Tedavi, Kumandaş Sefer, Cankurtaran Mehmet, Editor, AKADEMİSYEN KİTABEVİ, Ankara, pp.2973-2983, 2022

IV. **ÖN BOYNUZ MOTOR NÖRON HASTALIKLARI**
DEMİR E., ÖZBUDAK P.
in: Temel Pediatrik Nöroloji: Tanı ve Tedavi, Kumandaş Sefer, Cankurtaran Mehmet, Editor, AKADEMİSYEN KİTABEVİ, Ankara, pp.3003-3029, 2022

V. **40. Bölüm. DEHB ve Nörolojik Bozukluklar**
öztürk z., DEMİR E.
in: DİKKAT EKSİKLİĞİ HİPERAKTİVİTE BOZUKLUĞU EL KİTABI, A. Şebnem Soysal Acar, Editor, Nobel Akademik Yayıncılık Eğitim Danışmanlık Tic. Ltd. Şti., Ankara, pp.455-464, 2019

Refereed Congress / Symposium Publications in Proceedings

I. **18-36 ay arası sağlıklı çocuklarda ekran maruziyetinin dil gelişimi üzerine etkisinin araştırılması**
Kılıç R. K., Küçükünal I. S., Çuhacı Çakır B., Duyan Çamurdan A., Serdaroğlu E., Serdaroğlu A., Demir E., Arhan E., Hirfanoğlu T.
67. Türkiye Milli Pediatri Kongresi, Girne, Cyprus (Kktc), 18 - 22 October 2023, pp.61

II. **FIRST LATE ONSET CEREBRAL FOLATE TRANSPORTER DEFICIENCYWITH A NOVEL MUTATION MIMICKING SSPE AND WILSON'SDISEASE**
Kılıç R. K., Arhan E., Kayhan G., Demir E., Hırfanoğlu T.
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III. **SSPE yi Taklit Eden Geç Çocukluk Döneminde Klinik Bulgular İle Ortaya Çıkan Nadir Tedavi Edilebilen Bir Nörometabolik Hastalık: "Serebral Folat Transportu Eksikliği" (PS-033)**
Kılıç R. K., Hırfanoğlu T., Kayhan G., Demir E., Arhan E.
24. Ulusal Çocuk Nörolojisi Kongresi 17-21 Mayıs 2023, Muğla, Muğla, Turkey, 17 - 21 May 2023, vol.1, pp.187

IV. **KONJENİTAL CD59 EKSİKLİĞİ: KLİNİK BULGULAR, TANI VE TEDAVİ**
AKBAŞ S., ÇALIK M., KİPOĞLU O., ÖZCAN N., KILIÇ R. K., YILMAZ C., DEMİR E.
I. Türk Dünyası Erciyes Çocuk Nörolojisi Kongresi, Kayseri, Turkey, 15 - 19 March 2023

V. **ANTİ-NÖBET İLAÇLARIN ALERJİK REAKSİYONLARINA KARŞI ÇOCUK HEKİMLERİNİN FARKINDALIKLARI, BİLGİ DÜZEYLERİ VE TUTUMLARININ DEĞERLENDİRİLMESİ**
KILIÇ R. K., AKBAŞ S., SERDAROĞLU E., ARHAN E., HIRFANOĞLU T., DEMİR E.
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VI. **COVİD-19 ilişkili ADEM ve LETM kliniğinde Adenovirus koenfeksiyonlu bir olgu**
Akkuzu E., Uysal Yazıcı M., Azapağası E., Hirfanoglu T., Derinkuyu B. E., Tezer H., Demir E.
66. Türkiye Milli Pediatri Kongresi, Girne, Cyprus (Kktc), 5 - 09 October 2022, pp.59

VII. **PERONEAL NERVE PALSY DUE TO OSTEOCHONDROMA OF THE FİBULAR HEAD: A RARE CAUSE OF FOOT DROP**
Kılıç R. K., Akbaş S., Öztürk A. M., Demir E.

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VIII. **PRIDOSTIGMIN TREATMENT IN X-LINKED MYOTUBULAR MYOPATHY**
Akbaş S., Kılıç R. K., Gücüyener K., Serdaroğlu E., Demir E.
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IX. **Neonatal Amplitude Integrated EEG (aEEG): Contribution to Clinical, Etiologic, and Prognosis**
Cankurt İ., Arhan E., Kılıç R. K., Serdaroğlu E., Demir E., Hırfanoğlu T.
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X. **NEONATAL EEG: IMPORTANCE AND UTILITY IN DIFFERENTIAL DIAGNOSIS and PROGNOSIS**
Kılıç R. K., Arhan E., Cankurt İ., Serdaroğlu E., Demir E., Hırfanoğlu T.
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XI. **PRE-SURGERY PHASE 1 EVALUATION RESULTS AND CONTRIBUTION TO SURGERY IN PEDIATRIC PATIENTS WITH DRUG-RESISTANT EPILEPSY**
Cankurt İ., Hırfanoğlu T., Kılıç R. K., Serdaroğlu E., Demir E., Serdaroğlu A., Arhan E.
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Supported Projects

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Metrics

Publication: 90
Citation (WoS): 1027
Citation (Scopus): 1110
H-Index (WoS): 17
H-Index (Scopus): 16

Congress and Symposium Activities

VI. NÖROMUSKÜLER HASTALIKLAR KONGRESİ, Session Moderator, Antalya, Turkey, 2023

GENÇ MERAKLILAR İÇİN PEDİATRİK NÖROMUSKÜLER HASTALIKLAR KURSU, Invited Speaker, Antalya, Turkey, 2023

1. Uluslararası Katılımlı Erciyes Türk Dünyası Çocuk Nörolojisi Kongresi, Invited Speaker, Kayseri, Turkey, 2023

GENÇ MERAKLILAR İÇİN PEDİATRİK NÖROMUSKÜLER HASTALIKLAR KURSU, Invited Speaker, İstanbul, Turkey, 2023

23. ULUSAL ÇOCUK NÖROLOJİ KONGRESİ, Session Moderator, İzmir, Turkey, 2021
V. NÖROMUSKULER HASTALIKLAR KONGRESİ, Invited Speaker, Samsun, Turkey, 2021

Non Academic Experience

U 523, INSTİTUT DE MYOLOGİE, GROUPE HOSPİTALİER PİTİÉ-SALPÊTRIÈRE, PARİS, FRANSA
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