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Publons / Web Of Science ResearcherID: W-7266-2019

Yoksis Araştırmacı ID: 56671

Eğitim Bilgileri

Tıpta Yandal Uzmanlık, İstanbul Üniversitesi, İstanbul Tıp Fakültesi, Klinik Nörofizyoloji, Nöroloji, Türkiye 2011 - 2011

Doktora, İstanbul Üniversitesi, Sağlık Bilimleri Enstitüsü, Genetik, Türkiye 2002 - 2009

Tıpta Uzmanlık, İstanbul Üniversitesi, İstanbul Tıp Fakültesi, Nöroloji, Türkiye 1996 - 2001

Yabancı Diller

Fransızca, B2 Orta Üstü

İngilizce, B2 Orta Üstü

Yaptığı Tezler

Doktora, Temporal Lob Epileptogeneziyle İlişkili Genlerin Ekspresyon Analizi, İstanbul Üniversitesi, Deneysel Tıp Araştırmaları Enstitüsü, Genetik Anabilim Dalı, 2009

Tıpta Uzmanlık, İdyopatik jeneralize epilepsili ailelerde kalıtım özellikleri ve genetik inceleme, İstanbul Üniversitesi, Klinik Nörofizyoloji, Nöroloji, 2001

Araştırma Alanları

Tıp, Sağlık Bilimleri, Dahili Tıp Bilimleri, Nöroloji, Tıbbi Genetik

Akademik Unvanlar / Görevler

Prof. Dr., İstanbul Üniversitesi, İstanbul Tıp Fakültesi, Dahili, 2013 - Devam Ediyor

Doktor, İstanbul Üniversitesi, Deneysel Tıp Araştırmaları Enstitüsü, Genetik Anabilim Dalı, 2010 - Devam Ediyor

Doç. Dr., İstanbul Üniversitesi, İstanbul Tıp Fakültesi, Dahili, 2008 - 2013

Verdiği Dersler

Beyin sapı-kranyal sinirler, Doktora, 2013 - 2014
Normal EEG, Yüksek Lisans, 2013 - 2014
Bilinç Bozuklukları, Yüksek Lisans, 2013 - 2014
Mitokondriyal kalıtım, Doktora, 2013 - 2014
Bilinç Bozukluğunda EEG, Yüksek Lisans, 2013 - 2014
İlaçlara bağlı değişiklikler, Yüksek Lisans, 2013 - 2014
X'e bağlı kalıtım, Yüksek Lisans, 2013 - 2014

Yönetilen Tezler

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- I. **Exome sequencing of 20,979 individuals with epilepsy reveals shared and distinct ultra-rare genetic risk across disorder subtypes**
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- IV. **Acute effect of transcranial direct current stimulation on photoparoxysmal response.**
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- VI. **Diffusion tensor imaging in photosensitive and nonphotosensitive juvenile myoclonic epilepsy**
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- VII. **Exploring shared triggers and potential etiopathogenesis between migraine and idiopathic/genetic epilepsy: Insights from a multicenter tertiary-based study**
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- VIII. **GWAS meta-analysis of over 29,000 people with epilepsy identifies 26 risk loci and subtype-specific genetic architecture**

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- IX. **Ictal Blinking: Reappraisal of the Lateralization and Localization Value in Focal Seizures.**
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- X. **Birth outcomes in pregnant women with epilepsy: A Nationwide multicenter study from Türkiye**
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- XII. **Sex-based electroclinical differences and prognostic factors in epilepsy with eyelid myoclonia**
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- XIII. **Photosensitive occipital lobe epilepsy: Delineation of an under-recognized reflex epilepsy syndrome according to the new ILAE criteria and long-term follow-up**
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- XIV. **Elevated sTREM2 and NFL levels in patients with sepsis associated encephalopathy**
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- XV. **Epilepsy or neurodevelopmental disorders are associated with homozygous and pathogenic ELP2 variation in three siblings.**
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- XVI. **Status epilepticus in mesial temporal lobe epilepsy**
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- XVII. **Neurological effects of long-term diet on obese and overweight individuals: An electroencephalogram and event-related potential study**
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- XVIII. **Posterior Reversible Encephalopathy in Sepsis-Associated Encephalopathy: Experience from a Single Center.**
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- XIX. **The effect of fear of COVID-19 on quality of life in patients with epilepsy**
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- XX. **The rare rs769301934 variant in NHLRC1 is a common cause of Lafora disease in Turkey.**
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- XXI. **Idiopathic photosensitive occipital lobe epilepsy: an exceedingly rare reflex epilepsy syndrome**

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- XXII. **International case control study of Sudden Unexpected Death in Epilepsy (SUDEP)**
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- XXIV. **Clinical and Genetic Features in Patients With Reflex Bathing Epilepsy**
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- XXV. **Progressive Myoclonus Epilepsy Caused by a Homozygous Splicing Variant of SLC7A6OS.**
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- XXVI. **Depression is a major determinant of sleep abnormalities in patients with epilepsy.**
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- XXVII. **Slow repetitive transcranial magnetic stimulation in refractory juvenile myoclonic epilepsies**
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- XXVIII. **Peripheral blood expression levels of inflammasome complex components in two different focal epilepsy syndromes**
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- XXIX. **Functional connectivity analysis of patients with temporal lobe epilepsy displaying different ictal propagation patterns**
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- XXX. **Reflex epileptic features in patients with focal epilepsy of unknown cause**
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- XXXI. **Long-term follow-up of a large cohort with focal epilepsy of unknown cause: deciphering their clinical and prognostic characteristics**
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- XXXII. **Let's talk SUDEP**
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- XXXIII. **Relationship Between Sleep Characteristics and Sudden Unexplained Death Risk in Epilepsy**
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- XXXIV. **Investigation of Generalized EEG Paroxysms Accompanying Focal Epilepsies**
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- XXXV. **Ultra-Rare Genetic Variation in the Epilepsies: A Whole-Exome Sequencing Study of 17,606 Individuals**
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XXXVI. Impact of seizure-related injuries on quality of life

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XXXVIII. Identification of epilepsy related pathways using genome-wide DNA methylation measures: A trio-based approach

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XXXIX. Investigation of SLC2A1 gene variants in genetic generalized epilepsy patients with eyelid myoclonia

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XL. Investigation of the Video-EEG Findings and Clinical Data in Patients Diagnosed With Epilepsy and Psychosis

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XLI. Unusual ictal propagation patterns suggesting poor prognosis after temporal lobe epilepsy surgery: Switch of lateralization and bilateral asynchrony

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- XLIX. The Characteristics and Long-Term Course of Epilepsy in Lipoid Proteinosis: A Spectrum From Mild to Severe Seizures in Relation to ECM1 Mutations.**
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- L. Genital automatisms: Reappraisal of a remarkable but ignored symptom of focal seizures**
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- LI. Stereotyped high-frequency oscillations discriminate seizure onset zones and critical functional cortex in focal epilepsy**
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- LII. INVESTIGATIONAL CLUES ABOUT INSOMNIA IN PATIENTS WITH EPILEPSY**
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- LIII. SLEEP CHARACTERISTICS OF EPILEPSY PATIENTS AND THE RELATION OF SUDEP RISK**
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- LIV. Investigation of anti-neuronal antibodies in status epilepticus of unknown etiology: a prospective study**
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- LVI. Analysis of heart rate variability and risk factors for SUDEP in patients with drug-resistant epilepsy**
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- LVII. How Different Are the Patients With Bilateral Hippocampal Sclerosis From the Unilateral Ones Clinically?**
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- LIX. Teratogenicity of Antiepileptic Drugs.**
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- LX. Vagus Nerve Stimulation in Adults With Drug-Resistant Epilepsy: Efficacy, Adverse Effects and Outcomes**
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- LXI. Investigation of neuronal auto-antibodies in systemic lupus erythematosus patients with epilepsy**
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- LXII. Analysis of the tremor in juvenile myoclonic epilepsy**
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- LXIV. LONG TERM FOLLOW-UP OF NEUROPSYCHOLOGICAL OUTCOME IN PATIENTS WITH TEMPORAL LOBE EPILEPSY**
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- LXV. THE EFFECTS OF FIXATION-OFF SENSITIVITY ON EYE CLOSURE AND EYE CLOSED STATUS-RELATED EPILEPTIFORM DISCHARGES**
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- LXVI. NEURONAL AUTO- ANTIBODIES IN MESIAL TEMPORAL LOBE EPILEPSY WITH HIPPOCAMPAL SCLEROSIS: EPIPHENOMENON OR MARKER FOR AN AUTOIMMUNE SUBTYPE?**
Vanli-Yavuz E. N., Tuzun E., Ekizoglu E., Erdag E., Baysal-Kirac L., Ulusoy C., Peach S., Gundogdu G., Sencer S., Sencer A., et al.
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- LXVII. THE CLINICAL OUTCOME OF LENNOX-GASTAUT SYNDROME IN ADULTS**
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- LXX. LATERALIZED PERIODIC DISCHARGES ASSOCIATED WITH STATUS EPILEPTICUS IN THE FIRST YEAR AFTER STROKE**
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- XXIII. **Socio-Demographic, Clinical Characteristic And Social Life Of Patient With Epilepsy**
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- XXIV. **Epilepsy Knowledge, Attitude, Stigma, Anxiety and Depression and Quality of Life: A Structural Equation Modelling**
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13th European Congress on Epileptology, Vienna, Avusturya, 26 - 30 Ağustos 2018, cilt.59
- XXV. **The Predictors of Drug-Resistance in Patients with MRI-Negative Focal Epilepsies of Unknown Cause**
Atalar A. C., Yavuz E. N. V., Bebek N., Baykan B.
13th European Congress on Epileptology, Vienna, Avusturya, 26 - 30 Ağustos 2018, cilt.59
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XLIX. VALIDITY AND RELIABILITY OF TURKISH VERSION OF THE EPILEPSY SELF MANAGEMENT SCALE

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Dede H. O., Gurses C., Bebek N., Baykan B., Gokyigit A.

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LIV. THE RETROSPECTIVE EVALUATION OF ELEVEN EPILEPTIC PATIENTS WITH A DIAGNOSIS OF TUBEROUS SCLEROSIS COMPLEX

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LX. TWONovel AND THREE KNOWN EPM2A AND NHLRC1 (EPM2B) GENE VARIANTS LEADING TO LAFORA DISEASE IN TURKISH PATIENTS

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XXI. Long-term follow-up of genetic generalized epilepsy with typical absence seizures and generalized paroxysmal fast activity in adulthood

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Taskiran E., Gurses C., Matur Z., Bebek N., Baykan B., Gul G., Gokyigit A.

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LXIX. CLINICAL, GENETIC, AND EEG FEATURES OF TURKISH PROGRESSIVE MYOCLONIC EPILEPSY PATIENTS

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LXXII. INVESTIGATING THE EXPRESSION OF INFLAMMATION RELATED GENES IN TEMPORAL LOBE EPILEPSY PATHOGENESIS

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LXXV. Investigation of Bromodomain-Containing Protein 2 (BRD2) Gene Mutations in Photosensitive Epilepsy

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- LXXVII. **Evaluation of seizure and sleep characteristics of patients with pure nocturnal seizures**
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- LXXVIII. **RECURRENT FORM OF COMPLEX PARTIAL STATUS EPILEPTICUS WITH FRONTAL LOBE ORIGIN: TWO CASE REPORTS WITH LONG-TERM FOLLOW-UP**
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- LXXIX. **Morphological alterations in the blood-brain barrier, glia and neuron in the hippocampus of patients with mesial temporal lobe epilepsy**
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- LXXX. **Anti-neuronal antibodies in Hashimoto's thyroiditis patients with central nervous system involvement**
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- LXXXII. **Electrophysiological evaluation of children from mothers with epilepsy using anti-epileptic drugs**
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- LXXXIII. **Myoclonic seizures subside in the fourth decade in juvenile myoclonic epilepsy: A long-term observational study**
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- LXXXV. **Efficacy of levetiracetam as an add-on therapy in patients with startle epilepsy**
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- LXXXVIII. **Late onset and slow progressing Lafora Disease in four siblings with EPM2B mutation**
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- LXXXIX. **Brain derived neurotrophic factor (BDNF) and multidrug resistance (MDR) gene polymorphisms in patients with mesial temporal sclerosis (MTS)**
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- XC. **Two HIV (-) patients with neurosyphilis presenting convulsive and nonconvulsive status epilepticus in different stages of their disease**
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XCIV. Evaluation of four cases with electrical status epilepticus during slow sleep

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XCV. Clinical characteristics of bilateral mesial temporal sclerosis

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XCVIII. Clinical and genetic investigations of families with pure benign familial infantile convulsions (BFIC): Confirmation of linkage to chromosome 16

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