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Yoksis Araştırmacı ID: 126268

Eğitim Bilgileri

Tıpta Yandal Uzmanlık, İstanbul Üniversitesi, İstanbul Tıp Fakültesi, Nöroloji Ad, Çocuk Nörolojisi Bd, Türkiye 1999 - 2004

Tıpta Uzmanlık, Şişli Etfal Hastanesi Nöroloji, Nöroloji, Türkiye 1989 - 1994

Lisans, İstanbul Üniversitesi, Cerrahpaşa Tıp Fakültesi, Türkiye 1982 - 1988

Yabancı Diller

İngilizce, C1 İleri

Yaptığı Tezler

Tıpta Uzmanlık, Serebral Kortikal Gelişim Malformasyonları'nda Klinik ve Manyetik Rezonans Görüntüleme Özellikleri, İstanbul Üniversitesi, Nöroloji, Çocuk Nörolojisi, 2004

Tıpta Uzmanlık, İntraserebral Hematomlarda Akut Dönemde Prognozu Etkileyen Faktörler, --Seçiniz--, Nöroloji, 1994

Araştırma Alanları

Tıp, Sağlık Bilimleri, Dahili Tıp Bilimleri, Nöroloji

Akademik Unvanlar / Görevler

Prof.Dr., İstanbul Üniversitesi, İstanbul Tıp Fakültesi, Dahili Bilimler, 2012 - Devam Ediyor

Doç.Dr., İstanbul Üniversitesi, İstanbul Tıp Fakültesi, Dahili Bilimler, 2006 - Devam Ediyor

Akademik İdari Deneyim

İstanbul Üniversitesi, 2009 - 2013

Verdiği Dersler

Kranial felçler, Lisans, 2020 - 2021

SCI, SSCI ve AHCI İndekslerine Giren Dergilerde Yayınlanan Makaleler

- I. **Exome data of developmental and epileptic encephalopathy patients reveals de novo and inherited pathologic variants in epilepsy-associated genes**
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- XIV. **A Patient with Glucose Transporter Type 1 Deficiency Syndrome: Paroxysmal Choreoathetosis and Cerebral Positron-Emission Tomography Findings**
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- XV. **Adjunctive everolimus therapy for tuberous sclerosis complex-associated refractory seizures: Results from the postextension phase of EXIST-3**
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- XVI. **Novel Protein Biomarkers of Monoamine Metabolism Defects Correlate with Disease Severity**
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- XVII. **Immune alterations in subacute sclerosing panencephalitis reflect an incompetent response to eliminate the measles virus**
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- XVIII. **Novel mutations in ATP13A2 associated with mixed neurological presentations and iron toxicity due to nonsense-mediated decay**
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- XX. **Non-convulsive status epilepticus in two patients with tuberous sclerosis**
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- XXV. **Clinical and genetic spectrum of an orphan disease MPAN: a series with new variants and a novel phenotype.**

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- XXVI. **Investigation of neuronal auto-antibodies in children diagnosed with epileptic encephalopathy of unknown cause**

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- XXVII. **A new splice-site mutation in SLC12A6 causing Andermann syndrome with motor neuronopathy**

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- XXVIII. **DBS in pediatric patients: institutional experience**

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- XXIX. **Adjunctive everolimus for children and adolescents with treatment-refractory seizures associated with tuberous sclerosis complex: post-hoc analysis of the phase 3 EXIST-3 trial**

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- XXXI. **Megalencephalic leukoencephalopathy with subcortical cysts Characterization of disease variants**

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- XXXIII. **Pallidal Stimulation in an 11-Year-Old Boy with Treatment-Resistant Tourette Syndrome**

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- XLI. **A novel gene mutation in *PANK2* in a patient with severe jaw-opening dystonia**
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- XLV. **L-2-Hydroxyglutaric Aciduria: Report of Four Turkish Patients from the Same Family**
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Metrikler

Yayın: 105

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