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Uluslararası Araştırmacı ID'leri

Publons / Web Of Science ResearcherID: AAT-9501-2020

Yoksis Araştırmacı ID: 14464

Eğitim Bilgileri

Tıpta Uzmanlık, İstanbul Üniversitesi, İstanbul Tıp Fakültesi, Nöroloji Anabilim Dalı, Türkiye 1993 - 1998

Lisans, Anadolu Üniversitesi, Tıp Fakültesi, Tıp, Türkiye 1985 - 1991

Yabancı Diller

İngilizce, C1 İleri

Yaptığı Tezler

Tıpta Uzmanlık, AMYOTROFİK LATERAL SKLEROZ HASTALIĞINDA NÖROPSİKOLOJİK DEĞERLENDİRME VE OLAYA İLİŞKİN POTANSİYELLER, İstanbul Üniversitesi, Tıp Fakültesi, Nöroloji, 1998

Araştırma Alanları

Tıp, Sağlık Bilimleri, Dahili Tıp Bilimleri, Nöroloji

Akademik Unvanlar / Görevler

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

Doç.Dr., İstanbul Üniversitesi, İstanbul Tıp Fakültesi, Tıp, 2006 - 2012

Uzman, İstanbul Üniversitesi, İstanbul Tıp Fakültesi, Tıp, 1993 - 1998

SCI, SSCI ve AHCI İndekslerine Giren Dergilerde Yayınlanan Makaleler

- TREX1 p.A129fs and p.Y305C variants in a large multi-ethnic cohort of CADASIL-like unrelated patients**

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- II. **Identification of metabolic correlates of mild cognitive impairment in Parkinson's disease using magnetic resonance spectroscopic imaging and machine learning**
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- IV. **Decreased levels of cytokines implicate altered immune response in plasma of moderate-stage Alzheimer's disease patients**
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- VIII. **Kappa/Lambda Light-chain Typing in Alzheimer's Disease**
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- IX. **Genetic variants of vitamin D metabolism-related *DHCR7/NADSYN1* locus and *CYP2R1* gene are associated with clinical features of Parkinson's disease.**
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- XI. **Clinical and molecular genetic findings of hereditary Parkinson's patients from Turkey.**
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- XII. **Frequency of frontotemporal dementia-related gene variants in Turkey**
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- XIII. **Detection of visual and frontoparietal network perfusion deficits in Parkinson's disease dementia.**
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- XIV. **Cerebellar ataxia, neuropathy and vestibular areflexia syndrome (Canvas) is an important cause of late-onset ataxia**
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- XV. **A new alpha-synuclein missense variant (Thr72Met) in two Turkish families with Parkinson's disease**
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- XVI. **Neurological features and outcomes of Wilson's disease: a single-center experience.**
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- XIX. **A novel SACS p.Pro4154GlnfsTer20 mutation in a family with autosomal recessive spastic ataxia of Charlevoix-Saguenay**
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Mencacci N. E., Brockmann M. M., Dai J., Pajusalu S., Atasu B., Campos J., Pino G., Gonzalez-Latapi P., Patzke C., Schwake M., et al.
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- XXVII. **Functional Connectivity Analysis in Heterozygous Glucocerebrosidase Mutation Carriers**
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- XXXI. **Peripheral TREM2 mRNA levels in early and late-onset Alzheimer disease's patients.**
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- XXXII. **Caregiver Burden, Quality of Life and Related Factors in Family Caregivers of Dementia Patients in Turkey.**
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- XXXIV. **Clinico-Genetic, Imaging and Molecular Delineation of COQ8A-Ataxia: A Multicenter Study of 59 Patients**
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- XXXV. **The association of serum clusterin levels and Clusterin rs11136000 polymorphisms with Alzheimer disease in a Turkish cohort**
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- XXXVI. **Event-related potential changes due to early-onset Parkinson's disease in parkin (PARK2) gene mutation carriers and non-carriers**
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- XXXVII. **Turkish inappropriate medication use in the elderly (TIME) criteria to improve prescribing in older adults: TIME-to-STOP/TIME-to-START**
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XXXIX. Neuroinflammation Mediators are Reduced in Sera of Parkinson's Disease Patients with Mild Cognitive Impairment

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XL. Multicenter study of levodopa carbidopa intestinal gel in Parkinson's disease: the Turkish experience

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XLII. Peripheral GRN mRNA and Serum Progranulin Levels as a Potential Indicator for Both the Presence of Splice Site Mutations and Individuals at Risk for Frontotemporal Dementia.

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L. Peripheral GRN mRNA and Serum Progranulin Levels as a Potential Indicator for Both the Presence of Splice Site Mutations and Individuals at Risk for Frontotemporal Dementia

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- LII. **Which cognitive dual-task walking causes most interference on the Timed Up and Go test in Parkinson's disease: a controlled study.**
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- LV. **Clinical phenotype of hereditary spastic paraplegia due to KIF1C gene mutations across life span.**
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- LVII. **Nasu Hakola Disease: A Rare Cause of Dementia and Cystic Bone Lesions, Report of a New Turkish Family**
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- LVIII. **An assessment of Movement Disorder Society Task Force diagnostic criteria for mild cognitive impairment in Parkinson's disease**
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- LXI. **The p.Thr11Met mutation in c19orf12 is frequent among adult Turkish patients with MPAN.**
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- LXIII. **Impact of Neuro-Behcet Disease Immunoglobulin G on Neuronal Apoptosis**
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Metrikler

Yayın: 277

Atıf (WoS): 4495

Atıf (Scopus): 4881

H-İndeks (WoS): 27

H-İndeks (Scopus): 29