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Education Information

Expertise In Medicine, Istanbul University, Istanbul Medical Faculty, Nöroloji Anabilim Dalı, Turkey 1993 - 1998

Undergraduate, Anadolu University, School Of Medicine, Tıp, Turkey 1985 - 1991

Foreign Languages

English, C1 Advanced

Dissertations

Expertise In Medicine, AMYOTROFİK LATERAL SKLEROZ HASTALIĞINDA NÖROPSİKOLOJİK DEĞERLENDİRME VE OLAYA İLİŞKİN POTANSİYELLER, Istanbul University, Tıp Fakültesi, Nöroloji, 1998

Research Areas

Medicine, Health Sciences, Internal Medicine Sciences, Neurology

Academic Titles / Tasks

Professor, Istanbul University, Istanbul Medical Faculty, Tıp, 2012 - Continues

Associate Professor, Istanbul University, Istanbul Medical Faculty, Tıp, 2006 - 2012

Expert, Istanbul University, Istanbul Medical Faculty, Tıp, 1993 - 1998

Published journal articles indexed by SCI, SSCI, and AHCI

- 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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- III. **Anticholinergic Burden, Polypharmacy, and Cognition in Parkinson's Disease Patients with Mild Cognitive Impairment: A Cross-Sectional Observational Study**
SÜMBÜL ŞEKERCİ B., Bilgic B., PAŞIN Ö., Demir M. E., Hanagasi H. A.
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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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- V. **Genotype-Phenotype correlations of SCARB2 associated clinical presentation: a case report and in-depth literature review**
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- VI. **Medication management and treatment adherence in Parkinson's disease patients with mild cognitive impairment**
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- VII. **Intensive voice treatment (the Lee Silverman Voice Treatment [LSVT[®]LOUD]) for individuals with Wilson's disease and adult cerebral palsy: two case reports.**
Ertan E., GÜRVİT İ. H., Hanagasi H. A., Bilgic B., Tuncer M. A., Yilmaz C.
Logopedics, phoniatrics, vocology, vol.47, no.4, pp.262-270, 2022 (SCI-Expanded)
- VIII. **Kappa/Lambda Light-chain Typing in Alzheimer's Disease**
Kaya Z. Z., Tuzuner M. B., Sahin B., Akgun E., AKSUNGAR F., Koca S., SERDAR M. A., ŞAHİN Ş., Cinar N., Karsidag S., et al.
CURRENT ALZHEIMER RESEARCH, vol.19, no.1, pp.84-93, 2022 (SCI-Expanded)
- IX. **Genetic variants of vitamin D metabolism-related *DHCR7/NADSYN1* locus and *CYP2R1* gene are associated with clinical features of Parkinson's disease.**
ALAYLIOĞLU M., DURSUN E., Genc G., Sengul B., Bilgic B., GÜNDÜZ A., APAYDIN H., KIZILTAN G., GÜRVİT İ. H., Hanagasi H. A., et al.
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- X. **A comprehensive analysis of copy number variation in a Turkish dementia cohort**
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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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- XVI. **Neurological features and outcomes of Wilson's disease: a single-center experience.**
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- XVII. **TREM2 variants as a possible cause of frontotemporal dementia with distinct neuroimaging features.**
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European journal of neurology, vol.28, no.8, pp.2603-2613, 2021 (SCI-Expanded)
- XVIII. **A new alpha-synuclein missense variant (Thr72Met) in two Turkish families with Parkinson's disease.**
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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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- XX. **The Complex Genetic Landscape of Hereditary Ataxias in Turkey and Implications in Clinical Practice**
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- XXI. **The ARCA Registry: A Collaborative Global Platform for Advancing Trial Readiness in Autosomal Recessive Cerebellar Ataxias.**
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- XXII. **A novel PSEN2 p.Ser175Phe variant in a family with Alzheimer's disease**
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- XXIII. **A novel PNPLA6 mutation in a Turkish family with intractable Holmes tremor and spastic ataxia.**
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- XXVII. **Functional Connectivity Analysis in Heterozygous Glucocerebrosidase Mutation Carriers**
Sezgin M., Kicik A., Bilgic B., Kurt E., Bayram A., Hanagasi H. A., Tepgec F., Toksoy G., Gurvit H., Uyguner O., et al.
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- XXVIII. **Cognition of the mothers of patients with Duchenne muscular dystrophy.**
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- XXIX. **Characterization of Recessive Parkinson Disease in a Large Multicenter Study**
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- XXX. **The cerebral blood flow deficits in Parkinson's disease with mild cognitive impairment using arterial spin labeling MRI.**
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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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- XXXIII. **Revisiting the complex architecture of ALS in Turkey: Expanding genotypes, shared phenotypes, molecular networks, and a public variant database**
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- XXXIV. **Clinico-Genetic, Imaging and Molecular Delineation of COQ8A-Ataxia: A Multicenter Study of 59 Patients**
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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**
Bahat G., İlhan B., Erdogan T., Halil M., Savas S., Ulger Z., Akyuz F., Bilge A., Cakir S., Demirkan K., et al.
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- XXXVIII. **Small ubiquitin-related modifier (SUMO) 3 and SUMO4 gene polymorphisms in Parkinson's disease**
Küçükali C. İ., Salman B., Yüceer H., Ulusoy C. A., Abacı N., Ekmekci S., Tüzün E., Bilgiç B., Hanağası H. A.

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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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- XLIII. **A patient with early-onset Alzheimer's disease with a novel PSEN1 p.Leu424Pro mutation**
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- XLVII. **Patients with Lately Diagnosed Cerebrotendinous Xanthomatosis**
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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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- LIV. **HPCA Confirmed as a Genetic Cause of DYT2-Like Dystonia Phenotype**
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- LVI. **Role of LRRK2 and SNCA in autosomal dominant Parkinson's disease in Turkey**
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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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- LXIII. **Impact of Neuro-Behcet Disease Immunoglobulin G on Neuronal Apoptosis**
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- LXIV. **GC and VDR SNPs and Vitamin D Levels in Parkinson's Disease: The Relevance to Clinical Features**
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Clinical neurology and neurosurgery, vol.154, pp.34-42, 2017 (SCI-Expanded)
- LXVI. **Cognitive and anatomical correlates of anosognosia in amnesic mild cognitive impairment and early-stage Alzheimer's disease**
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Metrics

Publication: 277

Citation (WoS): 4495

Citation (Scopus): 4881

H-Index (WoS): 27

H-Index (Scopus): 29