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

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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

- Responsiveness of the Scale for the Assessment and Rating of Ataxia and Natural History in 884 Recessive and Early Onset Ataxia Patients**

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- II. **Validity and Reliability of the Pictures of Facial Affect (POFA) in Healthy People and Patients with Essential Tremor in the Turkish Population.**
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- III. **Medication management and treatment adherence in Parkinson's disease patients with mild cognitive impairment**
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- IV. **TREX1 p.A129fs and p.Y305C variants in a large multi-ethnic cohort of CADASIL-like unrelated patients**
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- VI. **Eating Difficulties and Relationship With Nutritional Status Among Patients With Dementia.**
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- VII. **Anticholinergic Burden, Polypharmacy, and Cognition in Parkinson's Disease Patients with Mild Cognitive Impairment: A Cross-Sectional Observational Study**
SÜMBÜL ŞEKERCİ B., Bilgic B., PAŞİN Ö., Demir M. E., Hanagasi H. A.
DEMENTIA AND GERIATRIC COGNITIVE DISORDERS, vol.51, no.5, pp.386-395, 2023 (SCI-Expanded)
- VIII. **Establishing an online resource to facilitate global collaboration and inclusion of underrepresented populations: Experience from the MJFF Global Genetic Parkinson's Disease Project**
Vollstedt E., Madoev H., Aasly A., Ahmad-Annur A., Al-Mubarak B., Alcalay R. N., Alvarez V., Amorin I., Annesi G., Arkadir D., et al.
PloS one, vol.18, no.10, 2023 (SCI-Expanded)
- IX. **Quantitative Gait and Balance Outcomes for Ataxia Trials: Consensus Recommendations by the Ataxia Global Initiative Working Group on Digital-Motor Biomarkers**
Ilg W., Milne S., Schmitz-Hübsch T., Alcock L., Beichert L., Bertini E., Mohamed Ibrahim N., Dawes H., Gomez C. M., Hanagasi H., et al.
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- X. **Identification of metabolic correlates of mild cognitive impairment in Parkinson's disease using magnetic resonance spectroscopic imaging and machine learning**
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- XII. **Genotype-Phenotype correlations of SCARB2 associated clinical presentation: a case report and in-depth literature review**
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BMC NEUROLOGY, vol.22, no.1, 2022 (SCI-Expanded)
- XIII. **Kappa/Lambda Light-chain Typing in Alzheimer's Disease**
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al.

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- XIV. **Intensive voice treatment (the Lee Silverman Voice Treatment [LSVT[®]LOUD]) for individuals with Wilson's disease and adult cerebral palsy: two case reports.**
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- XV. **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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- XVI. **A comprehensive analysis of copy number variation in a Turkish dementia cohort**
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- XVII. **Clinical and molecular genetic findings of hereditary Parkinson's patients from Turkey.**
Emekli I., Tepgeç F., Samanci B., Toksoy G., Hasanoğulları Kına G., Tüfekçioğlu Z., Başaran S., Bilgiç B., Gürvit I. H., Emre M., et al.
Parkinsonism & related disorders, vol.93, pp.35-39, 2021 (SCI-Expanded)
- XVIII. **Frequency of frontotemporal dementia-related gene variants in Turkey**
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- XIX. **Detection of visual and frontoparietal network perfusion deficits in Parkinson's disease dementia.**
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- XXV. **A novel SACS p.Pro4154GlnfsTer20 mutation in a family with autosomal recessive spastic ataxia of Charlevoix-Saguenay**
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NEUROLOGICAL SCIENCES, vol.42, no.7, pp.2969-2973, 2021 (SCI-Expanded)
- XXVI. **The Complex Genetic Landscape of Hereditary Ataxias in Turkey and Implications in Clinical Practice**
Vural A., Simsir G., Tekgul S., Kocoglu C., Akcimen F., Kartal E., Sen N. E., Lahut S., Omur O., Saner N., et al.
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- XXVII. **The ARCA Registry: A Collaborative Global Platform for Advancing Trial Readiness in Autosomal Recessive Cerebellar Ataxias.**
Traschuets A., Reich S., Adarmes A. D., Anheim M., Ashrafi M. R., Baets J., Basak A. N., Bertini E., Brais B., Gagnon C., et al.
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- XXVIII. **A novel PSEN2 p.Ser175Phe variant in a family with Alzheimer's disease**
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- XXX. **Biallelic variants in TSPOAP1, encoding the active-zone protein RIMBP1, cause autosomal recessive dystonia**
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- XXXII. **Inflammation and regulatory T cell genes are differentially expressed in peripheral blood mononuclear cells of Parkinson's disease patients.**
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- XXXIII. **Functional Connectivity Analysis in Heterozygous Glucocerebrosidase Mutation Carriers**
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- XXXV. **Characterization of Recessive Parkinson Disease in a Large Multicenter Study**
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- XXXVI. **The cerebral blood flow deficits in Parkinson's disease with mild cognitive impairment using arterial spin labeling MRI.**
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- XXXVII. **Revisiting the complex architecture of ALS in Turkey: Expanding genotypes, shared phenotypes, molecular networks, and a public variant database**
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XXXVIII. **Clinico-Genetic, Imaging and Molecular Delineation of COQ8A-Ataxia: A Multicenter Study of 59 Patients**

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XL. **Caregiver Burden, Quality of Life and Related Factors in Family Caregivers of Dementia Patients in Turkey.**

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XLI. **Event-related potential changes due to early-onset Parkinson's disease in parkin (PARK2) gene mutation carriers and non-carriers**

USLU A., ERGEN M., Demirci H., Lohmann E., Hanagasi H., DEMİRALP T.

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XLVII. **A patient with early-onset Alzheimer's disease with a novel PSEN1 p.Leu424Pro mutation**

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L. **Using global team science to identify genetic parkinson's disease worldwide**

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- LII. Association between selected cholesterol-related gene polymorphisms and Alzheimer's disease in a Turkish cohort**
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- LIII. Patients with Latently Diagnosed Cerebrotendinous Xanthomatosis**
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- LIV. Amyloid Beta Adsorption Problem with Transfer Plates in Amyloid Beta 1-42 IVD Kits**
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- LXI. Clinical phenotype of hereditary spastic paraplegia due to KIF1C gene mutations across life span.**
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- LXIII. Nasu Hakola Disease: A Rare Cause of Dementia and Cystic Bone Lesions, Report of a New Turkish Family**

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- LXIV. **An assessment of Movement Disorder Society Task Force diagnostic criteria for mild cognitive impairment in Parkinson's disease**
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- LXV. **SYNE1 related cerebellar ataxia presents with variable phenotypes in a consanguineous family from Turkey**
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