Dr. Öğr. Üyesi Arman ÇAKAR

Kişisel Bilgiler

İş Telefonu: +90 0212 414 2000 Dahili: 32571
E-posta: arman.cakar@istanbul.edu.tr
Web: http://aves.istanbul.edu.tr/143/

Uluslararası Araştırmacı ID'leri
ORCID: 0000-0002-8129-4172
Publons / Web Of Science ResearcherID: AAE-5475-2020
Yoksis Araştırmacı ID: 190348

Araştırma Alanları

Nöroloji

Akademik Unvanlar / Görevler

Dr. Öğr. Üyesi, İstanbul Üniversitesi, İstanbul Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2020 - Devam Ediyor

SCI, SSCI ve AHCI İndekslerine Giren Dergilerde Yayınlanan Makaleler

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IV. **Phenotypic features of RETREG1-related hereditary sensory autonomic neuropathy**
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V. **Thymoma patients with or without myasthenia gravis have increased Th17 cells, IL-17 production and ICOS expression**
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MUSCLE & NERVE, cilt.66, sa.6, ss.736-743, 2022 (SCI-Expanded)

VIII. **Cerebellar ataxia, neuropathy and vestibular areflexia syndrome (canvas): an important cause of late-onset ataxia with unique clinical features**
ÇAKAR A., Sahin E., Tezel S., Candayan A., Samanci B., BATTALOĞLU E., Basak A. N., Bilgic B., Hanagasi H. A., Durmus H., et al.
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IX. **NOVEL VARIANTS BROADEN THE MUTATIONAL SPECTRUM OF HEREDITARY SENSORY AND AUTONOMIC NEUROPATHY DISORDERS**
Lischka A., Eggermann K., Cakar A., Record C., Elbracht M., Hornemann T., Senderek J., Parman Y., Auer-Grumbach M., Reilly M., et al.
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XIII. **Genetics of Pain: Novel variants identified by the European Network on Inherited Sensory Neuropathies and Insensitivity to Pain (ENISNIP)**
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XIV. **Lumbar Spinal Stenosis: A Rare Presentation of Hereditary Transthyretin Amyloidosis**
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XV. **An Exploratory Study of Cognitive Involvement in Hereditary Transthyretin Amyloidosis**
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XVI. **Cerebellar ataxia, neuropathy and vestibular areflexia syndrome (Canvas) is an important cause of late-onset ataxia**
Cakar A., Sahin E., Tezel S., Candayan A., Samanci B., BATTALOĞLU E., Basak A. N., Bilgic B., Hanagasi H., Durmus H., et al.
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XVII. **Thymoma associated myasthenia gravis patients have increased Th17 Cells, IL-17 production and ICOS expression on CD4 T cells**
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XVIII. **The Complex Genetic Landscape of Hereditary Ataxias in Turkey and Implications in Clinical Practice**
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XIX. **SOD1 Mutation: A Single Center Experience**
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XX. **Late-onset TK2-Deficiency Patients from Turkey**
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XXI. **Clinical Features of the Patients with Neuromyelitis Optica Spectrum Disorder**
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XXIV. **Coronavirus Disease 2019 (COVID-19) From the Point of View of Neurologists: Observation of Neurological Findings and Symptoms During the Combat Against a Pandemic**
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XXV. **CD4+ T Cells of Myasthenia Gravis Patients Are Characterized by Increased IL-21, IL-4, and IL-17A Productions and Higher Presence of PD-1 and ICOS**
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XXXI. **Orbital Myositis: An Underrecognized Clinical Syndrome with a Need of Management Guidelines**
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Diğer Dergilerde Yayınlanan Makaleler

I. **LETTER TO THE EDITOR Novel and nano-rare genetic causes of paediatric-onset motor neuronopathies**
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II. **Botulinum Toxin Treatment in Blepharospasm: Single-center Experience Blefarospazm Tedavisinde Botulinum Toksin: Tek-merkez Deneyimi**
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III. **Comparison of Mitoxantrone versus Cyclophosphamide Treatment in Patients with Secondary Progressive Multiple Sclerosis**
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IV. **Aquaporin-4 Gene Polymorphisms in Neuromyelitis Optica and Recurrent Optic Neuritis**
Çakar A., Ulusoy C. A., İplik E. S., Küçükali C. İ., Tüzün E., Kürtüncü M.
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V. **The COVID-19 from Neurological Overview**
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VII. **Transthyretin-Related Familial Amyloid Polyneuropathy: In the Light of New Developments**
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Kitap & Kitap Bölümleri

I. **Familial Amiloid Polinöropati - Dünya ve Türkiye Deneyimi Yeni Gelişmeler Işığında Transtiretin İlişkili Ailevi Amiloid Polinöropatisi**
ÇAKAR A., DURMUŞ TEKÇE H., DEYMEER F., OFLAZER P., PARMAN F. Y.
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Hakemli Kongre / Sempozyum Bildiri Kitaplarında Yer Alan Yayınlar

I. **Long-term follow-up of five families from Turkey with UBQLN2 variants**
Durmus H., ÇAKAR A., Aysal F., Ertas M., Basak N., Parman Y.
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II. **Paraplegia following lumbar puncture: a rare complication of spinal dural arteriovenous fistula**
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III. **An Exploratory Study of Cognitive Involvement in Hereditary Transthyretin Amyloidosis**
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IV. **YAYGIN HİPERPİGMENTASYON İLE BAŞVURAN BİR DERMATOMİYOZİT OLGUSU**
GEZEGEN H., ALTINKAYNAK M., ÇAKAR A., DURMUŞ TEKÇE H., AKPINAR T. S., PARMAN F. Y.
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V. **Pandemi Öncesi ve Pandemi Döneminde Guillain-Barré Sendromunun Klinik ve Elektrofizyolojik Özellikleri: Çok Merkezli İstanbul Çalışması**
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57. Ulusal Nöroloji Kongresi, Antalya, Türkiye, 27 Kasım - 01 Aralık 2021

VI. **Cognitive involvement in transthyretin-related familial amyloid polyneuropathy (TTR-FAP)**
DURMUŞ TEKÇE H., ÇAKAR A., Demirci H., PARMAN F. Y.
Virtual Annual Meeting of the American-Academy-of-Neurology, ELECTR NETWORK, 17 - 22 Nisan 2021

VII. **Expanding the phenotypical spectrum of SACS mutation: A Single Center Experience**
ÇAKAR A., İNCİ M., ÖZDAĞ ACARLI A. N., DURMUŞ TEKÇE H., PARMAN F. Y.
Virtual Annual Meeting of the American-Academy-of-Neurology, ELECTR NETWORK, 17 - 22 Nisan 2021

VIII. **WRITER'S CRAMP: A SINGLE-CENTER EXPERIENCE**
Cakar A., Tufekcioglu Z., Hanagasi H., Parman Y.
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IX. **Clinical features of a homozygous missense mutation in the FXN gene resulting in a Charcot-Marie-Tooth-like phenotype**
ÇAKAR A., Candayan A., Yunisova G., Battaloglu E., DURMUŞ TEKÇE H., PARMAN F. Y.
Virtual Conference of Peripheral-Nerve-Society, ELECTR NETWORK, 01 Ocak 2020, ss.490

X. **Cognitive involvement in ATTR amyloidosis (TTR-FAP)**
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XI. **Clinical and genetic features of SPG11: A Single Center Experience**
Cakar A., Gezegen H., Tunca C., Bayraktar E., Basak N., Durmus-Tekce H., Parman F. Y.
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XII. **Circulating B Cell Subsets and Cytokine Gene Expression Levels in Peripheral Blood and Skin Biopsy in Chronic Inflammatory Demyelinating Polyneuropathy**
Acarli A. N. O., Yilmaz V., Sirin N. G., ÇAKAR A., Soysal A., Aysal F., DURMUŞ TEKÇE H., Tuzun E., Parman Y.
Annual Meeting of the American-Academy-of-Neurology, Toronto, Kanada, 25 Nisan - 01 Mayıs 2020

XIII. **Skin Biopsy as a Biomarker in Chronic Inflammatory Demyelinating Polyneuropathy**
Parman Y., Acarli A. N. O., ÜNVERENGİL G., Sirin N. G., ÇAKAR A., DURMUŞ TEKÇE H.
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XIV. **Studying Clinical and Genetic Characteristics of Emery-Dreifuss Muscular Dystrophy**
Yunisova G., Oflazer P., Deymeer F., ÇAKAR A., PARMAN F. Y., DURMUŞ TEKÇE H.
Annual Meeting of the American-Academy-of-Neurology, Toronto, Kanada, 25 Nisan - 01 Mayıs 2020, cilt.94

XV. **Autosomal Recessive Charcot-Marie-Tooth Disease in Turkey**
Parman Y., Cakar A., Candayan A., Akcay H. I., Yunisova G., Ulukan Ç., Durmus-Tekce H., BATTALOĞLU E.
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XVI. **Comparison of Mitoxantrone versus Cyclophosphamide in Patients with Secondary Progressive Multiple Sclerosis**
GÜNDÜZ T., ÖZCAN G., ÇAKAR A., AKÇAY H. İ., ÖZDAĞ ACARLI A. N., KÜRTÜNCÜ M., ERAKSOY M.
2nd Congress of European Academy of Neurology, Kopenhag, Danimarka, 28 - 31 Mayıs 2016, cilt.23, ss.601-879

XVII. **Switching From Glatiramer Acetate: Comparison of Disease Modifying Treatments**
Kurtuncu M., Akcay H. I., Cakar A., Gunduz T., Tuzun E., Eraksoy M.
ACTRIMS Forum, Louisiana, Amerika Birleşik Devletleri, 18 - 20 Şubat 2016, cilt.22, ss.18-19

XVIII. **Clinical features of patients with relapsing inflammatory optic neuritis**
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Metrikler

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Akademi Dışı Deneyim

UCL Queen Square Institute of Neurology