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International Researcher IDs

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Education Information

Doctorate, Hacettepe University, Sağlık Bilimleri Enstitüsü, Tıbbi Biyoloji A.B.D., Turkey 2018 - Continues

Postgraduate, Hacettepe University, Sağlık Bilimleri Fakültesi, Nörolojik Bilimler Ve Psikiyatri Ens., Turkey 2013 - 2016

Expertise In Medicine, Hacettepe University, Tıp, Nöroloji, Turkey 2006 - 2011

Undergraduate, Hacettepe University, Tıp, Turkey 1999 - 2006

Foreign Languages

French, C1 Advanced

English, C1 Advanced

Research Areas

Medicine, Health Sciences, Internal Medicine Sciences, Neurology

Academic Titles / Tasks

Associate Professor, Hacettepe University, Tıp Fakültesi (Türkçe), Dahili Tıp Bilimleri Bölümü, 2018 - Continues

Researcher, Ohio State University, Columbus, Nationwide Children's Hospital, Center for Gene Therapy, 2021 - 2022

Assistant Professor, Hacettepe University, Dahili Tıp Bilimleri Bölümü, 2015 - 2018

Research Assistant, Hacettepe University, Nörolojik Bilimler ve Psikiyatri Enstitüsü, 2014 - 2015

Researcher, Hacettepe University, 2011 - 2011

Researcher, Bayerische Julius-Maximilians-Universitaet Würzburg, Medicine, Neuropathology Laboratory, 2011 - 2011

Research Assistant, Hacettepe University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2006 - 2011

Academic and Administrative Experience

Ethics Committee Member, Hacettepe University, Tıp Fakültesi (Türkçe), Dahili Tıp Bilimleri Bölümü, 2016 - Continues

Courses

Motor muayene, Undergraduate, 2017 - 2018, 2016 - 2017
Peripheral nerve disorders, Undergraduate, 2017 - 2018
motor examination, Undergraduate, 2017 - 2018, 2016 - 2017
good clinical practice, Undergraduate, 2016 - 2017
mental muayene, Undergraduate, 2016 - 2017

Published journal articles indexed by SCI, SSCI, and AHCI

- I. **Nusinersen for adults with spinal muscular atrophy**
ARSLAN D., İNAN B., KILINÇ M., Bekircan-Kurt C. E., ERDEM ÖZDAMAR S., Tan E.
Neurological Sciences, vol.44, no.7, pp.2393-2400, 2023 (SCI-Expanded)
- II. **The evaluation of small fibers in multiple sclerosis**
Bekircan-Kurt C. E., Jahanroshan J., Tuncer A., ERGÜL ÜLGER Z., Gunes G., ERDEM ÖZDAMAR S., Tan E.
Multiple Sclerosis and Related Disorders, vol.72, 2023 (SCI-Expanded)
- III. **Two distinct skeletal muscle microRNA signatures revealing the complex mechanism of sporadic ALS**
AKSU MENGEŞ E., Balci-Hayta B., Bekircan-Kurt C. E., Aydinoglu A. T., ERDEM ÖZDAMAR S., Tan E.
ACTA NEUROLOGICA BELGICA, vol.122, no.6, pp.1499-1509, 2022 (SCI-Expanded)
- IV. **Comprehensive evaluation of velopharyngeal function in myasthenia gravis patients**
BAŞTUĞ DUMBAK A., KULAK KAYIKCI M. E., ŞAHİN M. İ., KUŞCU O., Bekircan-Kurt C. E., ERDEM ÖZDAMAR S.
ACTA NEUROLOGICA BELGICA, vol.122, no.5, pp.1229-1236, 2022 (SCI-Expanded)
- V. **The functional and structural evaluation of small fibers in asymptomatic carriers of TTR p.Val50Met (Val30Met) mutation**
Bekircan-Kurt C. E., YILMAZ E., ARSLAN D., Yıldız F. G., DİKMETAŞ Ö., ERGÜL ÜLGER Z., KOCABEYOĞLU S., Irkec M., HEKİMSOY V., Tokgozoglu L., et al.
NEUROMUSCULAR DISORDERS, vol.32, no.1, pp.50-56, 2022 (SCI-Expanded)
- VI. **Eculizumab in refractory generalized myasthenia gravis previously treated with rituximab: subgroup analysis of REGAIN and its extension study**
Siddiqi Z. A., Nowak R. J., Mozaffar T., O'Brien F., Yountz M., Patti F., Mazia C. G., Wilken M., Wilken M., Barroso F., et al.
Muscle and Nerve, vol.64, no.6, pp.662-669, 2021 (SCI-Expanded)
- VII. **Biologia Futura: the importance of 3D organoids-a new approach for research on neurological and rare diseases**
AKBABÄ T. H., Bekircan-Kurt C. E., Balci-Peynircioglu B., Balci-Hayta B.
BIOLOGIA FUTURA, vol.72, no.3, pp.281-290, 2021 (SCI-Expanded)
- VIII. **The evaluation of small fibers in asymptomatic carriers of Val30Met mutation: Results of three years follow-up**
Bekircan-Kurt C. E., YILMAZ E., Aslan D., Yıldız F. G., DİKMETAŞ Ö., ERGÜL ÜLGER Z., KOCABEYOĞLU S., Irkec M., HEKİMSOY V., Tokgozoglu L., et al.
JOURNAL OF THE PERIPHERAL NERVOUS SYSTEM, vol.26, no.3, pp.379, 2021 (SCI-Expanded)
- IX. **One Disease with two Faces: Semidominant Inheritance of a Novel HTRA1 Mutation in a Consanguineous Family**
Bekircan-Kurt C. E., ÇETİNKAYA A., GÖÇMEN R., KOŞUKCU C., Soylemezoglu F., ARSAVA E. M., Tuncer A., ERDEM ÖZDAMAR S., Akarsu N. A., TOPÇUOĞLU M. A.
JOURNAL OF STROKE & CEREBROVASCULAR DISEASES, vol.30, no.9, 2021 (SCI-Expanded)
- X. **Reliability and Validity of Turkish Myasthenia Gravis-Activities of Daily Living Scale**
KARANFİL E., SALCI Y., FİL BALKAN A., Bekircan-Kurt C. E., ERDEM ÖZDAMAR S., ARMUTLU K.
OTJR-OCCUPATION PARTICIPATION AND HEALTH, vol.41, no.2, pp.101-107, 2021 (SSCI)
- XI. **Consistent improvement with eculizumab across muscle groups in myasthenia gravis**

- Mantegazza R., O'Brien F. L., Yountz M., Howard J. F., Gabriel Mazia C., Wilken M., Barroso F., Saba J., Rugiero M., Bettini M., et al.
Annals of Clinical and Translational Neurology, vol.7, no.8, pp.1327-1339, 2020 (SCI-Expanded)
- XII. Trigeminal sensory-motor neuropathy in a patient with mixed connective tissue disease and review of the literature**
 Bekircan-Kurt C. E., TEMUÇİN Ç. M., APRAŞ BİLGEN Ş. Ş., ERDEM ÖZDAMAR S.
NEUROLOGICAL SCIENCES AND NEUROPHYSIOLOGY, vol.37, no.3, pp.148-151, 2020 (SCI-Expanded)
- XIII. A Novel Amplification-Refractory Mutation System-PCR Strategy to Screen MT-TL1 Pathogenic Variants in Patient Repositories**
 Bulduk B. K., Kilic H. B., Bekircan-Kurt C. E., Haliloglu G., ERDEM ÖZDAMAR S., Topaloglu H., KOCAEFE Y. Ç.
GENETIC TESTING AND MOLECULAR BIOMARKERS, vol.24, no.3, pp.165-170, 2020 (SCI-Expanded)
- XIV. Giant cell myositis associated with myasthenia gravis and thymoma**
 İNAN B., KURT C. E., Yıldız F. G., ERDEM ÖZDAMAR S., TEMUÇİN Ç. M., Tan E.
NEUROLOGICAL SCIENCES AND NEUROPHYSIOLOGY, vol.36, no.4, pp.236-237, 2019 (SCI-Expanded)
- XV. Laboratory diagnosis of metachromatic leukodystrophy requires more than arylsulfatase A assay**
 Pekgül F., Bekircan-Kurt C. E., Konuskan B., Erdem-Ozdamar S., Tan E., Akarsu N., Topcu M., Anlar B., Ozkara H. A.
FEBS OPEN BIO, vol.9, pp.199, 2019 (SCI-Expanded)
- XVI. Two sisters with anti-MuSK-positive myasthenia gravis**
 KURT E., Bekircan-Kurt C. E., KONUŞKAN B., ERKENT ERCAN İ., Tan E., Anlar B.
CLINICAL NEUROLOGY AND NEUROSURGERY, vol.182, pp.17-18, 2019 (SCI-Expanded)
- XVII. Long-term safety and efficacy of eculizumab in generalized myasthenia gravis**
 Muppidi S., Utsugisawa K., Benatar M., Murai H., Barohn R. J., Illa I., Jacob S., Vissing J., Burns T. M., Kissel J. T., et al.
Muscle and Nerve, vol.60, no.1, pp.14-24, 2019 (SCI-Expanded)
- XVIII. Establishment of primary myoblast cell cultures from cryopreserved skeletal muscle biopsies to serve as a tool in related research & development studies**
 Balci-Hayta B., Bekircan-Kurt C. E., AKSU MENGEŞ E., DAYANGAÇ ERDEN D., Tan E., ERDEM ÖZDAMAR S.
JOURNAL OF THE NEUROLOGICAL SCIENCES, vol.393, pp.100-104, 2018 (SCI-Expanded)
- XIX. The histopathological evaluation of small fiber neuropathy in patients with vitamin B12 deficiency**
 Gunes H. N., Bekircan-Kurt C. E., Tan E., ERDEM ÖZDAMAR S.
ACTA NEUROLOGICA BELGICA, vol.118, no.3, pp.405-410, 2018 (SCI-Expanded)
- XX. Ocular surface alterations and in vivo confocal microscopic characteristics of corneas in patients with myasthenia gravis**
 ERKAN TURAN K., KOCABEYOĞLU S., Bekircan-Kurt C. E., BEZCİ F., ERDEM ÖZDAMAR S., Irkec M.
EUROPEAN JOURNAL OF OPHTHALMOLOGY, vol.28, no.5, pp.541-546, 2018 (SCI-Expanded)
- XXI. ERLIN1 mutations cause teenage-onset slowly progressive ALS in a large Turkish pedigree**
 Tunca C., Akcimen F., Coskun C., Gundogdu-Eken A., Kocoglu C., ÇEVİK B., Bekircan-Kurt C. E., Tan E., Basak A. N.
EUROPEAN JOURNAL OF HUMAN GENETICS, vol.26, no.5, pp.745-748, 2018 (SCI-Expanded)
- XXII. Safety and efficacy of eculizumab in anti-acetylcholine receptor antibody-positive refractory generalised myasthenia gravis (REGAIN): a phase 3, randomised, double-blind, placebo-controlled, multicentre study**
 Howard J. F., Utsugisawa K., Benatar M., Murai H., Barohn R. J., Illa I., Jacob S., Vissing J., Burns T. M., Kissel J. T., et al.
The Lancet Neurology, vol.16, no.12, pp.976-986, 2017 (SCI-Expanded)
- XXIII. Myophosphorylase (PYGM) mutations determined by next generation sequencing in a cohort from Turkey with McArdle disease**
 Inal-Gultekin G., Toptas-Hekimoglu B., Gormez Z., Gelisin O., Durmus H., Erguner B., Demirci H., Sagiroglu M. S., Parman Y., Deymeer F., et al.
NEUROMUSCULAR DISORDERS, vol.27, no.11, pp.997-1008, 2017 (SCI-Expanded)
- XXIV. CELLULAR INFILTRATES IN SKIN AND SURAL NERVE OF PATIENTS WITH POLYNEUROPATHIES**
 Ueçeyler N., Braunsdorf S., Kunze E., Riediger N., Scheytt S., Divisova S., Bekircan-Kurt C. E., Toyka K. V., Sommer C.
MUSCLE & NERVE, vol.55, no.6, pp.884-893, 2017 (SCI-Expanded)
- XXV. New mutations and genotype-phenotype correlation in late-onset Pompe patients**

- Bekircan-Kurt C. E., Gunes H. N., Yildiz F. G., SAKA TOPÇUOĞLU E., Tan E., ERDEM ÖZDAMAR S.
 ACTA NEUROLOGICA BELGICA, vol.117, no.1, pp.269-275, 2017 (SCI-Expanded)
- XXVI. Single-pulse Transcranial Magnetic Stimulation in Amyotrophic Lateral Sclerosis: Experience of a single institution**
 Yildiz F. G., Bekircan-Kurt C. E., VARLI K.
 JOURNAL OF NEUROLOGICAL SCIENCES-TURKISH, vol.34, no.1, pp.70-75, 2017 (SCI-Expanded)
- XXVII. The evaluation of small fibres in asymptomatic patients with Val30Met mutation**
 Bekircan-Kurt C. E., Yildiz F. G., Gunes H. N., Erdem-Ozdamar S., Tan E.
 EUROPEAN JOURNAL OF NEUROLOGY, vol.23, pp.231, 2016 (SCI-Expanded)
- XXVIII. Autoimmune storm**
 Aslan S., Bekircan-Kurt C. E., Kurne A. T., Erdem-Ozdamar S.
 EUROPEAN JOURNAL OF NEUROLOGY, vol.23, pp.437, 2016 (SCI-Expanded)
- XXIX. Spinal muscular atrophy type III: Molecular genetic characterization of Turkish patients**
 BORA-TATAR G., YESBEK-KAYMAZ A., BEKIRCAN-KURT C. E., ERDEM-OZDAMAR S., Erdem-Yurter H.
 EUROPEAN JOURNAL OF MEDICAL GENETICS, vol.58, no.12, pp.654-658, 2015 (SCI-Expanded)
- XXX. Three Turkish families with different transthyretin mutations**
 Bekirean-Kurt C. E., GUNES N., Yilmaz A., ERDEM-OZDAMAR S., TAN E.
 NEUROMUSCULAR DISORDERS, vol.25, no.9, pp.686-692, 2015 (SCI-Expanded)
- XXXI. The Activation of RAGE and NF-KB in Nerve Biopsies of Patients with Axonal and Vasculitic Neuropathy**
 Bekircan-Kurt C. E., TAN E., OZDAMAR S. E.
 NOROPSİKIYATRI ARSIVİ-ARCHIVES OF NEUROPSYCHIATRY, vol.52, no.3, pp.279-282, 2015 (SCI-Expanded)
- XXXII. Voltage gated calcium channel antibody-related neurological diseases.**
 Bekircan-Kurt C. E., Derle Ç., KURNE A., ANLAR B.
 World journal of clinical cases, vol.3, pp.293-300, 2015 (SCI-Expanded)
- XXXIII. CUTANEOUS ACTIVATION OF RAGE IN NONSYSTEMIC VASCULITIC AND DIABETIC NEUROPATHY**
 Bekircan-Kurt C. E., Ueceyler N., SOMMER C.
 MUSCLE & NERVE, vol.50, no.3, pp.377-383, 2014 (SCI-Expanded)
- XXXIV. The Course of Myasthenia Gravis with Systemic Lupus Erythematosus**
 Bekircan-Kurt C. E., Kurne A. T., Erdem-Ozdamar S., Kalyoncu U., Karabudak R., Tan E.
 EUROPEAN NEUROLOGY, vol.72, pp.326-329, 2014 (SCI-Expanded)
- XXXV. 'Jaw clenching' in anti-Ri - Antibody-associated paraneoplastic syndrome**
 Bekircan-Kurt C. E., TEMUÇİN Ç. M., ELİBOL B., SAKA E.
 PARKINSONISM & RELATED DISORDERS, vol.19, no.1, pp.132-133, 2013 (SCI-Expanded)
- XXXVI. Myasthenia Gravis; Single Entity, Variable Clinical Features: Ten Years of Clinical Experience in a Tertiary Care Center Ten Years Clinical Experience of a Tertiary Care Center**
 BEKIRCAN-KURT C. E., KURNE A., ERDEM ÖZDAMAR S., KARABUDAK R., KANSU T., TAN E.
 JOURNAL OF NEUROLOGICAL SCIENCES-TURKISH, vol.30, no.1, pp.135-143, 2013 (SCI-Expanded)

Articles Published in Other Journals

- I. Identifying a Cut-off Point for Timed Up and Go Test in Neuromuscular Diseases
 Demirci C. S., SÜTÇÜ G., AYVAT F., ONURSAL KILINÇ Ö., DOĞAN M., AYVAT E., KURT C. E., ERDEM ÖZDAMAR S., Yıldırım S. A., KILINÇ M., et al.
 TURKISH JOURNAL OF NEUROLOGY, vol.28, no.1, pp.6-9, 2022 (ESCI)
- II. A Rare Cause of Dystonia: Spinal Meningioma
 Sarayli A. A. K., Bekircan-Kurt C. E., GÖÇMEN R., ÖGE H. K., Tan E.
 TURKISH JOURNAL OF NEUROLOGY, vol.27, no.2, pp.215-216, 2021 (ESCI)
- III. Organelle Positioning in Neurons and Skeletal Muscle Cells
 AKSU MENGEŞ E., KURAL MANGIT E., Bekircan-Kurt C. E., BORA G.

- GAZI MEDICAL JOURNAL, vol.32, pp.341-347, 2021 (ESCI)
- IV. Psychosocial Adjustment and Adherence to Medication in Patients with Myasthenia Gravis**
 AŞİRET G. D., KAPUCU S., KAYMAZ T. T., Bekircan-Kurt C. E.
 GAZI MEDICAL JOURNAL, vol.32, no.3, pp.371-376, 2021 (ESCI)
- V. Dysferlinopathy: A Case Report and Literature Update**
 Kutlu O., KURT C. E., Unsal I., Aribas Z., Renkliyildiz B., Eruzun H., Karagulmez A. D., ERDEM ÖZDAMAR S.
 İSTANBUL MEDICAL JOURNAL, vol.17, no.4, pp.136-140, 2016 (ESCI)

Refereed Congress / Symposium Publications in Proceedings

- I. NUSİNERSEN TEDAVİSİ ALAN ERİŞKİN SPİNAL MUSKÜLER ATROFİ HASTALARINDA İNFLAMASYON VE GLİAL HÜCRE İLİŞKİSİ BİYOBELİRTEÇLERİN TESPİTİ**
 ARSLAN D., ERGÜL ÜLGER Z., gökşen s., ESENDAĞLI G., ERDEM ÖZDAMAR S., TAN M. E., KURT C. E.
 59. Ulusal Nöroloji Kongresi, Antalya, Turkey, 13 December 2023
- II. MİYASTENİK KRİZ İLE BAŞVURAN HASTALARDA YARDIMCI FOLİKÜLER T HÜCRE (TFH) VE YARDIMCI T HÜCRE 17 (TH17) İLİŞKİSİ SİTOKİNLERİN TEDAVİ ÖNCESİ VE SONRASI DEĞİŞKENLİĞİ**
 ARSLAN D., ERGÜL ÜLGER Z., gökşen s., ESENDAĞLI G., ERDEM ÖZDAMAR S., TAN M. E., KURT C. E.
 59. Ulusal Nöroloji Kongresi, Antalya, Turkey, 13 December 2023
- III. Evaluation of the Role of Glial Factors in the Pathogenesis of Spinal Muscular Atrophy**
 ARSLAN D., ERGÜL ÜLGER Z., gökşen s., İNAN B., ESENDAĞLI G., ERDEM ÖZDAMAR S., TAN M. E., KURT C. E.
 2023 Neuromuscular Study Group Meeting, Orlando, United States Of America, 22 September 2023
- IV. Serum Cytokines Levels in Patients with Myasthenia Gravis and Their Changes After Treatment**
 ARSLAN D., ERGÜL ÜLGER Z., gökşen s., ESENDAĞLI G., ERDEM ÖZDAMAR S., TAN M. E., KURT C. E.
 75th Annual Meeting of the American-Academy-of-Neurology (AAN), Boston, United States Of America, 22 April 2023
- V. MME MUTASYONUNA BAĞLI DEMİYELİNİZAN NÖROPATİ**
 Demirel Özbeğ E., ERGÜL ÜLGER Z., TAN M. E., ERDEM ÖZDAMAR S., KURT C. E.
 58. Ulusal Nöroloji Kongresi, Antalya, Turkey, 19 - 24 November 2022
- VI. The evaluation of small fibers in asymptomatic carriers of Val30Met mutation: Results of three years follow-up**
 KURT C. E., YILMAZ E., aslan d., YILDIZ SARIKAYA F. G., DİKMETAŞ Ö., ERGÜL ÜLGER Z., KOCABEYOĞLU S., irkec m., HEKİMSOY V., TOKGÖZÖĞLU S. L., et al.
 2021 PNS Annual Meeting, United States Of America, 12 - 26 June 2021, vol.26, pp.379-179
- VII. Multimodal Assessment of Intensive Care Unit- Acquired Weakness in Severe Stroke Patients**
 İNAN B., KURT C. E., ERGÜL ÜLGER Z., yılmaz m.
 2021 American Academy of Neurology, 2 - 10 April 2021
- VIII. The effect and challenges of nusinersen treatment in adult spinal muscular atrophy patients - preliminary results**
 Inan B., Bekircan-Kurt C. E., Kilinc M., Erdem-Ozdamar S., Tan E.
 6th Congress of the European-Academy-of-Neurology (EAN), ELECTR NETWORK, 23 - 26 May 2020, vol.27, pp.625
- IX. Erişkin Kas Hastalarında Fiziksel Aktiviteyi Objektif ve Subjektif Olarak Değerlendiren Yöntemlerin Karşılaştırılması**
 AYVAT F., AYVAT E., ONURSAL KILINÇ Ö., SÜTCÜ G., DOĞAN M., KILINÇ M., AKSU YILDIRIM S., KURT C. E., ERDEM ÖZDAMAR S., TAN M. E.
 TND II. Nöromusküler Hastalıklar Kongresi, İzmir, Turkey, 19 - 21 April 2019
- X. Nusinersen İlacı İçin Başvuruda Bulunan Spinal Musküler Atrofi Tip 2 ve Tip 3 Tanılı Hastaların Profili**
 ONURSAL KILINÇ Ö., AYVAT F., SÜTCÜ G., DOĞAN M., AYVAT E., KILINÇ M., AKSU YILDIRIM S., KURT C. E., ERDEM ÖZDAMAR S., TAN M. E.
 TND II. Nöromusküler Hastalıklar Kongresi, İzmir, Turkey, 19 - 21 April 2019

- XI. **Kas Hastalarında Farklı Yüksekliklerden Oturmadan Ayağa Kalkma Sırasında Uyluk Kaslarının Aktivasyon Seviyelerinin İncelenmesi**
SÜTÇÜ G., DOĞAN M., ONURSAL KILINÇ Ö., AYVAT F., AYVAT E., AKSU YILDIRIM S., KILINÇ M., KURT C. E., ERDEM ÖZDAMAR S., TAN M. E.
TND II. Nöromusküler Hastalıklar Kongresi, İzmir, Turkey, 19 - 21 April 2019
- XII. **Erişkin Nöromusküler Hastalarda Düşme Öyküsünün AktiviteLimitasyonu ve Gövde Bozukluk Düzeyi ile İlişkisinin İncelenmesi**
DOĞAN M., ONURSAL KILINÇ Ö., AYVAT F., SÜTÇÜ G., AYVAT E., KILINÇ M., AKSU YILDIRIM S., KURT C. E., ERDEM ÖZDAMAR S., TAN M. E.
TND II. Nöromusküler Hastalıklar Kongresi, İzmir, Turkey, 19 - 21 April 2019
- XIII. **Erişkin Kas Hastalarında Fiziksel Aktivite ile İlişkili Faktörlerin Belirlenmesi**
AYVAT F., AYVAT E., ONURSAL KILINÇ Ö., SÜTÇÜ G., DOĞAN M., KILINÇ M., AKSU YILDIRIM S., KURT C. E., ERDEM ÖZDAMAR S., TAN M. E.
TND II. Nöromusküler Hastalıklar Kongresi, İzmir, Turkey, 19 - 21 April 2019
- XIV. **Differentially Expressed microRNA Profile in Skeletal Muscle Tissue of Sporadic ALS Patients**
Bekircan-Kurt C. E., AKSU MENGEŞ E., Balcı-Hayta B., Aydinoglu A. T., Erdem-Ozdamar S., Tan E.
71st Annual Meeting of the American-Academy-of-Neurology (AAN), Pennsylvania, United States Of America, 4 - 10 May 2019, vol.92
- XV. **The expression of epidermal growth factor (EGF) and keratinocyte growth factor (KGF) in skin biopsy of amyotrophic lateral sclerosis patients**
KURT E., Bekircan-Kurt C. E., ERDEM ÖZDAMAR S., Tan E.
70th Annual Meeting of the American-Academy-of-Neurology (AAN), Los-Angeles, Chile, 21 - 27 April 2018, vol.90
- XVI. **Myastenia gravis hastalarında altı dakika yürüyüş testinin kullanımı**
SALCI Y., KARANFİL E., ÇALIK KÜTÜKCÜ E., FİL BALKAN A., CEREN A. N., AYVAT F., KURT C. E., ARMUTLU K.
1. Kardiyopulmoner Rehabilitasyon Kongresi, Turkey, 22 - 25 November 2017, vol.29
- XVII. **Myastenia gravis hastalarının hastalığa psikososyal uyumu ve ilaç uyumlarının belirlenmesi.**
DURU AŞİRET G., KAPUCU S., TÜRTEM KAYMAZ T., KURT C. E.
. 5. Uluslararası, 16. Ulusal Hemşirelik Kongresi, Ankara, Turkey, 5 - 08 November 2017
- XVIII. **VECTOR-BORNE VIRAL INFECTIONS IN GUILLAIN BARRE SYNDROME PATIENTS**
Okar S., ERGÜNAY K., Bekircan-Kurt C. E., Erdem-Ozdamar S., Tan E.
Peripheral-Nerve-Society Meeting, Sitges, Spain, 8 - 12 July 2017, vol.22, pp.349-350
- XIX. **Acute myopathy in patients who are using short-term and low-dose systemic glucocorticoids: Preliminary findings**
İLBAY A., DURUSU TANRIÖVER M., KURT C. E., YILDIZ SARIKAYA F. G., ÖZİŞİK L., ERDEM ÖZDAMAR S.
16th European Congress of Internal Medicine, 31 August - 02 September 2017
- XX. **Recurrent peripheral and central demyelination in a seronegative patient**
KURT C. E., YILDIZ SARIKAYA F. G., TEMUÇİN Ç. M., TUNCER M. A., TAN M. E., ERDEM ÖZDAMAR S.
peripheral Nerve Society Meeting, 8 July - 13 September 2017
- XXI. **Low levels of SMN cause alterations in microtubule associated protein 1b**
BORA G., Rademacher S., YEŞBEK KAYMAZ A., Hensel N., KURT C. E., ERDEM ÖZDAMAR S., TOPALOĞLU H. A., Claus P., YURTER H.
21th Annual Spinal Muscular Atrophy Researcher Meeting, Orlando, United States Of America, 29 June - 02 July 2017
- XXII. **The Histopathological Evaluation of Small Fiber Neuropathy in Patients with Vitamin B12 Deficiency**
Bekircan-Kurt C. E., Gunes N., DEMİRCİ M., ERDEM ÖZDAMAR S., Tan E.
68th Annual Meeting of the American-Academy-of-Neurology (AAN), Vancouver, Canada, 15 - 21 April 2016, vol.86
- XXIII. **Myastenia Gravis de Velofaringeal Fonksiyonun Araştırılması**
BAŞTUĞ A., KULAK KAYIKCI M. E., KUŞCU O., KURT C. E., ERDEM ÖZDAMAR S.
4. YUTMA BOZUKLUKLARI KONGRESİ, Turkey, 19 - 21 November 2015
- XXIV. **A HEREDITARY AMYLOID POLYNEUROPATHY FAMILY DUE TO VAL30MET MUTATION WITH AN**

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Bekircan-Kurt C. E., ERDEM-OZDAMAR S., TAN E.

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- XXV. The incidence of Pompe disease in patients with vacuolar myopathy: a pilot study of a tertiary care center**

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- XXVI. A RARE CAUSE OF LUMBOSACRAL PLEXOPATHY: BLADDER DIVERTICULUM**

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- XXVII. A novel mutation in the HTRA1 gene in a Turkish family with CARASIL**

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- XXIX. Three families with hereditary amyloid polyneuropathy with three different mutations**

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Supported Projects

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ERGÜNAY K., ERDEM ÖZDAMAR S., KURT C. E., Project Supported by Higher Education Institutions, Guillain-Barre Hastalarında Vektör Kaynaklı Viral Enfeksiyonlar, 2017 - 2020

BALCI B., KURT C. E., AKSU E., AYDINOĞLU A. T., PEYNİRCİOĞLU B., Project Supported by Higher Education Institutions, İnfamasyon ile İlişkili Nöromusküler Hastalıklarda Dolaşımındaki Mitokondriyal DNA'nın Araştırılması, 2018 - 2019

ÖZKARA H. A., KURT C. E., ERDEM ÖZDAMAR S., TAN M. E., KONUŞKAN B., KARAAĞAOĞLU A. E., PEKGÜL F., TOPÇU M., ANLAR B., AKARSU A. N., Project Supported by Higher Education Institutions, Metakromatik lökodistrofi hastlığının alt tiplerinin tanımlanması, patojenik mutasyonların belirlenmesi, patogenezde inflamasyonun incelenmesi, 2017 - 2019

TAN M. E., ERDEM ÖZDAMAR S., KURT C. E., KURT E., Project Supported by Higher Education Institutions, Amiyotrofik Lateral Skleroz hastalarının deri biyopsilerinde Epidermal Büyüme Faktörü (EGF) ve Keratinosit Büyüme Faktörünün (KGF) ekspresyonunun değerlendirilmesi, 2016 - 2017

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Scholarships

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