Res. Asst. MEHMET BERKAY AKCAN

Personal Information

Email: berkay.akcan@comu.edu.tr

International Researcher IDs
ORCID: 0000-0003-0160-0377
Yoksis Researcher ID: 313197

Research Areas

Medicine

Academic Titles / Tasks

Research Assistant PhD, Canakkale Onsekiz Mart University, Tıp Fakültesi, Dahili Tıp Bilimleri, 2019 - Continues

Published journal articles indexed by SCI, SSCI, and AHCI

I. **Evaluating of colchicine use patterns and attack frequency of familial Mediterranean fever patients in the COVID-19 pandemic**
AKCAN M. B., Albuz B., Ozdemir O., SILAN F.
International Journal of Rheumatic Diseases, vol.26, no.5, pp.988-991, 2023 (SCI-Expanded)

II. **Anemia and thrombocytopenia due to a novel BRPF1 variant in a family from Çanakkale with intellectual disability and dysmorphic facies: Case report and review of the literature**
Ceylan Köse C., Kaya D., Akcan M. B., Sılan F.
AMERICAN JOURNAL OF MEDICAL GENETICS, PART A, no.6, pp.1-6, 2023 (SCI-Expanded)

III. **A New Case of Rare Microdeletion 10q22.3q23 along with Mosaic Klinefelter Syndrome Associated with Facial Dysmorphic Finding, Atrial Ventricular Septal Defect, and Motor Retardation**
Dincsoy Bir F., SILAN F., Velickovic J., Berkay Akcan M. B., ÖZDEMİR Ö.
MOLECULAR SYNDROMOLOGY, vol.13, no.3, pp.254-260, 2022 (SCI-Expanded)

Articles Published in Other Journals

I. **Familial intragenic X-linked OPHN1 gene deletion in a newborn male infant with low birth weight and distinctive facial appearance that diagnosed by advanced microarray-CGH method**
Aylanc H., Sılan F., Çokyaman T., Akcan M. B., Özdemir Ö.
Cumhuriyet Tıp Dergisi (ELEKTRONİK), vol.44, no.1, pp.125-130, 2022 (Peer-Reviewed Journal)

Refereed Congress / Symposium Publications in Proceedings

I. **Oculocutaneous albinism caused by a compound heterozygous mutations in TYR gene: a case report**
AKCAN M. B., BAKIOĞLU KAYA D., SÖNMEZ V., KABLAN A., ÖZDEMİR Ö., SILAN F.
Ulusal tıbbi genetik kongresi 2020, Online, Turkey, 20 November 2020

II. **Multiple pterygia, joint contractures, hypoplastic lungs and hydrops: A rare case of lethal multiple pterygium syndrome**
BAKIOĞLU KAYA D., AKCAN M. B., SÖNMEZ V., KABLAN A., ÖZDEMİR Ö., SILAN F.
Ulusal tıbbi genetik kongresi 2020, Online, Turkey, 20 November 2020

III. **A novel variant for SQSTM1 gene in a patient with frontotemporal dementia.**
OCAK Ö., KARAKAYA T., ALBUZ B., AKBAŞ N. E., SÖNMEZ V., AKCAN M. B., SILAN F., ÖZDEMİR Ö.
European Biotechnology Congress 2020, Prag, Czech Republic, 24 September 2020

IV. **A Case Of Coffin-Siris Syndrome With Atypical Phenotype Caused By A Novel De Novo Mutation In ARID1B Gene.**
ALBUZ B., AKBAŞ N. E., KARAKAYA T., SÖNMEZ V., AKCAN M. B., SILAN F., ÖZDEMİR Ö.
European Biotechnology Congress 2020, Prag, Czech Republic, 24 September 2020

V. **Chromatin remodelling dysfunction and CHD2 related epilepsy:Reports of two unrelated child.**
KABLAN A., ALBUZ B., AKBAŞ N. E., KARAKAYA T., SÖNMEZ V., AKCAN M. B., SILAN F., ÖZDEMİR Ö.
European Biotechnology Congress 2020, Prag, Czech Republic, 24 September 2020

VI. **A rare lethal multiple pterygium syndrome caused by a homozygous point mutation In CHRNG gene:a case report.**
AKCAN M. B., ALBUZ B., KABLAN A., ÖZDEMİR Ö., SILAN F.
European Biotechnology Congress 2020, Czech Republic, 24 September 2020

Metrics

Publication: 13