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Education Information

Doctorate, Baskent University, Institute Of Health Scıences, Tıbbi Genetik (Dr), Turkey 2018 - Continues
Undergraduate, Anadolu University, Faculty Of Busıness Admınıstratıon, Department Of Busıness, Turkey 2009 - 2012
Associate Degree, Anadolu University, Open Educatıon Faculty, Department Of Management And Organızatıon, Turkey 2007 - 2009
Associate Degree, Anadolu University, Open Educatıon Faculty, Department Of Management And Organızatıon, Turkey 2007 - 2009
Doctorate, Bursa Uludağ University, TIP FAKÜLTESİ, TEMEL TIP BİLİMLERİ, Turkey 1996 - 2002

Foreign Languages

German, B1 Intermediate
English, C1 Advanced
Italian, B1 Intermediate

Research Areas

Health Sciences, Natural Sciences, Engineering and Technology

Academic Titles / Tasks

Professor, Bursa Uludağ University, TIP FAKÜLTESİ, DAHİLİ TIP BİLİMLERİ, 2022 - Continues
Associate Professor, Bursa Uludağ University, TIP FAKÜLTESİ, DAHİLİ TIP BİLİMLERİ, 2013 - 2022

Academic and Administrative Experience

Head of Department, Bursa Uludağ University, TIP FAKÜLTESİ, 2018 - 2021
Head of Department, Bursa Uludağ University, TIP FAKÜLTESİ, DAHİLİ TIP BİLİMLERİ, 2017 - 2019

Courses

Courses

Translasyonel Tıp, Doctorate, 2017 - 2018
Tıbbi Genetik, Undergraduate, 2017 - 2018

Published journal articles indexed by SCI, SSCI, and AHCI

I. **Homozygous missense variants in YKT6 result in loss of function and are associated with developmental delay, with or without severe infantile liver disease and risk for hepatocellular carcinoma.**
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XI. **An Endocrinological Perspective on 22q11.2 Deletion Syndrome: A Single-center Experience.**
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XXVII. **Psoriasis and 5HT-R2C Gene Polymorphism: Association between Clinical, Demographic and Therapeutic Parameters in the Turkish Population.**
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XXVIII. **BRCA Variations Risk Assessment in Breast Cancers Using Different Artificial Intelligence Models**
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XXXVIII. **Biallelic mutations in M1AP are associated with meiotic arrest, severely impaired spermatogenesis and male infertility**
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XXXIX. **Identification of unsolved rare genetic cases of North Cyprus**
Ergoren M. C., Manara E., Paolacci S., Tuncel G., TEMEL Ş. G., Mocan G., DÜNDAR M., Bertelli M.
EUROPEAN JOURNAL OF HUMAN GENETICS, vol.28, pp.944, 2020 (SCI-Expanded)

XL. **Functional coding/non-coding variants in EGFR, ROS1 and ALK genes and their role in liquid biopsy as a personalized therapy.**
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XLI. **CC2D1A AS A NOVEL CILIOPATHY GENE**
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XLII. **Unique combination and in silico modeling of biallelic POLR3A variants as a cause of Wiedemann-Rautenstrauch syndrome**
TEMEL Ş. G., Ergoren M. C., Manara E., Paolacci S., Tuncel G., Gul S., Bertelli M.
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XLIII. **Diagnostic Efficiency of Clinical Exome Solution Panel in patients with Hearing loss/Hereditary Deafness by using Next Generation Sequencing**
Temel Ş. G., Alemdar A., Yılmaz M., Aliyeva L., Sag S. O.
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Pirim D., Kaya N., Yıldırım E., Sag S., Temel Ş. G.
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