

## Res. Asst. MERT COŞKUN

### Personal Information

Office Phone: [+90 242 249 6427](tel:+902422496427)

Office Phone: [+90 242 249 6536](tel:+902422496536)

Email: [mertcoskun@akdeniz.edu.tr](mailto:mertcoskun@akdeniz.edu.tr)

Web: <https://avesis.akdeniz.edu.tr/mertcoskun>

### Education Information

Expertise In Medicine, Akdeniz University, Faculty Of Medicine, Tıbbi Genetik Anabilim Dalı, Turkey 2019 - Continues  
Postgraduate, Ege University, Faculty Of Medicine, Turkey 2012 - 2018

### Research Areas

Medicine, Health Sciences, Internal Medicine Sciences, Medical Genetics

### Academic Titles / Tasks

Research Assistant PhD, Akdeniz University, Faculty of Medicine, Department of Internal Medicine, 2019 - Continues

### Published journal articles indexed by SCI, SSCI, and AHCI

- I. **Real-Life Experience of Dose-Adjusted Venetoclax in Acute Myeloid Leukemia Patients Concomitantly Using Posaconazole for Antifungal Prophylaxis: A Single-Center Experience**  
ILTAR U., Vural E., ALHAN F. N., Vurgun S., Atas U., YÜCEL O. K., SALİM O., Ulas T., COŞKUN M., TOYLU A., et al.  
UHOD - Uluslararası Hematoloji-Onkoloji Dergisi, vol.33, no.2, pp.75-82, 2023 (SCI-Expanded)
- II. **Clinical and molecular evaluation of MEFV gene variants in the Turkish population: a study by the National Genetics Consortium**  
Dündar M., Fahrioglu U., Yildiz S. H., Bakır-Güngör B., Temel Ş. G., Akın H., Artan S., Cora T., Şahin F. İ., Dursun A., et al.  
FUNCTIONAL & INTEGRATIVE GENOMICS, vol.22, no.3, pp.291-315, 2022 (SCI-Expanded)

### Refereed Congress / Symposium Publications in Proceedings

- I. **NADİR BİR BOY KISALIĞI NEDENİ: 3M SENDROMLU 2 OLGU**  
Öztürk N., Coşkun M., Karamık G., Yılmaz Bayer Ö., Parlak M., Nur B., Mıhçı E.  
5. EGE ENDOKRİN HASTALIKLAR VE GENETİK SEMPOZYUMU, İzmir, Turkey, 9 - 11 March 2023
- II. **TMC6 GENİNDE NADİR MİSSENSE VARYANT SAPTANAN EPİDERMODİSPLAZİ VERRÜSİFORMİS OLGUSU**  
Coşkun M., Toyulu A., Altıok Clark Ö., Nur B., Alpsoy E.  
15.ULUSAL TIBBİ GENETİK KONGRESİ, Muğla, Turkey, 9 - 13 November 2022
- III. **Manifestations of a heterozygous CDK13 mutation as a rare entity; congenital heart defects,**

**dysmorphic facial features, and intellectual developmental disorder**

PEKER A., Karamık G., COŞKUN M., Öztürk N., NUR B., MIHÇI E.

EURODYSMORPHO, Barcelona, Spain, 14 - 17 September 2022

**IV. NOONAN SENDROMLU ÇOCUKLARDA GENOTİP-FENOTİP İLİŞKİSİ**

Nacak G., Coşkun M., Toylu A., Altıok Clark Ö., Mihçi E., Nur B.

57.TPK, Gazimagusa, Cyprus (Kktc), 22 - 26 May 2022

## **Metrics**

Publication: 17

Citation (WoS): 4

Citation (Scopus): 5

H-Index (WoS): 1

H-Index (Scopus): 1

## **Congress and Symposium Activities**

6. Ulusal Çocuk Genetik Kongresi, Attendee, Aydın, Turkey, 2023

15. Ulusal Tıbbi Genetik Kongresi , Attendee, Muğla, Turkey, 2022

European Human Genetics Conference (ESHG 2022), Attendee, Vienna, Austria, 2022

17. Ulusal Tıbbi Biyoloji ve Genetik Kongresi (online katılımlı), Attendee, Antalya, Turkey, 2021

5. Ulusal Çocuk Genetik Kongresi (online katılımlı), Attendee, Antalya, Turkey, 2021

14. Ulusal Tıbbi Genetik Kongresi (online katılımlı), Attendee, Antalya, Turkey, 2020

4. Ulusal Çocuk Genetik Kongresi, Attendee, Ankara, Turkey, 2019