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

I. **Brittle cornea syndrome with a novel pathogenic variant of PRDM5 gene**
SUSAM E., YILDIRIM N., KOCAGİL S., ÇİLİNGİR O.
EUROPEAN JOURNAL OF HUMAN GENETICS, vol.30, no.SUPPL 1, pp.119-120, 2022 (SCI-Expanded)

## Refereed Congress / Symposium Publications in Proceedings

I. Özbakır D. H., Susam E., Bütün Z., Artan S.
15. Ulusal Tıbbi Genetik Kongresi, Muğla, Turkey, 9 - 13 November 2022, pp.90
II. **A Brittle cornea syndrome with a novel pathogenic variant of PRDM5 gene**
Susam E., Yıldırım N., Kocagil S., Çilingir O.
54th European Society of Human Genetics (ESHG) Conference:Virtual Conference, 28 - 31 August 2021, vol.30, pp.119-120
III. **A Rare Form Of Constitutional Chromoanasynthesis: Ring Chromosome 18**
Susam E., Erzurumluoğlu Gökalp E., Tosumoğlu E., Kocagil S., Çilingir O., Durak Aras B., Artan S.
14. ULUSAL TIBBİ GENETİK KONGRESİ'xx'xxULUSLARAARSI KATILIMLI'xx'xx, Ankara, Turkey, 20 - 22 November 2020, vol.31, pp.15
IV. **A Novel Mutation of DYSF Gene in A Patient with Limb Girdle Muscular Dystrophy Type 2b**
KOCAGİL S., KAPLAN E., SUSAM E., DURAK ARAS B., ARTAN S., ERZURUMLUOĞLU GÖKALP E., ÇİLİNGİR O.
European Human Genetics Virtual Conference, 06 June 2020
V. **MACROORCHIDISM AS A UNIQUE SIGN IN 3q13.31 DELETION SYNDROME**
SUSAM E., ÇİLİNGİR O., DURAK ARAS B., ERZURUMLUOĞLU GÖKALP E., KOCAGİL S., ARTAN S.
ESHG 2019, Gothenburg, Sweden, 15 - 19 June 2019, vol.27, pp.1843
VI. **The 15q11.2 BP1–BP2 Microdeletion Syndrome with Variable Expressivity**
ERZURUMLUOĞLU GÖKALP E., ÇİLİNGİR O., DURAK ARAS B., KOCAGİL S., SUSAM E., ARTAN S.
13th Balkan Congress of Human Genetics, Edirne, Turkey, 17 - 20 April 2019, vol.22, pp.95
VII. **Macroorchidism as a Uniq Sign in 3q13.31 Deletion Syndrome.**
SUSAM E., ÇİLİNGİR O., DURAK ARAS B., ERZURUMLUOĞLU GÖKALP E., KOCAGİL S., ARTAN S., ŞİMŞEK E.
European Human Genetics Conference, Gothenburg, Sweden, 15 June 2019, vol.27, pp.1843
VIII. **A patient with a balanced inversion of chromosome 11 and unbalanced inversion of chromosome 2**
SUSAM E., ÇİLİNGİR O., BAŞ H., DURAK ARAS B., ERZURUMLUOĞLU GÖKALP E., ARTAN S.
Erciyes Meidcal Genetics Days 2019, Kayseri, Turkey, 21 - 23 February 2019, vol.41, pp.20
IX. **A Patient with a Balanced Inversion of Chromosome 11 and Unbalancded Inversion of Chromosome 2.**
SUSAM E., ÇİLİNGİR O., BAŞ H., DURAK ARAS B., ERZURUMLUOĞLU GÖKALP E., ARTAN S.
Erciyes Medical Genetics Days, Kayseri, Turkey, 21 February 2019, vol.21, pp.20

## Supported Projects

Durak Aras B., Erzurumluoğlu Gökalp E., Çilingir O., Artan S., Köşger P., Kıztanır H., Temena A., Çınar D., Susam E., Project Supported by Higher Education Institutions, Primer Kardiyomiyopatilerde mtDNA Varyantlarının Değerlendirilimesi, 2019 - 2020

## Metrics

Publication: 12