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Published journal articles indexed by SCI, SSCI, and AHCI

- I. **Further Evidence for *RFWD3* Gene Causing Fanconi Anemia Complementation Group W: Detailed Clinical Report of the Second Case in the Literature**
KOCAGİL S., ŞAFKİ İ. N., SARAÇ E., AYDIN C., ARTAN S., KİREL B.
MOLECULAR SYNDROMOLOGY, vol.14, pp.509-515, 2023 (SCI-Expanded)
- II. **Clinical and molecular evaluation of MEFV gene variants in the Turkish population: a study by the National Genetics Consortium**
Dundar M., Fahrioglu U., Yildiz S. H., Bakir-Gungor B., Temel S. G., Akin H., Artan S., Cora T., Sahin F. I., Dursun A., et al.
FUNCTIONAL & INTEGRATIVE GENOMICS, vol.22, no.3, pp.291-315, 2022 (SCI-Expanded)

Refereed Congress / Symposium Publications in Proceedings

- I. **Two Siblings Harboring Two Nonsense Variants: Kaufman Oculocerebrofacial Syndrome With Variable Intrafamilial Expression**
Saraç E., Çarman K. B., Durak Aras B., Artan S.
EuroDysmorpho 2023, Lisbon, Portugal, 13 - 16 September 2023, pp.1
- II. **Complete paternal isodisomy of chromosome 15 in a patient with atypical presentation of Angelman syndrome**
Çilingir O., Saraç E., Aynacı S., Kocagil S.
14th European Cytogenomics Conference, Montpellier, France, 1 - 04 July 2023, pp.48
- III. **Fanconi Aplastik Anemili 5 Olgu: FANCA, ERCC4, RFWD3 Genlerinde 3 Novel/Nadir Varyant**
Özbakır D. H., Saraç E., Susam E., Kocagil S., Aynacı S., Kirel B., Çilingir O.
2. Ulusal HematoOnkoGenetik Kongresi, İskele, Cyprus (Kktc), 4 - 07 May 2023, pp.52
- IV. **Trizomi 21 Tanılı Bir Yenidoğanda Geçici Anormal Miyelopoez ve GATA1 Geninde İki Somatik Varyant**
Saraç E., Özdemir Z. C., Tosumoğlu E., Çilingir O.
2. Ulusal HematoOnkoGenetik Kongresi, Gazimagusa, Cyprus (Kktc), 4 - 07 May 2023, pp.95
- V. **Farklı Yapısal Y Kromozom Anomalisi Saptanan Olgularda Klinik ve Genetik Sonuçların İncelenmesi**
Tosumoğlu E., Saraç E., Üre İ., Kocagil S.
15. Ulusal Tıbbi Genetik Kongres, Muğla, Turkey, 9 - 13 November 2022, pp.240
- VI. **A novel LRP5 gene variant in a patient with Osteoporosis-Pseudoglioma syndrome**
Saraç E., Kirel B., Çilingir O., Artan S.
7.Uluslararası Erciyes Tıp Tıbbi Genetik Kongresi, Kayseri, Turkey, 26 - 28 May 2022, pp.104-105
- VII. **Atipik bulguları olan Smith-Magenis sendromlu bir olgu sunumu**
Saraç E., Özbakır D. H., Kocagil S., Erzurumluoğlu Gökalp E., Çilingir O., Durak Aras B., Artan S.
14. ULUSAL TIBBİ GENETİK KONGRESİ'xx'xxULUSLARAARSI KATILIMLI'xx'xx, Eskişehir, Turkey, 20 - 22 November 2020, vol.31, pp.92

Metrics

Publication: 10

Citation (WoS): 4

Citation (Scopus): 5

H-Index (WoS): 1

H-Index (Scopus): 1