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Refereed Congress / Symposium Publications in Proceedings

- I. **Report of first Endocrine-cerebro-osteodysplasia patient to reach childhood age**
KOCAGİL S., ÖZBAKIR D. H., Kabaoglu U., ERZURUMLUOĞLU GÖKALP E., ÇİLİNGİR O.
56th Annual Conference of the European-Society-of-Human-Genetics (ESHG), Glasgow, England, 10 - 13 June 2023, pp.145
- II. **REPORT OF THE FIRST ENDOCRINE-CEREBRO-OSTEODYSPLASIA PATIENT TO REACH CHILDHOOD AGE**
Özbakır D. H., Kocagil S., Erzurumluoğlu Gökalp E., Çilingir O.
EURODYSMORPHO 2023, Lisbon, Portugal, 13 - 16 September 2023, pp.28
- III. **A de novo small marker chromosome that causes Trisomy 9p in a patient with failure to develop, microcephaly and normal neuromotor development**
ARTAN S., ÖZBAKIR D. H., TOSUMOĞLU E., HARMANCI K., PANAL G., ERZURUMLUOĞLU GÖKALP E.
14th European Cytogenomics Conference, France, 1 - 04 July 2023, pp.48
- IV. **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
- V. **Özbakır D. H., Susam E., Bütün Z., Artan S.**
15. Ulusal Tıbbi Genetik Kongresi, Muğla, Turkey, 9 - 13 November 2022, pp.90
- VI. **CRPPA gene-associated Walker-Warburg syndrome: A rare prenatal case with intragenic homozygous deletion**
Özbakır D. H., Bütün Z., Kocagil S.
7. Uluslararası Erciyes Tıp Tıbbi Genetik Kongresi, Kayseri, Turkey, 26 - 28 May 2022, pp.102-103
- VII. **Saraç E., Özbakır D. H., Kocagil S., Erzurumluoğlu Gökalp E., Çilingir O., Durak Aras B., Artan S.**
14. ULUSAL TIBBİ GENETİK KONGRESİ, 20 - 22 November 2020, pp.91

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