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Articles Published in Other Journals

- I. **A report of two siblings diagnosed with Cutis Laxa**
GÜNDOĞDU ÖĞÜTLÜ Ö. B., SEZER A., DEMİRBAŞ M. H., KAYHAN G., PERÇİN F. E.
Gazi Medical Journal, vol.31, no.2, 2020 (ESCI)
- II. **International Participated Erciyes Medical Genetics Days 2019A novel mutation in HECW2 gene resulting neurodevelopmental disorder with hypotonia, seizures, and absent language**
DEMİRBAŞ M. H., ÖZBUDAK P., SERDAROĞLU A., PERÇİN F. E.
Erciyes Medical Journal, pp.25, 2019 (Peer-Reviewed Journal)
- III. **Deletion of the SOX3 Gene Causes Panhypopituitarism: A Case Report**
KAYHAN G., DEMİRBAŞ M. H., PERÇİN F. E.
Gazi Medical Journal, vol.30, no.1, pp.1-101, 2019 (Scopus)

Refereed Congress / Symposium Publications in Proceedings

- I. **A mosaic double aneuploidy: mos 45,X/47,XX,18 with mild phenotype**
DEMİRBAŞ M. H., HABİLOĞLU E., ERGÜN M. A., PERÇİN F. E., YİRMİBEŞ KARAOĞUZ M.
European Human Genetics Conference, Milano, İtalya, 16 - 19 June 2018

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Publication: 6