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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 (Refereed Journals of Other Institutions)
- 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 (Refereed Journals of Other Institutions)
- 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 (Refereed Journals of Other Institutions)

### 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