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Refereed Congress / Symposium Publications in Proceedings

- I. **Importance of genetic diagnosis in demyelinating diseases; adult-onset Alexander Disease case with novel GFAP mutation**
CEYLAN KÖSE C., OCAK Ö., GÜNDÜZ O. R., SILAN F.
15. Ulusal Tıbbi Genetik Kongresi, Muğla, Turkey, 9 - 13 November 2022
- II. **A rare Malpuech, Mignarelli, Michels, Carnevale Syndrome (3MC Syndrome); Novel variant in the MASP1 gene**
GÜNDÜZ O. R., CEYLAN KÖSE C., SILAN F.
EUROPEAN BIOTECHNOLOGY CONGRESS 2022, PRAG, Czech Republic, 5 - 07 October 2022
- III. **A rare syndrome from Canakkale; Kindler Syndrome with loss of heterozygosity of FERMT1 gene region**
GÜNDÜZ O. R., IŞIK MERMUTLU S., CEYLAN KÖSE C., SILAN F.
7. Uluslararası Erciyes Tıp Tıbbi Genetik Kongresi, Kayseri, Turkey, 26 - 28 May 2022

Metrics

Publication: 4