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Refereed Congress / Symposium Publications in Proceedings

- I. **De novo GABRB3 c.103G>A mutation detected in a patient with epilepsy and speech retardation**
AKBAŞ N. E., SÖNMEZ V., AKCAN M. B., MUTLUER Y. E., CELİK K. M., SILAN F., ÖZDEMİR Ö.
6. Uluslararası Katılımlı Erciyes Tıp Tıbbi Genetik Kongresi, Kayseri, Turkey, 16 September 2021

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