In the Clinical/Scientific Note “KCNQ2 encephalopathy manifesting with Rett-like features: A follow-up into adulthood” by Mastrangelo et al., the first sentence of the fifth paragraph should read, “A next-generation sequencing panel that included 140 genes involved in genetic epilepsies revealed the de novo heterozygous KCNQ2 variant c.629 G > C (p.Arg210Pro), which was not present in the Human Gene Mutation Database (portal.biobase-international.com/).” The authors regret the error.

Reference
KCNQ2 Encephalopathy Manifesting With Rett-like Features: A Follow-up Into Adulthood

Neurol Genet 2021;7;
DOI 10.1212/NXG.0000000000000604

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