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Corrigendum to "A single-center, retrospective analysis of genotype–phenotype correlations in children with Dravet syndrome"

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The authors regret that several of the SCN1A variants reported in the supplementary table were annotated incorrectly per chart reports, and appreciate the assistance of the HGMD editors in correcting these variants. In the Results section, p.Ala1442Val should be noted as p.Ala1440Val, and p.Phe1761Thrfs*8 should be noted as p.Phe1671Thrfs*8. Appropriate corrections to all other variants are provided in updated supplementary tables (see attachment) verified through sequence alignment to the cDNA RefSeq for transcript variant 1 of *SCN1A* (NM_001165963.4) and original genetic testing reports wherever possible.

The authors would like to apologise for any inconvenience caused.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Appendix A. Supplementary data

Supplementary material related to this article can be found, in the online version, at doi:https://doi.org/10.1016/j.seizure.2020.04.004.