

CORRECTION



Correction: Novel compound heterozygous *ABCA2* variants cause IDPOGSA, a variable phenotypic syndrome with intellectual disability

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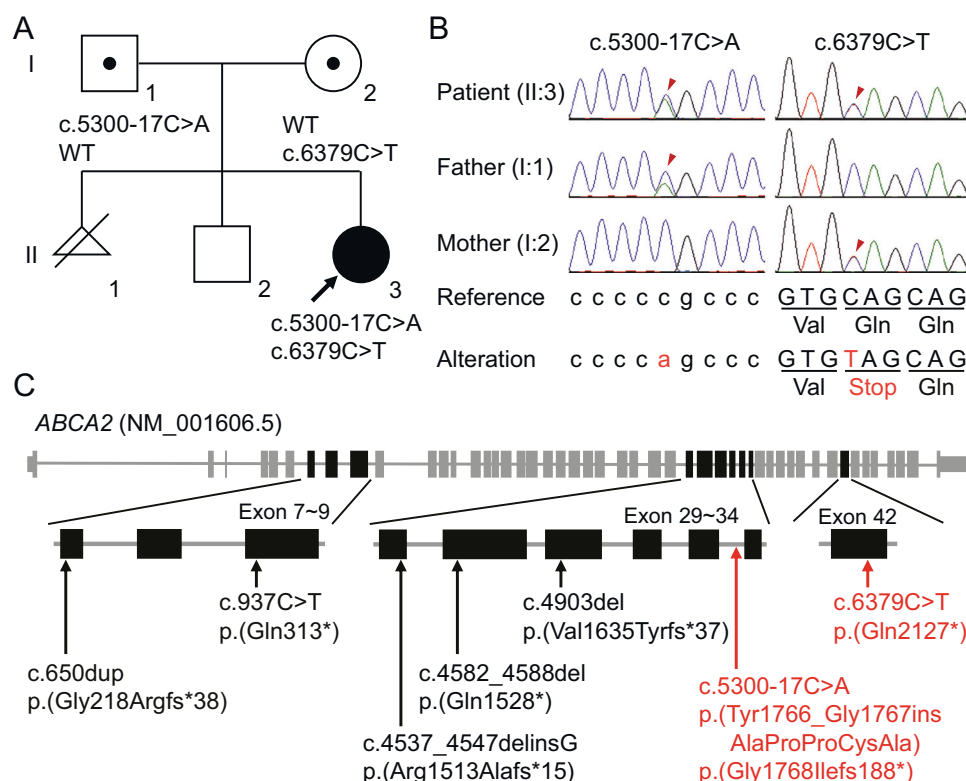
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Correction to: *Journal of Human Genetics* <https://doi.org/10.1038/s10038-024-01219-8>, published online 17 January 2024

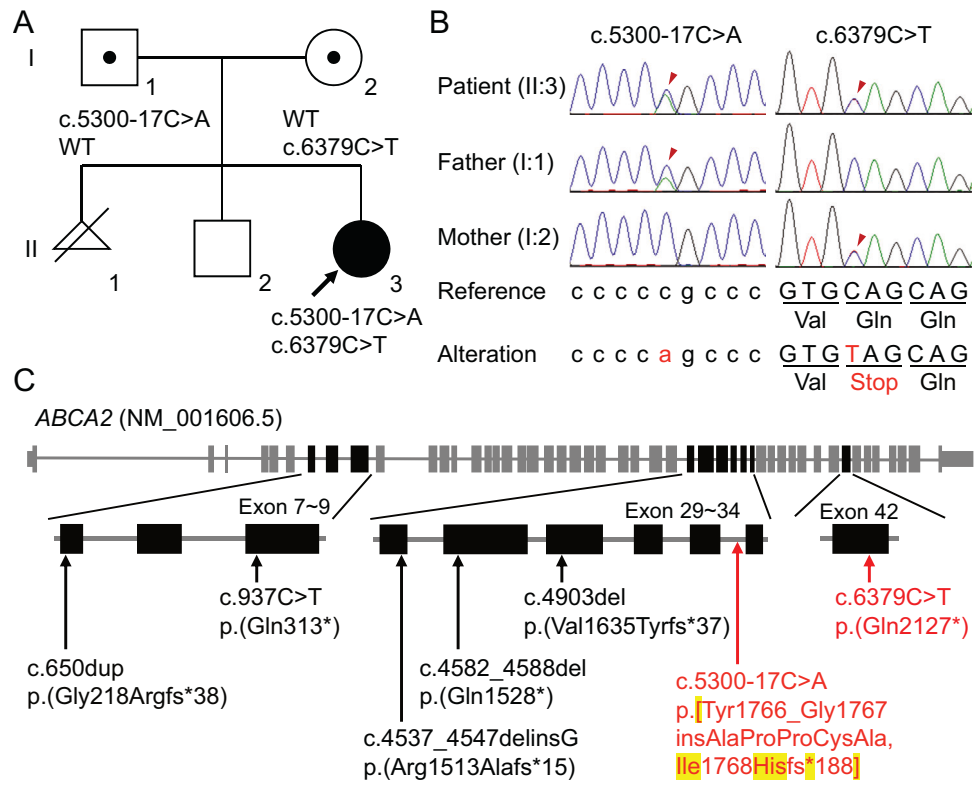
The updated figures 1 and 2 are as follows.

The patient variant described on pages 144–146 is accurately stated as follows: compound heterozygous *ABCA2* variants, NM_001606.5:c.[5300–17C>A];[6379C>T] p.[Tyr1766_Gly1767insAlaProProCysAla,Ile1768Hisfs*188];[(Gln2127*)]

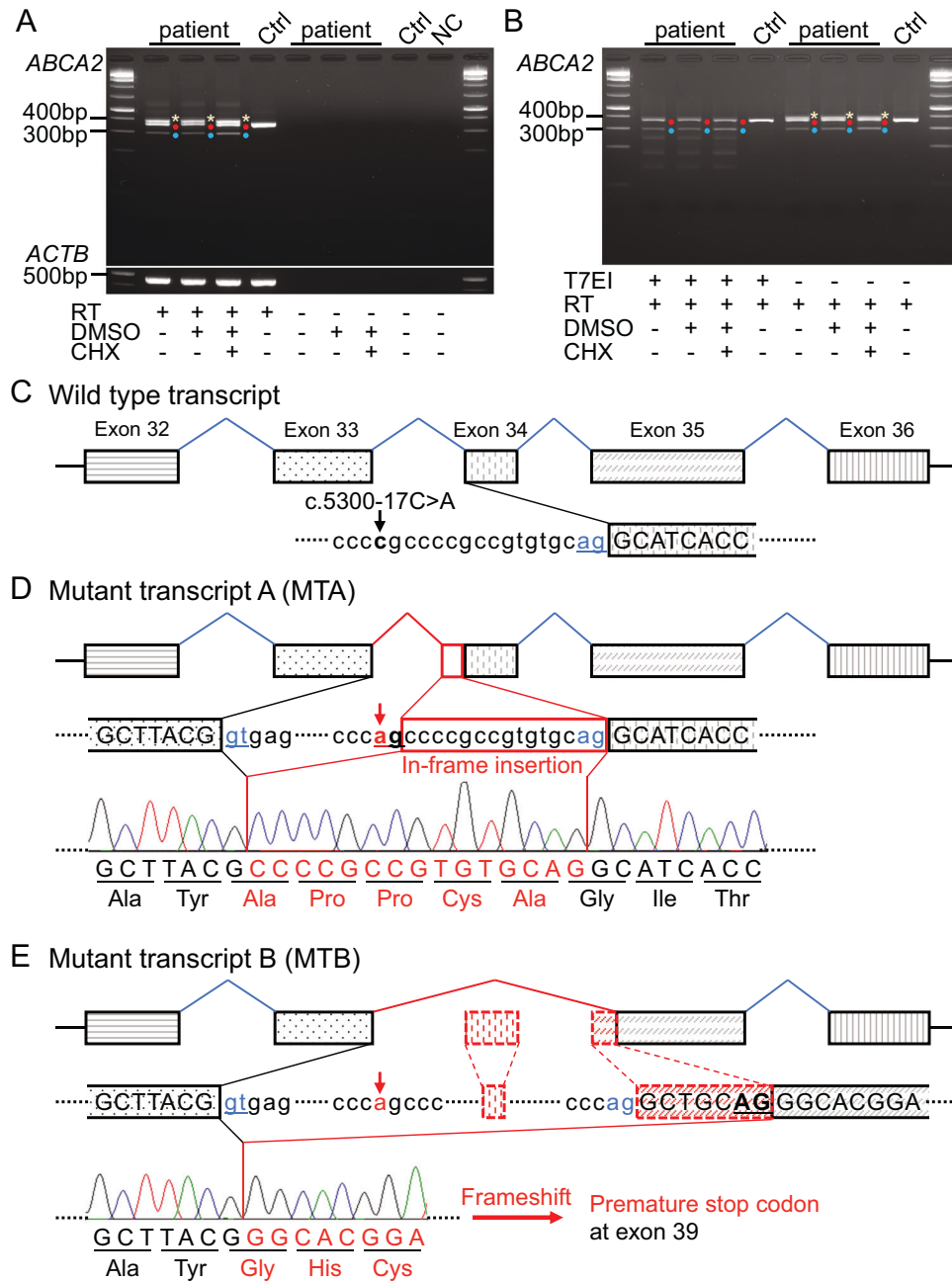
Original Fig. 1



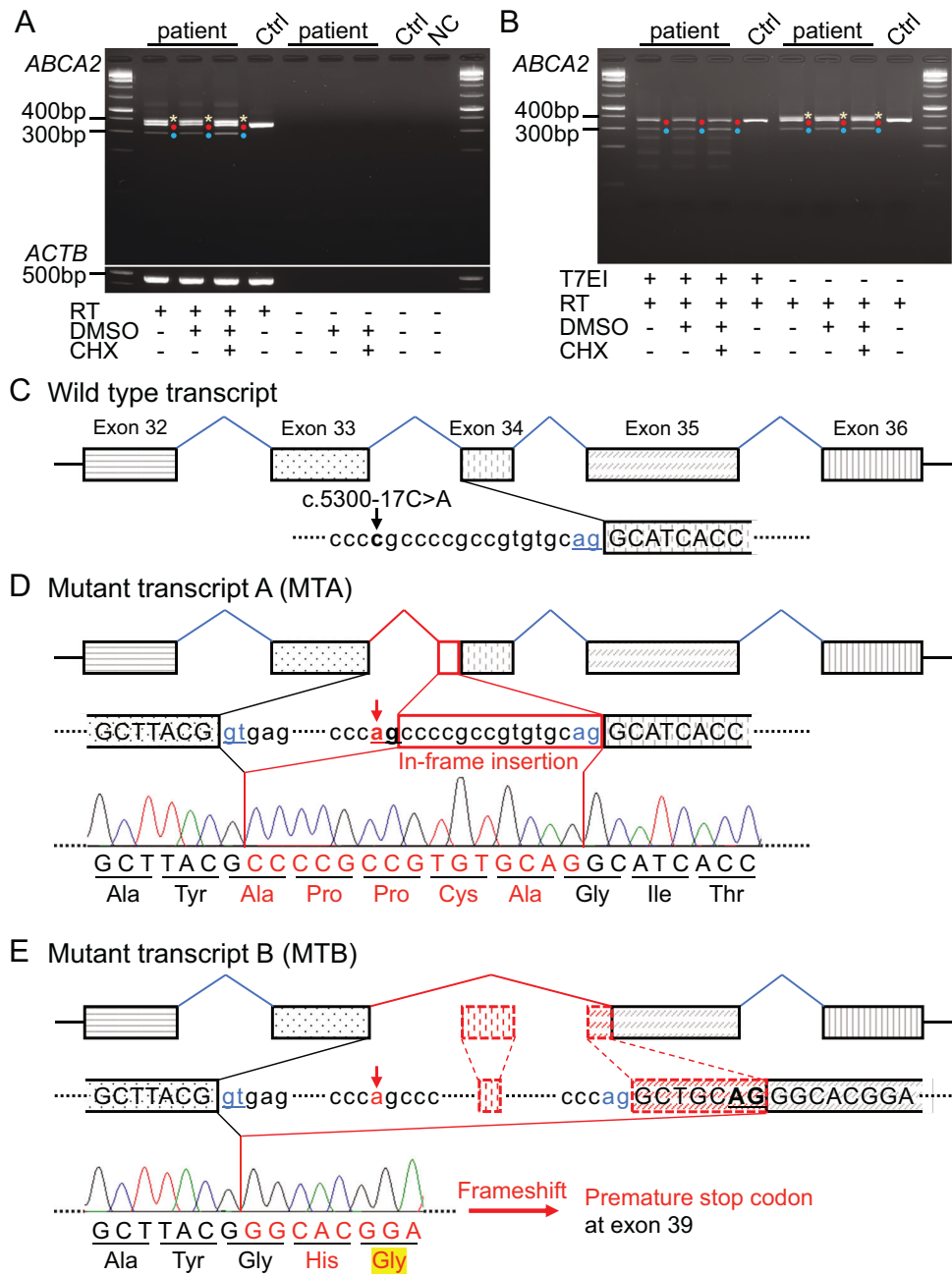
Updated Fig. 1



Original Fig. 2



Updated Fig. 2



The original article has been corrected.

ADDITIONAL INFORMATION

The online version contains supplementary material available at <https://doi.org/10.1038/s10038-025-01422-1>.

Supplementary information The online version contains supplementary material available at <https://doi.org/10.1038/s10038-025-01422-1>.

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