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CORRECTION

Variants in *PHF8* cause a spectrum of X-linked neurodevelopmental disorders and facial dysmorphology

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The variant description c.1030C>T p.(Gln343*) should have been c.1027C>T p.(Gln343*) three times in the originally published version of this article. In Table 1 for variant 5 (individuals 6, 7, and 8), in Figure 5, and for individuals 6, 7, and 8 in Table S1. The article has been corrected online. The authors regret the errors.

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