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CORRECTION

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Correction to: Phenotype, genotype and long-term prognosis of 40 Chinese patients with isobutyryl-CoA dehydrogenase deficiency and a review of variant spectra in ACAD8

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Correction to: Orphanet J Rare Dis (2021) 16:392
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Following the publication of the original article [1] the authors asked to revise the following sentence of the "Discussion" section:

"Another special IBDD patient, with a severe lack of speech development and lack of social interactions, was reported to be associated with autism that was genetically confirmed in the *DNA2* gene [18]."

The correct sentence should read:

"Another special IBDD patient, with a severe lack of speech development and lack of social interactions, was reported to be associated with autism [18]."

The original article has already been corrected as above.

The original article can be found online at https://doi.org/10.1186/s13023-021-02018-6.

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