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CORRECTION

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Correction: Prader-Willi syndrome patient with atypical phenotypes caused by mosaic deletion in the paternal 15q11-q13 region: a case report

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Correction: Ital J Pediatr 48, 204 (2022)
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The original article [1] mistakenly mis-numbered a section of citations in the main body; this has since been rectified.

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Reference

1. Jinying W, et al. Prader-Willi syndrome patient with atypical phenotypes caused by mosaic deletion in the paternal 15q11-q13 region: a case report. Ital J Pediatr. 2022;48:204. https://doi.org/10.1186/s13052-022-01398-0.

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