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Corrigendum: Spondyloocular syndrome: A novel *XYLT2* variant with description of the neonatal phenotype

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A Corrigendum on

Spondyloocular syndrome: A novel XYLT2 variant with description of the neonatal phenotype

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In the published article, there was an error. There was a sentence missing in the **Acknowledgments** section.

A correction has been made to the **Acknowledgments**. This sentence previously stated: "The authors thank the family for participating in the study."

The corrected sentence appears below:

"The authors thank the family for participating in the study. Two of the authors of this publication are members of the European Reference Network for rare malformation syndromes and rare intellectual and neurodevelopmental disorders, ERN-ITHACA."

The authors apologize for this error and state that this does not change the scientific conclusions of the article in any way. The original article has been updated.

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