

CORRECTION

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# Correction: Multilocus pathogenic variants contribute to intrafamilial clinical heterogeneity: a retrospective study of sibling pairs with neurodevelopmental disorders

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**Correction:** BMC Med Genomics 17, 85 (2024)  
<https://doi.org/10.1186/s12920-024-01852-4>

Following the publication of the Original Article [1], the authors reported an error in the placement of the plots during the manuscript preparation step. The authors accidentally placed the plots for Family HOU1842 in the Family of HOU4131, which resulted the duplication of Figure 1 Plots B and H and Supplementary Figure 2B and 5B.

The original article can be found online at <https://doi.org/10.1186/s12920-024-01852-4>.

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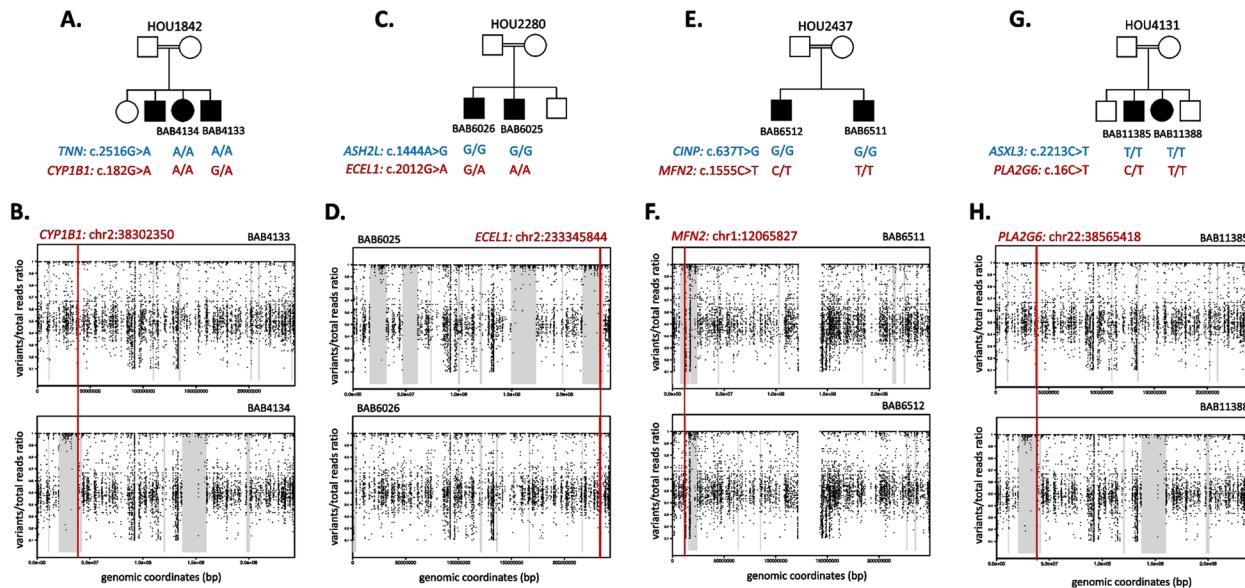
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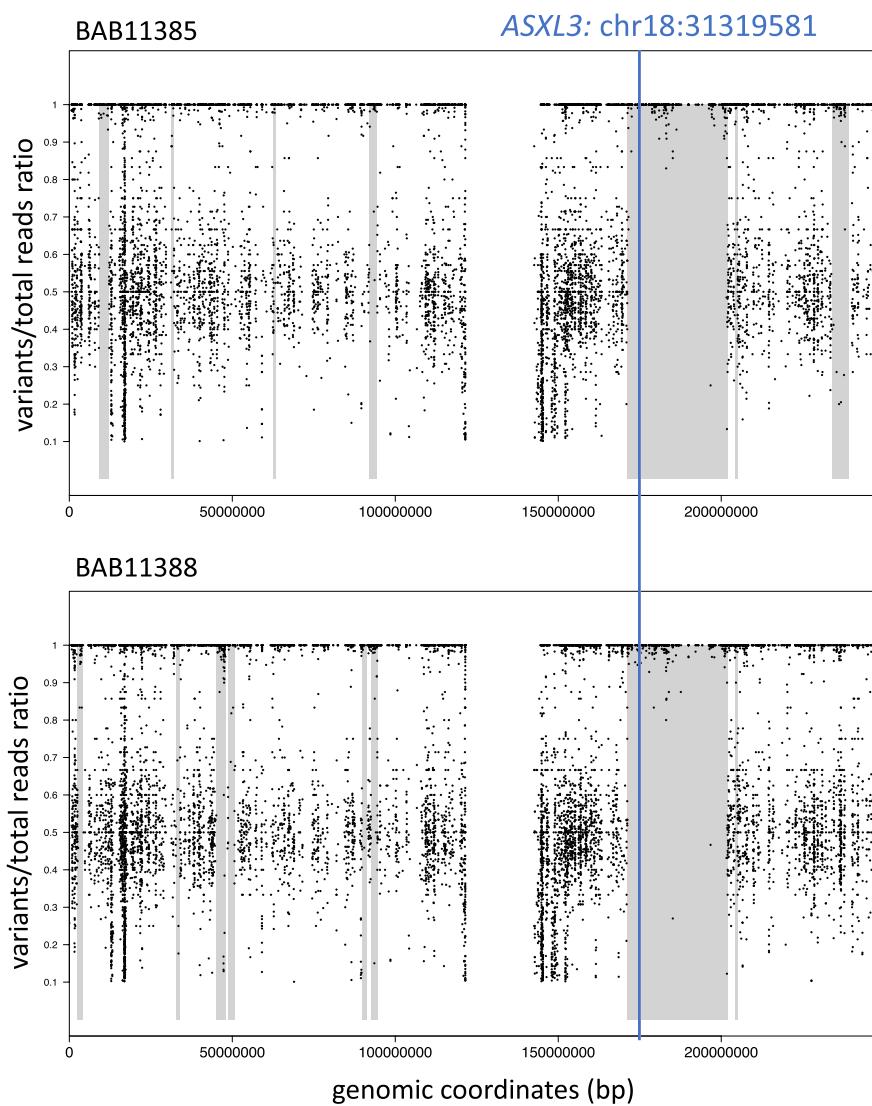
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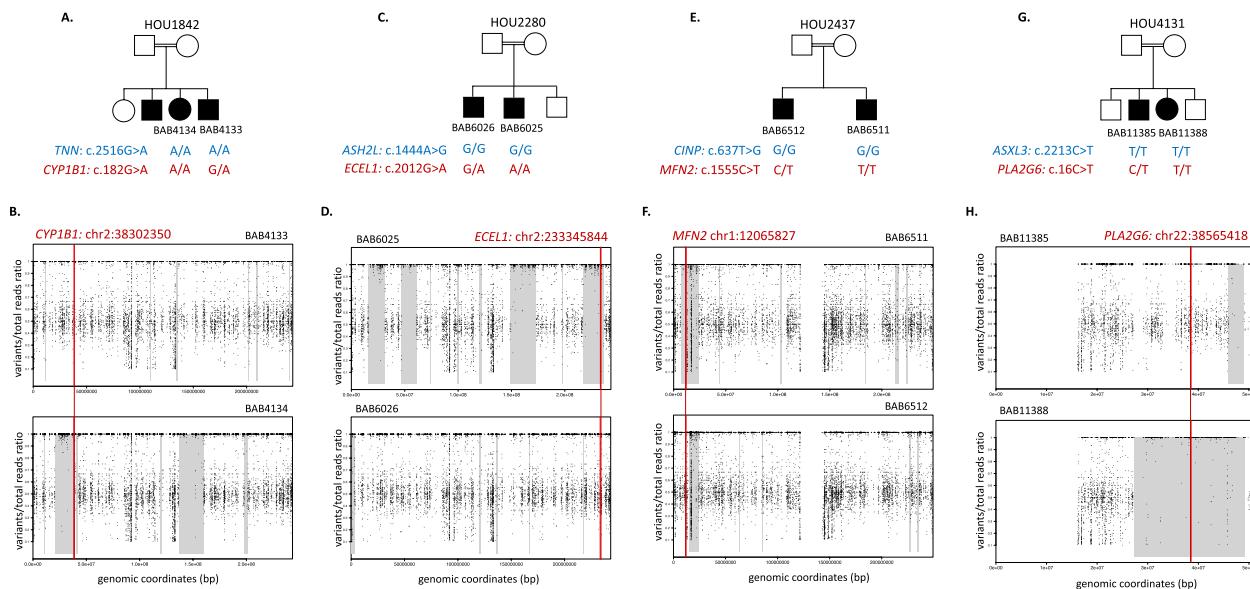
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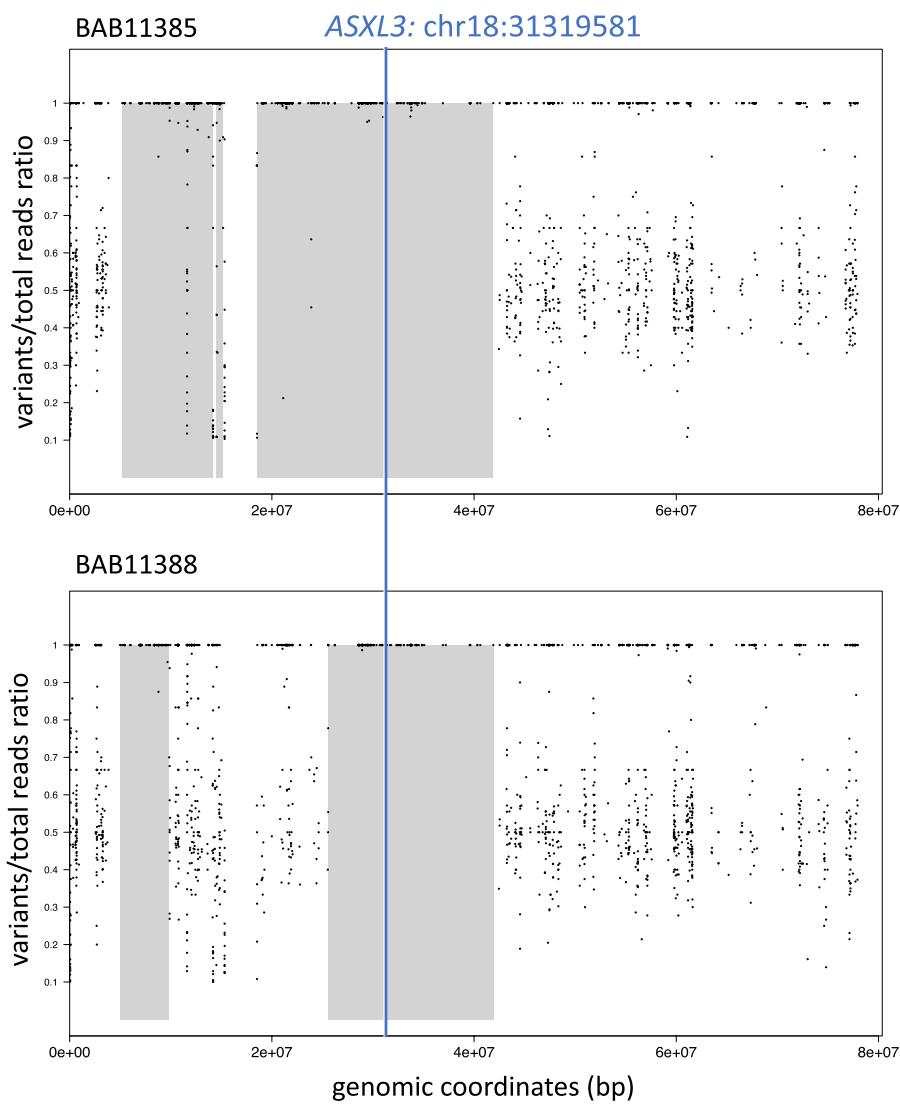
**Incorrect**  
**Figure 1**



**Supplementary Figure 5****A.****B.**

**Correct**  
**Figure 1**



**Supplementary Figure 5****A.****B.**

The authors indicate that these errors do not affect any of the results nor the conclusions presented in the text of the study.

The Original Article has been corrected.

Published online: 07 August 2024

**Reference**

1. Bozkurt-Yozgatli, et al. Multilocus pathogenic variants contribute to intra-familial clinical heterogeneity: a retrospective study of sibling pairs with neurodevelopmental disorders. *BMC Med Genomics*. 2024;17:85. <https://doi.org/10.1186/s12920-024-01852-4>.