



**Correction:** The phenotypic spectrum of *WWOX*-related disorders: 20 additional cases of WOREE syndrome and review of the literature

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The article has been corrected to account for one patient being investigated through genome sequencing rather than exome sequencing as originally published; thus amendments to the Abstract and Methods have been made as well as addition of the relevant authors and acknowledgment.



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These authors contributed equally: Juliette Piard and Lara Hawkes.

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**Correction:** Analysis of fetal DNA in maternal plasma with markers designed for forensic DNA mixture resolution

Amandine Moriot, BA<sup>1</sup> and Diana Hall, PhD<sup>1</sup>

*Genetics in Medicine* (2019) 21:1668; <https://doi.org/10.1038/s41436-019-0465-6>

Correction to: *Genetics in Medicine* <https://doi.org/10.1038/s41436-018-0102-9>; published online 03 August 2018

In the original version of this Article, the data for Mother ID 8; 13; 25; 31; and 47 of Table 2 was merged. This has now been corrected in both the PDF and HTML versions of the Article.

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Published online: 11 March 2019

**Correction:** Toward automation of germline variant curation in clinical cancer genetics

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*Genetics in Medicine* (2019) 21:1668; <https://doi.org/10.1038/s41436-019-0490-5>

Correction to: *Genet Med* advance online publication 21 February 2019; <https://doi.org/10.1038/s41436-019-0463-8>.

An update to the original published conflict of interest for author Liying Zhang, PhD. L.Z. received compensation from Future Technology Research LLC (seminar on precision medicine), Roche Diagnostics Asia Pacific, BGI, Illumina (speaking activities at conferences/workshop). L.Z.'s family member has a leadership position and ownership interest of Shanghai Genome Center. This correction has been made.

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## Open

**Correction:** DNA breakpoint assay reveals a majority of gross duplications occur in tandem reducing VUS classifications in breast cancer predisposition genes

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Correction to: *Genetics in Medicine*; <https://doi.org/10.1038/s41436-018-0092-7>; published online 28 July 2018.

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**Correction:** The Gen-Equip Project: evaluation and impact of genetics e-learning resources for primary care in six European languages

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**Correction:** Comparative assessment of gene-specific variant distribution in prenatal and postnatal cohorts tested for Noonan syndrome and related conditions

N. T. Leach, PhD<sup>1</sup>, D. R. Wilson Mathews, MS, LCGC<sup>1</sup>, L. S. Rosenblum, PhD<sup>1</sup>, Z. Zhou, PhD<sup>1</sup>, H. Zhu, PhD<sup>1</sup> and R. A. Heim, PhD<sup>1</sup>

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The original version of this article contained an error in the spelling of the author N. T. Leach, which was incorrectly given as N. L. Leach. This has been corrected in both the PDF and HTML versions of the article.

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Open

**Correction:** Opportunities to implement a sustainable genomic medicine program: lessons learned from the IGNITE Network

Kenneth D. Levy, PhD, MBA<sup>1</sup>, Kathryn Blake, PharmD<sup>2</sup>, Colette Fletcher-Hoppe, BA<sup>3</sup>, James Franciosi, MD, MS<sup>4</sup>, Diasuke Goto, PhD<sup>5</sup>, James K. Hicks, PharmD, PhD<sup>6</sup>, Ann M. Holmes, PhD<sup>7</sup>, Sri Harsha Kanuri, MBBS, PhD<sup>1</sup>, Ebony B Madden, PhD<sup>8</sup>, Michael D. Musty, BA<sup>9</sup>, Lori Orlando, MD<sup>9</sup>, Victoria M. Pratt, PhD<sup>1</sup>, Michelle Ramos, MPH<sup>10</sup>, Ryanne Wu, MD, MHS<sup>9</sup> and Geoffrey S. Ginsburg, MD, PhD<sup>9</sup>

*Genetics in Medicine* (2019) 21:1670; <https://doi.org/10.1038/s41436-018-0280-5>

Correction to: *Genetics in Medicine*; <https://doi.org/10.1038/s41436-018-0080-y>; published online 12 July 2018.

The original version of this Article contained an error in the spelling of the author Geoffrey S. Ginsburg, which was incorrectly given as Geoffrey Ginsburg. This has now been corrected in both the PDF and HTML versions of the Article.



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**Correction to:** The landscape of epilepsy-related GATOR1 variants

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The original version of this Article contained an error in the author list where the corresponding author Stéphanie Baulac was repeated twice. This has now been corrected in the HTML version; the PDF was correct at the time of publication.

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