Correction: Nonrandom occurrence of multiple de novo coding variants in a proband indicates the existence of an oligogenic model in autism

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Genetics in Medicine (2019) 21:2662; <https://doi.org/10.1038/s41436-019-0629-4>

Correction to: *Genetics in Medicine*; <https://doi.org/10.1038/s41436-019-0610-2>; published online 23 July 2019

In the original version of the published Article the word “mutation” was used throughout the text. The ACMG changed its terminology from “mutation” to “variant” prior to the publication of the Article and the requested text changes were not incorporated by the typesetter. The PDF and HTML versions of the Article have now been modified accordingly.

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Published online: 1 August 2019

Correction: The value of diagnostic testing for parents of children with rare genetic diseases

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Genetics in Medicine (2019) 21:2662; <https://doi.org/10.1038/s41436-019-0615-x>

Correction to: *Genetics in Medicine*; <https://doi.org/10.1038/s41436-019-0583-1>, published online 26 June 2019

In the original version of this Article, financial support was not fully acknowledged. The PDF and HTML versions of the Article have now been corrected to include the following:

S.H. acknowledges that the University of Aberdeen's Health Economics Research Unit is funded by the Chief Scientist Office of the Scottish Government Health and Care Directorates.

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Published online: 18 July 2019

Correction: Inaccuracies and shortcomings in “Adherence of cell-free DNA noninvasive prenatal screens to ACMG recommendations”

Katherine Johansen Taber¹, Carrie Haverty¹, Susan Hancock¹ and James Goldberg¹

Genetics in Medicine (2019) 21:2663; <https://doi.org/10.1038/s41436-019-0589-8>

Correction to: *Genetics in Medicine*; <https://doi.org/10.1038/s41436-019-0555-5>, published online 03 June 2019

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Published online: 19 June 2019

Correction: Variants in *MED12L*, encoding a subunit of the Mediator kinase module, are responsible for intellectual disability associated with transcriptional defect

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Genetics in Medicine (2019) 21:2663; <https://doi.org/10.1038/s41436-019-0590-2>

Correction to: *Genetics in Medicine* 2019; <https://doi.org/10.1038/s41436-019-0557-3>, published online 03 June 2019

In the Acknowledgements section of the paper the authors neglected to mention that the study was supported by a grant from the National Human Genome Research Institute (NHGRI) UM1HG007301 (S.H., M.L.T.). In addition, the award of MD was associated with the authors Michelle L. Thompson and Susan Hiatt instead of PhD. The PDF and HTML versions of the Article have been modified accordingly.

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Published online: 3 July 2019