Genetics inMedicine | CORRECTION

Correction: Genetic evaluation of cardiomyopathy: a clinical practice resource of the American College of Medical Genetics and Genomics (ACMG)

Ray E. Hershberger[1], Michael M. Givertz[2], Carolyn Y. Ho[3], Daniel P. Judge[4], Paul F. Kantor[5], Kim L. McBride[6], Ana Morales[1], Matthew R. G. Taylor[7], Matteo Vatta[8,9,10] and Stephanie M. Ware[9,11] on behalf of the ACMG Professional Practice and Guidelines Committee

Genetics in Medicine (2019) 21:2406; https://doi.org/10.1038/s41436-019-0521-2

Correction to: *Genetics in Medicine*; https://doi.org/10.1038/s41436-018-0039-z, published online 14 June 2018

In the original version of this Article, the Statement on Conflict of Interest originally submitted by the authors was not published and should have read:

Disclosure: R.E.H. has served as a consultant for Array Biopharma. C.Y.H. has served as a consultant and has received research support from MyoKardia. D.P.J. has served as a consultant for Array Biopharma, Eidos Therapeutics, Glaxo Smith Kline, Invitae, MyoKardia, and Pfizer. M.R.G.T. has served as a consultant for Array Biopharma, Guidepoint Global, and Wellpoint, Inc., and has served as a speaker for GeneDx. M.V. is employed by Invitae Corporation. The other authors declare no conflicts of interest.

Furthermore, in the Acknowledgements section we neglected to state that:

This article is an abbreviated version of "Genetic Evaluation of Cardiomyopathy—a Heart Failure Society of America Practice Guideline," published in *Journal of Cardiac Failure*, and on which the American College of Medical Genetics and Genomics and the Heart Failure Society of America collaborated.

The PDF and HTML versions of the Article have now been corrected with the above.

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Correction: Variant classification changes over time in *BRCA1* and *BRCA2*

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In the original version of this Article, the affiliation details for Drs. Jordan Lerner-Ellis and George Charames did not include the Department of Pathology and Laboratory Medicine at the University of Toronto. In addition, Drs. Jordan Lerner-Ellis and George Charames were incorrectly affiliated with the institute of Health Policy, Management and Evaluation at the University of Toronto. These errors have now been corrected in both the PDF and HTML versions of the Article.

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Correction: Evaluation of the cost and effectiveness of diverse recruitment methods for a genetic screening study

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Correction to: *Genet Med* 2019; https://doi.org/10.1038/s41436-019-0497-y published online 1 April 2019

The original version of this Article contained an error in the undergraduate degree awarded to the author Ian Halim, which was incorrectly given as BS. This has now been corrected to BA in both the PDF and HTML versions of the Article.

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These authors contributed equally: Hila Milo Rasouly and Julia Wynn

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Open

Correction: Sequencing as a first-line methodology for cystic fibrosis carrier screening

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Genetics in Medicine (2019) 21:2407–2408; https://doi.org/10.1038/s41436-019-0543-9

Correction to: *Genetics in Medicine;* https://doi.org/10.1038/s41436-019-0525-y, published online 30 April 2019

The original version of this Article contained an error in Fig. **3**. Specifically, the result "3 (67%) TOP" should read "2 (67%) TOP." This has now been corrected in both the PDF and HTML versions of the Article.

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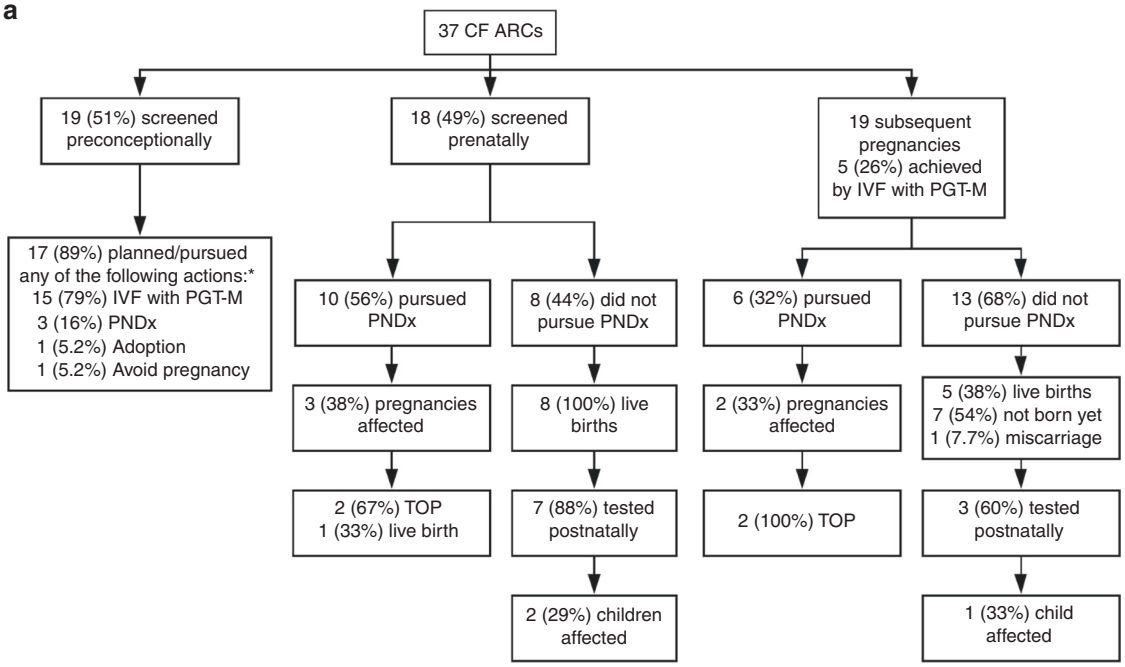
a



b

	"CF23 ARCs", N (%)	"Non-CF23 ARCs", N (%)	*P*-value
Screened preconceptionally	13	6	
Planned/Pursued action to avert affected pregnancy (a)	12 (92)	5 (83)	1
All pregnancies (b)	25	12	
Pursued prenatal diagnosis	9 (36)	7 (58)	0.29
Pregnancies affected	2 (22)	3 (43)	
Affected pregnancy Outcome	1 (50) live birth 1 (50) TOP	3 (100) TOP	

(a) Actions include IVF with PGT-M, prenatal diagnostic testing if/when pregnancy occurs, adoption, or avoidance of pregnancy

(b) Current and subsequent pregnancies in those screened prenatally, and subsequent pregnancies that occured in those screened preconceptionally.

Fig. 3

Correction: ClinGen expert clinical validity curation of 164 hearing loss gene–disease pairs

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In the original version of this Article, the address for affiliation 8, *Division of Hearing and Balance Research, National Hospital Organization Tokyo Medical Center, Tokyo, Japan*, was incorrectly given as *Laboratory of Auditory Disorders and Division of Hearing and Balance Research, Tokyo, Japan*.

Furthermore, the address for affiliation 9, *Medical Genetics Center, National Institute of Sensory Organs, National Hospital Organization Tokyo Medical Center, Tokyo, Japan*, was incorrectly given as *Medical Genetics Center, National Institute of Sensory Organs, National Tokyo Medical Center, Tokyo, Japan*.

They have now been corrected in both the PDF and HTML versions of the Article.

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Correction: *SMAD6* is frequently mutated in nonsyndromic radioulnar synostosis

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

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