CORRECTION | Genetics inMedicine

Correction: Putting genome-wide sequencing in neonates into perspective

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The original version of this Article contained an error in the spelling of the author Pleuntje J. van der Sluijs, which was incorrectly given as Eline (P. J.) van der Sluijs. This has now been corrected in both the PDF and HTML versions of the Article.

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Correction: A new microdeletion syndrome involving *TBC1D24*, *ATP6V0C*, and *PDPK1* causes epilepsy, microcephaly, and developmental delay

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

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Correction: The ARID1B spectrum in 143 patients: from nonsyndromic intellectual disability to Coffin–Siris syndrome

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The original version of this Article contained an error in the spelling of the author Pleuntje J. van der Sluijs, which was incorrectly given as Eline (P. J.) van der Sluijs. This has now been corrected in both the PDF and HTML versions of the Article.

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Correction: Estimating the burden and economic impact of pediatric genetic disease

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Genetics in Medicine (2019) 21:2161–2162; https://doi.org/10.1038/s41436-019-0458-5

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Correction: Genomic mosaicism in the pathogenesis and inheritance of a Rett syndrome cohort

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The second author Jiarui Li is now listed as a co–first author according to her contribution to this paper. The list of authors who contributed equally now reads: Qingping Zhang, Xiaoxu Yang, Jiaping Wang, and Jiarui Li. This has now been corrected in both the PDF and HTML versions of the Article.

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Correction: Population genomic screening of all young adults in a health-care system: a cost-effectiveness analysis

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In the original version of this Article, the affiliation details for Lei Zhang were given as Monash University. While working on the Article Dr. Zhang was also affiliated with the Department of Epidemiology and Biostatistics, School of Public Health, Xi'an Jiaotong University Health Science Center, Xi'an, Shaanxi, PR China. This has now been corrected in both the PDF and HTML versions of the Article.

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Correction: Clinical and genetic spectrum of children with congenital diarrhea and enteropathy in China

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Correction to: *Genetics in Medicine*; https://doi.org/10.1038/s41436-019-0488-z; published online 21 March 2019.

There is an error in the figure legend of Fig. 1(a). The correct figure legend for this figure should be "Fig. 1(a) Hematoxylin and eosin (H&E) analysis of the descending duodenum shows the loss of goblet cells and Paneth cells and the presence of apoptotic cells in patient 48."

The authors apologize for these errors and state that this does not change the scientific conclusions of the article in any way. The PDF and HTML versions of the Article have been modified accordingly.

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Correction: Mitochondrial oxodicarboxylate carrier deficiency is associated with mitochondrial DNA depletion and spinal muscular atrophy–like disease

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Correction to: *Genetics in Medicine* **20**:1224–1235; https://doi.org/10.1038/gim.2017.251; Article published online 08 March 2018

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