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CORRIGENDUM

The diagnostic utility of exome sequencing in Joubert syndrome and related disorders

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Correction to: *Journal of Human Genetics* (2013) **58,** 113–115; doi: 10.1038/jhg.2012.117; published online 4 October 2012

Since the publication of the above article, the authors have noticed an error in the description of mutation.

The mutation described as c.6012-12A>T in the main text and Figure 1a should have been c.6012-12T>A.

This correction does not alter the results and their interpretation as discussed in the paper.

The authors would like to apologize for this mistake.