Erratum to: Concordance rates of Wilson's disease phenotype among siblings

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The Results section of this article contained errors. The third paragraph of the subsection headed “Genetic analysis” should have read as follows:

Aside from H1069Q, the most frequent missense mutations included T977M (c.2930C>T), T1220M (c.3659C>T), S653Y (c.1958C>A), and G988R (c.2962G>C). Among severe mutations, the Q1351X (c.4051C>T), A1135fs (c.3402delC), D1047fs (c.3140delA), and M769fs (c.2304insC) were identified.

The authors apologize for any inconvenience caused.

The online version of the original article can be found at <http://dx.doi.org/10.1007/s10545-013-9625-z>.

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