

## Erratum to: Barth syndrome without tetralinoleoyl cardiolipin deficiency: a possible ameliorated phenotype

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The original version of this article unfortunately contained a mistake. The reference by Whited is incomplete. The corrected reference is:

Whited K, Baile MG, Currier P, Claypool SM (2013) Seven functional classes of Barth syndrome mutation. *Human Molecular Genetics* 22:483–492.

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The online version of the original article can be found at <http://dx.doi.org/10.1007/s10545-014-9747-y>.

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