


CORRECTION

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Correction to: Comprehensive genetic characteristics of dystrophinopathies in China

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Correction to: Ma et al. *Orphanet Journal of Rare Diseases*
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The authors are grateful to the Editor of *Orphanet Journal of Rare Diseases* for allowing them the opportunity to publish this Correction, and apologize to the readership for any inconvenience caused.

A number of the variant descriptions in Additional file 1 of the original article [1] were not correct because of using different Refseq. The authors have now revised all the variant descriptions by Reference Sequence Database (NM_004006.2) in “Additional file 1” accompanying this Correction, as well as the original article.

All the authors agree to this Correction.

Supplementary Information

The online version contains supplementary material available at <https://doi.org/10.1186/s13023-021-01853-x>.

Additional file 1. Clinical and genetic information of patients with small mutation.

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1. Ma, et al. Comprehensive genetic characteristics of dystrophinopathies in China. *Orphanet J Rare Dis*. 2018;13:109. <https://doi.org/10.1186/s13023-018-0853-z>.

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