Correction to: Genetic analysis in Chinese patients with familial or young-onset amyotrophic lateral sclerosis

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In our recently published article, we quoted our previously published work [Shi J, Zhao F, Pang X, Huang S, Wang J, Chang X, Zhang J, Liu Y, Guo J, Zhang W. Whole-exome sequencing identifies a heterozygous mutation in SLC12A6 associated with hereditary sensory and motor neuropathy. *Neuromuscul Disord*. 2021 Feb;31(2):149–157. <https://doi.org/10.1016/j.nmd.2020.11.002>. Epub 2020 Nov 18. PMID: 33323309] in a part of the method of genetic testing but we forgot to fully acknowledge the source of the overlap among references. We wish to supplement this article in the references.

The authors apologize for this error and state that this does not change the scientific conclusions of the article in any way.

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The original article can be found online at <https://doi.org/10.1007/s10072-021-05634-z>.

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