

CORRECTION

Correction: Correction: Wild-Type Mouse Models to Screen Antisense Oligonucleotides for Exon-Skipping Efficacy in Duchenne Muscular Dystrophy

The PLOS ONE Staff

There is an error in the Correction published on November 15, 2018. In the last sentence of the caption for \$2 Fig, "mdx" should be "C57BL6." The publisher apologizes for the error. The correct sentence is: No difference was observed between treated and untreated C57BL6 mice. Please view \$2 Fig and the complete, correct \$2 Fig caption below.

Supporting information

S2 Fig. Routine hematoxylin and eosin staining for examining muscle morphology . Hematoxylin and eosin staining of TA tissue sections from treated C57BL6 mice with 2 μ g PMO, 5 μ g PNA and 5 μ g 2′Ome PS by local injection at different time-points e.g. 48 hr, 2 and 4 weeks after injection, and C57BL6 normal controls. Scale Bar = 100 μ m. No difference was observed between treated and untreated C57BL6 mice. (TIF)

References

- Cao L, Han G, Gu B, Yin H (2014) Wild-Type Mouse Models to Screen Antisense Oligonucleotides for Exon-Skipping Efficacy in Duchenne Muscular Dystrophy. PLoS ONE 9(11): e111079. https://doi.org/ 10.1371/journal.pone.0111079 PMID: 25365558
- Cao L, Han G, Gu B, Yin H (2018) Correction: Wild-Type Mouse Models to Screen Antisense Oligonucleotides for Exon-Skipping Efficacy in Duchenne Muscular Dystrophy. PLoS ONE 13(11): e0207817. https://doi.org/10.1371/journal.pone.0207817 PMID: 30440030



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