

CORRECTION

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

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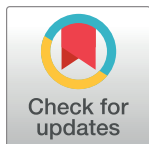
[S2 Fig](#) contains incorrectly duplicated panels. The authors have provided a corrected version here.

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 µg PMO, 5 µg PNA and 5 µg 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 µm. No difference was observed between treated and untreated *mdx* mice. (TIF)

Reference

1. 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



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