Supplementary materials

Genes and primer sequences used for this study are as follows:

Fragile X mental retardation gene 1, autosomal homolog (*Fxr1***)**: *Fxr1* encodes for Fragile X mental retardation syndrome-related protein 1, which is an RNA binding protein that regulates mRNA translation and is necessary for striated muscle function ^{39,40}.

Forward: GATAATACAGAATCCGATCAG

Reverse: CTGAAGGACCATGCTCTTCAATCAC

ATPase Ca²⁺ -transporting plasma membrane (*Atp2b1***):** *Atp2b1* encodes for plasma membrane Ca⁺² transporting ATPase 1 protein. This protein plays a critical role in calcium homeostasis, which is necessary for muscle contraction ⁴¹.

Forward: GTGGCCAGATCTTGTGGTTT

Reverse: CATCAATAAGGGGGATGTGC

Myotubularin-related protein 3 (*Mtmr3*): *Mtmr3* gene is related to myotubularin phosphatase implicated in myopathy ^{42,43}.

Forward: GTTGGCTACCTGACCACCTG

Reverse: CTCGACTGGGTTCAAAGAGC

Protein Phosphatase 3, catalytic subunit, alpha isoform (*Ppp3ca***)**: *Ppp3ca* encodes for the catalytic subunit alpha of calcineurin, which is a calcium dependent serine/threonine phosphatase. Calcineurin has functions in muscle growth, adaptation, and pathology specifically regulating cell differentiation and muscle fiber specification ⁴⁴⁻⁴⁷.

Forward: CTGGCGGGAAACAGACTCTG

Reverse: GTGGCATCCTCTCGTTAATTCG

Glutathione peroxidase 8 (*Gpx8***)**: *Gpx8* is involved in the anti-oxidant response ^{48,49} that is blunted under diabetic conditions contributing to oxidative stress ^{50,51}. Forward: GAAGGACCGTGTCTCTGGAA

Reverse: ATGGCTTCAAGGGGTTCTTC