

SUPPLEMENTARY FIG. S5. Increased mitochondrial oxidant levels in hypoxic myotubes upon miR-210 blocking. C2C12 mouse myoblasts were transfected with SCR or anti-miR-210. The next day, cells were switched to differentiation medium and allowed to differentiate for 48 h, followed by incubation in 1% O_2 hypoxic conditions for 24 h. Then, myotubes were stained with MitoSOX and Hoechst 33342 dyes, and fluorescence was revealed by fluorescence microscopy. (A) Representative pictures display MitoSOX and Hoechst 33342 fluorescences, either separate or merged. Calibration bar = 50 μ m. (B) The bar graph indicates MitoSOX fluorescence intensity normalized for the number of Hoechst 33342-positive nuclei (**p<0.002; n=6) MitoSOX-associated fluorescence was increased when miR-210 was inhibited. (C) The bar graph shows miR-210 levels in C2C12 cells transfected with anti-miR-210 or SCR. miR-210 levels were efficiently decreased by anti-miR-210 (n=6; *p<0.03).