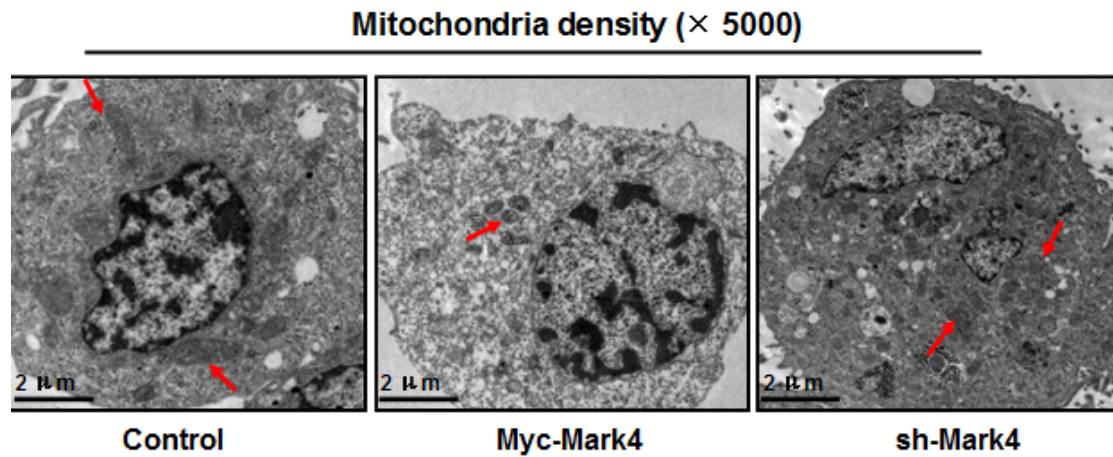
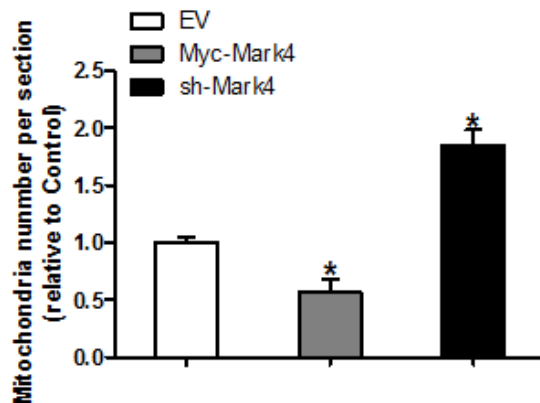


Figure S2

A



B



C

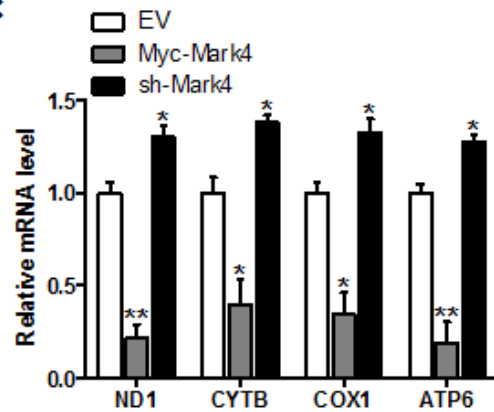


Figure S2. MARK4 decreases mitochondrial content in pig primary trophoblast cells challenged with 400 μ M NEFA. (a) Mitochondrial density assessed by electron microscopy in cytotrophoblasts after transfection with Myc-MARK4 and sh-MARK4 for 48h. Cells were then incubated with 400 μ M NEFA for 36h. Original magnification, $\times 5,000$; (b) Quantification of mitochondrial number per image area in (a) relative to control group (analysis of 10 random images per cell section, $n = 3$); (c) The relative mRNA level of various mitochondria-encoded genes was determined by quantitative RT-PCR in pig trophoblast cells ($n = 3$). Results were expressed as fold change relative to the values of control cells set to 1 unit. Values are expressed as mean \pm SEM. ** $P < 0.01$; * $P < 0.05$ compared with the control group. NEFA: non-esterified fatty acid; Myc-MARK4 group: over expression of MARK4 group; sh-MARK4 group: knock down of MARK4 group; Control: empty vector (EV) group; Red arrow: mitochondria.