

CARDIAC PROTEIN KINASE D1 ABLATION ALTERS THE MYOCYTES β -ADRENERGIC RESPONSE

SUPPLEMENTARY MATERIALS

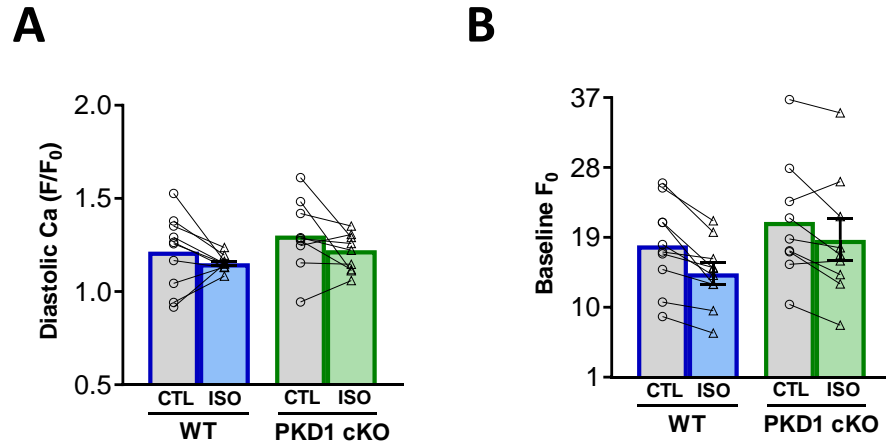


FIGURE S1. Diastolic Ca²⁺ changes were comparable in both WT and PKD1 cKO myocytes. **A**, Diastolic Ca²⁺ during pacing (minimum during pacing, F/F₀). **B**, baseline fluorescence (at rest, non-paced, F₀=F_{cell}-F_{background}). Same recordings as in Figure 1. Data points represent cells (WT, n = 9; KO, n = 8). Mice: WT, N = 5; KO, N = 4. Data are presented as mean \pm SEM. Two-way ANOVA, followed by Tukey's multiple comparisons test. Differences were considered statistically significant if P < 0.05.

FULL BLOTS USED ON FIGURE 5A-D

WT= wild type, KO= PKD1 cKO, Sal= saline, Iso= isoproterenol.

Blots were cut to probe separately with different antibodies and imaged in the ChemiDoc™ MP Imaging System (Bio-Rad) (see methods section in the manuscript).

The samples used in Figure 5A-D are labelled in **bold red font**.

MW marker: Precision Plus Protein Dual Color Standards (Bio-Rad).

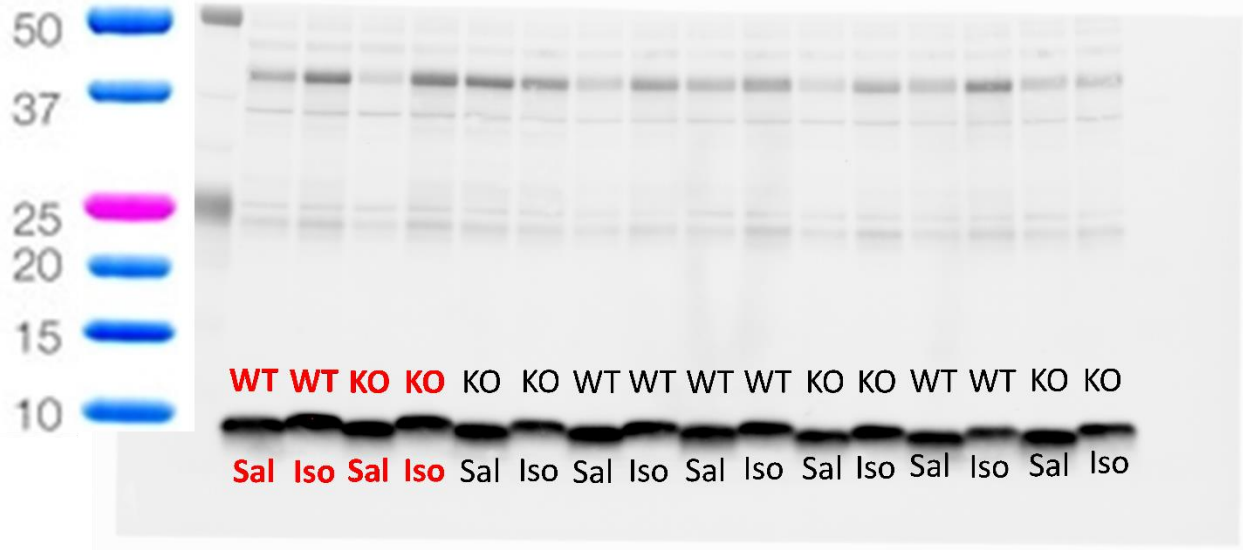


Full blots used on Figure 5Aa

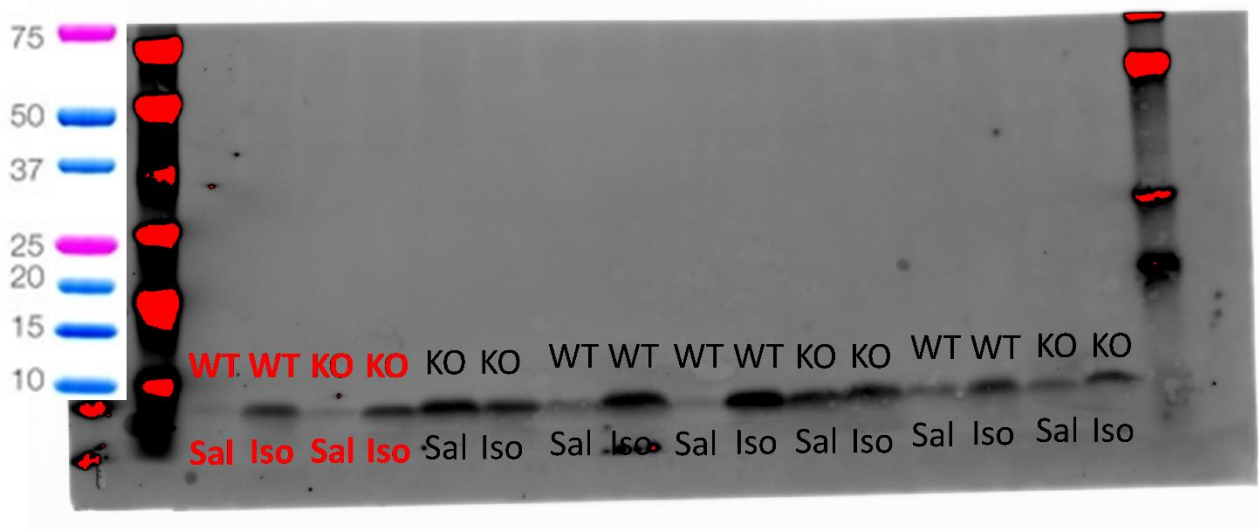
GAPDH



Total PLB

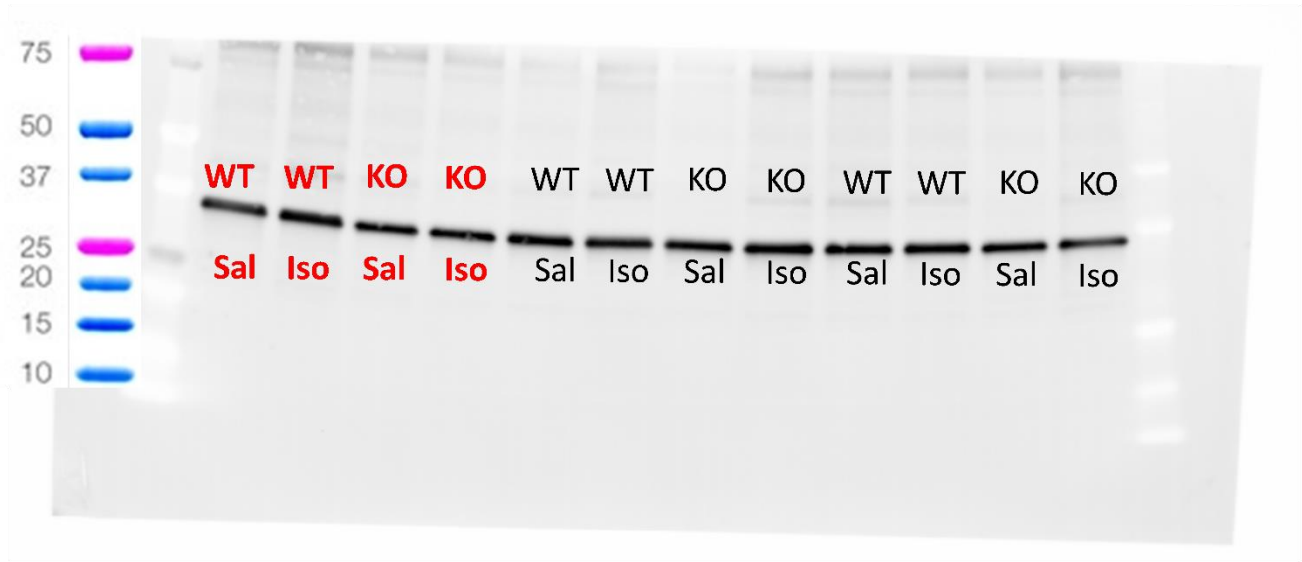


pPLB

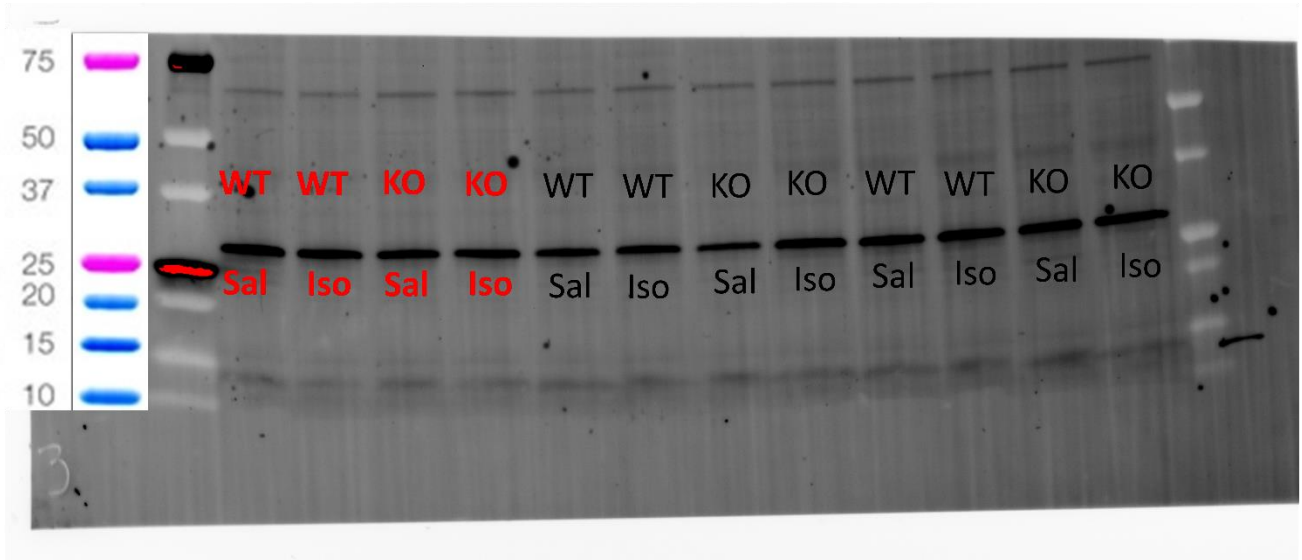


Full blots used on Figure 5Ba

GAPDH



Total Tnl

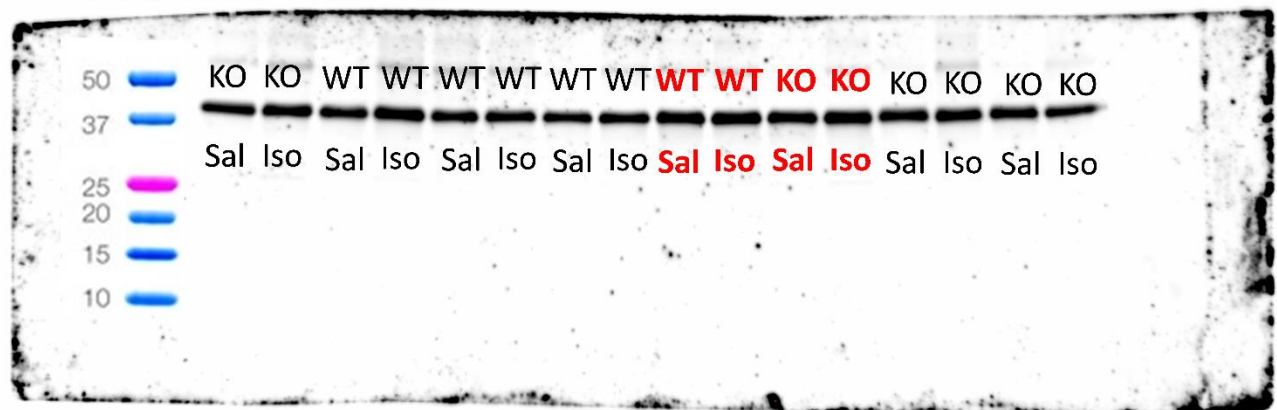


pTnl

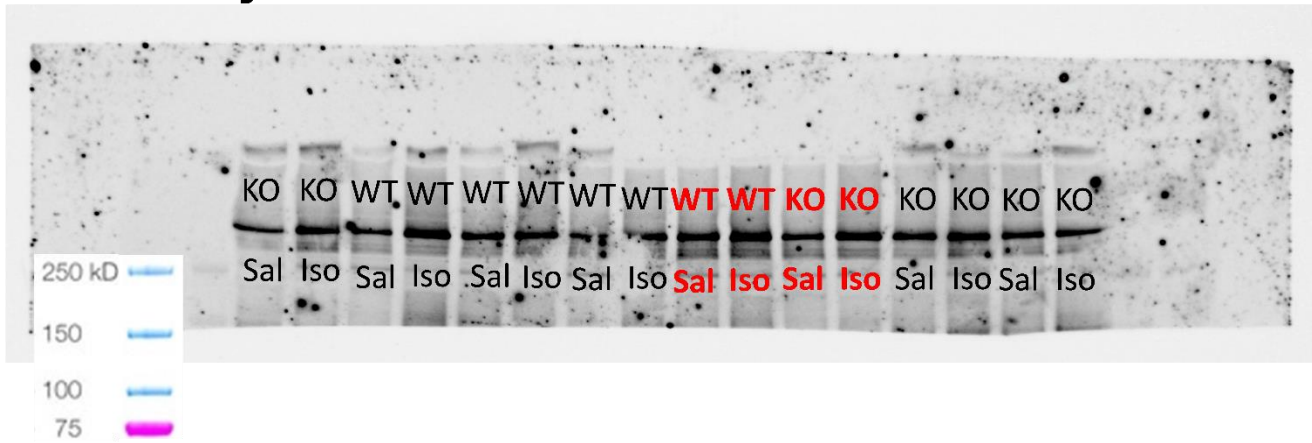


Full blots used on Figure 5Ca

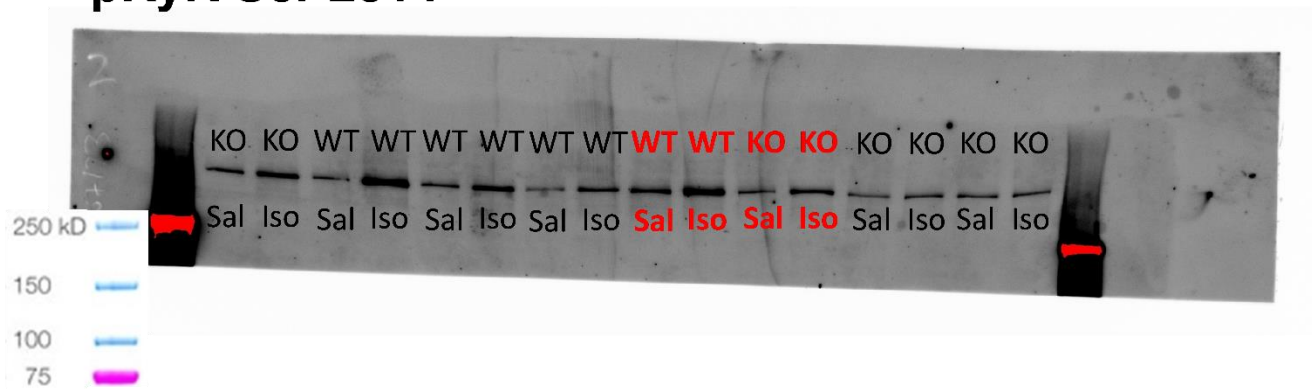
GAPDH



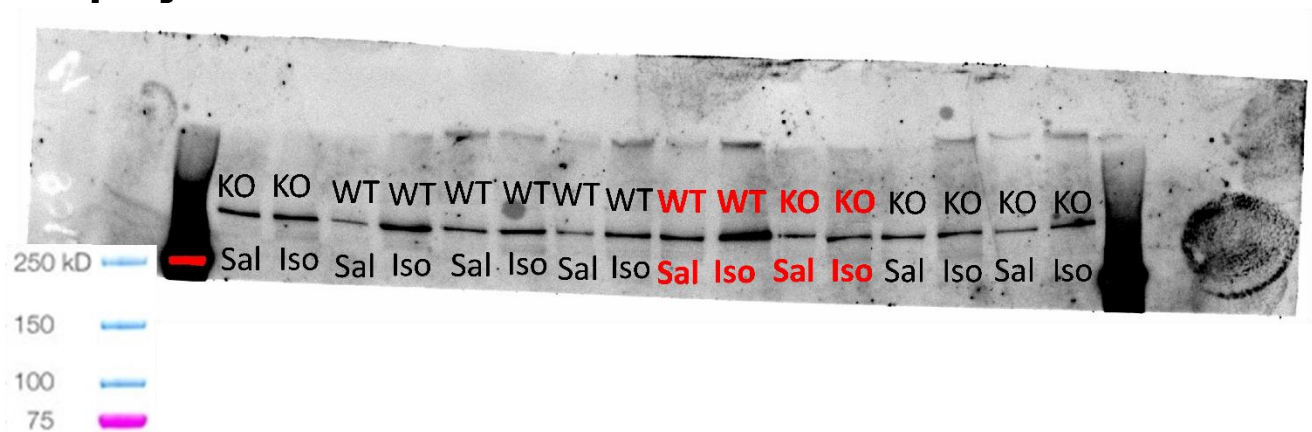
Total RyR



pRyR Ser 2814

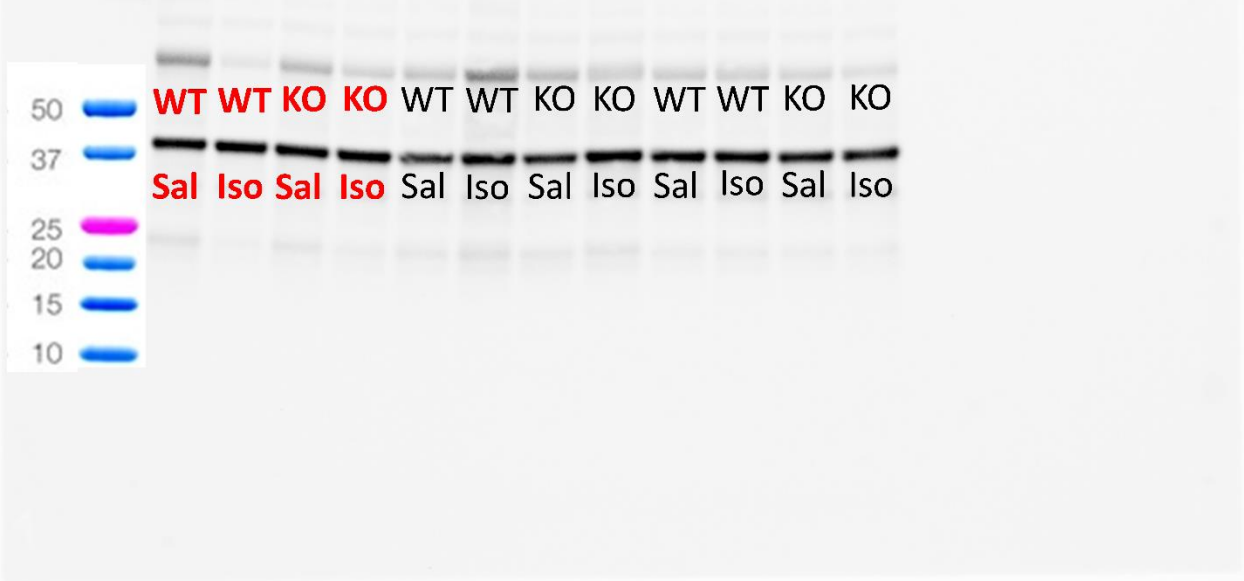


pRyR Ser2808



Full blots used on Figure 5Da

GAPDH



Total SERCA2

