

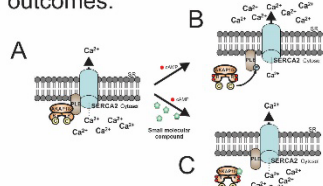
New therapeutic strategy targeting regulation of the SERCA2 complex protects from myocardial ischemia-reperfusion injury

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INTRODUCTION

The β -adrenergic receptor-cAMP-protein kinase A (PKA) signaling pathway regulates heart rate and contractility. Central in this regulation is the supramolecular complex PKA/AKAP18 δ /PLB/SERCA2 (Fig 1A). This complex controls the adrenergic effect on Ca²⁺ re-uptake and heart relaxation (Fig 1B). Discrete control of PLB phosphorylation is facilitated by the AKAP18 δ , which holds PKA and PLB in close proximity. We aimed to find small molecular compounds that disrupt the AKAP18 δ -PLB protein-protein interaction (PPI) (Fig 1C) as this may protect from ischemia reperfusion injury (IRI) in the treatment of acute myocardial infarction. Total infarct size is a key indicator of post MI outcomes.



CONCLUSION

Compound 13M is the first small molecule to target the AKAP18 δ -PLB interaction and is now in preclinical drug development phase. We propose that specifically blocking the adrenergic regulation of SERCA2-activity is beneficial and provide evidence that small molecular PPI disruptors that have such a mechanism-of-action reduce infarct size and preserve cardiac function.

RESULTS

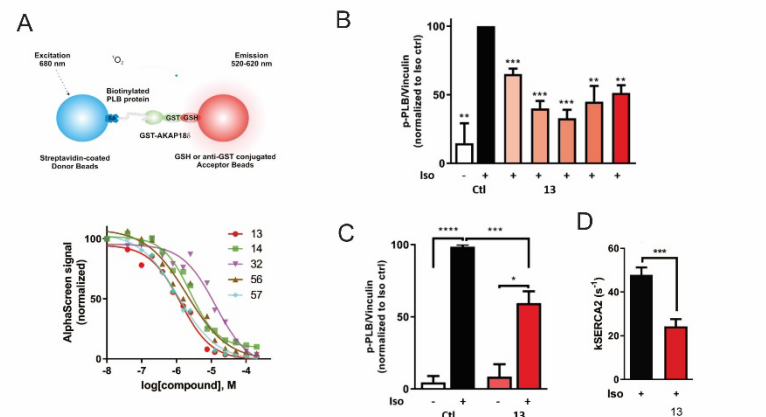


Figure 2: *In vitro* testing of compound 13. A) Principle of the bead-based AlphaScreen assay. AKAP18 and PLB binding with increasing concentration of different compounds. B) PLB phosphorylation (p-PLB) on H9C2 cell line. C) p-PLB on rat adult cardiomyocytes. D) SERCA2 activity determined by field stimulated in adult cardiomyocytes.

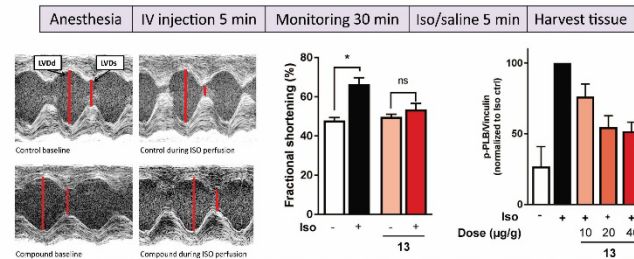


Figure 3: *In vivo* testing of compound 13 - Initial efficacy data on rats. Illustration of the protocol. Compound blocks Isoproterenol effect on the heart contractility measured by echocardiography (fractional shortening; FS=(LVdD-LVDs)/LVdD*100) and phospholamban phosphorylation.

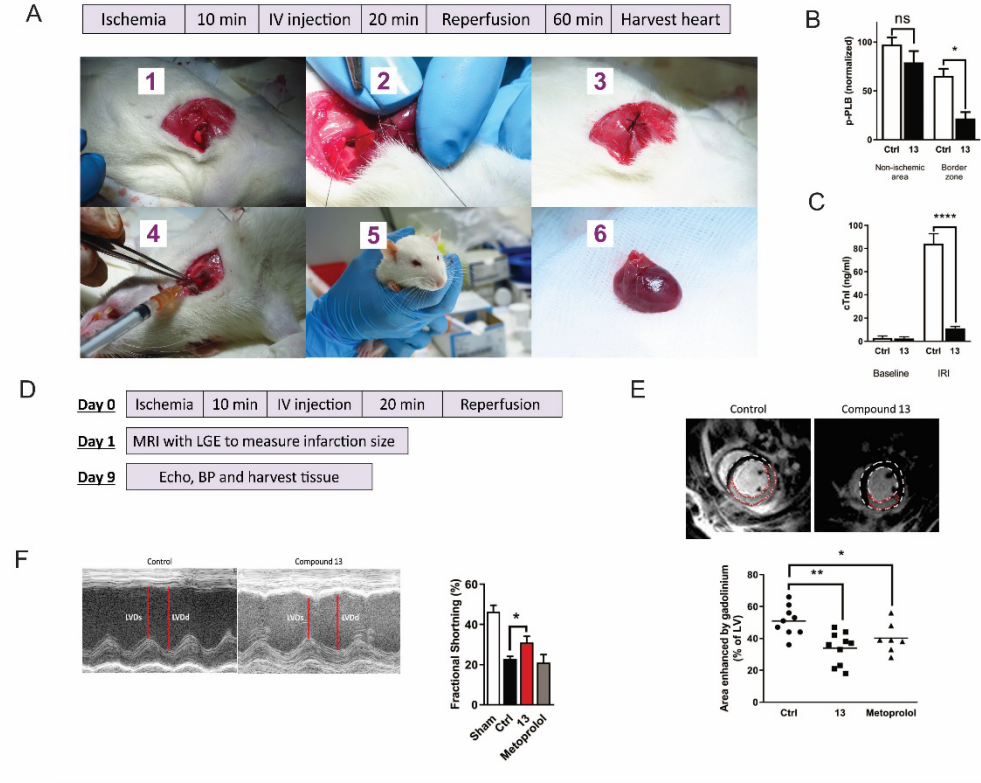


Figure 4: IRI disease model . A) Illustration of the protocol for the ischemia reperfusion injury (IRI) model. B) Compound 13 decrease p-PLB in the border area, but not in the non-ischemic region. C) Compound decreases troponin in blood. D) Illustration of the protocol for the IRI model. E) MRI analysis showed a decrease in area enhanced by gadolinium in animals treated with compound 13 and with Metoprolol. F) Echocardiography on day 9 show that treated animals have better contractility.

DISCLOSURE

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