



Activation of ϵ PKC reduces reperfusion arrhythmias and improves recovery from ischemia: Optical mapping of activation patterns in the isolated guinea-pig heart

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ABSTRACT

Pervious biochemical and hemodynamic studies have highlighted the important role of ϵ PKC in cardioprotection during ischemic preconditioning. However, little is known about the electrophysiological consequences of ϵ PKC modulation in ischemic hearts. Membrane permeable peptide ϵ PKC selective activator and inhibitor were used to investigate the role of ϵ PKC modulation in reperfusion arrhythmias. **Methods:** Protein transduction domain from HIV-TAT was used as a carrier for peptide delivery into intact Langendorff perfused guinea pig hearts. Action potentials were imaged and mapped (124 sites) using optical techniques and surface ECG was continuously recorded. Hearts were exposed to 30 min stabilization period, 15 min of no-flow ischemia, followed by 20 min reperfusion. Peptides (0.5 μ M) were infused as follows: (a) control (vehicle-TAT peptide; TAT-scrambled $\psi\epsilon$ RACK peptide); (b) ϵ PKC agonist (TAT- $\psi\epsilon$ RACK); (c) ϵ PKC antagonist (TAT- ϵ V1). **Results:** Hearts treated with ϵ PKC agonist $\psi\epsilon$ RACK had reduced incidence of ventricular tachycardia (VT, 64%) and fibrillation (VF, 50%) compared to control (VT, 80%, $P < 0.05$) and (VF, 70%, $P < 0.05$). However, the highest incidence of VT (100%, $P < 0.05$) and VF (80%) occurred in hearts treated with ϵ PKC antagonist peptide ϵ V1 compared to control and to ϵ PKC agonist $\psi\epsilon$ RACK. Interestingly, at 20 min reperfusion, 100% of hearts treated with ϵ PKC agonist $\psi\epsilon$ RACK exhibited complete recovery of action potentials compared to 40% ($P < 0.05$) of hearts treated with ϵ PKC antagonist peptide, ϵ V1 and 65% ($P < 0.05$) of hearts in control. At 20 min reperfusion, maps of action potential duration from ϵ PKC agonist $\psi\epsilon$ RACK showed minimal dispersion (48.2 ± 9 ms) compared to exacerbated dispersion (115.4 ± 42 ms, $P < 0.05$) in ϵ PKC antagonist and control (67 ± 20 ms, $P < 0.05$). VT/VF and dispersion from hearts treated with scrambled agonist or antagonist peptides were similar to control. **Conclusion:** The results demonstrate that ϵ PKC activation by $\psi\epsilon$ RACK peptide protects intact hearts from reperfusion arrhythmias and affords better recovery. On the other hand, inhibition of ϵ PKC increased the incidence of arrhythmias and worsened recovery compared to controls. The results carry significant therapeutic implications for the treatment of acute ischemic heart disease by preconditioning-mimicking agents.

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1. Introduction

Recently, the identification of novel therapeutic targets for the prevention of ischemia/reperfusion-induced cardiac injury has gained considerable attention due to the importance of cardiovascular disease as a public healthcare problem worldwide [1]. Ischemic preconditioning is a powerful phenomena which confers protection to hearts subjected to brief episodes of ischemia prior

to the subsequent ischemic insult [2,3]. There is a strong evidence supporting the involvement of ϵ PKC activation in preconditioning [4–6]. If indeed activation of ϵ PKC is required for cardioprotection, an ϵ PKC selective peptide activator would be expected to act as a preconditioning mimetic. Although hemodynamic and biochemical data using an ϵ PKC-selective agonist in mouse [5,7] and porcine [8] hearts demonstrated that ϵ PKC translocation is required for protection from ischemic injury, the detailed electrophysiological effects of ϵ PKC isozyme modulation in intact hearts is lacking. In this study, we delivered selective membrane permeable peptide ϵ PKC modulators (activator and inhibitor) into intact whole hearts to investigate the electrophysiological role of PKC isozymes in

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reperfusion arrhythmias. Protein transduction domain from HIV-TAT peptide [8,9] was used as a carrier for peptide delivery and optical mapping technique was used to record action potentials in Langendorff perfused guinea pig hearts.

2. Material and methods

2.1. Surgical procedure and experimental setup

All experimental protocols were approved by the Animal Studies Subcommittee of the Department of Veterans Affairs, New York Harbor Healthcare System and carried out in accordance with EU Directive 2010/63/EU for animal experiments.

The details of the surgical procedure have been described elsewhere [10,11]. Briefly, Dunkin–Hartley guinea pigs of either sex, weighing between 300 and 400 g, were anesthetized by intraperitoneal injection of sodium pentobarbital (30 mg/kg) and heparinized (1000 U/kg). A mid-thoracotomy was performed and the hearts were rapidly excised and placed in cold oxygenated Tyrode's solution containing 10,000 U/l heparin. The excised heart was rapidly cannulated at the aorta and retrogradely perfused in a modified Langendorff apparatus. The Tyrode's solution contained (in mM): NaCl 130, KCl 4.75, CaCl₂ 1.0, MgSO₄ 1.2, NaHCO₃ 12.5, and glucose 15.0. The solution was continuously bubbled with 95% O₂/5% CO₂ through a fritted glass tube. Temperature was maintained at 36 ± 0.3 °C by monitoring the temperature of the perfusate within the closed chamber.

2.2. Langendorff preparation and perfusion chamber

Experiments were started after the heart was stabilized and a steady perfusion pressure of 60–70 mmHg during sinus rhythm was maintained. A custom designed perfusion chamber was used for studying intact hearts using optical recording methods [10,12–14]. Bipolar surface electrograms were recorded using Teflon-coated silver wires (250-μm diameter) exposed at the tip and chlorided with an interpolar distance of 500 μm and positioned separately on the epicardial surface.

2.3. Fluorescent dye staining

A dye, di-4-ANEPPS (Molecular Probes; Eugene, OR), was used as the potentiometric fluorescent probe. Dye fluorescence was measured at wavelengths above the 645-nm cutoff filter when excited with a 520 ± 20 nm interference filter [10–14]. Hearts were

stained by gradual injection of 40–60 μl from a 2.5 mM stock solution of dye into a 5 ml bubble trap situated directly above the aortic cannula. The final dye concentration was approximately 1.8 μM; 10–15 min was allowed for the staining to complete. For longer protocols for which photobleaching and/or dye washout may reduce the optical signal amplitudes, hearts were restrained with smaller amounts of dye (5–10 μl) to restore the original signal-to-noise ratio.

2.4. Instrument setup

Details of the optical and recording apparatus have been described elsewhere [10,13,14]. The epicardial surface was illuminated with light from two 45 W tungsten halogen lamps, which was collimated and passed through 520 ± 20 nm interference filters. A 45° mirror in the optical apparatus was used to focus the grid pattern on the region of interest using a 35 mm camera lens (50 mm, F1:1.4, Nikon). Epi-fluorescent light from the stained heart was collected, projected through a 645 nm cutoff filter, and focused to form an image of the heart on a 12 × 12 element photodiode array. 124 diodes were current to voltage converted and sampled.

2.5. Experimental design

Hearts were subjected to a 30-min control stabilization period, followed by 15-min no-flow ischemia and then 20-min reperfusion. Optical recordings were performed every 2–5 min during the stabilization/perfusion period, every 2–3 min during ischemia, immediately following reperfusion and every 2–3 min thereafter. Permeable peptide (0.5 μM) modulators of εPKC isozyme, their scrambled peptides or vehicles were administered for 30-min prior to ischemia. The membrane permeable peptides, ψεRACK; (εV1–7 [HDAPIGYD]), which activates εPKC translocation and function [5,15] and εV1–2 [EAVSLKPT], which inhibits the translocation and function [5,15] of εPKC were conjugated to TAT peptide [YGRKKRRQRRR] via cysteine bond at the N-terminus for permeability [6,8]. For controls, we used scrambled peptides for ψεRACK [CPDYHDAGI], εV1–2 [LSETKPAV] and vehicle alone (TAT-peptide). Effective delivery of εPKC modulator peptides to intact hearts has been tested and demonstrated [8]. The peptides were synthesized from Genemed Synthesis, South San Francisco, CA and at the Protein and Nucleic Acid Facility, Stanford University, Stanford, CA. All peptides used were over 90% pure.

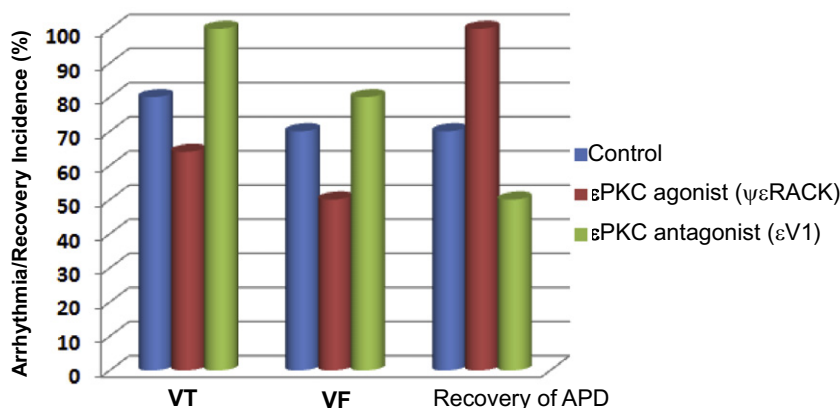


Fig. 1. Incidence of ventricular arrhythmia and recovery of action potentials during reperfusion in guinea-pig hearts. The incidence of ventricular tachycardia (VT), ventricular fibrillation (VF) and recovery of action potential duration at 90% (APD₉₀) at 20 min reperfusion are shown for untreated control, εPKC agonist ψεRACK (0.5 μM) and εPKC antagonist εV1 (0.5 μM) hearts. Hearts treated with εPKC agonist ψεRACK had the lowest incidence of VT/VF and complete recovery of APD₉₀ compared with hearts treated with εPKC antagonist εV1 and control.

Table 1
Summary of arrhythmic events.

	N	VT (%)	VF (%)	Recovery of APD90 (%)
Control	11	80	70	65
ϵ -PKC agonist $\psi\epsilon$ RACK	8	64	50	100
ϵ -PKC antagonist ϵ V1	8	100	80	40
Scrambled ϵ -PKC agonist $\psi\epsilon$ RACK	6	85	65	60
Scrambled ϵ -PKC antagonist ϵ V1	4	90	70	80
TAT-vehicle	4	75	70	70

VT: Ventricular Tachycardia; VF: Ventricular Fibrillation; APD90: Action Potential Duration at 90% repolarization.

2.6. Signal acquisition and data analysis

Activation time was defined as the peak temporal derivative of the action potential upstroke and recovery was defined as the point of maximum second derivative during repolarization. VT was defined as a successive run of at least 10 premature ventricular contractions. Data are presented as means \pm SE. The number of experiments (n) indicates the number of hearts used. Arrhythmia and optical mapping data from peptides treated and controls are compared by paired or unpaired Student's t -test as appropriate. A value of $P < 0.05$ was considered significant.

3. Results

A total of 41 hearts were used as follows: 11 control hearts which did not receive any treatment; 8 hearts were treated with

ϵ PKC agonist peptide, $\psi\epsilon$ RACK; 8 hearts were treated with ϵ PKC antagonist peptide, ϵ V1; 6 hearts were treated with scrambled ϵ PKC agonist $\psi\epsilon$ RACK (negative control); 4 hearts were treated with scrambled ϵ PKC antagonist peptide (negative control) and 4 hearts were treated with TAT alone (vehicle).

Hearts treated with ϵ PKC agonist $\psi\epsilon$ RACK had significantly reduced incidence of ventricular tachycardia, VT (64%) and fibrillation, VF (50%) compared to control VT (80%, $P < 0.05$) and VF (70%, $P < 0.05$). However, the highest incidence of VT (100%, $P < 0.05$) and VF (80% ($P < 0.05$)) occurred in hearts treated with ϵ PKC antagonist peptide ϵ V1 compared to control and to ϵ PKC agonist $\psi\epsilon$ RACK (Fig. 1). At the end of reperfusion (20 min), hearts treated with ϵ PKC agonist $\psi\epsilon$ RACK had 100% recovery of action potential duration measured at 90% repolarization (APD₉₀) compared to 40% ($P < 0.05$) of hearts treated with ϵ PKC antagonist peptide ϵ V1 and 65% ($P < 0.05$) in control hearts (Fig. 1). The incidence of arrhythmias and APD₉₀ recovery in hearts treated with scrambled peptides or vehicle was similar to control hearts (Table 1).

Fig. 2 illustrates selected action potentials and surface ECG recorded from control, ϵ PKC agonist $\psi\epsilon$ RACK and ϵ PKC antagonist ϵ V1 during stabilization period (panels A, E, I), 15 min of ischemia (panels B, F, J), during early reperfusion (panels C, G, K) and at the end of 20 min reperfusion (panels D, H, L). Ischemia resulted in action potential shortening in control, ϵ PKC agonist $\psi\epsilon$ RACK and ϵ PKC antagonist ϵ V1 (panels B, F and J, respectively). Early reperfusion caused VF in both control (panel C) and ϵ PKC antagonist ϵ V1 (panel K), but only a self terminating VT run (panel G) in the ϵ PKC agonist $\psi\epsilon$ RACK. At the end of reperfusion (20 min), complete recovery of action potential (panel H) was observed in the ϵ PKC agonist $\psi\epsilon$ RACK, however a slow VT was still present in the ϵ PKC

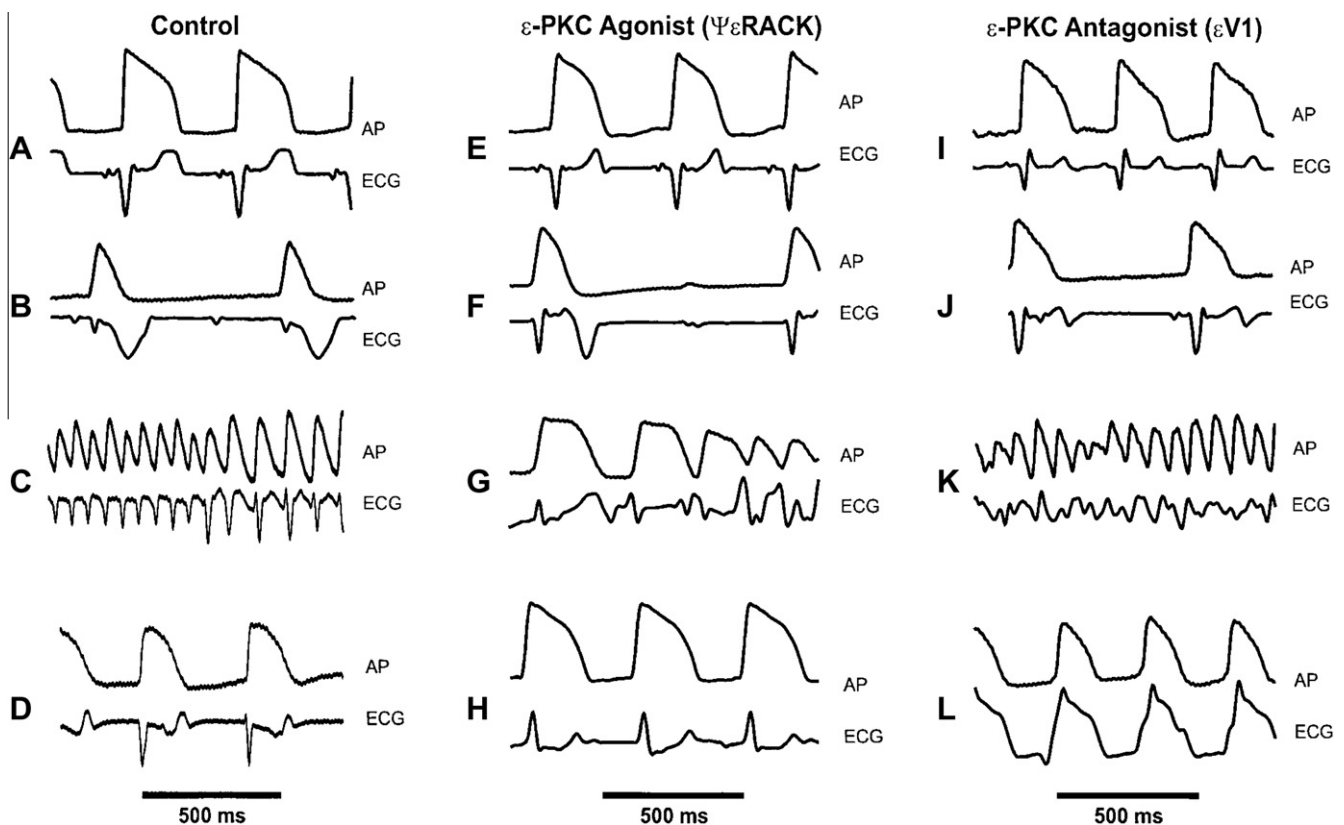


Fig. 2. Action potentials and electrocardiograms from guinea pig hearts. Action potentials and ECGs were recorded from untreated control, ϵ PKC agonist $\psi\epsilon$ RACK and ϵ PKC antagonist hearts. In the control, (A) represents recordings at 30 min stabilization period, (B) at 15 min global ischemia, (C) at 10 s of reperfusion with an onset of ventricular fibrillation (VF) and (D) at 20 min reperfusion demonstrating partial recovery of action potentials. In the ϵ PKC agonist $\psi\epsilon$ RACK (0.5 μ M), (E) represents recordings at 30 min stabilization period, (F) at 15 min global ischemia, (G) at 30 s reperfusion with self terminated ventricular tachycardia (VT) and (H) at 20 min reperfusion demonstrating full recovery of action potentials. In the ϵ PKC antagonist ϵ V1 (0.5 μ M), (I) represents recordings at 30 min stabilization period, (J) at 15 min global ischemia, (K) at 20 s reperfusion showing VT and (L) at 20 min reperfusion demonstrating poor recovery of action potential and ECG with VT still present.

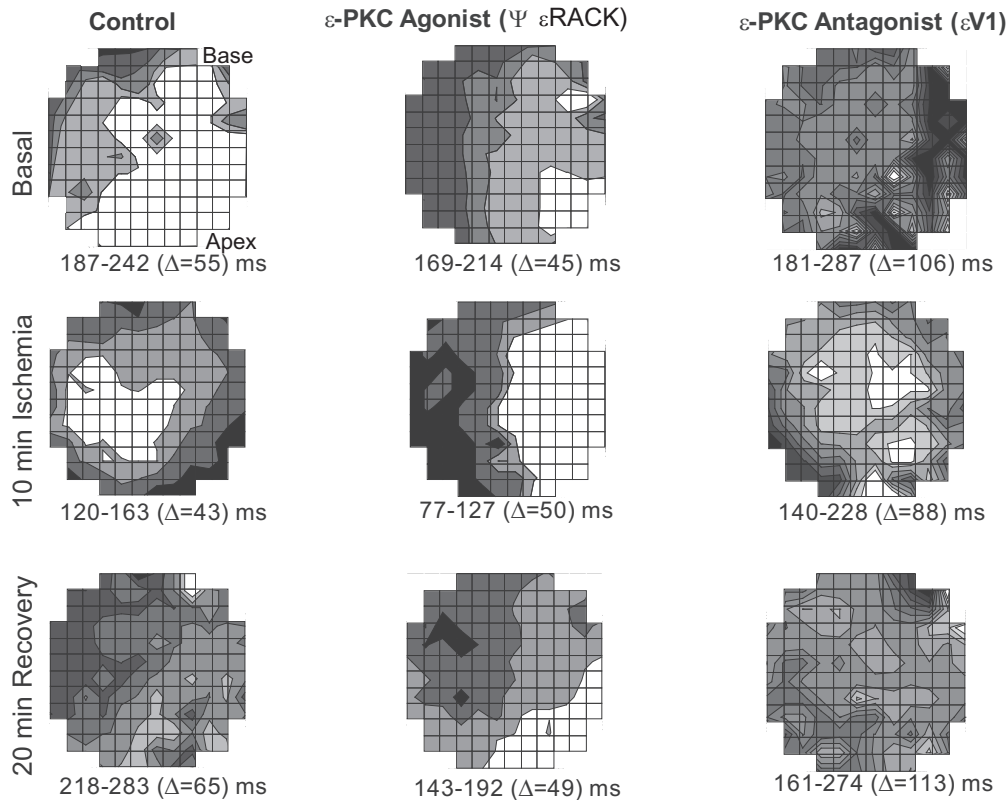


Fig. 3. Maps illustrating spatial distribution of action potentials recorded from guinea pig hearts. The maps were obtained at 10 ms isochrones of action potential duration at 90% (APD_{90}) during basal conditions, 10 min ischemia and 20 min reperfusion. During basal conditions (prior to ischemia), the control untreated hearts and ϵ PKC agonist $\psi\epsilon$ RACK treated hearts, APD_{90} had less dispersion of 55 ms and 45 ms, respectively compared with ϵ PKC antagonist ϵ V1 treatment (106 ms). During ischemia, the ϵ PKC antagonist ϵ V1 treatment resulted in the worst spatial heterogeneity of the shortening of APD_{90} with the highest dispersion (88 ms) compared to control (43 ms) and ϵ PKC agonist $\psi\epsilon$ RACK (50 ms). At the end of the 20 min reperfusion, ϵ PKC agonist $\psi\epsilon$ RACK treatment resulted in full recovery of APD_{90} dispersion (49 ms) similar to the dispersion prior to ischemia under basal conditions (45 ms). However, in the ϵ PKC antagonist ϵ V1 treated hearts, APD_{90} dispersion was the worst (113 ms) compared to untreated control (65 ms) and to ϵ PKC agonist ϵ V1 (49 ms).

antagonist ϵ V1 (panel L) and incomplete recovery of action potentials in control (panel D) was noted.

To further characterize the electrophysiologic properties of ϵ PKC modulation, action potential duration activation maps were obtained. In these maps (Fig. 3), each shaded zone represents an isochronal region activated at successive 10 ms intervals. The high incidence of VT/VF in the control and ϵ PKC antagonist ϵ V1 correlated with the high dispersion of APD_{90} during reperfusion compared to ϵ PKC agonist $\psi\epsilon$ RACK which showed minimal dispersion. At 20 min reperfusion, the average dispersion of APD_{90} in the ϵ PKC agonist $\psi\epsilon$ RACK, ϵ PKC antagonist ϵ V1 and control was 48.2 ± 9 , 115.4 ± 42 and 67 ± 20 ms respectively ($P < 0.05$ when compared to control). Fig. 3 shows that APD_{90} dispersion at 20 min reperfusion was 65, 49 and 113 ms during control, ϵ PKC agonist $\psi\epsilon$ RACK and ϵ PKC antagonist ϵ V1, respectively. Dispersion APD_{90} was the highest in the ϵ PKC antagonist ϵ V1 compared to control and ϵ PKC agonist $\psi\epsilon$ RACK during not only basal condition, but also during ischemia and reperfusion. Fig. 3 shows ADP_{90} dispersion of 106, 88 and 113 ms during basal condition, 10 min ischemia and at 20 min reperfusion respectively. On the other hand, APD_{90} dispersion in control and ϵ PKC agonist $\psi\epsilon$ RACK were 55, 43 and 65 vs. 45, 50 and 49 ms during basal condition, 10 min ischemia and 20 min reperfusion respectively.

4. Discussion

The primary finding of this study is that in the intact isolated heart, ϵ PKC activation by $\psi\epsilon$ RACK peptide (ϵ PKC agonist) resulted in a significant reduction of the incidence of reperfusion induced

VT/VF likely by reducing APD dispersion and by affording better action potential recovery. This novel anti-arrhythmic effect of the $\psi\epsilon$ RACK peptide has significant therapeutic implications for patients with ischemic heart diseases and in the development of new preconditioning mimicking agents.

4.1. Comparison with previous studies

The findings from this study are consistent with biochemical data demonstrating that $\psi\epsilon$ RACK peptide is cardioprotective against ischemic injuries when delivered in vivo before the ischemic period [8]. Specifically, we showed that $\psi\epsilon$ RACK peptide treatment resulted in significant reduction in infarct size, CPK release and improved ejection fraction. More importantly, $\psi\epsilon$ RACK peptide continuous delivery conferred sustained ϵ PKC activation without desensitization as seen in the settings of adenosine agonists or brief period of ischemic preconditioning [8]. The present data are also consistent with previous work in transgenic mice where moderate in vivo activation of ϵ PKC protects the ischemic heart from reperfusion arrhythmias [7] and moderate in vivo inhibition of ϵ PKC exacerbates the incidence of reperfusion arrhythmia [7]. Collectively, the present electrophysiological data demonstrating that post-reperfusion arrhythmias are significantly reduced by the $\psi\epsilon$ RACK peptide ϵ PKC agonist together with the above biochemical data showing cardioprotection from myocardial injuries point to a unique and potential novel pharmacological approach in treating both arrhythmias and myocardial injuries related to acute ischemic heart disease.

4.2. Potential mechanism (s) of anti-arrhythmic effect of ϵ PKC activation

Although, previous *in vitro* and *in vivo* studies using ϵ PKC modulating peptides and ϵ PKC transgenic mice have shown that ϵ PKC activation significantly improved the hemodynamic of the heart, attenuated cellular damage, and reduced infarction size [5,8,16], the role of ϵ PKC in ischemia–reperfusion related ventricular arrhythmias is poorly understood. Using optical maps, we have previously demonstrated that spontaneous Ca oscillations during ischemia/reperfusion resulted in premature ventricular beats that initiated runs of polymorphic VT/VF in a guinea-pig heart [12]. This tachyarrhythmia results in further increase in the level of Ca transient (Ca_T). Tachyarrhythmia-induced increase in Ca_T and the degeneration of the arrhythmia to VF may be related to the development of fast spontaneous Ca oscillations and/or Ca induced cell-to-cell uncoupling [12]. Using transgenic mice with specific cardiac constitutive activation of ϵ PKC, we found that sustained ϵ PKC resulted in reduced basal Ca current density as well as blunted β -adrenergic activation of Ca current [17] both of which are beneficial in the settings of undesirable catecholamine release during ischemia [18,19].

PKC regulates several ion channels including Na, K and Ca channels by phosphorylation [20–22]. Because these channels play critical role in action potential genesis and conduction, alteration in these channels function lead to abnormal electrical activity and arrhythmias. Previous data showed that ϵ PKC activation consistently inhibited Ca and Na channels [20,22] but increased the K channel (IKs). The combined effect of ϵ PKC activation on these channels may limit the amount of Ca entering the cell during ischemia especially the inhibition of the Ca current (less Ca entering the cell) combined with the activation of the K current, IKs (shortening of action potential thus limiting amount of Ca entry to the cell). Thus intracellular Ca accumulation during ischemia may play an important role in reperfusion arrhythmias through after depolarizations [11,12,23]. Consistent with the concept that interventions which limit Ca entry to the cell during ischemia, may reduce or eliminate reperfusion arrhythmias, earlier clinical and experimental studies have shown that verapamil, a Ca channel blocker, terminated reperfusion related arrhythmia [24,25]. However the negative hemodynamic side undermined its usefulness in the clinical settings. Unlike verapamil, $\psi\epsilon$ RACK peptide does not affect the hemodynamic parameters, heart rate and does not cause desensitization or downregulation of ϵ PKC in a porcine myocardial infarction model [8] making $\psi\epsilon$ RACK peptide a useful therapeutic agent for patients with ischemic heart disease. Altogether, this study demonstrates that in addition to the cardioprotective effects, ϵ PKC activation by $\psi\epsilon$ RACK peptide also suppressed the ischemia–reperfusion related ventricular tachyarrhythmia, thus providing a new therapeutic approach using preconditioning mimetic agents to manage patients with ischemic heart diseases.

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