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Sidorenko GI, Gelis LG, Medvedeva EA,

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artery bypass grafting in patients with  
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The present studies have demonstrated that  
protection with PP by activation of  $\beta$   
1-adrenoreceptors or adenosine receptors



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is identical to that of IP in the human myocardium. In addition, they have shown that mitoK ATP channels, PKC and p38MAPK are an integral part of the cellular signal transduction involved in this cardioprotection in which mitoK ATP channels are placed upstream and p38MAPK placed downstream of PKC.

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pharmacological research and

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risk factors and coronary artery disease on  
pharmacological protection the large  
majority of data demonstrating

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myocardium during myocardial infarction  
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of the institution of myocardial anoxia or ischemia may protect the myocardium by preventing atp depletion structural damage to cell membranes and organelles and comprehensive pharmacological protection of the myocardium section 1

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## ~~Pharmacological Protection of the Myocardium and ...~~

It is a technique of cardiac protection in  
which repeated brief interruptions of blood  
flow are performed before restoring final

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Of The Myocardium, improving  
myocardial metabolic recovery [1, 2].

~~Pharmacological protection of reperfusion  
injury in ST...~~

It protects against both myocardial  
infarction and stunning. This phenomenon  
involves the development of an altered

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myocardial cell structure which is dependent on stress responsive gene transcription leading to the synthesis of cardioprotective proteins, including most importantly cyclooxygenase-2 and inducible nitric oxide synthase.

~~Perioperative myocardial protection | BJA~~

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Overall, our results show that angiotensin-(1-9) confers pharmacological post-conditioning of the myocardium against I/R injury and this effect is mediated by AT<sub>2</sub> R and Akt. Thus, this peptide may be considered a valuable potential therapeutic agent in the setting of

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acute ischemia and post MI and future studies are required to thoroughly explore its cardioprotective power and translational value.

## Advances

~~Protection of the myocardium against ischemia/reperfusion ...~~

Pharmacological protection of acute

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ischemic myocardial injury was studied using diltiazem, bunazosin, coenzyme-Q10 (Co-Q), and nicorandil, in dogs of which the left anterior descending coronary artery was ligated for 60 min. Drugs were given intravenously prior to and/or during coronary ligation.

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~~Protection of acute ischemic myocardial  
injury by ...~~

Pharmacological protection of reperfusion  
injury in ST-segment elevation myocardial  
infarction. Gone with the wind? By

Elisabetta Tonet, Davide Bernucci,  
Giampaolo Morciano and Gianluca  
Campo. Cite . BibTex; Full citation

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Abstract. Primary percutaneous coronary  
intervention (PCI) represents the greatest  
progress in the treatment of ST-elevation  
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~~Pharmacological protection of reperfusion  
injury in ST ...~~

The rational approach to pharmacological



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protection of the myocardium has been to prevent the metabolic and structural changes that take place during and immediately after anoxic cardiac arrest.

Advanced

Within 7 to 10 minutes after arrest, tissue  $P_{O_2}$  levels in the myocardium drop to essentially zero, and anaerobic metabolism then proceeds at a nearly uniform pace

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until reoxygenation [ 11].  
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