Interaction of Insulin and AMPK in the Ischemic Heart
Another Chapter in the Book of Metabolic Therapy?

Rong Tian, James A. Balschi

There has been a longstanding interest in the development of metabolic modulations to improve the outcome of myocardial ischemia. The glucose-insulin-potassium (GIK) regimen presents the classic example of such a quest. For more than four decades since GIK was first described, substantial evidence from animal research and small size clinical studies has supported the efficacy of GIK treatment for acute myocardial ischemia. However, a recent large clinical trial, CREATE-ECLA, showed no effect of GIK in patients with acute ST-segment elevation myocardial infarction. Although this is extremely disappointing for supporters of metabolic treatment, careful evaluations of why GIK got lost in translation will benefit future development of therapeutic strategies based on metabolic principles. The study by Folmes et al published in this issue of Circulation Research sheds some light on this matter.

Decreased perfusion during myocardial ischemia limits the delivery of oxygen and nutrients and decreases oxidative metabolism. Under these conditions, increased glycolytic ATP production becomes a critical energy source (Figure). Several mechanisms have been proposed to account for the increases in glucose uptake and glycolysis during myocardial ischemia. These include: α-adrenergic mechanisms, p38 activation, and, more recently, stimulation of AMPK-activated protein kinase (AMPK). AMPK acts as an intracellular energy sensor. Its activity is highly sensitive to the cellular AMP to ATP ratio. AMPK activity markedly increases in response to energy depletion (increased AMP/ATP) during myocardial ischemia. Previous studies suggested that the increased AMPK activity promoted glycolytic ATP production and, thereby, protected against ischemic injury. It was also suggested, however, that stimulation of fatty acid oxidation by AMPK during reperfusion was detrimental to posts ischemic recovery. This is because increased fatty acid oxidation would suppress glucose oxidation, whereas continued glycolysis will result in cytosolic proton production and intracellular acidosis (Figure). This uncoupling between the rates of glycolysis and glucose oxidation could be exaggerated when enhanced glycolysis occurred in the presence of high circulating fatty acids, because the latter would facilitate fatty acid oxidation, which suppresses glucose oxidation, thereby increasing the "uncoupling" and predisposing the reperfused myocardium to further injury.

Multiple mechanisms have been proposed to explain the benefit of providing insulin and glucose to ischemic hearts (Figure). First, this regimen enhances glycolytic ATP production during restricted perfusion and impaired oxidative ATP production. Second, high insulin will suppress the high blood fatty acid levels that are commonly observed in patients with acute ischemic syndrome. Third, activation of PI3K/Akt signaling by insulin promotes cell survival and protects against apoptosis. Finally, recent studies suggest that insulin, via activation of Akt, inhibits AMPK activity by phosphorylation of Ser 485 of α-AMPK. This interesting finding provides a mechanism by which insulin could antagonize the adverse effects of AMPK in posts ischemic hearts. The study by Folmes et al critically tested this hypothesis in isolated mouse hearts. They reported that although insulin inhibited AMPK activity and improved posts ischemic recovery in hearts perfused with glucose as the only substrate, the inhibitory effect of insulin on AMPK activity was not observed when palmitate was present in the perfusate at either low (0.2 mmol/L) or high (1.2 mmol/L) concentration. Furthermore, in contrast to the improved function observed in glucose-perfused hearts treated with insulin, insulin impaired the post-ischemic recovery in hearts perfused with the high concentration of palmitate.

Although this study was performed in vitro it illustrates several important considerations regarding metabolic modulation for the management of myocardial ischemia. One critical issue is the level of circulating fatty acids to which the heart is exposed during ischemia and reperfusion. The study reports that high levels of fatty acids, commonly observed in patients during acute stress, abolished the benefit of insulin. This result suggests that the insulin effects observed in ex vivo animal studies may not extend to clinical situations. However, a potential effect of GIK treatment is the insulin-mediated suppression of circulating fatty acid levels. If so, the peripheral effect of insulin may also serve or contribute to its cardiac action. Such an effect was only tested in small groups of patients in early studies of GIK treatment when the management of myocardial ischemia was drastically different from that of today. Low-molecular weight heparin used during reperfusion therapy releases lipoprotein lipase into circulation that could counteract the lowering of free fatty acids by insulin, thus the effect of insulin on plasma levels of free fatty acids should be reevaluated in today’s clinical studies.

The study by Folmes et al also suggests that the amount of insulin supplied to the heart might contribute to the differen-
...and suggests that reperfusion with insulin can eliminate the adverse effects of AMPK-mediated myocardial protection and suggests that reperfusion...low-flow ischemia, a simulation of under-perfusion, contributes significantly to postischemic recovery.11,14 Another interesting yet unanswered question raised by the study of Folmes et al. is why insulin exerts opposite effects on postischemic recovery in hearts perfused with and without high concentrations of fatty acid. In hearts perfused with high concentration of fatty acid, insulin suppressed the fatty acid oxidation rate without an effect on AMPK activity. The net proton production from uncoupled glycolysis was essentially identical in reperfused heart receiving insulin regardless of whether they were reperfused with glucose or glucose plus palmitate, suggesting additional mechanism(s) may yet be discovered. In this regard, Folmes and colleagues have met the criteria for carrying out an excellent study; that is, to raise new questions for investigation. Defying predictions that the CREATE-ECLA Trial would put the issue of metabolic modulation therapy to rest, the study by Folmes has likely just opened a new chapter.

Acknowledgments

We thank Linda Johnson for her professional assistance in preparing the manuscript.

Sources of Funding

The authors’ work is supported by research funds from the National Institute of Health HL46033 (to J.A.B.), HL67970 and HL59246 (to R.T.). R.T. is an Established Investigator of the American Heart Association.

Disclosures

None.

References


**Key Words:** insulin □ AMPK □ glucose □ myocardial ischemia □ metabolic modulation
Interaction of Insulin and AMPK in the Ischemic Heart: Another Chapter in the Book of Metabolic Therapy?
Rong Tian and James A. Balschi

_Circ Res._ 2006;99:3-5
doi: 10.1161/01.RES.0000233142.26369.f6

_Circulation Research_ is published by the American Heart Association, 7272 Greenville Avenue, Dallas, TX 75231
Copyright © 2006 American Heart Association, Inc. All rights reserved.
Print ISSN: 0009-7330. Online ISSN: 1524-4571

The online version of this article, along with updated information and services, is located on the World Wide Web at:
http://circres.ahajournals.org/content/99/1/3

Permissions: Requests for permissions to reproduce figures, tables, or portions of articles originally published in _Circulation Research_ can be obtained via RightsLink, a service of the Copyright Clearance Center, not the Editorial Office. Once the online version of the published article for which permission is being requested is located, click Request Permissions in the middle column of the Web page under Services. Further information about this process is available in the Permissions and Rights Question and Answer document.

Reprints: Information about reprints can be found online at:
http://www.lww.com/reprints

Subscriptions: Information about subscribing to _Circulation Research_ is online at:
http://circres.ahajournals.org/subscriptions/