The early cardiac physiologists discovered that depriving the heart of its blood supply led to an acceleration of glycolysis which could provide anaerobic energy. Richard Bing\(^1\) in the 1950s used the novel tool of coronary sinus catheterisation to study the energy sources of human heart muscle in patients with heart failure. He showed that for its energy sources the heart uses mostly fatty acids and to a lesser extent glucose. Circulating free-fatty acids (FFA) could inhibit myocardial glucose oxidation.\(^2\) In 1962 Sodi-Pallares\(^3\) launched the concept of metabolic therapy using glucose-insulin-potassium (GIK) solutions that reduced acute electrocardiographic signs of ischaemic damage.

The mechanism whereby GIK provides benefit is in part by reducing the toxic high FFA levels that are found in early acute myocardial infarction (AMI) as a result of major catecholamine stimulation with acute adipose tissue lipolysis (Fig. 1).\(^4,5\) In heart failure (HF), beta-blockade and GIK also reduce circulating FFA, while the newer drugs trimetazidine and ranolazine act metabolically to improve the condition of patients with ischaemia or HF.\(^6\)

**Metabolic therapy must be given early**

Almost all of the prior GIK studies for patients with acute coronary syndromes (ACS) started infusions far too late after the onset of symptoms. During the critical first hour, GIK infusions could potentially reduce major energy depletion. Crucial experimental support for this concept came from relevant work at the University of Cape Town (UCT) supported by the Chris Barnard Fund,\(^7\) thus supporting the rationale for the positive IMMEDIATE study in which GIK was infused by paramedics to patients in the prehospital emergency ambulance setting.\(^8\) Started immediately after the onset of symptoms, and continued thereafter, GIK reduced by 40% the rate of the combined cardiovascular endpoint, cutting in-hospital mortality or cardiac arrest by half, besides reducing infarct size measured at 30 days.\(^8\)

**Molecular cardioprotection**

The extraordinary complexity of the molecular protective pathways must, like the metabolic paths, have evolved millions of years ago when rapid cardioprotection was required after the hyperadrenergic stresses and blood losses experienced when hunting and escaping from wild animals.\(^9\) Today these paths can be brought into action during prompt therapy for acute coronary occlusion by rapid reperfusion, as in AMI. Although saving many cells otherwise threatened with ischaemic cell death, rapid reperfusion kills a significant percentage of cells that could have been saved by the appropriate intervention. Indeed, the time has come to take reperfusion injury seriously!\(^10\)

Reperfusion damage, with the sudden return of oxygen and reversal of tissue pH changes, is inevitable during the optimal therapy of AMI by prompt percutaneous coronary intervention (PCI). Working on this problem, the group of Hausenloy and Yellon\(^10\) in London discovered the RISK (Reperfusion Injury Salvage Kinases) path and the SAFE (Survival Activating Factor Enhancement) path, on which our group in Cape Town has focused. These paths help to lessen ischaemic-perfusion damage, and may, hypothetically, also be activated by intense exercise.

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**Corresponding author:** L H Opie (lionel.opie@uct.ac.za)
the proposal that insulin therapy, a promoter of optimal cardiac metabolic protection, can directly promote cardiac cell survival during reperfusion. From the evolutionary point of view it would have been inadequate to have only one molecular cardioprotective path. Thus another path was discovered, namely the SAFE (Survival Activating Factor Enhancement) pathway, that is the focus of our group under Sandrine Lecour. This path involves cytokine protection by low levels of tumour necrosis factor-alpha (TNF-alpha) acting on a specific series of molecular events and leading to activation of transcription factor signal transducer and activator of transcription-3 (STAT-3).

The jump from experiments to clinical certainty

Postconditioning by balloon inflation-relation

Can the basic laboratory data be clinically applied? In animals such as the baboon, the major part of myocardial metabolic damage occurs in the first hour. The first human studies with early intervention were with ischaemic postconditioning. That means that after coronary flow had been abruptly restored to the previously ischaemic myocardium by PCI, the intra-coronary balloon was blown up again to cause temporary ischaemia. The balloon was then released only to be briefly blown up again, a procedure repeated for five cycles, thereby causing a very significant reduction in infarct size as in the basic observations made by Thibault et al. With late reperfusion, after the crucial first 3 hours, the extent of reperfusion damage can be expected to be slight and the benefit of therapy aimed at reperfusion damage likewise slight or totally absent.

Remote conditioning

An even more interesting approach is remote conditioning, whereby intermittent occlusion-reperfusion of a limb acts at a distance to reduce cardiac reperfusion damage. First shown in patients with coronary bypass operations by Yellon’s group, the concept has now been widely extended to involve other organs such as kidney and brain. This approach has found human application by the simple procedure of pumping up and down an ordinary blood pressure sphygmomanometer in the ambulance taking patients to hospital. The primary endpoint was myocardial salvage index at 30 days after primary PCI, measured, by myocardial perfusion imaging, as the proportion of the area at risk of cell death but salvaged by treatment. The area saved from cellular death was about 20 - 25%. This procedure has no known side-effects and reduced the extent of the infarcted tissue; benefit was seen even in infarcts sustained by patients who were very rapidly transferred to the emergency room for PCI, following the onset of symptoms, by the efficient ambulance service that Denmark enjoys. The vexing question of how remote ischaemic conditioning works (cardioprotection at a distance) is still a matter of dispute; there are arguments for both humoral and nervous mechanisms.

From near-certainty to wishful thinking

Wouldn't it be wonderful if we could imagine that certain foodstuffs or beverages contain protective agents that promote our molecular ischaemic pathways? Although this is fanciful, nonetheless there is a possibility that certain components of wine, namely resveratrol and melatonin, may protect during reperfusion injury during the therapy of heart attacks. The impact on myocardial infarct size after coronary occlusion and reperfusion following administration of red wine, resveratrol, and melatonin to rats has been examined in our laboratory. Both resveratrol and melatonin gave substantial protection. The group of Sandrine Lecour has in the past suggested that the resveratrol content of red wine may, with only modest intake, reach blood levels high enough to be cardioprotective although that conclusion is only inferential and hypothetical. More recently melatonin has been discovered in red wine and also in white. Melatonin is commonly used by persons threatened by jet lag during air travel because it promotes a natural sleep rhythm (and may be one of the reasons why red wine with meals goes with a good night’s sleep).

Speculatively, red wine is part of the French paradox with, however, many other factors contributing to the relative protection from myocardial infarction that the French enjoy, because of their traditional lifestyle (which regretfully is now fading). Overall, modest wine intake is of itself no panacea but rather only a small part of part of the healthy five-point lifestyle in which non-smoking and daily vigorous exercise, for more than 30 minutes, are the two top criteria. As fully argued in my recent book, Living Longer, Living Better, intense exercise not only acts by promoting preponderance of the vagus anti-adrenergic protective nervous system at rest, but also by stimulating the protective molecular paths.

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8. Sodhi-Pathan D, Behrmann, J, Sheshin PL, et al. A double-blind randomized controlled trial of intravenous glucose, insulin, and potassium (GIK) for acute coronary syndromes in emergency medical services. The IMMEDIATE (Immediate Myocardial Metabolic Enhancement during Initial Assessment and Treatment in Emergency Care) Trial. JAMA 2012; online 27 March 2012.
17. Opie LH. The five-point healthy lifestyle. SA Heart 2011;8:154-163.

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