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Nicorandil in myocardial ischaemia

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Studies of contemporary populations show that patients with angina pectoris have significantly impaired quality of life [1, 2]. Despite taking one or more conventional antianginal drugs, patients frequently experience three or more anginal episodes each week, and increased angina frequency, which is associated with decreased overall feeling of well-being. Ideally, antianginals should not only relieve symptoms with little or no side-effects, but they should also improve patient outcome [3]. The deficiencies of current medical therapy for angina are well recognised [4] and with the exception of aspirin, beta blockers and statins, it makes little or no impact on disease progression. In order to favourably improve outcome, aggressive cardiovascular risk factor management should be undertaken in parallel with stringent efforts to optimise and tailor medical treatment to the individual patient.

This review looks at a novel agent, the potassium adenosine triphosphate-(K\textsubscript{ATP}-)sensitive channel opener, nicorandil and its role in the treatment of myocardial ischaemia.

Background

Potassium channels, particularly those regulated by intracellular ATP are ubiquitous in the heart and blood vessels and are important modulators of cardiovascular function. Opening potassium channels, there is an increased efflux of potassium ions from the cell and the resting membrane potential is shifted towards more negative values (hyperpolarisation); this leads to an inhibition of calcium influx or indirect calcium antagonism, causing a fall in intracellular calcium concentration, relaxation of vascular smooth muscle cells and vasodilation.

Nicorandil was developed as the first oral K\textsubscript{ATP} channel opener in Japan where it became available in the early 1980s, since when it has been widely used in that country. It was introduced in the UK in 1994, and became available in Austria in 1997. Nicorandil is described as a hybrid compound, consisting of a nicotinamide vitamin group and an organic nitrate. Similarly to nitrates, nicorandil also leads to activation of the enzyme guanylate cyclase which in turn induces an increase in intracellular cyclic guanosine monophosphate (cGMP) and a decrease in intracellular calcium ions. Ultimately this results in reduction in vascular tone, predominantly in the venous vascular bed, so that mainly the venous capacitance vessels are dilated. Because of this dual mechanism of action, both preload and afterload are reduced producing a dose-related improvement in haemodynamics (Figure 1).

Nicorandil causes reduction in total peripheral resistance and blood pressure, at therapeutic concentrations this effect is small [5] and transient, and does not normally lead to first-dose hypotension. It improves resting coronary blood flow without changing cardiac or stroke volume indices and without causing reflex tachycardia or an increase in oxygen consumption. Nicorandil is not negatively inotropic at therapeutic doses and contractility is largely unchanged, it may therefore be used with relative safety in patients with left ventricular dysfunction, but it does not have a licence for the treatment of heart failure. Angiographic studies have shown that the drug dilates both stenotic and non-stenotic coronary arteries.

Unlike classical nitrates there appears to be an absence of haemodynamic tolerance to nicorandil. This was demonstrated in a study [6] of 25 patients with congestive heart failure who were infused with either glyceryl trinitrate (GTN) or nicorandil. The mechanism of action of nicorandil (KCO, Potassium channel opener)

**Figure 1.**
Nicorandil in myocardial ischaemia

has also been shown to prevent platelet aggregation. It has been of coronary ligation it may cause limitation of infarct size. It cardioprotective properties in a number of models of ischaemia/reperfusion, myocardial stunning, and in canine models used. It has been proposed that nicorandil may possess The term cardioprotection, although ill-defined, is now widely not totally dependent on its activation of cGMP to be effective.

Clinical experience

Nicorandil is indicated for the prevention and long-term treatment of angina, and is available (in the UK) as 10 and 20 mg tablets; the recommended starting dose is 10 mg twice daily which can be titrated upwards to a maximum of 30 mg twice daily, although the usual therapeutic range is 10–20 mg twice daily. It may be appropriate when starting therapy to advise patients who are susceptible to nitrate-induced or non-specific headache to divide the 10 mg scored tablet in two and to give 5 mg twice daily.

Clinical trials

Nicorandil has been extensively studied in Europe in more than 1500 patients with angina, both in placebo-controlled studies and in comparative studies with other antianginals. The majority of studies show that nicorandil is superior to placebo in reducing angina episodes and increasing exercise duration [7]. While one placebo-controlled study [8] did not show any significant improvement in angina frequency, total exercise duration, or time to ischaemia with nicorandil, another placebo-controlled study of 46 patients, two hours post-dosing, showed nicorandil improved time to onset of angina by 38%, without significant alteration in heart rate or blood pressure [9]. Nicorandil also appears to have comparable efficacy to nitrates, beta blockers and calcium antagonists.

In a comparative study of oral doses of nicorandil 20 mg twice daily and sossorbide-5-mononitrate 20 mg twice daily, both drugs showed comparable improvement in exercise duration and time to onset of angina [10]. In a comparative study with atenolol 50–100 mg twice daily, propranolol 40–50 mg twice daily and metoprolol 100 mg twice daily, nicorandil in doses 10 and 20 mg twice daily produced equivalent improvement in angina attack rate and exercise tolerance with chronic dosing [11], whereas atenolol significantly reduced heart rate, this did not change significantly in patients receiving nicorandil.

Nicorandil has also been compared to a number of calcium antagonists. In a randomized study of nicorandil 10 mg twice daily, increasing to 20 mg twice daily versus nifedipine 20 mg twice daily, both agents showed equal efficacy in terms of increased exercise duration and time to onset of angina, but, unlike nicorandil, nifedipine significantly increased resting heart rate [12].

Is nicorandil cardioprotective?
The term cardioprotection, although ill-defined, is now widely used. It has been proposed that nicorandil may possess cardioprotective properties in a number of models of ischaemia/reperfusion, myocardial stunning, and in canine models of coronary ligation it may cause limitation of infarct size. It has also been shown to prevent platelet aggregation. It has been suggested that nicorandil may mimic ischaemic preconditioning [13], a condition which offers powerful protection against ischaemic necrosis (see Yellon review in this issue, p: 8–11). The clinical value of these apparent properties of nicorandil remains to be established, however, by attenuation of ischaemia, nicorandil may play an important role in bypass surgery and confer protection in patients undergoing percutaneous transluminal coronary angioplasty (PTCA).

Findings from two recent studies in unstable angina [14] and acute myocardial infarction [15] support the view that nicorandil is safe and may have a protective role in acute ischaemic syndromes. A randomised, double-blind, placebo-controlled trial was conducted to assess the safety and efficacy of oral nicorandil in unstable angina. The study in 200 patients with unstable angina [14] showed that nicorandil, when added to aspirin, unfractionated heparin, nitrates, atenolol and diltiazem, significantly further reduced the numbers of episodes of transient myocardial ischaemia, supraventricular tachycardia and ventricular tachycardia, compared to placebo (Figure 2).

Beneficial effects of nicorandil were also seen in a small, placebo-controlled, double-blind safety study in 45 patients with acute myocardial infarction [15]. This showed a reduction in the incidence of arrhythmic events and the overall incidence of adverse events was lower in patients with acute myocardial infarction who received nicorandil orally. The drug also appeared to attenuate the progression of infarction in patients without Q-wave abnormalities at the time of inclusion. But nicorandil did not appear to attenuate infarct progression in those with Q-wave changes on entry to the study. While it is difficult to make firm conclusions on the cardioprotective effects of nicorandil, further studies to evaluate the safety and efficacy of nicorandil as an additive therapy in acute ischaemic syndromes are justified.

Outcome study

The first major outcome study, Impact of Nicorandil in Angina (IONA) is currently underway in the UK. The primary aim of the study is to show that treatment with nicorandil can reduce cardiovascular events in patients with stable angina. IONA is planned to recruit 5000 patients, men and women, and is being co-ordinated by the University of Glasgow. Patients will consist of existing or newly diagnosed angina pectoris, and will include those who have had a myocardial infarction or coronary artery bypass graft surgery. Patients will receive either nicorandil 20 mg twice daily or placebo, in ad-
Nicorandil may also have a role in individuals with obstructive or have profound ankle oedema with a calcium antagonist. Agents, particularly those who feel lethargic on beta blockade, are beneficial in patients who cannot tolerate either of these other 'add-on' to beta blockers and calcium antagonists. It may be effective as monotherapy in some, perhaps more mildly symptomatic, patients. It may be less useful for those who are headache-prone. It can be used safely as an alternative agent with comparable efficacy and tolerability to existing antianginals. Nicorandil may be particularly useful where some of these antianginals are contraindicated or ineffective. While the role of nicorandil in acute ischaemic syndromes remains to be defined, the results of small recently conducted studies give cause for optimism.

**References**


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