EDITORIAL

Should primary angioplasty be available for all patients with an ST elevation myocardial infarction?

A de Belder

For the acute myocardial infarction patient, percutaneous coronary intervention is clearly superior to thrombolysis for many clinical end points, yet widespread availability of PCI services is still far from being realised.

A patient with cardiac chest pain calls for the emergency services.

Scenario 1—Within minutes a trained paramedic crew has established the diagnosis of acute myocardial infarction (AMI), and transmits an ECG electronically to a myocardial infarction centre, where a coordinator mobilises the catheter laboratory staff to prepare for angioplasty. On instruction from the cardiologist coordinator, the trained staff administer drugs (aspirin, clopidogrel, perhaps thrombolysis, or abciximab, or bivalirudin) and consents the patient for coronary intervention. The patient does not go to the accident and emergency department of the nearest hospital.

Scenario 2—An ambulance arrives and the patient is taken to the nearest hospital, where an ECG establishes the diagnosis of an AMI. Intravenous streptokinase is given, but after 90 minutes chest pain continues and the ST segments have not shifted. A decision is made to transfer the patient to a percutaneous coronary intervention (PCI) centre. Because of a heavy workload the ambulance facility cannot give the patient urgent priority, but is eventually taken to another hospital where, nine hours after the onset of chest pain, an angioplasty is performed.

What would you want?

Many studies of animal and human AMI have shown that the duration of a coronary occlusion is the main determinant of final infarct size. The matters are biologically complex involving endothelial oedema, oxidative stress, reperfusion injury, embolisation of thromboresistant clots, and formation of leucocyte aggregates within the microcirculation. What is important is to establish normal flow in the infarct related vessel, as soon as is feasible.

The two strategies available to achieve this aim, thrombolysis and primary angioplasty, have been pitched against each other as to which is the best treatment. An evaluation of the 23 randomised trials comparing primary PCI with intravenous thrombolytic treatment have shown that the PCI strategy:

- saves lives in the short and long term
- leads to better TIMI III flow in the infarct related artery
- has less reinfarction
- leads to a shorter hospital stay
- has less hospital readmission
- causes less heart failure
- leads to less angina
- causes less strokes.

The overall combined end point (death, reinfarction, and stroke) was 8% for primary PCI versus 14% for thrombolysis ($p < 0.0001$).

WHAT HAPPENS TO PATIENTS OVER THE LONG TERM?

The pioneering Zwolle group whose randomised trial between PCI and thrombolysis in non-cardiogenic shock ST elevation myocardial infarction (STEMI) patients reported five year mortality rates of 13% versus 24%, respectively ($p < 0.01$), with a reduction in non-fatal MI (6% v 22%), recurrent ischaemia, and heart failure with no differences in overall costs. These Kaplan-Meier curves were divergent showing an increased benefit with time.

In this issue of Heart, Parodi and colleagues publish their five year outcome data for patients undergoing primary coronary intervention for acute ST elevation MI (all ages and including cardiogenic shock) in a single centre. The door to balloon time was a remarkable mean (SD) 22 (15) minutes, which shows what can be achieved with good organisation and communication. The five year mortality of 20% after five years is all the more extraordinary when a comparison is made with a thrombolysis driven service giving comparable mortality at 30 days.

Why has there been such reluctance to implement what seems to be a superior treatment for a common life threatening condition? A cursory glance of the medical literature will find many examples whereby evidence is ignored in favour of accepted, though flawed, dogma.

PCI VERSUS EARLY THROMBOLYSIS

Pre-hospital thrombolysis delays the time of onset of symptoms by 33° to 55° minutes, and the mortality benefits for patients treated within 60 minutes of onset of chest pain was significant.

The CAPTIM investigators compared a strategy of pre-hospital thrombolysis with transfer to an interventional facility for primary PCI. Patients randomised < 2 hours from symptom onset.

Abbreviations: AMI, acute myocardial infarction; PCI, percutaneous coronary intervention; STEMI, ST elevation myocardial infarction; TIMI, thrombolysis in myocardial infarction

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showed a trend toward lower 30 day mortality in the prehospital thrombolysis group (2.2% v 5.7%) (p = 0.05), and the mortality for patients treated > 2 hours was similar (5.9% v 3.7%) (p = 0.47). It is fair to conclude that PCI does not confer any mortality benefit over thrombolysis that is given within < 2 hours of chest pain onset.

WHAT ABOUT IN THE REAL WORLD?
This is fine for those places that have the facilities, but what about the real world where scarce resources make this option an impossibility?

The DANAMI-2 study examined whether transfer of patients to a PCI centre would still offer patient benefit—1572 patients were randomised to PCI (n = 790) or intravenous alteplase given on admission (n = 782). All transfers were made within three hours. The primary end point (mortality, reinfarction, disabling stroke at 30 days) was 14.2% in the thrombolysis group versus 8.5% in the primary PCI group (p = 0.002), the majority of the benefit being from prevention of reinfarction. The Czech Republic have harnessed their resources and now provide a 24 hour PCI service for AMI throughout the country, and yet they only have 22 catheter laboratories. Their management strategy is based on the PRAGUE 2 trial—a randomised controlled study comparing long distance travel for PCI versus thrombolysis in the nearest available hospital (n = 850). The study was stopped prematurely because of a 2.5-fold excess mortality in the thrombolysis group among patients treated > 3 hours after the onset of symptoms (15.3% thrombolysis v 6% PCI); however there was no significant difference between the strategies if the patient was treated < 3 hours from symptom onset.

WHY SHOULD THROMBOLYSIS BE MUTUALLY EXCLUSIVE OF A PCI SERVICE?
The PRAGUE study examined three strategies for PCI for patients presenting with STEMI to a district hospital without on-site catheterisation facilities—immediate thrombolysis, thrombolytic treatment during transfer for PCI, and transfer for primary angioplasty. The combined end point of death, reinfarction, or stroke was reached in 23%, 15%, and 8% of patients, respectively.

The timing and delivery of glycoprotein IIb/IIIa inhibitors before PCI for STEMI has improved results, but the delivery of these agents is not straightforward in the prehospital setting. Perhaps, this so-called facilitated angioplasty with thrombolysis ± glycoprotein IIb/IIIa inhibitors may have a role, particularly for patients who may have to travel long distances for their PCI. The outcome of trials such as FINESSE and ASSENT IV will provide the answer, but I would be surprised if these strategies will provide such striking differences that the investment of expensive drugs combined with intervention will become routine.

THROMBOLYSIS WITH BACKUP PCI
What about a strategy of thrombolysis with backup PCI for those patients in whom chest pain or ECG changes persist? This is the default position of many units, which dramatically reduces the number of patients requiring PCI in the short term. The randomised trials are not fully clear on the best course of action—there is a suggestion that late PCI may have some benefit, but only if the intervention is successful.

Most patients that survive a STEMI subsequently undergo angiography and revascularisation at later date; why not do it at the time the data strongly shows is the best time for PCI to occur—as soon after the acute occlusion as possible?

PRIMARY PCI: MOVING FORWARDS
Indeed for those involved in primary PCI, the debate has moved on. Previous studies have established the benefit of stenting over plain old balloon angioplasty, mainly for the benefit of recurrent angina. Parodi and his colleagues have analysed their data to discuss the role of glycoprotein IIb/IIIa inhibitors, appropriateness and timing of non-culprit coronary lesion treatment, and the provision of a risk stratification score based on simple clinical criteria. In most catheter laboratories patients are undergoing PCI for lesions causing stable angina—a procedure that confers little or no prognostic benefit; yet the patients with STEMI for whom this procedure has proven prognostic benefit are denied it. Our priorities need to refocus.

ARE THERE ENOUGH INTERVENTIONAL CARDIOLOGISTS?
To reproduce Parodi’s results, primary PCI should be performed at facilities that undertake a sufficient volume of work to develop and maintain skills. The American College of Cardiology/American Heart Association guidelines suggest that institutions performing PCI should be doing at least 400 cases a year with a 24 hour capability. To my mind, if a patient can reach such a PCI facility within two hours of arrival of the emergency services, reorganisation of regional strategies should take place to facilitate this. For those areas where this is currently an impossibility, regional discussions should take place to decide where such a centre could be developed.

Many centres are embracing this challenge, but there remain manpower issues which need to be grasped, understood, and dealt with if PCI is to become a routine procedure.

‘WE CAN’T DO AN ANGIOPLASTY—IT WILL AFFECT OUR DOOR TO NEEDLE TIMES’
Most units are highly efficient in the delivery of thrombolysis to patients coming through the hospital door, yet paradoxically, protocols such as this can become barriers to change. The measurement of door to needle times has some bearing on the patient’s outcome, but unless the crucial time measurement of onset of symptoms becomes the yardstick, the patient may well end up with the wrong treatment—all decisions should be based on the onset of the infarct, and not the time medical care starts.

WHAT IS THERE TO DO?
• The most significant delay contributing to delays in treatment for STEMI is the time it takes for the emergency services to be called by the patient and their family. The average time for pain to call time is one hour. Media campaigns have made little difference in reducing this, but the message that chest pains require immediate contact with the emergency services needs to be hammered home.
• Establish a local high volume PCI centre if one is not available.
• Once contact is made with the patient, the main purpose is to establish an ECG diagnosis and make an immediate strategy for treatment by contacting the local AMI coordinator.
• Data from the various travelling trials for STEMI PCI have shown that the visit to the local hospital engenders a 30–50 minute delay. Therefore, it is no longer acceptable for patients with STEMI to be transferred to the most local hospital, but to one that delivers the best treatment.

REFERENCES
An unusual cause of syncope

A 65 year old man was admitted to our hospital for recurrent syncope. Two years before presentation, he had placement of a Le Veen shunt for an intractable ascite complicating liver cirrhosis. On physical examination he appeared well. His blood pressure was 130/80 mm Hg and heart rate was 85 beats/min. No sign of congestion or cardiac murmur was noticed and an ECG showed a normal sinus rhythm without abnormalities. Transthoracic echocardiogram detected a 3 x 8 cm floating fluid filled mass in the right atrium that prolapsed into the right ventricle at each cardiac cycle. Transoesophageal echocardiography indicated that the mass was an intra-cardiac pseudocyst and was attached to the venous tip of the shunt (panel A). Pulmonary perfusion lung scan did not reveal signs of pulmonary embolism. The patient underwent open heart surgery and the cavitary ascitic fluid filled mass was removed completely (panel B). Histologic examination confirmed that the wall of the pseudocyst comprised fibrin, red blood cells, mononuclear cells, and capillaries.

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