Using simulation to estimate the cost effectiveness of improving ambulance and thrombolysis response times after myocardial infarction

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Methods
Coronary heart disease computer simulation model
This paper describes results derived from a discrete event simulation of the treatment of coronary heart disease (CHD), from the first onset of angina, or MI, to age 85 years or death. A further simulation to model interventions for preventing of CHD was developed in parallel. Both simulations were written using patient orientated simulation technique software (POST) with a Delphi interface. They were used to test the cost, effect, and overall cost effectiveness of treatments and interventions, based on the most recent published data available.

Two proved lifesaving treatments in the management of acute myocardial infarction (MI) are resuscitation from out of hospital cardiac arrest and restoration of blood flow to the blocked coronary artery (using thrombolysis and aspirin and/or primary angioplasty). In a cardiac arrest resulting from ventricular fibrillation or pulseless ventricular tachycardia, rapid defibrillation is required to return the heart to spontaneous output before brain and cardiac death occurs. Thrombolysis limits damage to the heart muscle and its consequences. Standards of care for these treatments in the National Service Framework (NSF) for Coronary Heart Disease for England in 2000 include an ambulance response within 8 minutes of the call to the national emergency number.2

The aim of this study is to quantify, using a computer simulation model, the health gains and the costs associated with moving from practice in the mid-1990s to meeting the NSF ambulance and thrombolysis targets.

In the treatment model, simulated patients transfer between stable and unstable angina, MI, and sudden cardiac death. There is an arbitrary cut-off point of 85 years because older patients have multiple diseases and it is difficult to isolate the data relating to CHD. In discrete event simulation, an event is a point in time at which a patient's state changes (for example, a patient joins a waiting list or has an MI). The transitions between events are based on risks such as age, severity of coronary artery disease, and previous history of CHD. Times between events are sampled from the relevant distributions for each patient, depending on the events and risk group. Parameter estimates are based on best evidence through searching the Cochrane database, MEDLINE and EMBASE, checking grey literature, and in discussion with experts. The simulation output has been validated against current routine sources of data. In this research, we are looking at implications of faster access to treatment for MI and so MI is in the centre of the simplified diagram of patient flows in the model shown as fig 1. For more details of the simulation model construction and assessment of model validity please see www.emjonline.com/supplemental/ and/or reference 5.

The simulation (see fig 1) describes the flow of patients from disease or treatment state to the next. Separate spreadsheet models described:

Abbreviations: MI, myocardial infarction; CHD, coronary heart disease; DTN, door-to-needle; NICE, National Institute for Clinical Excellence; NSF, National Service Framework; UKHAS, United Kingdom Heart Attack Study

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the effects of faster ambulance arrival times on death rates
both before and after admission to hospital with MI
the effect of varying the provision of thrombolysis and the
timing of thrombolysis on the relative risk of death for
patients after admission to hospital.

These relative risks were then incorporated into the simulation
model in order to assess the life years saved under
different policies.

Sources of data
To address the cost effectiveness of the NSF ambulance and
thrombolysis targets, all parameter estimates not directly
affected by changing ambulance and thrombolysis timings
were fixed. For more details, please see www.emjonline.com/
supplemental/. Ambulance and thrombolysis data were
extracted from three datasets: Heartstart Scotland (1991–
98), the United Kingdom Heart Attack Study
(UKHAS) (1994–95), and the West Midlands Thrombolysis

Ambulance response time and survival
In the UKHAS, 61% of patients had witnessed cardiac arrests
or MIs. The Heartstart study provided data by ambulance
response times on the survival rate to hospital admission for
patients where resuscitation was attempted. The 8 minute
ambulance response target was estimated by moving the 75%
quartile in Heartstart Scotland of 10.4 to 8.0 minutes. To
achieve this, all calls that occurred between 8.0 and
10.4 minutes were added to the baseline frequency of calls
in each time band (0–8 min) in proportion to the number
of calls in these bands. The 25% of baseline calls that occurred
after 10.4 minutes were each decreased by a simple down-
ward shift of 2.4 minutes (table 1). To obtain a new survival
time to hospital arrival for patients experiencing a witnessed
cardiac arrest or MI, the percentage frequency in each time
band was multiplied by the survival rate for that time band
and summed over all time bands.7 The overall estimated
change in survival rate for all cardiac arrests or MIs was
applied to the survival rate in each age/sex band.

UKHAS showed that cardiac arrest survivors were four
times more likely to die prior to hospital discharge than other
patients with an acute MI. We have assumed therefore that
the additional patients who survive as a result of faster
ambulance times have this poorer in-hospital survival rate,
thus reducing the overall hospital survival rate. We explored
the impact of changing this mortality rate in a sensitivity
analysis using a hypothetical assumption: that cardiac arrest
survivors have an in-hospital survival rate similar to those
who do not experience an out of hospital cardiac arrest.

Thrombolysis timing
In the UKHAS about 50% of acute MI patients at hospital
arrival were diagnosed as having experienced a definite MI

| Table 1 Ambulance response time distribution for cardiac arrest patients at baseline and for achieving the National Service Framework target |
|----------------------------------|-----------------|-----------------|-----------------|
| Response time interval (min)     | Percentage of patients discharged (survival rate to hospital discharge) | Baseline, percentage of patients responded to in time interval* | If 75th percentile response time is 8 minutes, percentage of patients |
| 0–4                              | 14.7            | 12.2            | 22.1            |
| 4–8                              | 9.2             | 42.9            | 52.9            |
| 8–12                             | 5.1             | 28.8            | 17.1            |
| >12                              | 4.3             | 16.1            | 7.9             |

*Data extracted from the Heartstart Scotland dataset.
and received thrombolysis. We assessed the effect of reducing the median time from hospital arrival to thrombolysis for such patients—that is, door-to-needle (DTN) time. Another 10% of patients had a definite MI and indications for thrombolysis but did not receive it. The effect of providing thrombolysis for such patients was also assessed.

Estimates of thrombolysis efficacy were taken from a meta-analysis. Data for efficacy were related to the time interval from onset of symptoms to administration of thrombolysis—that is, a relative mortality risk of 0.70 was associated with a time interval of up to 1 hour; 0.74 for 2–3 hours; 0.79 for 4–6 hours; 0.85 for 7–12 hours and 0.98 for more than 12 hours. The effect of using efficacy estimates from a more recent meta-analysis, which showed greater efficacy in the first few hours, was estimated in a sensitivity analysis. Long term follow up data show that following thrombolysis, there is little evidence that further lives are saved after 45 days. We have therefore assumed that thrombolysis only affects in-hospital mortality rates.

In the West Midlands Thrombolysis Project, which provided data on event times for patients with definite MIs diagnosed at hospital arrival, the 75th percentile DTN time was 80 minutes. To derive a frequency distribution in which the 75th percentile DTN time was 30 minutes, we multiplied each individual patient’s DTN time by three-eighths and calculated new symptom-to-needle times. It was thus possible to derive the change in efficacy arising from the faster DTN times (see table 2) from the symptom-to-needle times. A similar approach was used to derive the distribution for the 20 minute DTN target. The overall change in hospital survival, taking into account both those who require thrombolysis and those who do not, was applied to the survival for each age/sex group.

**Costs**

The Review of Ambulance Performance Standards estimated that the cost associated with attaining a 75% ambulance response time within 8 minutes would be £15 million for England (excluding London). This cost was scaled to the total population being modelled, up-rated to year 2000 costs and applied as a fixed cost in the scenarios which used a more recent meta-analysis, which showed greater efficacy in the first few hours, was estimated in a sensitivity analysis. Long term follow up data show that following thrombolysis, there is little evidence that further lives are saved after 45 days. We have therefore assumed that thrombolysis only affects in-hospital mortality rates.

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**Table 2** Distribution of time from hospital arrival to thrombolysis (door-to-needle time) for eligible patients (excluding the 10% of patients who fail to receive treatment), baseline and proposed for achieving the National Service Framework 30 minute target

<table>
<thead>
<tr>
<th>Door-to-needle time (min)</th>
<th>Baseline, percentage of patients*</th>
<th>75th percentile in 30 minutes, percentage of patients</th>
</tr>
</thead>
<tbody>
<tr>
<td>0–29</td>
<td>29.8</td>
<td>75.3</td>
</tr>
<tr>
<td>30–59</td>
<td>33.7</td>
<td>18.1</td>
</tr>
<tr>
<td>59–119</td>
<td>4.9</td>
<td>4.9</td>
</tr>
<tr>
<td>120–239</td>
<td>8.8</td>
<td>1.1</td>
</tr>
<tr>
<td>&gt;240</td>
<td>2.5</td>
<td>0.6</td>
</tr>
</tbody>
</table>

*Data extracted from the West Midlands Thrombolysis Project dataset.

83% of patients received streptokinase and 17% alteplase (as found in the UKHAS) with just one administration of the relevant drug. Retepase and tenecteplase were not widely available at the time and we did not include them in the model.

In attempting to reduce time to thrombolysis, a number of strategies have been tried such as improving communication between ambulances crew and receiving hospitals, and moving thrombolysis to the accident and emergency department. Some studies have shown that increasing deployment of nurses to coordinate assessment and initiation of thrombolysis can achieve the 30 minute DTN target. In each study, more than four experienced cardiac nurses were trained to undertake this new role. However, there was little information on the extent of additional staff requirements. We assumed that one to five additional F or G grade nurses would be needed, with an annual salary of between £22 500 and £25 000 (plus 40% overheads, at 2000 prices) for a population of about 250 000 (the average catchment of a hospital providing thrombolysis, based on information from the Myocardial Infarction National Audit Project (MINAP). These costs were scaled up to a population of one million, and applied in the scenarios about faster access to thrombolysis.

**Derivation of cost per life year saved**

The estimated life years saved each year was the difference between the total number of patient life years in the simulated scenario, with the changed relative risks, compared to the base run. The costs in the model, which were the estimated NHS costs, for the treatment of coronary heart disease were compared between the scenarios. The annual incremental cost per life years saved of each scenario compared to baseline were accumulated over 20 years; costs were discounted at 3.5% and benefits were discounted at 3.5%, as recommended by the National Institute for Clinical Excellence (NICE). The discounted costs per life year saved were averaged over the 20 years.

**Simulations and sensitivity analysis**

A simulation was run for all CHD patients arising from a population of 0.5 million people, for 20 years with 100 replications. The results were scaled to a population of one million. The main scenarios were as follows:

1. 75% of “life-threatening” calls to receive an ambulance response within 8 minutes.
2. 75th percentile of distribution of DTN time for thrombolysis is 30 minutes (with the existing proportion of patients receiving thrombolysis).
3. 75th percentile of distribution of DTN time for thrombolysis is 20 minutes (with the existing proportion of patients receiving thrombolysis).
4. 75th percentile of distribution of DTN time for thrombolysis is 30 minutes (including the additional eligible patients (10% of the total) who currently fail to receive thrombolysis).
5. Combination of scenarios 1 and 2.

The results from each scenario were compared with the baseline values, shown in the relevant tables, which relate to the position in England in the mid-1990s. We explored the sensitivity of the results to the following scenarios:

6. The mortality rate for cardiac arrest survivors. For the ambulance response scenario 1, those cardiac arrest survivors were assumed to have a similar in-hospital survival rate to those who did not experience an out of hospital cardiac arrest.
the 30 minute DTN scenario.

Boersma et al's data improved the life years saved to 24 for thrombolysis scenario 2, and we used Boersma et al's relative mortality risk estimates.

RESULTS

The estimated number of deaths prevented per million population per year ranged from two to seven across the five main scenarios, and life years saved from 15 to 70 (table 3). The standard deviations of these results show that in different populations the outcome may vary considerably.

Based on the average values, the ambulance target generated 42 more life years saved than were saved by achieving the 30 minute DTN target. Little difference was found between the 30 minute and 20 minute DTN targets. Increasing the use of thrombolysis to a further 10% of patients had similar benefits to improving DTN times to a 75th percentile of 30 minutes. Figure 2 shows that the difference in the cumulative number of lives saved between these scenarios increases over the 20 year simulation.

Sensitivity analysis

The results for the sensitivity analyses are shown in table 4. Assuming that cardiac arrest survivors had the same inhospital mortality rate as those who did not have an out of hospital cardiac arrest (scenario 6) increased the estimated life years saved per million population by 48. The results are shown to be sensitive to the ambulance response profile and the worse and best scenarios ranged between 35 and 81 (scenario 8 and 7). Increasing the ambulance response to 90% within 8 minutes (scenario 9) would result in a further 28 life year saved over scenario 1. The benefits of thrombolysis were sensitive to the efficacy estimates; the use of Boersma et al’s data improved the life years saved to 24 for the 30 minute DTN scenario.

Ambulance response time cost effectiveness

The marginal annual cost of achieving the 8 minute target was estimated at £448 100 for a population of one million. The estimated discounted incremental cost per life year saved, to age 85, accumulated over 20 years and divided by 20 to get the average per year, was £8540. The cost effectiveness ratios are much less favourable in the short term with an accumulated value of £19 110 per year at 5 years and £12 400 per year at 10 years.

Thrombolysis time cost effectiveness

The annual cost of nurse initiated thrombolysis to meet the 30 minutes DTN time for thrombolysis was assumed to be between £135 000 and £752 500 per million population. The discounted incremental cost per life year saved, to age 85, accumulated over 20 years and divided by 20 to get the average per year, was estimated to be between £10 150 and £54 230. Again, the cost effectiveness ratios are less favourable in the short term with the cumulative values for the lower cost at £20 560 per year in 5 years and at £13 800 per year in 10 years.

DISCUSSION

This simulation model has indicated that it may be cost effective to improve both ambulance response and thrombolysis DTN times with results from a 20 year period or more. The costs per live years saved were well below the putative threshold set by NICE.25 Much greater benefits are achieved by reaching the ambulance response target; particularly with the higher 90% response target. However, the benefit of ambulance timing was sensitive to the assumption about the poorer inhospital survival rate for cardiac arrest survivors; though the weight of evidence suggests a poorer prognosis for these patients than MI patients who have not experienced a cardiac arrest.9 Further research is needed on the impact of faster ambulance times on inhospital mortality rates. Very
little additional gain is achieved by moving from the 30 minute to the 20 minute NSF thrombolysis target, whereas ensuring that all eligible patients receive thrombolysis would be beneficial.

Although our model is based on UK data, our results would be applicable to other developed countries with similar emergency services. Since the 1990s, changes in practice have led to ambulance response times and speed of delivery of thrombolysis (sometimes involving use of newer agents) that are now more in line with the NSF targets.

The cost effectiveness of achieving the ambulance target might be greater if non-CHD lives saved were included, but there were no data to estimate this. Furthermore, there are likely to be local variations in the feasibility and cost of achieving the target, particularly between urban and rural populations. The Review of Ambulance Performance Standards estimated that moving to the 75% target would gain 530 life years per million population per year at a cost per life year saved of £1100 which is more favourable than our estimates. This is likely to be optimistic because long-term mortality risk associated with MI was not taken into account. Pell et al showed that shorter ambulance response times increased the proportion of cardiac arrest victims receiving defibrillation and increased survival to hospital discharge; they estimated that a 90% 8 minute response time would save about 5–6 deaths per million population per year. These are close to our estimates.

Our thrombolysis results may have underestimated the absolute benefit by taking an optimistic baseline time distribution from the West Midlands Thrombolysis Project in which hospitals volunteered for the audit to promote improvement. There is wide variation in the cost effectiveness estimates for the 30 minute DTN target depending on the numbers of new nursing staff needed. There are other costs, which we have not taken into account, such as a possible increased need for coronary care unit (CCU) staff which would reduce the cost effectiveness. In common with other studies, we have not taken account of the cost of treating patients whose lives are saved but who acquire other diseases. These are likely to be considerable in the older population.

It is important to consider other approaches, to reducing time from symptom onset to starting thrombolysis. Public education on identifying heart attack symptoms might be expected to further increase lives saved. The SHARP study found that delays to hospital arrival were greater in those who contacted their general practitioners directly. However, studies evaluating educational campaigns designed to reduce delay between symptom onset and treatment of suspected MI have generated inconclusive results. Campaigns aimed at ensuring calls are made directly to the ambulance service may save some lives but may increase inappropriate calls and costs. Prehospital thrombolysis is increasingly available in the UK and could further increase lives saved especially in remote areas.

In conclusion, improving ambulance response and inhospital time to thrombolysis appears to be cost effective. The absolute gain with improving ambulance response is much greater than that associated with further reducing DTN time for thrombolysis.

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