Myocardial perfusion scintigraphy and cost effectiveness of diagnosis and management of coronary heart disease

S R Underwood, L J Shaw

Myocardial perfusion scintigraphy (MPS) is a well validated, non-invasive imaging technique that has a valuable role in the diagnosis, management, and assessment of prognosis of coronary heart disease (CHD). The diagnostic accuracy of MPS allows reliable risk stratification and guides the selection of patients for further interventions, such as revascularisation. MPS also has particular advantages over alternative techniques in the management of a number of patient subgroups, including women, the elderly, and those with diabetes. Increased use of MPS will improve patient outcomes and be associated with greater cost effectiveness of treatment, in terms of life-years saved, particularly in these special patient groups. Currently, however, access to MPS in the UK is limited, with inappropriately long waiting times, and MPS activity levels fall short of estimated need.

COST EFFECTIVENESS OF MYOCARDIAL PERFUSION SCINTIGRAPHY

Principles of cost effectiveness

Several principles underlie why a more accurate diagnostic test with additional prognostic information, such as MPS, can be more cost effective even if it is more expensive than an alternative test such as the exercise ECG (table 1).

For example, fig 1 shows that when a patient with presenting likelihood of CHD of 50% has an abnormal exercise ECG the post-test likelihood of disease is 73%, which is not sufficiently high to be confident of the diagnosis. A subsequent abnormal MPS gives a likelihood of 96%, but if the same patient had gone directly to MPS the post-test likelihood would have been 90%, which should be sufficiently high to diagnose the presence of CHD depending upon the clinical circumstances.

Costs

True costs, reflecting consumption of resources, are difficult to estimate. Some analyses of cost effectiveness have used prices as a surrogate for cost, which may be valid from the perspective of the referring clinician, but are less helpful from the perspective of the National Health Service. Two studies have estimated cost, one using UK figures, but the findings are similar at £220 in the UK and £179 (range £159–348) in the USA (table 2). These costings are sensitive to throughput. A reasonable throughput per camera is 2000 patients per year, but this can be increased to as much as 4000 in high volume centres by running the camera 12 hours per day and six days per week.

Cost effectiveness of MPS

Although randomised controlled and blinded trials are often used to evaluate the cost effectiveness of treatment, this study design is difficult or impossible in the case of diagnostic testing. However, some data have been derived from decision analytical models and these have demonstrated the cost effectiveness of MPS both in patients presenting with stable chest pain syndromes and in pre-operative risk assessment. Garber and colleagues showed that in 55 year old men presenting with chest pain, a strategy of MPS proceeding to angiography compared with exercise ECG proceeding to angiography cost £25 000 per quality adjusted life year (QALY), a figure that is generally regarded as acceptable. The benefit of MPS would have been even greater if a more realistic model had been used, since it was assumed that all patients with disease proceed to angiography, whereas, as demonstrated above, it is possible to treat low risk patients with CHD without the need for angiography. Patterson and colleagues also found initial testing with MPS to be more cost effective than angiography cost £25 000 per QALY. However, despite the attractive cost effectiveness of MPS, access to MPS in the UK is limited, with inappropriately long waiting times.
effective than the exercise ECG, although again the model assumed that all patients with disease undergo angiography. Kuntz and colleagues found a higher incremental cost effectiveness of initial MPS over the exercise ECG of £34 250 in a 55 year old man with atypical chest pain, but they used an unrealistically low specificity of 64% for MPS and again assumed that all patients with disease undergo angiography. It would be valuable to repeat these models assuming that patients with abnormal but low risk non-invasive tests are treated medically in the first instance, since the prognostic power of MPS is likely to lead to even greater cost effectiveness. Interpretation of these numbers should be placed in the context of other generally accepted medical interventions. For example, coronary artery bypass grafting for patients with triple vessel coronary artery disease and severely impaired left ventricular function costs £25 500 per QALY gained, and cholesterol lowering therapy in a 60 year old man with cholesterol of 7.5 mmol/l costs approximately £30 000 per QALY. Thus, MPS appears to be a cost effective use of resources compared with other generally accepted medical procedures.

Mathematical simulations have the advantage of using data from diverse sources or from expert opinion, but they are limited by a lack of real world effectiveness data. However, two controlled clinical studies have demonstrated savings for similar outcomes using MPS in patients presenting with chest pain. The economics of myocardial perfusion imaging in Europe (EMPIRE) study compared patients presenting with stable chest pain syndromes to centres that routinely use MPS and those that do not, in each of four European countries. Diagnostic strategies using MPS were cheaper and equally effective than those that did not, both for the costs of diagnosis and for overall two year management costs, but patient outcome was the same (fig 2).

The economics of non-invasive diagnosis (END) study was a larger (11 372 patients) registry study with very similar findings of 30–40% savings in costs over 2.5 years in patients with stable chest pain syndromes undergoing initial MPS and selective angiography (fig 3). An additional series of 9521 patients showed that MPS was more cost effective than stress echocardiography in patients with known CHD and stable chest pain at $39 347 per life-year saved. Unfortunately, modelling studies have provided conflicting results on the comparative cost effectiveness of strategies using MPS and stress echocardiography and there are insufficient clinical data to make a firm statement on this.

Other clinical studies indirectly support the cost effectiveness of MPS. In a US practice making routine use of MPS the number of patients with normal MPS proceeding to angiography is 1%, whereas in a UK rapid access chest pain clinic that makes little use of MPS, coronary angiography rates are high and 56% of coronary angiograms in women are normal. It has been estimated that routine use

<table>
<thead>
<tr>
<th>Table 2 Cost of common diagnostic tests calculated using principles that estimate the true consumption of resources.</th>
<th>EMPIRE4</th>
<th>USA average3</th>
<th>USA range3</th>
</tr>
</thead>
<tbody>
<tr>
<td>Rest ECG</td>
<td>£20</td>
<td>£55</td>
<td>£39–£207</td>
</tr>
<tr>
<td>Exercise ECG</td>
<td>£70</td>
<td></td>
<td></td>
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<tr>
<td>Rest echocardiography</td>
<td>£100</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Stress echocardiography</td>
<td>£1644</td>
<td>£114–£2134</td>
<td></td>
</tr>
<tr>
<td>Computed x ray tomography</td>
<td>£172</td>
<td>£55–£288</td>
<td></td>
</tr>
<tr>
<td>MPS</td>
<td>£220</td>
<td>£179</td>
<td>£139–£348</td>
</tr>
<tr>
<td>MRI</td>
<td>£529</td>
<td>£318–£739</td>
<td></td>
</tr>
<tr>
<td>PET</td>
<td>£771</td>
<td>£582–£891</td>
<td></td>
</tr>
<tr>
<td>Coronary angiography</td>
<td>£1100</td>
<td>£1097</td>
<td>£516–£2873</td>
</tr>
</tbody>
</table>

EMPIRE, economics of myocardial perfusion imaging in Europe; MRI, magnetic resonance imaging; MPS, myocardial perfusion scintigraphy; PET, positron emission tomography.
of MPS in this setting would reduce cost by between £30 000 and £65 000 per year, mainly by reducing the normal coronary angiography rate. Similarly in the USA it has been estimated that about one third of referrals to coronary angiography are inappropriate.

Clinical outcomes
Modelling studies indicate that clinical outcome is improved by MPS at a cost per QALY that is acceptable and that life expectancy is increased by between seven days and two years. Because of the scale and nature of the studies that would be required, it has been more difficult to confirm this finding in clinical studies. However, in a reanalysis of the END data, Shaw and colleagues have shown that testing with initial MPS adds between one and two years of life, and in a separate study they showed that MPS added 0.5 years compared with stress echocardiography.

CONCLUSION
MPS is a well established, non-invasive imaging technique with a large body of evidence to support its effectiveness in the diagnosis and management of angina and myocardial infarction. It is more accurate than the exercise ECG in detecting myocardial ischaemia and it is the single most powerful technique for predicting future coronary events. The high diagnostic accuracy of MPS allows reliable risk stratification and guides the selection of patients for further interventions, such as revascularisation. This in turn allows more appropriate utilisation of resources, with the potential for both improved clinical outcomes and greater cost effectiveness. Evidence from modelling and observational studies supports the enhanced cost effectiveness associated with MPS use. In patients presenting with stable or acute chest pain, strategies of investigation involving MPS are more cost effective than those not using the technique. MPS also has particular advantages over alternative techniques in the management of a number of patient subgroups, including women, the elderly, and those with diabetes, and its use will have a favourable impact on cost effectiveness in these groups.

ACKNOWLEDGEMENTS
This submission is based upon the deliberations of a consensus conference on the evidence for the clinical and cost effectiveness of MPS, organised by the British Nuclear Cardiology Society in January 2003. Members of the writing group were C Anagnostopoulos (co-chair), M Cerqueira, PJ Ell, J Flint, M Harbinson, A Kelion, A Al Mohammad, EM Pturolich, LJ Shaw, AC Tweeddle, SR Underwood (co-chair) and L J Shaw, Atlanta Cardiovascular Research Institute, Atlanta, Georgia, USA

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REFERENCES