Multicentre evaluation of the diagnostic value of cardiac troponin T, CK-MB mass, and myoglobin for assessing patients with suspected acute coronary syndromes in routine clinical practice

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Objective: To assess the diagnostic efficiency of the third generation cardiac troponin T assay in routine clinical practice.

Design: Prospective observational study of unselected consecutive admissions.

Setting: Multicentre study in 43 teaching and non-teaching hospitals in 13 countries.

Subjects: 1105 hospital admissions, median age 67 years (range 15–96 years, 63.7% male) with suspected acute coronary syndromes (72.3% of cases) or other non-specific symptoms where cardiac disease required exclusion (27.7%).

Interventions: Over the study period, myoglobin, creatine kinase MB isoenzyme (CK-MB), and cardiac troponin T where measured in parallel with conventional diagnostic tests. Final diagnostic classification involved standard ECG changes and CK-MB mass exceeding 5.0 µg/l.

Main outcome measures: Diagnostic efficiency was assessed by receiver operator characteristic curve analysis including and excluding patients with unstable angina.

Results: Measurement of cardiac troponin T was diagnostically equivalent to CK-MB and both were better than myoglobin, with areas under the curve at 12 hours of 0.94, 0.99, and 0.80, respectively. Diagnostic criteria using CK-MB were inadequate and showed bias when patients with unstable angina were included. Elevations of cardiac troponin T did not occur when cardiac disease could be categorically excluded but were found in clinical conditions other than suspected acute coronary syndromes.

Conclusions: CK-MB is unsuitable as a diagnostic gold standard even at the proposed lower threshold. A lower cut off for cardiac troponin T of 0.05 µg/l should be used for diagnosis of acute myocardial infarction. Diagnosis of acute myocardial infarction cannot be made solely on the basis of a cardiac troponin T result.
cardiospecificity of cTnT measurement might be expected to confer an advantage (myositis, rhabdomyolysis, multiple trauma).

Data were recorded prospectively on all patients, using a structured proforma including relevant clinical details, ECG data, routine biochemical results with reference intervals, and final diagnosis. Care was taken to record the duration of chest pain before admission where possible. As timing of chest pain can be unreliable, time from admission to venepuncture was recorded in all cases and was used as the index time for assessment.

Data pro formas were then independently reviewed centrally in a blinded fashion by two of the investigators (POC and PJS) to check whether the data were internally consistent and compatible with the diagnosis supplied. When there was a discrepancy, further information was obtained from the participating centre. Where a final diagnosis could not be agreed, third party adjudication was used to assign the case to the appropriate patient group.

In all, 1105 patients admitted to 44 hospitals in 13 countries with suspected acute coronary syndrome, or in whom acute coronary syndrome was considered as a secondary diagnosis, were enrolled in the study. Of these, 799 (72.3%) had a primary diagnosis of suspected acute coronary syndrome, and 306 (27.7%) had investigations for other reasons. The median age of the subjects was 67 years (range 15–96 years; lower quartile 57, upper quartile 76); 63.7% of the patients were male, median age 66 years (range 15–96 years; lower quartile 53.5, upper quartile 74), and 36.3% were female, median age 72 years (range 27–96 years, lower quartile 63, upper quartile 80). The median duration of chest pain was 3.9 hours (range very short to 12.5 hours).

**Diagnostic categorisation**

In order to allow comparison with previous studies, final diagnostic categorisation of patients on hospital discharge was made using WHO criteria for ischaemic heart disease but with CK-MB mass as the biochemical gold standard.

Acute myocardial infarction was diagnosed when there were characteristic ECG changes—either ST segment elevation or the development of new Q waves—confirmed by significant changes in serial cardiac enzymes. ST segment elevation was judged to be present if it was ≥ 0.1 mV at the J point in one or more extremity leads or ≥ 0.2 mV in one or more precordial leads. Significant changes in cardiac enzymes were considered to occur when creatine kinase mass exceeded twice the upper reference limit of the local laboratory, or if CK-MB mass exceeded 5 µg/l.

Unstable angina pectoris was diagnosed in patients with an episode of chest pain within the previous 48 hours and without characteristic ECG changes (ST segment elevation as defined above or development of new Q waves), who had non-specific ECG changes (ST depression or T wave inversion) or continuing chest pain with or without non-specific ECG changes and without significant changes in cardiac enzymes. Stable angina pectoris was diagnosed in patients with known ischaemic heart disease but without acute chest pain and without significant changes in cardiac enzymes.

Non-ischaemic chest pain was diagnosed when follow up investigations allowed coronary artery disease to be excluded or when a definite source of non-cardiac chest pain could be identified.

Other diagnostic groups were taken from the final discharge diagnosis where a positive diagnosis could be made and ischaemic heart disease excluded.

**Analytical methods**

Blood samples for measurement of cTnT, CK-MB mass, and myoglobin were drawn along with blood for routine investigations according to the local investigation protocol. The choice of the use of serum or plasma was left to the local participating centre. For serum, blood samples were taken, allowed to clot fully, and then centrifuged. For plasma, blood samples were taken, mixed by inversion, and then centrifuged. Plasma or serum samples (as appropriate) were then separated for analysis by the laboratory for cTnT, CK-MB mass, and myoglobin determinations. Measurement of other cardiac enzymes in the local laboratories were done by automated analysis according to the manufacturers’ recommended methods.

Measurements of cTnT, myoglobin, and CK-MB mass were done using an Elecsys analyser by the troponin T STAT third generation, myoglobin STAT, and CK-MB STAT methods (Roche Diagnostics, Furtwangen, Germany). These assays are based on electrochemiluminescence immunoassay technology (ECLIA) using two mouse monoclonal antibodies in a sandwich format, two step assay. They were done on Elecsys 1010 and 2010 immunoassay analysers according to the manufacturer’s instructions. For cTnT, the detection limit was 0.01 µg/l with an upper limit of 25 µg/l; for CK-MB the detection limit was 0.1 µg/l with an upper limit of 500 µg/l; and for myoglobin, the detection limit was 15 µg/l with an upper limit of 3000 µg/l.

The analytical performance of the assays was good, with total imprecision less than 10% in, respectively, 97% (cTnT, range 0.15–5.77 µg/l), 94% (CK-MB, range 5.95–53 µg/l), and 97% (myoglobin, 90.4–1133 µg/l) of participating centres.

**Data handling and statistical methods**

Parametric statistics were used for the derivation of all estimates of imprecision. Baseline demographics are expressed as medians and range, with upper and lower quartiles where appropriate. Diagnostic performance was assessed by construction of receiver operating characteristic (ROC) curves with areas compared for significance at the p = 0.05 level by the normal approximation method (as the numbers were large). All statistical analysis was done using the SAS statistical package (SAS Institute Inc, Cary, North Carolina, USA). It has been documented that when diagnosis is based on creatine kinase and CK-MB, an elevation of cTnT is seen in approximately 33% of patients with unstable angina; thus the effect of variation in the diagnostic categorisation on the underlying population was studied by doing an ROC analysis including and excluding patients with a final diagnosis of unstable angina.

**RESULTS**

**Patients with a primary diagnosis of suspected acute coronary syndrome**

In all, 799 patients were studied: 467 with acute myocardial infarction, 164 with unstable angina, 100 with stable angina, and 68 with non-ischaemic chest pain. The data from the ROC analysis are summarised in table 1.

When patients with unstable angina were excluded, measurement of cTnT in samples taken less than two hours from admission was better than either myoglobin or CK-MB mass. Measurement of cTnT was significantly better than myoglobin (p < 0.05) at all other time points, most notably at 12–24 hours from admission. CK-MB and cTnT showed equivalent diagnostic efficiency at between 2–6 hours and at all time points thereafter. Inclusion of patients with unstable angina resulted in an apparent fall in the diagnostic efficiency of both myoglobin and cTnT. At less than 2 hours, 2–6 hours, and 6–12 hours, CK-MB was better than cTnT, but diagnostic efficiencies converged at 12–24 hours from admission. CK-MB was superior to myoglobin at all time points. Myoglobin and cTnT showed comparable diagnostic efficiencies until 6–12 hours from admission, when there was a progressive decrease in the efficiency of myoglobin but an increase in the value for cTnT. The ROC curves are shown in figs 1 and 2.

The third generation assay has both an improved imprecision profile and a lower detection limit. A detailed evaluation of the ROC curve was therefore performed. The clinical
sensitivity for cTnT was examined using cut off values for cTnT from 0.01–0.1 µg/l (table 2). Reduction in the cut off value resulted in improvement in sensitivity but loss of specificity, with optimal trade off in the range of 0.04–0.06 µg/l.

Patients without a primary diagnosis of acute coronary syndrome

Patients with ischaemic heart disease

There were 122 patients with cardiac failure: 56 with congestive heart failure, 49 with arrhythmias producing heart failure, and 17 with underlying ischaemic heart disease. cTnT exceeded 0.05 µg/l in 52 (45.6%) of these, and was raised to 0.1 µg/l or above in 28. These data are summarised in fig 3.

Patients with chest pain without suspected acute coronary syndrome.

In all, 124 patients were admitted with a final diagnosis that excluded suspected acute coronary syndrome as a primary diagnosis. In one there was inadequate documentation so that patient was excluded from further analysis. In 75, cardiac disease could be categorically excluded (39 with rheumatological disorders, five with carcinoma, 13 with gastrointestinal tract disorders, four with viraemia, and 14 with other miscellaneous diagnoses); cTnT was less than 0.05 µg/l in all these cases. There were 37 patients with pulmonary disease (10 with pulmonary emboli, 27 with pulmonary interstitial disease). Four of the 10 patients with a pulmonary

<table>
<thead>
<tr>
<th>Population</th>
<th>Time (hours)</th>
<th>Troponin T</th>
<th>Myoglobin</th>
<th>CK-MB mass</th>
</tr>
</thead>
<tbody>
<tr>
<td>MI/NICP + SAP</td>
<td>&lt;2</td>
<td>461</td>
<td>364</td>
<td>245</td>
</tr>
<tr>
<td>MI/NICP + SAP</td>
<td>2 to &lt;6</td>
<td>296</td>
<td>254</td>
<td>203</td>
</tr>
<tr>
<td>MI/NICP + SAP</td>
<td>6 to &lt;12</td>
<td>349</td>
<td>293</td>
<td>242</td>
</tr>
<tr>
<td>MI/NICP + SAP</td>
<td>12 to &lt;24</td>
<td>332</td>
<td>251</td>
<td>213</td>
</tr>
<tr>
<td>MI/NICP + SAP + UAP</td>
<td>&lt;2</td>
<td>746</td>
<td>624</td>
<td>295</td>
</tr>
<tr>
<td>MI/NICP + SAP + UAP</td>
<td>2 to &lt;6</td>
<td>681</td>
<td>514</td>
<td>253</td>
</tr>
<tr>
<td>MI/NICP + SAP + UAP</td>
<td>6 to &lt;12</td>
<td>734</td>
<td>553</td>
<td>292</td>
</tr>
<tr>
<td>MI/NICP + SAP + UAP</td>
<td>12 to &lt;24</td>
<td>717</td>
<td>511</td>
<td>263</td>
</tr>
</tbody>
</table>

CK-MB, MB isoenzyme of creatine kinase, MI, acute myocardial infarction; NICP, non-ischaemic chest pain; SAP, stable angina pectoris; UAP, unstable angina pectoris.
embolus had a cTnT concentration of $\geq 0.05 \mu g/l$; in one of these the cTnT value was above 0.1 $\mu g/l$. Ten of the 27 patients with pulmonary interstitial disease had a cTnT concentration of $\geq 0.05 \mu g/l$ and in six of these the value exceeded 0.1 $\mu g/l$. There were two unusual cases—one with thyroiditis and one with systemic lupus erythematosus. In neither of these could myocardial involvement in the primary pathology be excluded. These data are summarised in table 3.

**Rhabdomyolysis and multiple trauma**

There were 30 patients with rhabdomyolysis. Their creatine kinase values were in the range of 1655–10600 U/l and the cTnT concentration exceeded 0.05 $\mu g/l$ in two; however, all the cTnT values in these patients were less than 0.1 $\mu g/l$.

Thirty patients were admitted with multiple trauma: 25 of these had cTnT concentrations less than 0.05 $\mu g/l$ and two had values above 0.1 $\mu g/l$ (0.108 and 0.129 $\mu g/l$). Data comparing the markers in patients with multiple trauma are shown fig 4.

**Figure 2** Receiver operating characteristic (ROC) plots of acute myocardial infarction versus stable angina pectoris plus unstable angina pectoris and non-ischaemic chest pain according to sample time from admission.

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**Table 2** Effect on sensitivity and specificity of reducing the cut off value for troponin T

<table>
<thead>
<tr>
<th>Time window (hours)</th>
<th>Cut off point</th>
<th>0.01 $\mu g/l$</th>
<th>0.03 $\mu g/l$</th>
<th>0.04 $\mu g/l$</th>
<th>0.05 $\mu g/l$</th>
<th>0.06 $\mu g/l$</th>
<th>0.1 $\mu g/l$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sensitivity [CI]</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>0</td>
<td>82.4 (74.8 to 88.6)</td>
<td>69.5 (60.7 to 77.2)</td>
<td>67.9 (59.4 to 76.0)</td>
<td>63.4 (54.5 to 71.5)</td>
<td>60.3 (51.3 to 68.9)</td>
<td>53.4 (44.4 to 62.3)</td>
<td></td>
</tr>
<tr>
<td>&gt;0 to &lt;2</td>
<td>99.0 (96.3 to 99.9)</td>
<td>77.5 (71.0 to 83.1)</td>
<td>73.8 (67.0 to 79.9)</td>
<td>69.6 (62.5 to 76.0)</td>
<td>67.5 (60.3 to 74.2)</td>
<td>60.7 (53.6 to 67.5)</td>
<td></td>
</tr>
<tr>
<td>2 to &lt;4</td>
<td>93.7 (85.8 to 97.9)</td>
<td>83.5 (73.6 to 90.9)</td>
<td>83.5 (73.6 to 90.9)</td>
<td>82.3 (72.2 to 89.9)</td>
<td>79.7 (69.1 to 88.0)</td>
<td>73.4 (62.3 to 82.7)</td>
<td></td>
</tr>
<tr>
<td>4 to &lt;6</td>
<td>94.9 (87.4 to 98.6)</td>
<td>92.3 (84.0 to 97.1)</td>
<td>92.3 (84.0 to 97.1)</td>
<td>91.0 (82.4 to 96.3)</td>
<td>88.5 (79.2 to 94.6)</td>
<td>85.9 (76.2 to 92.7)</td>
<td></td>
</tr>
<tr>
<td>6 to &lt;12</td>
<td>97.6 (94.5 to 99.2)</td>
<td>95.7 (92.0 to 98.0)</td>
<td>93.8 (89.7 to 96.7)</td>
<td>93.8 (89.7 to 96.7)</td>
<td>92.9 (88.5 to 95.9)</td>
<td>92.4 (87.9 to 95.6)</td>
<td></td>
</tr>
<tr>
<td>&gt;12</td>
<td>99.0 (98.0 to 99.6)</td>
<td>96.0 (94.3 to 97.3)</td>
<td>95.0 (93.1 to 96.5)</td>
<td>94.9 (93.0 to 96.4)</td>
<td>94.2 (92.2 to 95.8)</td>
<td>90.7 (88.4 to 92.7)</td>
<td></td>
</tr>
</tbody>
</table>

Specificity [CI]

<table>
<thead>
<tr>
<th>Time window (hours)</th>
<th>Cut off point</th>
<th>0.01 $\mu g/l$</th>
<th>0.03 $\mu g/l$</th>
<th>0.04 $\mu g/l$</th>
<th>0.05 $\mu g/l$</th>
<th>0.06 $\mu g/l$</th>
<th>0.1 $\mu g/l$</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>60.0 (32.3 to 83.7)</td>
<td>73.3 (44.9 to 92.2)</td>
<td>73.3 (44.9 to 92.2)</td>
<td>73.3 (44.9 to 92.2)</td>
<td>73.3 (44.9 to 92.2)</td>
<td>80.0 (51.9 to 95.7)</td>
<td></td>
</tr>
<tr>
<td>&gt;0 to &lt;2</td>
<td>51.6 (43.5 to 59.6)</td>
<td>77.4 (70.0 to 83.7)</td>
<td>88.0 (73.5 to 86.5)</td>
<td>82.6 (75.8 to 88.4)</td>
<td>87.1 (80.7 to 92.0)</td>
<td>75.8 (75.8 to 97.1)</td>
<td></td>
</tr>
<tr>
<td>2 to &lt;4</td>
<td>35.9 (21.1 to 52.8)</td>
<td>71.8 (53.4 to 85.0)</td>
<td>74.4 (58.0 to 87.0)</td>
<td>76.9 (60.7 to 88.9)</td>
<td>76.9 (60.7 to 88.9)</td>
<td>89.7 (75.8 to 97.1)</td>
<td></td>
</tr>
<tr>
<td>4 to &lt;6</td>
<td>43.8 (19.8 to 70.1)</td>
<td>68.8 (41.3 to 89.0)</td>
<td>81.3 (54.4 to 96.0)</td>
<td>81.3 (54.4 to 96.0)</td>
<td>81.3 (54.4 to 96.0)</td>
<td>81.3 (54.4 to 96.0)</td>
<td></td>
</tr>
<tr>
<td>6 to &lt;12</td>
<td>31.0 (19.3 to 44.5)</td>
<td>56.9 (41.3 to 70.8)</td>
<td>62.1 (48.3 to 74.6)</td>
<td>63.8 (49.9 to 75.9)</td>
<td>67.2 (53.7 to 79.0)</td>
<td>74.1 (61.1 to 84.7)</td>
<td></td>
</tr>
<tr>
<td>&gt;12</td>
<td>38.6 (32.4 to 45.1)</td>
<td>56.8 (50.1 to 63.2)</td>
<td>58.9 (52.5 to 65.1)</td>
<td>60.6 (54.2 to 66.7)</td>
<td>63.1 (56.6 to 69.2)</td>
<td>71.4 (65.2 to 77.0)</td>
<td></td>
</tr>
</tbody>
</table>

CI, confidence interval.
DISCUSSION

The results of this study show that a lower diagnostic cut off for cTnT using the third generation assay is appropriate. It also illustrates the flaw in using CK-MB as the gold standard. There is an apparent fall in diagnostic efficiency when a less well defined diagnostic group, unstable angina, is included in the population studied. Test evaluations are typically performed on selected patient populations or on samples collected as part of a clinical trial. The patients involved have a high probability of ischaemic heart disease (typically > 80%) and the majority have a clear diagnosis of acute myocardial infarction. Test evaluations are typically performed on selected patient populations or on samples collected as part of a clinical trial. The patients involved have a high probability of ischaemic heart disease (typically > 80%) and the majority have a clear diagnosis of acute myocardial infarction. In routine clinical practice we found the prevalence of ischaemic heart disease to be 50%, with acute myocardial infarction at presentation in only 10%. The majority of patients required cardiac enzyme measurement for diagnosis. The situation is worse in the emergency department population, where there is a significant incidence of missed acute myocardial infarction.24

Realistic evaluation requires usual measurement, an unselected population, cases that are diagnostically challenging, and technology available for routine clinical use.

The diagnostic sensitivity of CK-MB mass for acute myocardial infarction was close to 100%, consistent with the kinetics of CK-MB release22 and its inclusion in the conventional diagnostic classification. In patients with unstable angina, 70% had detectable cTnT, exceeding 0.1 µg/l in 31.3%. The confounding effect of CK-MB on the diagnostic classification is illustrated by exclusion of patients with unstable angina, when the diagnostic efficiency of cTnT measurement improved. Myoglobin did not prove to be a consistently better marker than CK-MB in the first six hours, and thereafter its performance declined. Myoglobin has been considered to be one of the earliest markers to rise following acute myocardial infarction. However, early studies did not compare myoglobin with sensitive assays for CK-MB mass and cTnT.10–12 25 The short time window for myoglobin means that early peak values could be missed. Myoglobin is probably useful in the emergency department for early exclusion of myocardial infarction, but is less useful in patients admitted to hospital, as blood samples tend to be taken later.

It has been proposed that the definition of acute myocardial infarction should include cardiac troponins7 and that treatment recommendations should be revised accordingly.26 Selection of an appropriate cut off point is important. In a “normal” group, cardiac troponins are undetectable. Detailed analysis of the diagnostic cut off point from the ROC data suggested that the value should be reduced to 0.05 µg/l for an unselected patient population. The FRISC study showed that at a value above 0.06 µg/l there is an increased risk of a cardiac event.27 Long term follow up of that study group over four years showed a very low event rate at cTnT values less than 0.06 µg/l.28 A recent study suggested a value of 0.03 µg/l for risk stratification but used highly selected patients in a clinical trial of unstable angina. Recently published data have shown that cTnT values exceeding 0.01 µg/l indicate higher risk in the unstable angina population.29 However, cut off points at this level are difficult to measure reliably. There is therefore good evidence to support the use of a cut off value in the range 0.03–0.06 µg/l. We feel that a value of 0.05 µg/l is

### Table 3 Values for troponin T, myoglobin, and CK-MB in non-cardiac populations

<table>
<thead>
<tr>
<th>Disease</th>
<th>n</th>
<th>Troponin T (µg/l)</th>
<th>Myoglobin (µg/l)</th>
<th>CK-MB (µg/l)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Rheumatoid joint disease</td>
<td>39</td>
<td>&lt;0.01 (&lt;0.01 to 0.041)</td>
<td>34.4 (25 to 127.0)</td>
<td>0.8 (0.2 to 6.1)</td>
</tr>
<tr>
<td>Carcinoma</td>
<td>5</td>
<td>&lt;0.01 (&lt;0.01 to 0.02)</td>
<td>41 (30 to 120)</td>
<td>1.3 (0.3 to 5.6)</td>
</tr>
<tr>
<td>GIT</td>
<td>13</td>
<td>&lt;0.01 (&lt;0.01 to 0.041)</td>
<td>58 (38.7 to 147), n=10</td>
<td>2.1 (1.0 to 7.5), n=12</td>
</tr>
<tr>
<td>Viraemia</td>
<td>4</td>
<td>&lt;0.01 (&lt;0.01 to 0.01)</td>
<td>49 (25 to 77)</td>
<td>1.4 (0.3 to 1.4)</td>
</tr>
<tr>
<td>Pneumonia</td>
<td>28</td>
<td>0.016 (&lt;0.01 to 0.72)</td>
<td>101 (36 to 3100), n=26</td>
<td>4.1 (0.3 to 235), n=25</td>
</tr>
<tr>
<td>PE</td>
<td>9</td>
<td>0.026 (&lt;0.01 to 4.06)</td>
<td>48.5 (33 to 84), n=8</td>
<td>1.85 (0.8 to 5.7), n=8</td>
</tr>
<tr>
<td>Other</td>
<td>26</td>
<td>&lt;0.01 (&lt;0.01 to 0.27)</td>
<td>59 (28 to &gt;3000), n=22</td>
<td>2.5 (0.6 to 72), n=25</td>
</tr>
</tbody>
</table>

Values are median (range).

CK-MB, MB isoenzyme of creatine kinase; GIT, gastrointestinal tract disease; PE, pulmonary embolus.
pragmatically the most appropriate. It is in the range for accurate measurement, useful in the general chest pain population, and consistent with our ROC data. The use of a lower cut off may be more appropriate in patients with a definitive diagnosis of acute coronary syndrome, but further studies are required. Clinical studies have shown that the magnitude of cTnT release is a reflection of the degree of actual or functional myocardial damage. The definition of acute myocardial infarction remains clinical, and where cardiac disease could be categorically excluded, no cTnT elevations were seen. In patients with rhabdomyolysis, cTnT was marginally raised if at all, despite substantial elevations of creatine kinase. Elevations of cTnT were only seen where a possibility of myocardial damage existed or where that would be consistent with data reported elsewhere. Clearly, cTnT measurements cannot be used uncritically for diagnosis. Elevation of cTnT is pathognomonic of myocardial cell damage, but only in the presence of appropriate clinical or ECG findings does this constitute acute myocardial infarction. The definition of acute myocardial infarction remains clinical, but cTnT is a core diagnostic test. The measurement of cTnT has been shown to identify patients who will benefit from revascularisation, or from low molecular weight heparin and glycoprotein IIb/IIIa antagonists. In order to be clinically useful a rapid and precise cTnT assay is required, with results comparable from both laboratory and point of care testing. These criteria are met by this assay.

Conclusions

In routine clinical practice, measurement of cTnT by the third generation assay on the 1010 analyser at a diagnostic cut off of 0.05 μg/l allows rapid accurate differential diagnosis of patients with suspected acute coronary syndrome, and of patients with other causes of chest pain. Myoglobin can serve as a useful short term marker but was not superior to cTnT or CK-MB in this study.

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A 61 year old man presented with a history of chest pain of 13 years’ duration. The frequency and severity of the chest pain had started to increase recently and was associated with shortness of breath at rest. There was no history of hyperlipidaemia and cigarette smoking. The patient was taking aspirin, a diuretic, an angiotensin converting enzyme inhibitor, digoxin, nitrates, and mexiletine. On examination his blood pressure was 130/80 mm Hg and pulse rate 80 beats/min. Auscultation revealed a mild systolic murmur on the apex, with slightly harsh vesicular breathing in the lower zone. A chest x ray showed cardiomegaly, and ECG revealed atrial fibrillation with left ventricular hypertrophy; in the strip trace a few ventricular extrasystoles were seen. Biochemical investigation revealed only hyperlipidaemia with no other pathology. Echocardiography showed that the left ventricular internal diameters were dilated (79 mm/57 mm); ejection fraction was 51%, and a third degree mitral insufficiency was evident. Radio-nuclide ventriculography was subsequently undertaken which showed significant hypokinesia in the septum and an ejection fraction of 20%.

The patient was admitted and coronary arteriography and ventriculography undertaken. Coronary arteriography showed a normal left anterior descending coronary artery, but in the circumflex (Cx) region neither normal vascular structure nor collateral circulation was evident, suggesting Cx occlusion (below left). Upon selective right coronary arteriography, the Cx artery arose as a terminal extension of the right coronary artery, supplying the Cx region; however, no stenosis was seen (below right). Left ventriculography showed diffuse hypokinesia. The patient was transferred to surgery for mitral valve replacement because of severe left ventricular dysfunction; however, the operation was cancelled and the patient was treated medically.

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