Detection of myocardial infarction by immunohistological staining for C9 on formalin fixed, paraffin wax embedded sections

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Abstract

Aims—To evaluate an immunohistological stain for complement component C9 as a method of detecting early myocardial infarction and to compare this with (1) an enzyme histochemical method and (2) conventional histological staining.

Methods—(1) Eight hearts taken at necropsy were stained using the nitroblue tetrazolium/phenazine methosulphate method and an immunohistological stain for C9. (2) Twenty five hearts from cases of suspected or confirmed myocardial infarction and 25 from cases without conventional evidence of infarction were stained for C9 and by haematoxylin and eosin.

Results—(1) The histochemical method indicated myocardial necrosis in five hearts and the C9 method in seven, all of which had clinical evidence of myocardial damage or a reason for it. The histochemical method required fresh myocardium, was difficult to use and was difficult to interpret. (2) Of 25 hearts with suspected or confirmed infarction, 24 were stained by the C9 method. Staining with haematoxylin and eosin showed infarction in 16 of these, all with infarcts at least 24 hours old; the other eight had clinical evidence of infarction less than 24 hours old. The heart not stained by C9 was from a patient who, on review, had no evidence of infarction. Of the 25 control hearts, none had infarction on staining with haematoxylin and eosin, but three were stained by the C9 method. These three were from patients with septicaemia or another reason for myocardial damage.

Conclusions—The immunohistological method for C9 is a simple, reliable and sensitive method for the detection of early myocardial necrosis that could be used on formalin fixed, paraffin wax embedded necropsy material. This had advantages over a histochemical method and conventional staining with haematoxylin and eosin.

Methods

Histochemical study
Transverse slices, about 1–2 cm thick, were taken from the left ventricle of eight patients at necropsy (table 1).

Nitroblue tetrazolium/phenazine methosulphate method
The slices of myocardium were rinsed and stored at 4°C in Sorensen’s phosphate buffer (0.1 M, pH 7·4) containing 0·1 M ascorbic acid. Fresh slices were cut just before staining. These were incubated in Sorensen’s buffer containing nitroblue tetrazolium, 50 mg/100 ml, and phenazine methosulphate, 2·5 mg/100 ml, for 15 minutes at 37°C. The slices were rinsed, immersed in 10% formal saline, and photographed.

Immunohistological method for C9
After fixation for at least 18 hours, histological blocks were cut from the myocardial slices, using the photographs as guides to the sites of blocks. Blocks were embedded in paraffin wax on a Shandon Hypercenter with a routine processing schedule. Sections were cut at 5 μm, dried on glass slides at 60°C for 30 minutes, rehydrated, and immersed in 1% hydrogen peroxide in methanol to block endogenous peroxidase. Sheep antiserum to C9 (The Binding Site, Birmingham, UK) was used at a dilution of 1 in 500 in 0·15 M phosphate buffered saline.
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<table>
<thead>
<tr>
<th>Patient (sex/age in years)</th>
<th>Clinical details</th>
<th>Haematoxylin and eosin</th>
<th>Ninhydrin tetrazolium/phenazine methosulphate staining</th>
<th>C9 immunohistological staining</th>
</tr>
</thead>
<tbody>
<tr>
<td>F/33</td>
<td>Died 24 hours after heart transplant</td>
<td>One group of contraction band necrosis</td>
<td>Patchy pink/red</td>
<td>Extensive patchy staining of single cells and groups of cells</td>
</tr>
<tr>
<td>M/44</td>
<td>Died two months after heart transplant</td>
<td>Severe acute rejection with patchy necrosis</td>
<td>Patchy pink/red</td>
<td>Extensive patchy staining of single cells and groups of cells</td>
</tr>
<tr>
<td>F/50</td>
<td>Died one hour after collapse from dysrhythmia: no clinical evidence of myocardial infarct: coronary arteries patent</td>
<td>No infarction identified</td>
<td>Dark blue</td>
<td>None</td>
</tr>
<tr>
<td>F/55</td>
<td>Died from septicaemic shock in liver failure</td>
<td>No infarction identified</td>
<td>Dark blue</td>
<td>A few scattered cells</td>
</tr>
<tr>
<td>M/62</td>
<td>Died seven days after myocardial infarct</td>
<td>Infarcted area</td>
<td>White area</td>
<td>Heavily stained area</td>
</tr>
<tr>
<td>M/68</td>
<td>Died two days after myocardial infarct</td>
<td>Infarcted area</td>
<td>White area</td>
<td>Heavily stained area</td>
</tr>
<tr>
<td>M/73</td>
<td>Died 24 hours after myocardial infarct</td>
<td>No infarction identified</td>
<td>Patchy pink/red</td>
<td>A few groups of cells</td>
</tr>
<tr>
<td>M/83</td>
<td>Died six hours after myocardial infarct</td>
<td>No infarction identified</td>
<td>Dark blue</td>
<td>Scattered single cells</td>
</tr>
</tbody>
</table>

Figure 1  Slice of left ventricle from a 62 year old man stained by the ninhydrin tetrazolium/phenazine methosulphate method seven days after a myocardial infarct. Non-infarcted myocardium appears dark and infarcted myocardium appears white. The junction between the dark and white areas is irregular, producing a mottled pattern at the edges of the main white area.

Figure 2  Histological section taken from the heart in fig 1 at the junction of macroscopically dark and white areas, stained by the immunohistological method for C9. Infarcted myocardial cells appear dark.

and applied to sections for 60 minutes at room temperature on a Shandon Sequenza immunostaining centre. After washing, peroxidase conjugated donkey antiserum to sheep IgG at a dilution of 1 in 100 was applied for 45 minutes. After another wash, peroxidase was detected with hydrogen peroxide and tetra- amino-biphenyl hydrochloride (diaminobenzidine), and sections were counterstained with haemalum, dehydrated and mounted.

Results

Histochemical study

Myocardium without obvious infarction stained dark blue with the ninhydrin tetrazolium/phenazine methosulphate method (table 1). Unstained areas, virtually white, were seen in two hearts with well established infarction (fig 1). Intermediate staining with patches of pink to red coloration was seen in three hearts.

Intracellular deposition of C9 was detected immunohistologically in the areas of infarction as defined by the enzymatic method, with

Study of conventional staining

Necropsy records were reviewed and 25 consecutive cases with clinically suspected or confirmed myocardial infarction were identified. Another 25 cases without clinical or conventional pathological evidence of myocardial infarction were also identified. Sections of formalin fixed, paraffin wax embedded myocardium, stained with haematoxylin and eosin, were inspected. The immunohistological method for C9 was applied to sections cut from the same blocks.
greater intensity of staining at the edges of infarcts (fig 2). There was patchy staining with C9 in the hearts that had pink/red histochemical staining. Scattered single cells stained with C9 in two hearts that had dark blue histochemical staining. C9 was not deposited in muscle cells in one heart which had no clinical, conventional pathological or histochemical evidence of infarction. C9 was detected in arterial walls in viable myocardium and was a useful internal positive control in all hearts.

Table 2 Comparison of staining of myocardium with haematoxylin and eosin (H and E) with immunohistology for C9 in 50 hearts

<table>
<thead>
<tr>
<th></th>
<th>25 hearts with suspected/confirmed myocardial infarction</th>
<th>25 hearts without obvious myocardial infarction</th>
</tr>
</thead>
<tbody>
<tr>
<td>H and E showed infarct; C9 positive</td>
<td>16</td>
<td>0</td>
</tr>
<tr>
<td>H and E showed infarct; C9 negative</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>H and E did not show infarction; C9 positive</td>
<td>8</td>
<td>3</td>
</tr>
<tr>
<td>H and E did not show infarction; C9 negative</td>
<td>1</td>
<td>22</td>
</tr>
</tbody>
</table>

Figure 3 Immunohistological staining for C9 on a section of myocardium from a 64 year old man five days after a myocardial infarct. Subendocardial myocardium is viable and unstained, with immunostaining at the edge of a well established infarct.

Figure 4 Immunohistological staining for C9 on a section of myocardium from a 57 year old man who died of peritonitis. Two myocardial cells contain C9 and appear dark.

Discussion

Immunohistological staining for C9 was better than the nitroblue tetrazolium/phenazine methosulphate method for detecting myocardial infarction. The C9 method had the advantage that formalin fixed, paraffin wax embedded material could be used. Unlike the technical difficulties of the histochemical method, there was no need for special storage or treatment of samples from hearts. The histochemical method was unreliable on an infarct under 24 hours old, whereas the C9 method stained this infarct. The histochemical method is a macroscopic test whereas the C9 method is microscopic and permitted much finer resolution, detecting staining of single cells. The C9 method was so sensitive that it detected scattered single myocardial cell necrosis in three of the 25 hearts without other evidence of myocardial damage, but with an adequate clin-
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ical explanation for such damage. This finding was similar to that in the 55 year old woman with septicaemic shock in the histochemical study (table 1). The C9 method was superior to conventional staining for detecting infarcts less than 24 hours old. Hearts without evidence of infarction were not stained by the C9 method.

Activation of the complement system and deposited C5b-9 complex is known to accumulate in infarcted myocardium, both in humans and experimental animals. Complement deposition in infarcted areas occurs after a few hours, two to four in the rat and five to six in the rabbit, except when reperfusion occurs, when complement deposition is rapid. This observation shows that circulating blood is necessary for the activation and deposition of complement, and accords with our finding that C9 deposition was most noticeable at the edges of well established infarcts.

The immunohistological stain for C9 has many potential uses such as in retrospective studies of hearts to detect unsuspected myocardial infarction and in the study of diffuse non-regional necrosis of individual myocardial cells.

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