A Clinicopathologic Correlation Study of Thallium-201 
Myocardial Scintigraphy in Diagnosis of 
Myocardial Infarction

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SUMMARY
In a series of 1,000 consecutive autopsy cases, we evaluated the 
clinical utility of thallium-201 (Tl-201) myocardial scintigraphy 
and electrocardiography (ECG) in 101 patients who had been 
studied while alive. Fifty-five cases had myocardial infarctions 
(MI) at autopsy. The Tl-201 scintigram and ECG in diagnosis 
of MI showed sensitivities of 68% and 60%, specificities of 87% 
and 83%, and diagnostic accuracies of 76% and 70%, respectively. 
The sensitivity of the Tl-201 scintigram was 70% in anterior MI, 
80% in postero-inferior MI, 25% in lateral and subendocardial 
infarction. The sensitivity was 88% for large massive MI, but was 
low in scattered (50%) or middle-sized MI (17%). The diagnostic 
limit of the resolution of Tl-201 scintigrams was 4.5 cm in long 
diameter. All 8 cases with MI of less than 4 cm could not be 
diagnosed with the technique. There were 48 cases of large MI 
(more than 5 cm), but 8 cases could not be diagnosed by scinti-
graphy because of non-transmural or scattered MI. A comparison 
of the Tl-201 scintigram and ECG showed that 27 cases out of 60 
cases were diagnosed by both methods, 14 only by the Tl-201 
scintigram, 9 only by ECG and 10 by neither method.

Additional Indexing Words:
Autopsy Scintigraphic defect Transmural infarction 
Non-transmural infarction

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Received for publication May 26, 1986.
Electrocardiography, serum enzymes and ultrasonic echocardiography have been used in the diagnosis of myocardial infarction (MI). Myocardial perfusion scintigraphy with thallium-201 (Tl-201) is regarded as a recent advancement for diagnosing MI. However, clinicopathologic correlation studies between Tl-201 myocardial scintigraphy and pathologic findings are still scanty. In our previous clinicopathologic study of Tl-201 myocardial scintigraphy in 47 cases (26 with MI and 21 without MI), it was concluded that the size threshold for MI detection of Tl-201 was 5 cm in long diameter. The purpose of the present study was to assess the utility and limitations of Tl-201 scintigrams and electrocardiography on the basis of pathologic findings in 101 patients.

Materials and Methods

In a series of 1,000 consecutive autopsy cases (during 3.5 years from September, 1980 to February, 1984), 101 patients had been examined with Tl-201 scintigraphy while alive. This population consisted of 47 men and 54 women, ranging in age from 63 to 93 (average age of 79 years). Fifty-five patients had a total of 60 previous myocardial infarctions. Myocardial images were recorded 10 min after an intravenous injection of 2mCi of ionic thallium-201 at rest, using a Searle PHO Gamma V scintillation camera with a converging collimeter. Myocardial images were obtained in four different views (anterior, left oblique 30° and 60°, and left lateral), acquiring 400,000 counts for each view. The site of scintigraphic defects was diagnosed as anterior (including antero-septal), inferior (including infero-septal) and lateral. A lesion at the apical portion was included in either involved adjacent myocardial segment. Each scintigram was evaluated visually by 2 radiologists, without knowledge of the clinical and pathologic findings, and was divided into positive (definitely abnormal or probably abnormal) and negative (probably normal or definitely normal) categories.

Twelve-lead electrocardiograms were also reviewed. An old MI was diagnosed when pathologic Q waves of ≥0.04 sec in duration and the following R waves of at least 25% were present, and acute MI was diagnosed when new pathologic Q waves and ST-T changes developed. Subendocardial infarction was diagnosed when significant ST-T changes and a typical enzyme pattern were present. A MI in patients with left bundle branch block (LBBB) was diagnosed in the same way.

At autopsy, the heart specimens were cut into four or five transverse slices with a thickness of 0.8 to 1.5 cm, after fixation in 10% formaldehyde.
A MI was diagnosed by gross examination when myocardial necrosis or a scar was more than 1 cm long in diameter. Microscopic studies were also performed. A MI was classified as anterior (including antero-septal), posterior (including postero-septal), lateral or subendocardial infarction. MI was also classified according to the types of wall involvement: massive or scattered, and transmural or non-transmural.\(^8,9\) The size of the MI was expressed as large (>5 cm for longest diameter), middle-sized (2-5 cm) and small (1-2 cm).\(^8\) The MIs which occurred after Tl-201 scintigraphy were included in the control group. Unpaired t-tests and Chi-square tests were used to assess differences between the groups.

**Results**

I. Clinical course of MI

Fig. 1 shows the latencies from the onset of the MI to Tl-201 scintigraphy and latencies from Tl-201 examination to death in 55 patients with a pathologically verified MI. The intervals from the onset of MI to Tl-201 scintigraphy...
Table I. Diagnosis of Myocardial Infarction—Autopsy and $^{201}$Tl Scintigram—

<table>
<thead>
<tr>
<th>Size/Origin</th>
<th>Anterior</th>
<th>Postero-inferior</th>
<th>Lateral</th>
<th>Subendocardial</th>
<th>Sensitivity</th>
</tr>
</thead>
<tbody>
<tr>
<td>large mass.</td>
<td>○● ●● ○● ○● ●● ○● ○● ○● □● ○●</td>
<td>○● ●● ○● ○● ○● ○●</td>
<td>●● ○●</td>
<td>35/40 = 87.5%</td>
<td></td>
</tr>
<tr>
<td>scattered</td>
<td>○● ●● ○● ○● ○● ○● ○●</td>
<td>○● ●● ○● ○● ○● ○●</td>
<td>○●</td>
<td>4/8 = 50.0%</td>
<td></td>
</tr>
<tr>
<td>middle-sized</td>
<td>○● ●● ○● ○● ○● ○●</td>
<td>○● ●● ○● ○● ○● ○●</td>
<td>○●</td>
<td>2/6 = 25.0%</td>
<td></td>
</tr>
<tr>
<td>scattered</td>
<td>●● ●● ●● ●●</td>
<td>●● ●● ●● ●●</td>
<td>●●</td>
<td>0/4 = 0%</td>
<td></td>
</tr>
</tbody>
</table>

Sensitivity: 70.4% (27/38), 80.0% (19/24), 25.0% (1/4), 25.0% (1/4), 68.1% (41/60).

○ Defect (+)
● = (-)
* = same cases

ranged from 3 days to about 17 years in 30 patients (33 MI) and were unknown in 25 patients (27 MI). Nineteen of 33 MI were studied within 1 month after the onset of MI and 15 (79%) of them showed positive scintigrams. Thirteen (93%) of 14 MI which were studied more than 1 month after the onset of MI had positive scintigrams, but only 13 (48%) of 27 in which the onset of MI was not known had positive scintigrams. In the 55 patients with MI, the intervals from T1-201 scintigraphy to death ranged from 3 days to 6 years (average of 19 months). Nineteen patients with negative scintigrams were dispersed irrespectively of these intervals. In 46 patients without a MI at autopsy, the intervals from T1-201 scintigraphy to death ranged from 3 days to 6 years (average of 17 months).

II. T1-201 scintigram—Autopsy correlations

The results of a comparative study between pathologic and T1-201 scintigraphic findings for 60 infarctions are shown in Table I. The T1-201 scintigram correctly diagnosed the presence of MI in 19 (70%) of 27 cases with an anterior infarction at autopsy, in 20 (80%) of 25 with a posterior infarction, in 1 (25%) of 4 with a lateral infarction and in 1 (25%) of 4 with a subendocardial infarction. In addition, 35 (88%) of 40 cases with a massive large infarction, 4 (50%) of 8 with a scattered large infarction and 2 (17%) of 12 with a middle-sized infarction had T1-201 scintigraphic defects. Fig. 2 shows 3 representative infarctions which were correctly diagnosed with T1-201 scintigraphy; a massive anterior MI (Fig. 2A), a massive posterior MI (Fig. 2B) and a middle-sized antero-septal MI (Fig. 2C). These three infarctions were transmural and showed thinning of the walls. Nine (75%) of 12 MIs with cardiac aneurysms had T1-201 scintigraphic defects. As
Fig. 2. Transverse sections of the heart viewed from the base. (A) Man, aged 84, massive anterior MI (horizontal length 5 cm, thickness 0.5 cm, longitudinal length 6 cm). The distance of the section from the apex was 3.0 cm. (B) Man, aged 81, massive posterior MI (9 x 0.8 x 7 cm). The distance from the apex was 4.5 cm. (C) Man, aged 83, massive antero-septal MI (3.2 x 0.5 x 4.7 cm). The distance from the apex was 2.8 cm.

a whole, there was a good agreement in 41 (68%) of 60 infarctions between autopsy results and Tl-201 scintigraphic defects.

III. Diagnostic limit of Tl-201 scintigram in size of MI

The relationship between the size of the MI at autopsy and Tl-201 scintigraphic defects is shown in Fig. 3. Eight infarctions less than 4 cm in long diameter were not detected by the Tl-201 scintigram; 7 were non-transmural and/or scattered infarctions. Thus, the limit of detection of MIs by the Tl-201 scintigram was 4.5 cm (long diameter). Fig. 4 illustrates a
Fig. 3. Diagnostic sensitivity of Tl-201 scintigram on the basis of size of myocardial infarction. The correlation between Tl-201 scintigram and size of MI is illustrated, with horizontal length represented by "a" and longitudinal length of the MI by "b". Open circle: cases with a positive Tl-201 scintigram, closed circle: cases with a negative scintigram, upper oblique bar from the circle: non-transmural MI, and lower oblique bar: scattered MI. Seven cases numbered from 1 to 7 were shown pathologically; 1: Fig. 2A, 2: Fig. 2B, 3: Fig. 2C, 4: Fig. 5A, 5: Fig. 5B, 6: Fig. 5C, 7: Fig. 7A and 7B.

Fig. 4. Tl-201 images at rest in Case 3 (Fig. 2C). Note the perfusion defect in the anterior region. Abbreviation: ANT = anterior; LAO = left anterior oblique; L-LAT = left lateral.
Tl-201 scintigraphic defect (anterior), which proved to be the smallest infarction (3.2×0.5×4.5 cm) at autopsy (Fig. 2C). Of 43 infarctions more than 5 cm, 8 were not diagnosed by Tl-201 scintigraphy (Fig. 3). Seven of these cases were non-transmural infarctions and 5 of 7 were also scattered infarctions. As a whole, 31 (89%) of 35 transmural infarctions could be diagnosed but only 10 (40%) of 25 non-transmural infarctions could be diagnosed (p<0.001).
Fig. 6. TI-201 images in Case 7. Note the perfusion defect in the apico-posterior region (partially anterior).

Fig. 5 shows 3 representative infarctions which could not be diagnosed with TI-201 scintigraphy, a circumferential and subendocardial infarction (Fig. 5A), a scattered large non-transmural lateral infarction (Fig. 5B) and a middle-sized septal infarction (Fig. 5C). The scintigram in Fig. 6 shows an apico-inferior defect but no defect in the high antero-septal area. Pathologic examination showed a large antero-septal infarction with a ventricular aneurysm (Fig. 7A) and an apico-posterior infarction (Fig. 7B). Only one of the 2 infarctions was detected with TI-201 scintigraphy.

IV. The sensitivity, specificity and diagnostic accuracy

In Table II the sensitivity, specificity and diagnostic accuracy are shown for TI-201 scintigraphy and ECG on the basis of pathologic findings. The sensitivities of TI-201 scintigraphy and ECG were 68% and 60%, the specificities were 87% and 83% and the diagnostic accuracies were 76% and 70%, respectively. Three indices were higher for TI-201 scintigraphy than for the ECG. Six cases with false positive TI-201 scintigraphy included 2 cases of cardiac amyloidosis, 2 of marked cardiac fibrosis, one of rheumatic endomyocardial fibrosis (Figs. 7C and 8), and one of dilated cardiomyopathy (DCM). Of
Fig. 7. (A) and (B) represent Case 7 (Fig. 6), woman, aged 87, antero-septal MI ($5 \times 0.2 \times 6.5 \text{ cm}$) with aneurysm formation and an apico-posterior MI ($5 \times 0.5 \times 4.5 \text{ cm}$). The distances were 4.0 and 2.0 cm from the apex, respectively. (C) Woman, aged 76, fibrosis due to rheumatic endomyocarditis (postero-lateral region). The distance from the apex was 4.1 cm.
Fig. 8. TI-201 images in Case of Fig. 7C. Note the perfusion defect at the posterior region.

Table II. Diagnosis of Myocardial Infarction—Comparisons of the Sensitivity, Specificity and Diagnostic Accuracy between TI-201 Scintigraphy and ECG—

<table>
<thead>
<tr>
<th>201TI myocardial scintigraphy</th>
<th>ECG</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
</tr>
<tr>
<td>Autopsy MI (+)</td>
<td>MI (+)</td>
</tr>
<tr>
<td>41</td>
<td>36</td>
</tr>
<tr>
<td>19</td>
<td>24</td>
</tr>
<tr>
<td>Total</td>
<td>60</td>
</tr>
<tr>
<td>Autopsy MI (-)</td>
<td>MI (-)</td>
</tr>
<tr>
<td>6</td>
<td>8</td>
</tr>
<tr>
<td>40</td>
<td>38</td>
</tr>
<tr>
<td>Total</td>
<td>46</td>
</tr>
<tr>
<td>Autopsy</td>
<td>MI</td>
</tr>
<tr>
<td>Total</td>
<td>106</td>
</tr>
</tbody>
</table>

8 false-positive ECG cases, 2 cases had cardiac amyloidosis, 2 cases scattered scars, 1 case had aortic regurgitation, 1 case had amyotrophic lateral sclerosis and the 2 remaining cases had no significant findings.

V. Comparison of TI-201 scintigram and ECG in diagnosis of MI

Table III compares the TI-201 scintigram and the ECG in diagnosis of 60 incidents of myocardial infarction. Twenty-seven (45%) were diagnosed by both TI-201 scintigraphy and ECG, 14 (23%) by TI-201 scintigraphy, 9 (15%) by ECG and 10 (16.7%) by neither method. Thirteen (48%) of
Table III. Comparisons of 201TI Scintigrams and ECG in Diagnosis of Various Sites of MI

<table>
<thead>
<tr>
<th></th>
<th>Anterior</th>
<th>Posterior-inferior</th>
<th>Lateral</th>
<th>Subendocardial</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Open circles</td>
<td>13</td>
<td>14</td>
<td>0</td>
<td>0</td>
<td>27/60 45.0%</td>
</tr>
<tr>
<td>Crosses</td>
<td>6</td>
<td>6</td>
<td>1</td>
<td>1</td>
<td>14/60 23.3%</td>
</tr>
<tr>
<td>Crosses</td>
<td>4</td>
<td>2</td>
<td>1</td>
<td>2</td>
<td>9/60 15.0%</td>
</tr>
<tr>
<td>Crosses</td>
<td>4</td>
<td>3</td>
<td>2</td>
<td>1</td>
<td>10/60 16.7%</td>
</tr>
<tr>
<td>Total</td>
<td>13/27</td>
<td>14/25</td>
<td>0/4</td>
<td>0/4</td>
<td>0%</td>
</tr>
</tbody>
</table>

Open circles show the correct diagnosis and crosses represent failure of diagnosis.

Fig. 9. Comparison of the diagnostic sensitivity of Tl-201 scintigrams and electrocardiogram. The size of MI was presented as in Fig. 3. Open circles represent cases, diagnosed by both Tl-201 scintigraphy and ECG, the triangles only by Tl-201 scintigraphy, the squares only by ECG, and closed circles by neither method.

27 anterior infarctions and 14 (56%) of 25 posterior infarctions could be diagnosed by both methods, but none of the 4 lateral and 4 subendocardial infarctions could be diagnosed by either method.

From the view of the size and nature of the MI (Fig. 9), 26 of 27 infarctions diagnosed by both methods were more than 5 cm (long diameter); only 3 cases were non-transmural and scattered infarctions. Eight of 14 diagnosed only by Tl-201 scintigraphy had massive large infarctions. While 4 of 9 diagnosed only by ECG had a large MI, 3 of these cases were non-transmural
and/or scattered MI. All 10 which could not be diagnosed by either method were non-transmural and/or scattered MI.

**DISCUSSION**

T1-201 myocardial scintigraphy is a clinically useful method for diagnosis of acute and old MI and angina pectoris. The MI can be visualized as a perfusion defect in T1-201 scintigraphy, reflecting reduced blood flow and reduced myocardial mass in the infarcted area; partially due to replacement of dead myocardial tissues by scars.\(^{10}\) However, clinicopathologic correlation studies between T1-201 scintigraphy and pathologic findings have been infrequent.\(^{11} \text{–7}\) Particularly, studies in a variety of MI patients and control subjects have been unsatisfactory in a number of patients.\(^{1} \text{–6}\) Wackers et al\(^{1}\) examined 9 cases, Bulkley et al\(^{11}\) 11, Nagai et al\(^{12}\) 12, Murata et al\(^{13}\) 20 and Nishimura\(^{6}\) 10, respectively. In this study, we examined 55 patients with various kinds of MI and 46 control cases without a MI on the basis of pathologic findings.

Wackers et al\(^{11}\) reported that the frequency of positive scans was significantly higher (90 of 96) in patients studied within 24 hours of the onset of MI than in patients studied later (75 of 104) and that repeated scintiscans obtained later than 24 hours showed few changes in defect size. Holman\(^{10}\) reported that the sensitivity of the technique decreases with time after the onset of symptoms. Fletcher et al\(^{12}\) reported that patients with small infarctions showed a greater reduction in size with time. In our patients with infarctions at autopsy, the latency from the onset of MI to T1-201 scintigraphy ranged from 3 days to 16 years and the interval from T1-201 scintigraphy to death was from 3 days to 6 years. Both groups of patients with and without T1-201 scintigraphic defect were distributed independently of these intervals. Cases with a known MI onset had positive scintigrams more frequently than those in which the onset was not known. However, most cases in which the onset was unknown had middle-sized and/or non-transmural infarctions.

The present study indicates that the ability of T1-201 scintigraphic to detect defects was higher for posterior infarctions (80%) than for anterior infarctions (70%). There are reports\(^{13},14\) that the detection of defects was lower for postero-inferior infarctions than for anterior infarctions. This discrepancy might be explained by the fact that, in this study, large and scattered infarctions and middle-sized infarctions were more frequent in anterior infarction than postero-inferior infarction. In cases of massive large infarctions, 17 of 18 with anterior infarctions and 17 of 19 with posterior infarctions showed T1-201 scintigraphic defects. The detection of T1-201 scinti-
Several studies\textsuperscript{11,13,7} have shown that small infarctions are difficult to diagnose with Tl-201 scintigraphy. In this study, 8 cases of MI below 4 cm (long diameter) could not be diagnosed. Small lesions may be obscured by the surrounding viable myocardium.\textsuperscript{15} Tl-201 scintigrams did not consistently show perfusion defects in 9 cases with infarctions from 4 to 5 cm (long diameter). Therefore, the Tl-201 scintigram could reliably detect MIs greater than 4.5 cm in long diameter. In experimental studies using phantoms, Cook et al\textsuperscript{16} reported that lesions over 2.5 cm can be seen in any projection and Narita et al\textsuperscript{17} reported that lesions over 2.5 cm in diameter were accurately detected with Tl-201 scintigraphy. Mueller et al\textsuperscript{18} reported that abnormal scintigrams could be detected for smaller deficits in a phantom than in dogs. In reality, though, the diagnostic limit of Tl-201 scintigram in man could be larger (4.5 cm) than in phantom experiments due to the presence of surrounding tissues of the human body and the beating during the entire cardiac cycle.

Eight cases of large infarction were not diagnosed by Tl-201 scintigraphy; 7 had non-transmural and/or scattered infarction. The remaining false-negative case had a primarily non-transmural infarction with a partially transmural lesion; only the transmural lesion was detected. In 2 previous studies\textsuperscript{19,20} and this study, the detection of Tl-201 scintigraphic defects was lower for non-transmural infarctions (40\% to 80\%) than for transmural infarctions (88\% to 90\%). This may be due to the presence of viable myocardium in both the infarcted area (as in scattered MI) and surrounding regions. Thus, cases in which perfusion defects in Tl-201 scintigrams were detected had 3 common factors; i) transmural infarction, ii) thinning of the wall in the infarcted area and iii) large (more than 5 cm for longest diameter) infarction.

The sensitivity of Tl-201 scintigraphy in diagnosis of an old MI, on the basis of pathologic findings, was reported as 72\% by Wackers et al,\textsuperscript{11} 67\% by Takahashi et al,\textsuperscript{7} 59\% by Murata et al\textsuperscript{5} and 100\% by Nagai et al.\textsuperscript{4} By contrast, the sensitivity of ECG on the basis of pathologic findings was reported as 35\% by Alonso et al,\textsuperscript{21} 96\% by Savage et al\textsuperscript{22} and 60\% by Gunnar et al.\textsuperscript{23} Only Takahashi et al\textsuperscript{7} compared the sensitivity, specificity and diagnostic accuracy of Tl-201 scintigraphy with ECG. In acute MI, Wackers et al\textsuperscript{11} reported that there was a 91\% correspondence between scintigraphic defects and the postmortem location of infarctions and 70\% correspondence between the ECG and postmortem findings. In our study, the sensitivities of Tl-201 scintigraphy and ECG were 68\% and 60\%, the specificities were
87% and 83% and diagnostic accuracies were 76% and 70%, respectively. Furthermore, TI-201 scintigraphy was more useful than ECG in diagnosis of an old MI. In addition, TI-201 scintigraphy will be extremely helpful in diagnosis of MI in patients with abnormal conduction disturbances.11,24) In this study, 1 of 2 patients with LBBB and 1 with an intraventricular conduction disturbance showed TI-201 scintigraphic defects.

False-positive cases were reported in patients with coronary artery disease,25,26) dilated cardiomyopathy,27) aortic valve stenosis and sarcoidosis.28) There were 6 false-positive cases in this study, of which 2 had cardiac amyloidosis, 2 marked cardiac fibrosis, 1 rheumatic endomyocardial fibrosis and 1 dilated cardiomyopathy.

Twenty-seven of 60 cases (45%) could be diagnosed by both TI-201 scintigrams and ECG, 14 (23%) only by the TI-201 scintigram, and 9 (15%) only by ECG. A MI could be diagnosed more correctly and more frequently by a combination of methods. In this study, 5 of 9 patients who were diagnosed only by ECG had non-transmural and/or scattered infarctions. Fourteen who were diagnosed only by TI-201 scintigraphy had MIs of various sizes. Since 8 large infarctions were included in the group, the reason for the failure of ECG diagnosis was not known. Finally, 10 cases could not be diagnosed by either method.

References