Does technetium-99m diethylenetriaminepentaacetic acid clearance predict the clinical course of idiopathic pulmonary fibrosis?

Marco Mura MD, Gelmoma Belmonte MD, Stefano Fanti MD, Angela Maria Grazia Pacilli MD, Luca Fasano MD, Maurizio Zompatori MD, Mario Schiavina MD, Mario Fabbri MD

Clearance of inhaled technetium-99m diethylenetriaminepentaacetate (99mTc-DTPA) is a potential indicator of disease activity and progression in idiopathic pulmonary fibrosis (IPF). The objective of the present study was to evaluate the prognostic value of 99mTc-DTPA scans in IPF. A total of 22 patients (18 males), aged 33 to 80 years with IPF were followed for six to 20 months (mean 13 months). At diagnosis, high-resolution computed tomography (HRCT) scans showed a honeycomb pattern with bibasilar reticular opacities in all cases. At T0 (diagnosis) and T1 (follow-up), each patient had pulmonary function tests (forced vital capacity, diffusing capacity of the lung for carbon monoxide and partial arterial O₂ pressure), extension of fibrosis evaluated by HRCT visual score and 99mTc-DTPA lung clearance. Results at T0 and T1 were compared, taking into account the whole population and patients with relatively fast and slow 99mTc-DTPA wash-out. 99mTc-DTPA clearance did not show any significant correlation with functional tests or HRCT score. These findings indicate that clearance of inhaled 99mTc-DTPA is not of value in following the progress of IPF.

Key Words: 99mTc-DTPA clearance; High resolution computed tomography; Idiopathic pulmonary fibrosis; Lung epithelial permeability

I diopathic pulmonary fibrosis (IPF) is a chronic fibrotic pulmonary disease with poor prognosis (1,2). Currently, the recommended approach to stage and predict the progression of IPF is based on pulmonary function tests (PFTs) and high-resolution computed tomography (HRCT) (3-5).

Technetium-99m diethylentriaminepentaacetic acid (99mTc-DTPA) aerosol clearance is a noninvasive index of pulmonary epithelial permeability; increased permeability results from inflammation of the lung parenchyma in a variety of diffuse lung diseases (6). Test interpretation is based on the analysis of radionuclide clearance after the isotope is administered as a soluble aerosol, with a particle diameter of less than 2 µm, and after cigarette smoking has been suspended for at least one week (7-9). A clearance half-time of less than 50 min is considered to be pathological and suggests diffuse interstitial damage (10,11). Increased 99mTc-DTPA clearance has been cited as an early marker of inflammation and activity grade (12,13). The role of 99mTc-DTPA clearance in IPF is controversial; according to some authors (13), the procedure should not be recommended as a routine test in IPF; while other investigators (14,15) have identified this technique as a significant predictor of clinical course and survival.

The present study was designed to evaluate the potential of 99mTc-DTPA scans to assess disease activity and progression of IPF. To assess the usefulness of 99mTc-DTPA scans, we prospectively studied a group of IPF patients and compared the results of this nuclear medicine test with HRCT quantitative scanning and PFTs.
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TABLE 1
Pulmonary function studies, radiographic extent of idiopathic pulmonary fibrosis and technetium-99m diethylenetriaminepentaacetate (99mTc-DTPA) findings at diagnosis (T0) and follow-up (T1) in the whole population (n=22)

<table>
<thead>
<tr>
<th></th>
<th>T0</th>
<th>T1</th>
<th>P</th>
</tr>
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<tbody>
<tr>
<td>HRCT score (%)</td>
<td>46.5±12.8 (range 20.0 to 70.0)</td>
<td>51.8±13.4 (range 25.8 to 75.0)</td>
<td>0.0013</td>
</tr>
<tr>
<td>FVC (%) pred</td>
<td>69.8±18.6</td>
<td>64.8±16.7</td>
<td>0.0002</td>
</tr>
<tr>
<td>DLCO (%) pred</td>
<td>36.9±21.1</td>
<td>27.7±19.9</td>
<td>0.00003</td>
</tr>
<tr>
<td>PaO2 (mmHg)</td>
<td>75.0±14.5</td>
<td>68.7±15.2</td>
<td>0.0011</td>
</tr>
<tr>
<td>Clearance of 99mTc-DTPA (min)</td>
<td>29.5±11.3</td>
<td>35.2±15.2</td>
<td>0.033</td>
</tr>
</tbody>
</table>

Student’s t test for dependent variables. DLCO Diffusing capacity of the lung for carbon monoxide; FVC Forced vital capacity; HRCT High resolution computed tomography; PaO2 Arterial O2 pressure at rest; pred Predicted normal value

METHODS

Twenty-two patients, 18 males and four females, aged 33 to 80 years (mean 66±10.5 years) (19 former smokers), diagnosed with IPF on the basis of American Thoracic Society/European Respiratory Society criteria (6), were studied and followed for a period of six to 20 months (mean follow-up was 13 months). HRCT scans revealed a honeycomb pattern with basal and peripheral reticular opacities, and traction bronchiectasis in all patients (16).

Patients were examined when stable, without signs of heart failure or acute inflammation, on two occasions at presentation (T0) and at follow-up (T1). Patients had PFTs, and HRCT and 99mTc-DTPA scans. Informed consent was obtained from all patients.

During the T0 to T1 interval, nine patients were treated with corticosteroids (prednisolone 0.5 mg/kg/day for four weeks, followed by 0.25 mg/kg/day for eight weeks and then followed by 0.125 g/kg/day for four to 12 weeks).

HRCT scans were performed with a single breath hold in the prone position (17), without intravenous injection of contrast material and using a filter for the osseous tissue. One millimetre thick sections were taken, with a 1 s scan time and 10 mm intervals between sections. Both lungs were included in the field of view. The extent of the lesions was evaluated with a visual score and expressed as a percentage of total lung volume, as previously described (18).

PFTs were performed according to American Thoracic Society guidelines (19). The following parameters were examined in all patients: partial arterial O2 pressure (PaO2) was determined using a Radiometer ABL 520 gas analyzer (Diamond Diagnostic, USA) and expressed in mmHg; forced vital capacity (FVC) was determined using a Morgan Transflow 544 pneumotachograph (Morgan Scientific, USA); and diffusing capacity of the lung for carbon monoxide (DLCO) was determined using the single breath method, with correction for hemoglobin values. FVC and DLCO were expressed as percentages of the predicted normal values (20).

The 99mTc-DTPA scans were performed by administering the radionuclide as an aerosol from an Ultravent nebulizer (Mallinckrodt Medical Inc, USA) consisting of particles 0.89 µm in diameter. Scintigraphy was performed with a gammacamera, equipped with a low energy general purpose collimator interfaced to a computer. Patients were administered the aerosol for 3 min to 4 min in a closed circuit system and at normal tidal volume. At the end of aerosol administration, chest images were immediately recorded with the patient in the seated position. Each dynamic image was acquired for 10 s in posterior projection on a 64x64 matrix, with a total acquisition time of 20 min. Regions of interest were manually drawn around each lung and data were processed to obtain time-activity curves. Clearance half-time was calculated using a least squares curve fitting routine to find the best exponential fit for the first 7 min of data because the contribution of background activity was insignificant in this time interval. A clearance half-time less than 50 min was considered to be pathological.

Differences between T0 and T1 findings were investigated using the Student’s t test for dependent variables, while the Student’s t test for independent variables was used to compare the findings between different groups. Correlations among HRCT scores, pulmonary function studies and 99mTc-DTPA clearance were analyzed using the two-tailed Pearson’s correlation coefficient. P<0.05 was regarded as significant.

RESULTS AND DISCUSSION

At presentation (T0), irregular opacities, prevailing at the bases, were observed in all patients’ chest radiographs. All patients showed a restrictive ventilatory impairment (Table 1). DLCO values correlated significantly with both PaO2 (r=0.53, P=0.021) and FVC (r=0.57, P=0.009) at T0. The extent of parenchymal abnormalities evaluated with HRCT score showed an inverse and significant correlation with PaO2 (r=-0.64, P=0.001), FVC (r=-0.47, P=0.023) and DLCO (r=-0.46, P=0.028) at T0. Clearance of 99mTc-DTPA did not show any significant relationship with the other parameters considered, at either T0 or T1.

The extent of the lesions demonstrated by HRCT scan showed a significant increase during the T0 to T1 interval; similarly, a significant decrease in FVC, DLCO and PaO2 was observed. A decrease in the extent of reticular opacities was

TABLE 2
Pulmonary function studies and radiographic extent of idiopathic pulmonary fibrosis at diagnosis (T0) and follow-up (T1) in patients with markedly accelerated clearance (CI) of inhaled technetium-99m diethylenetriaminepentaacetate (99mTc-DTPA) (group I) and patients with moderately accelerated CI of inhaled 99mTc-DTPA (group II)

<table>
<thead>
<tr>
<th></th>
<th>Group I (Cl&lt;25 min; n=10)</th>
<th>Group II (Cl&gt;25 min; n=12)</th>
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<tbody>
<tr>
<td>T0</td>
<td></td>
<td></td>
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<tr>
<td>HRCT scan score (%)</td>
<td>48.7±13.3</td>
<td>44.9±14.0</td>
</tr>
<tr>
<td>FVC (%) pred</td>
<td>67.8±20.6</td>
<td>69.8±18.3</td>
</tr>
<tr>
<td>DLCO (%) pred</td>
<td>38.1±18.8</td>
<td>35.4±24.8</td>
</tr>
<tr>
<td>PaO2 (mmHg)</td>
<td>76.9±10.3</td>
<td>73.2±17.8</td>
</tr>
<tr>
<td>T1</td>
<td></td>
<td></td>
</tr>
<tr>
<td>HRCT scan score (%)</td>
<td>53.7±11.7</td>
<td>49.6±13.0</td>
</tr>
<tr>
<td>FVC (%) pred</td>
<td>62.3±13.2</td>
<td>65.5±19.3</td>
</tr>
<tr>
<td>DLCO (%) pred</td>
<td>24.8±14.1</td>
<td>30.3±23.2</td>
</tr>
<tr>
<td>PaO2 (mmHg)</td>
<td>66.5±15.6</td>
<td>70.1±13.1</td>
</tr>
</tbody>
</table>

Comparison of the two groups (Student’s t test for independent variables) showed no significant differences at T0 or T1. DLCO Diffusing capacity of the lung for carbon monoxide; FVC Forced vital capacity; HRCT High resolution computed tomography; PaO2 Arterial O2 pressure at rest; pred Predicted normal value
not observed in any patient. On the other hand, the clearance of inhaled 99mTc-DTPA showed a significant slowing during the T0 to T1 interval (higher and therefore less pathological values of half-time) (Table 1).

Clearance of inhaled 99mTc-DTPA was less than 50 min in all patients at T0. To investigate whether 99mTc-DTPA clearance at presentation could be considered a predictive index of clinical course, the patients were subdivided into two groups: patients with a half-time clearance less than 25 min at T0 (group I, n=10) and patients with a half-time clearance greater than 25 min at T0 (group II, n=12). Comparing the two groups at T0, PFTs and HRCT scores showed no significant differences (Table 2).

During the T0 to T1 interval, a worsening of both radiological and functional parameters was observed in both groups I and II, although in patients with half-time clearance less than 25 min (group I), only DLCO and PaO2 changes were statistically significant. In patients with a half-time clearance greater than 25 min (group II), all parameter changes were significant. Further, at T1 there were no significant differences between the two groups (Table 2). Therefore, the evolution of impairment was very similar between the two groups.

No correlations were found between 99mTc-DTPA clearance and PFTs or HRCT quantitative scores at baseline, and changes in HRCT and PFTs were not correlated with changes in 99mTc-DTPA clearance values. Patients with more rapid clearance at presentation did not show more functional or radiological deterioration. In fact, clearance tended to improve, while the other indexes worsened.

These results were not influenced by treatment because no significant difference, in any considered parameter, was observed between treated and untreated subjects, either at T0 or at T1 (data not shown).

Previous studies have investigated the role of 99mTc-DTPA scans in the management of IPF patients; according to some authors (14), a normal 99mTc-DTPA clearance at presentation may predict stable lung function, while accelerated clearance may identify those cases that will further progress. However, this study included a majority of patients with idiopathic interstitial pneumonias other than IPF.

Recently, Mogulkoc et al (15) analyzed the 99mTc-DTPA clearance in a considerable number of patients with significant ground-glass opacities in the HRCT scan, thought to represent IPF at an early stage. The authors identified 99mTc-DTPA clearance as an independent predictor of survival, together with age, total lung capacity and DLCO (15). However, clearance was the least significant among the parameters considered, and did not correlate with PFTs or HRCT scores (15). A prognostic role cannot be excluded, especially when the diagnosis of IPF is early; however, at present, these results may not justify the use of an expensive and time-consuming technique such as 99mTc-DTPA scans.

CONCLUSIONS
Although the small number of patients considered precludes any definitive conclusion, our results suggest that 99mTc-DTPA scans may be too expensive when compared with the information supplied by other, less costly techniques. We conclude that the use of 99mTc-DTPA scan is not of value in following the progress of IPF, due to its poor predictive value on the clinical course of this disease.

REFERENCES
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