Prognostic Value of the Carcinoembryonic Antigen Found in Pleural Lavage Fluid From Patients With Lung Carcinoma

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OBJECTIVE: To detect the tumor marker carcinoembryonic antigen (CEA) in pleural lavage fluid taken during surgery from patients with pulmonary carcinoma without associated pleural effusion and assess its possible prognostic implications.

PATIENTS AND METHODS: A prospective, observational study was undertaken to include consecutive patients who underwent surgical treatment for lung carcinoma in which pleural lavage was performed prior to closure of the thoracic cavity (study group). The same techniques and measurements were used in patients undergoing thoracotomy for benign disease (control group). The preoperative blood level of CEA was also quantified.

RESULTS: In the study group, the median CEA levels in blood and pleural lavage fluid were 2.90 ng/mL and 0.40 ng/mL respectively; these figures are higher than those corresponding to the control group. A CEA level of 0.30 ng/mL in pleural lavage fluid was established as a cutoff point, based on the corresponding receiver operating characteristic curve, with a sensitivity of 68.4% and a specificity of 35.7%. A graph of survival in relation to this cutoff point revealed a statistically significant effect (P<0.05).

CONCLUSIONS: It is possible to detect CEA in pleural lavage fluid from the thoracic cavity of patients with lung carcinoma. The values obtained are higher than those found in fluid from patients without neoplastic disease, and this parameter functions as an independent predictor of disease course.

Key words: Carcinoembryonic antigen. Pleural lavage. Pulmonary carcinoma.

Introduction

Lung cancer is the most common cause of death due to neoplastic disease in developed countries. Prognosis is poor, and the 5-year survival rate of less than 15% is attributed to the lack of early detection and the impossibility of cure in patients with extensive disease.

TNM classification is the principal guide to the prognosis of non-small cell lung cancer.1 Prognosis can be influenced by many additional biological factors, of which the most notable are the circulating tumor markers in blood. Numerous studies indicate the particular relevance of carcinoembryonic antigen (CEA) in the diagnosis and management of patients with non-small cell lung cancer.2

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CEA was one of the first markers to be evaluated in lung cancer.\textsuperscript{3} Although it is also elevated in patients with adenocarcinoma of the gastrointestinal tract, CEA is found at increased concentrations in the blood of 40% to 80% of lung cancer patients, and the antigen acts as a sensitive indicator of adenocarcinomas and advanced stages of the disease. In fact, the blood concentration of CEA has been correlated with the spread and/or progression of the primary disease.\textsuperscript{4}

Analysis of tumor markers in serum has been applied to the diagnosis, prognosis, and follow up of patients with lung cancer. Although the possibility of early diagnosis has been considered, analysis of tumor markers is currently most useful for the evaluation of therapeutic efficacy and as an indicator of relapse or disease progression.\textsuperscript{5}

Pleural fluid from patients with neoplastic pleural effusion is known to contain detectable levels of tumor markers, notably CEA.\textsuperscript{6} However, the literature does not contain descriptions of the detection of CEA in pleural lavage fluid from patients with lung carcinoma without concomitant pleural effusion. Thus, the aim of this study was to assess the possibility of isolating CEA in pleural lavage fluid, with the intention of establishing its possible significance as an independent prognostic variable.

Patients and Methods

\textbf{Patient Characteristics}

From January 2000 to January 2004, a total of 164 patients with potentially resectable lung carcinoma were identified for inclusion in a prospective study. A medical history was taken, and physical examination, complete laboratory workup (including preoperative CEA), radiography (chest radiograph, and computed tomography scan of the thoracic and abdominal region), and bronchoscopy were performed in all patients.

Inclusion criteria were as follows: a) presence of a primary lung tumor; b) thoracotomy performed with the aim of curing the cancer; c) determination of serum CEA levels on the morning of surgery; d) washings of the thoracic cavity performed following thoracotomy; e) absence of kidney failure; f) absence of positive cytology for malignancy in pleural lavage fluid; and g) determination of CEA levels in lavage fluid from the thoracic cavity.

Exclusion criteria were as follows: a) presence of pleural effusion (as determined by radiography and confirmed on opening the thoracic cavity), and b) presence of serosanguineous fluid in pleural washes.

Patients were assigned to 2 groups: a study group, comprising patients with lung neoplasms who met the specified criteria (n=164), and a control group, containing patients who underwent thoracotomy for nonneoplastic diseases and in whom pleural lavage was performed (n=30).

After the period for inclusion of subjects was closed on January 31, 2004, a follow up was undertaken to determine the condition of patients at 3 months (April 30, 2004), either by telephone interview or assessment of the patient’s chart.

\textbf{Pleural Lavage Technique}

Washes of the pleural cavity were performed during surgery in all patients. The lavage procedure entailed the introduction of 200 mL of physiologic saline (isotonic solution of 0.9% sodium chloride, Braun-Dexon, Spangenberg, Germany) into the cavity, followed by recovery of 30 mL with a syringe after 2 to 3 minutes. Aliquots of the samples were placed in test tubes to be sent to the pathology department and hospital laboratory for cytology and determination of CEA levels.

\textbf{Statistical Analysis}

Since the quantitative variables displayed a nonparametric distribution, they are presented as the median and the 25th and 75th percentiles. Between-groups comparisons were performed using the Mann-Whitney U test.

To estimate the cutoff point for the use of CEA concentration in pleural lavage fluid as a prognostic marker, a receiver operating characteristic curve was constructed. The Kaplan-Meier procedure was used to plot the survival curve, and between-groups comparisons of the functions were performed using the log-rank test.

Statistical significance for hypothesis testing was established at \( P < .05 \).

\textbf{Results}

In all of the patients included in the study, the macroscopic appearance of the pleural lavage fluid was clear, with no evidence of blood contamination.

Table 1 shows the characteristics of the patients studied. Neoadjuvant treatment was performed with chemotherapy in 28 patients and with radiotherapy in 3 patients.

\begin{table}[h]
\centering
\begin{tabular}{|l|c|}
\hline
Characteristics & Results \tabularnewline \hline
Sex & \textsuperscript{a}Male/female 149/15 \tabularnewline Age, years & 63 (55-69)\textsuperscript{a} \tabularnewline Procedure & \textsuperscript{a}Atypical resection 14 \tabularnewline & Lobectomy 107 \tabularnewline & Pneumonectomy 36 \tabularnewline & No resection 7 \tabularnewline Histology & Adenocarcinoma 71 \tabularnewline & Squamous 77 \tabularnewline & Large cells 7 \tabularnewline & Others 9 \tabularnewline Stage & 0 3 \tabularnewline & Ia 20 \tabularnewline & Ib 62 \tabularnewline & Iia 3 \tabularnewline & IIb 24 \tabularnewline & IIIa 26 \tabularnewline & IIIb 19 \tabularnewline & IV 7 \tabularnewline Visceral pleura involvement & Yes 49 \tabularnewline & No 115 \tabularnewline Mean size of the tumor, cm & 3.5 (2.4-5.5)\textsuperscript{a} \tabularnewline \hline
\end{tabular}
\caption{Patient Characteristics}
\end{table}

\textsuperscript{a}Data correspond to the median (25th-75th percentiles).
In the study group, the median CEA levels in blood and pleural lavage fluid were 2.90 ng/mL and 0.40 ng/mL, respectively; the concentrations found in the study group were higher than those corresponding to the control group (0.85 ng/mL and 0.20 ng/mL, respectively). A CEA level of 0.30 ng/mL in pleural lavage fluid was established as a cutoff point, based on the corresponding receiver operating characteristic curve, with a sensitivity of 68.4% and a specificity of 35.7% (Table 2). The survival curves for the study group (Figure 2) revealed survival to be significantly different above and below the cutoff point (P = .044).

The median time until patient death was 9.7 months (25-75 percentiles: 13.3-26.5) for the group in which the CEA concentration in pleural lavage fluid was 0.30 ng/mL or higher, and 23.9 months (25-75 percentiles: 13.3-26.5) in the group in which the level was less than 0.30 ng/mL.

Univariate analysis (Table 3) did not show statistically significant relationships between CEA level in pleural lavage fluid and either involvement of the visceral pleura, tumor stage (initial stages [I and II] or advanced stages [III and IV]), mediastinal node involvement (grouping N0-N1 or N2 involvement), histologic type, or size of the tumor (<5 cm or ≥5 cm).

Discussion

In the literature, more than 150 prognostic factors based on characteristics of the tumor or patient, or on the therapeutic interventions have been identified as associated with lung carcinoma. Analysis of tumor markers in serum has been applied to the diagnosis, prognosis, and treatment of patients with lung tumors. Furthermore, pleural fluid from patients with neoplastic pleural effusion is known to contain detectable levels of tumor markers.

Tumor markers can be defined as substances or cellular changes associated with the process of neoplastic transformation. It would be desirable to use CEA as a diagnostic and therapeutic tool in the management of lung carcinoma.

TABLE 2

<table>
<thead>
<tr>
<th>CEA in Pleural Lavage Fluid (ng/mL)</th>
<th>&lt;0.30</th>
<th>≥0.30</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dead (n=38)</td>
<td>68.4% (n=26)</td>
<td>31.6% (n=12)</td>
</tr>
<tr>
<td>Living (n=126)</td>
<td>64.3% (n=81)</td>
<td>35.7% (n=45)</td>
</tr>
<tr>
<td>Total (n=164)</td>
<td>107</td>
<td>57</td>
</tr>
</tbody>
</table>

TABLE 3

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Pleural Lavage Fluid</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>Histology</td>
<td></td>
<td>.642</td>
</tr>
<tr>
<td>Squamous</td>
<td>0.4 (0.200-0.500)</td>
<td></td>
</tr>
<tr>
<td>Aденocarcinoma</td>
<td>0.3 (0.200-0.600)</td>
<td></td>
</tr>
<tr>
<td>Staging</td>
<td></td>
<td>.268</td>
</tr>
<tr>
<td>Stages I-II</td>
<td>0.3 (0.200-0.600)</td>
<td></td>
</tr>
<tr>
<td>Stages III-IV</td>
<td>0.4 (0.200-0.700)</td>
<td></td>
</tr>
<tr>
<td>Visceral pleura involvement</td>
<td></td>
<td>.57</td>
</tr>
<tr>
<td>No (n=115)</td>
<td>0.350 (0.200-0.575)</td>
<td></td>
</tr>
<tr>
<td>Yes (n=49)</td>
<td>0.450 (0.200-1.125)</td>
<td></td>
</tr>
<tr>
<td>Node involvement</td>
<td></td>
<td>.57</td>
</tr>
<tr>
<td>N0-N1 (n=132)</td>
<td>0.3 (0.200-0.600)</td>
<td></td>
</tr>
<tr>
<td>N2 (n=32)</td>
<td>0.4 (0.200-0.700)</td>
<td></td>
</tr>
<tr>
<td>Size of the tumor, cm</td>
<td></td>
<td>.193</td>
</tr>
<tr>
<td>&lt;5 (n=124)</td>
<td>0.3 (0.200-0.600)</td>
<td></td>
</tr>
<tr>
<td>≥5 (n=40)</td>
<td>0.4 (0.200-0.700)</td>
<td></td>
</tr>
</tbody>
</table>

*Values are shown as the median concentration of carcinoembryonic antigen (ng/mL) obtained in pleural lavage fluid (25th-75th percentiles, weighted average).
begin to use all of this biological information to complement the prognostic capacity of the anatomic TNM staging. In addition, malignant cells can be detected in pleural fluid, with a diagnostic yield of around 60%. Some studies have taken a step further and assessed the value of cytologic findings in pleural lavage fluid from patients without associated pleural effusion. However, no data has been published on the detection and value of tumor markers in pleural lavage fluid, the objective of this study.

CEA encompasses a heterogeneous group of proteins that circulate at high concentration in the blood of patients with certain malignant tumors, especially those of epithelial origin. Although the sensitivity and specificity for the diagnosis of lung tumors is low, elevated levels of this marker at the time of diagnosis are associated with poor prognosis, even after presumably curative surgical excision of the tumor. This conclusion has also been extended to the measurement of CEA in pleural fluid, where it achieves a sensitivity of 49% and a specificity of 98%. In addition, patients with malignant pleural effusion have been demonstrated to present higher levels of CEA in pleural fluid than patients with other pleural exudates.

In this study, we found higher CEA levels in pleural lavage fluid from patients with lung carcinoma than in a control group of patients undergoing thoracotomy for nonneoplastic disease.

Blood contamination of the chest cavity was avoided following thoracotomy, and serosanguineous pleural lavage fluid was discarded. Furthermore, the finding that preoperative blood CEA levels (determined on the day of surgery) differed from those of pleural lavage fluid reinforces the conclusion that there was no blood contamination of the samples.

One of the criteria for inclusion of patients was negative cytology of pleural lavage fluid. We decided to include this criterion in order to avoid additional complicating factors whilst making the study group as homogeneous as possible.

Following analysis of the concentration of CEA in pleural lavage fluid, we established a cutoff point of 0.30 ng/mL (Figure 1). Comparison of the survival curves (Figure 2) for the 2 groups included in the study (CEA levels in pleural lavage fluid of ≥0.30 ng/mL or <0.30 ng/mL) revealed a statistically significant difference.

At the end of the study, 126 patients were alive and 38 had died as a result of neoplastic lung disease. In the patients who had died, the median concentration of CEA in pleural lavage fluid was 0.916 ng/mL, compared with 0.556 ng/mL in the living patients.

We believe that the CEA obtained in pleural lavage fluid from patients with lung carcinoma functions as additional objective data that is useful in establishing a more accurate prognosis of the neoplastic disease.

We have demonstrated that it is possible to detect CEA in pleural lavage fluid from the thoracic cavity of patients with pulmonary carcinoma, and that the values obtained are higher than those found in patients without neoplastic disease. This parameter functions as an independent predictor of disease course.

Acknowledgments

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REFERENCES