Original Article
Expression level of serum human epididymis 4 and its prognostic significance in human non-small cell lung cancer

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Abstract: The prognostic significance of serum human epididymis protein 4 (HE4) levels in human NSCLC among a Chinese population has not been investigated. The purpose of this study was to evaluate the prognostic significance of serum HE4 level in patients with NSCLC among a Chinese population. Serum HE4 expression levels were measured by enzyme-linked immunosorbent assay (ELISA). The overall survival (OS) analyzed by log-rank test, and survival curves was plotted according to Kaplan-Meier. The COX proportional hazards regression model was used to determine the joint effects of several variables on survival. Serum HE4 level was found to be significantly higher in patients with NSCLC than that of controls (13.76 ± 5.01 ng/ml vs. 5.09 ± 1.25 ng/ml, P < 0.01). High HE4 expression was correlated with TNM stage (P = 0.003), lymph node metastases (P = 0.007), and distant metastases (P < 0.001). Furthermore, patients with high serum HE4 level had a significantly lower 5-year OS rate (34.0% vs. 59.7%; P = 0.022) than those with low serum HE4 level. In a multivariate Cox model, we found that HE4 expression was an independent poor prognostic factor for 5-year OS (hazards ratio [HR] = 3.654, 95% confidence interval [CI] = 2.753-11.981, P = 0.019) in NSCLC. In conclusion, the detection of HE4 levels in the serum might serve as a new tumor biomarker in the prognosis of NSCLC among Chinese population.

Keywords: Lung cancer, prognosis, HE4, serum, marker

Introduction
Lung cancer still remains the most common cancer and the most common cause of cancer-related death worldwide [1, 2]. About 85% lung cancers were non-small cell lung cancers (NSCLC), and approximately two thirds of NSCLC cases were diagnosis at locally advanced or metastatic disease as the typically asymptomatic at early stages [3]. It is well known that pathologic TNM category, age, sex, and cell type are all important prognostic factors for the patients with NSCLC [4]. The advances in molecular biology have enabled researchers to focus on molecular or biological markers in NSCLC.

Human epididymis protein 4 (HE4) is an 11 kDa protein that is a precursor to the epididymal secretory protein E4 and has since been reported to be expressed in several normal tissues including the epithelia of respiratory and reproductive tissues [5, 6]. Previously, Paek et al. demonstrated that an elevated serum HE4 level was related to the advanced stage of epithelial ovarian cancer. An elevated serum level of HE4 was a poor prognostic factor for progression-free survival in patients with epithelial ovarian cancer who were treated with debulking surgery and adjuvant taxane and platinum-based chemotherapy [7]. Iwahori et al. found that serum HE4 levels were elevated in lung cancer, and it might be a potential prognostic marker for lung cancer patients [8]. However, the prognostic significance of serum HE4 levels in human NSCLC among Chinese population has not been investigated. The purpose of this study was to evaluate the prognostic significance of serum HE4 level in patients with NSCLC among Chinese population.
Committee of Provincial Hospital Affiliated to Shandong University). Tumor stage was determined according to the 2009 TNM staging classification system. The clinical characteristic of 100 patients with NSCLC was summarized in Table 1. Fasting blood was taken for all participants and serum was collected and stored at -80°C. Enzyme-linked immunosorbent assay was measured by enzyme-linked immunosorbent assay (ELISA) with immunoassay kit (Miltenyi, Germany) according to the manufacturer’s directions. The optical density (OD) at 450 nm was determined. The standard curves were established with OD450 as Y axis and the concentration of standard substance as X axis. The level of protein was obtained through standard curve. Results are reported as concentration of HE4 ng/ml in samples.

Statistical analysis was performed using SPSS 18.0 (SPSS, IBM, Chicago, IL, USA). For continu-

Materials and methods

Patients and serum samples

The selection criteria for patients with NSCLC were as follows: (1) pathologically confirmed patients with NSCLC; (2) the patients had no previous history of other cancers. 100 patients with recently diagnosed NSCLC in the Department of Thoracic Surgery, Provincial Hospital Affiliated to Shandong University between October 2007 and February 2013 were enrolled in the present study. Control serum samples were collected from 100 healthy volunteers. All subjects underwent clinical examination; plain chest radiograph; CT scan of the chest, upper abdomen, and brain; fiberoptic bronchoscopy; and bone scan. Blood samples were collected from the patients at the time of diagnosis, before any kind of treatment (surgery, radiation, or chemotherapy). The demographic and pathological data, including age, gender, and the tumor stage, were obtained by a review of the patients’ medical records (the data was used with the consent of the patients as well as the approval of the Ethics Committee of Provincial Hospital Affiliated to Shandong University). Tumor stage was determined according to the 2009 TNM staging classification system. The clinical characteristic of 100 patients with NSCLC was summarized in Table 1. Fasting blood was taken for all participants and serum was collected and stored at -80°C.

Enzyme-linked immunosorbent assay

HE4 was measured by enzyme-linked immunosorbent assay (ELISA) with immunoassay kit (Miltenyi, Germany) according to the manufacturer’s directions. The optical density (OD) at 450 nm was determined. The standard curves were established with OD450 as Y axis and the concentration of standard substance as X axis. The level of protein was obtained through standard curve. Results are reported as concentration of HE4 ng/ml in samples.

Statistical analysis

Statistical analysis was performed using SPSS 18.0 (SPSS, IBM, Chicago, IL, USA). For continu-
Results

Serum level of HE-4 in patients with NSCLC and controls

Serum HE4 level was found to be significantly higher in patients with NSCLC than that of controls. Mean serum HE4 level was 13.76 ± 5.01 ng/ml in the NSCLC group and 5.09 ± 1.25 ng/ml in the control group (P < 0.01, shown in Figure 1). 95% of HE4 values of the control group were 7.26 ng/ml, which was used as the cutoff value.

Increased serum HE4 expression level in NSCLC patients and its relationship with clinicopathological variables

In this study, NSCLC patients with values less than 95% of HE4 values of the control group (7.26 ng/ml) were assigned to the low expression group (n = 26), whereas those with values ≥ 7.26 ng/ml were assigned to the high expression group (n = 74). As shown in Table 1, high HE4 expression was correlated with TNM stage (P = 0.003), lymph node metastases (P = 0.007), and distant metastases (P < 0.001). However, high HE4 expression was not associated with other clinicopathological factors of patients, including age, sex, and smoking history (all P > 0.05).

Expression of HE4 in serum samples in relation to prognosis of patients with NSCLC

As shown in Figure 2, patients with high serum HE4 level had a significantly lower 5-year OS rate (34.0% vs. 59.7%; P = 0.022) than those with low serum HE4 level. Table 2 showed multivariate analyses of clinical variables and serum HE4 levels for their prognostic influence on OS. In a multivariate Cox model, we found that HE4 expression was an independent poor prognostic factor for 5-year OS (hazards ratio [HR] = 3.654, 95% confidence interval [CI] = 2.753-11.981, P = 0.019) in NSCLC.

Table 2. Multivariate analyses for prognostic factors in patients with NSCLC

<table>
<thead>
<tr>
<th>Variables</th>
<th>HR</th>
<th>95% CI</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age</td>
<td>0.981</td>
<td>0.287-1.826</td>
<td>0.387</td>
</tr>
<tr>
<td>Sex</td>
<td>0.822</td>
<td>0.376-2.118</td>
<td>0.467</td>
</tr>
<tr>
<td>Smoking history</td>
<td>1.782</td>
<td>0.763-3.734</td>
<td>0.101</td>
</tr>
<tr>
<td>TNM stage</td>
<td>4.945</td>
<td>2.345-9.344</td>
<td>0.014</td>
</tr>
<tr>
<td>Lymph node involvement</td>
<td>4.563</td>
<td>3.643-10.332</td>
<td>0.007</td>
</tr>
<tr>
<td>Distant metastases</td>
<td>5.981</td>
<td>3.643-12.355</td>
<td>0.001</td>
</tr>
<tr>
<td>Serum HE4 expression level</td>
<td>3.654</td>
<td>2.753-11.981</td>
<td>0.019</td>
</tr>
</tbody>
</table>

HR = hazard ratio; CI = confidence interval; HE4 = human epididymis protein 4.
Discussion

The best prognostic system for overall survival in NSCLC is still the TNM staging system. We are now in an era where personalized medicine and targeted therapies may give new hope for this patient group [9]. Identification of novel molecular markers which can improve diagnosis and prognostic stratification and serve as possible therapeutic targets will be of great importance in the near future. Circulating biomarkers are a promising means of prognosis, as serum and plasma samples are easily obtainable. A number of novel markers for lung cancer have been identified in recent years, such as carcinoembryonic antigen (CEA), serum cytokeratin 19 fragment (CYFRA 21-1) and progastrin releasing peptide (pro-GRP), however, they are not satisfactory for monitoring the disease because of their relatively low sensitivity and specificity [10-12].

HE4 is one of the most intensively studied of the novel biomarkers, which was first described as an epididymis specific gene using northern blot analysis and in situ transcript hybridization [13, 14]. HE4 is a member of the WAP domain family, and this domain shows fifty well-conserved amino acid motifs. This protein has a variety of functions, such as antiproteinases, SLPI, and elafin, which shows antibacterial activities and anti-inflammatory effects [15, 16]. In the previous reports, HE4 is overexpressed in ovarian cancer cells and secreted to the sera in the patients with ovarian cancer [17-19]. Moore and colleagues analyzed the serum samples for levels of CA125, soluble mesothelin-related peptide, HE4, CA72-4, activin, inhibin, osteopontin, and epidermal growth factor. As a single tumor marker, HE4 had the highest sensitivity for detecting ovarian cancer and combined CA125 and HE4 was a more accurate predictor of malignancy than either alone [20]. On the other hand, HE4 is less frequently elevated in patients with benign gynecologic disease compared with CA125 [21]. In addition, Andersen et al. demonstrated that combining the symptom index with a serum HE4 or CA125 test could improve the prediction of ovarian cancer in a prospective case-control study [22].

Previously, Iwahori et al. found that serum HE4 levels were elevated in lung cancer, and it might be a potential prognostic marker for lung cancer patients [8]. However, the prognostic significance of serum HE4 levels in human NSCLC among Chinese population has not been investigated. The purpose of this study was to evaluate the prognostic significance of serum HE4 level in patients with NSCLC among Chinese population. In the present study, we had three findings. Firstly, the serum HE4 level was significantly higher in patients with NSCLC than that of controls, and high HE4 expression was correlated with TNM stage, lymph node metastases, and distant metastases. Secondly, Kaplan-Meier method found that patients with high serum HE4 level had a significantly lower 5-year OS rate (34.0% vs. 59.7%; \( P = 0.022 \)) than those with low serum HE4 level. Thirdly, in a multivariate Cox model, we found that HE4 expression was an independent poor prognostic factor for 5-year OS in NSCLC. All the results suggested that serum HE4 level was befitting to predict prognosis of NSCLC patients.

In conclusion, our results indicated that serum HE4 was overexpressed in NSCLC and high expression of HE4 was associated with poor prognosis in patients with NSCLC. The data suggests that serum HE4 may be useful molecular markers in NSCLC among Chinese population.

Disclosure of conflict of interest

None.

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References


Serum epididymis and non-small cell lung cancer


