LUNG CANCER

Differential expression of VASP in normal lung tissue and lung adenocarcinomas

L Dertsiz, G Ozbilim, Y Kayisli, G A Gokhan, A Demircan and U A Kayisli


Background: Vasodilator stimulated phosphoprotein (VASP) is associated with focal adhesions and is thought to have an important role in actin filament assembly and cell motility. We hypothesise that an increase in the expression of VASP is involved in the progression and invasion of lung adenocarcinomas in parallel to tumour progression. A study was undertaken to analyse VASP expression in normal lung tissue and lung adenocarcinomas.

Methods: Human lung tissues with adenocarcinomas (n = 26) were used. Normal lung tissue specimens (n = 14) were taken from areas a standard distance (3 cm) from resected adenocarcinomas of patients who underwent surgical lung resection. Adenocarcinomas were classified according to pathological staging and histopathological grades. Tissues were stained for VASP using immunohistochemistry.

Results: Normal lung pneumocytes showed no VASP expression while alveolar macrophages had the strongest immunoreactivity for VASP. Bronchial epithelium (surface epithelium, goblet cells) and bronchial gland cells had a very weak immunoreactivity for VASP. Adenocarcinomas had significantly greater VASP expression than normal epithelium (p < 0.001). Moreover, VASP expression in adenocarcinomas increased significantly with more advanced tumour stage (p < 0.001).

Conclusions: The spatial and differential expression of VASP in normal lung tissue and lung adenocarcinomas suggests that it is likely to be involved in the differentiation of normal lung cells to adenocarcinomas. The significant increase in the expression of VASP in adenocarcinomas in parallel to pathological staging suggests that it may regulate the invasive behaviour of lung adenocarcinomas as adenocarcinoma invasion is increased in more advanced tumours.

Methods
Collection of tissues
Specimens of human lung tissue with adenocarcinomas (n = 26) were collected from surgically treated patients who gave their informed consent. Normal lung tissue samples (n = 14) were taken from areas a standard distance (3 cm) from resected adenocarcinomas of patients who underwent surgical lung resection. All tissue samples were collected during the 3 year period between October 2000 and May 2003. Normal lung tissue sections were histopathologically assessed for tumour cells and apparent pathological features and were only included in the study if they were morphologically normal. Consent forms and protocols were approved by Human Investigation Committees of Akdeniz University. Tissues were fixed in formalin fixative and embedded in paraffin for immunohistochemical analysis. They were then cut into 5 μm thick sections and mounted on poly-l-lysine coated slides. Immunohistochemical analysis was carried out.

Abbreviations: ECM, extracellular matrix; VASP, vasodilator stimulated phosphoprotein
using a standardised method based on the streptavidin-biotin technique described below.

None of the patients had undergone chemotherapy before surgery and all lung adenocarcinomas were the initial presentation. Pathological staging was performed according to the surgical resection materials and systematic mediastinal lymph node dissection: stage 1 (n = 7), stage 2 (n = 9), stage 3 (n = 8) and stage 4 (n = 2). The tumours were also graded histopathologically as well differentiated (grade I; n = 9), moderately differentiated (grade II; n = 7) and poorly differentiated (grade III; n = 10).

**Immunohistochemistry**

To detect VASP immunoreactivity a mouse monoclonal antibody (Alexis Platform, San Diego, CA, USA) was used. Following deparaffinization, tissues were rinsed twice in phosphate buffer saline (PBS) for 10 minutes. Endogenous peroxidase activity was quenched by 0.6 ml 3% hydrogen peroxide (H2O2) and 5.4 ml methanol for 10 minutes. Sections were then incubated with mouse anti-VASP (1:400 dilution) for 60 minutes at room temperature. Normal mouse antibody isotype was used as a negative control replacing the antibody with PBS. Following several PBS rinses, slides were incubated with streptavidin-peroxidase complex for 30 minutes (Vector Labs), then rinsed several times in PBS-T and incubated with DAB (Vector Labs) for 2 minutes. After slight staining with haematoxylin the slides were mounted using a permanent mounting medium.

**Statistical analysis**

Patients with pathological stages 1 and 2 tumours and those with stages 3 and 4 were combined for comparison. Semi-quantitative results of VASP were normally distributed as assessed by the Kolmogorov-Smirnov test. Analysis of variance (ANOVA) and the post hoc Tukey test for pairwise comparisons were used in statistical analysis. A p value of <0.05 was considered significant. Statistical calculations were performed using Sigmastat for Windows, Version 3.0 (Jandel Scientific Corporation, San Rafael, CA, USA).

**RESULTS**

Of the 26 patients with adenocarcinomas, 22 (84.6%) were men. The mean age of the patients was 59.7 years (range 39–79). Details of the clinical evaluation of the adenocarcinomas such as nodal or distant metastases are shown in table 1. Adenocarcinomas were classified as stage I–2 (n = 16) or stage 3–4 (n = 10) after clinical and pathological examination (table 1).

**VASP expression in normal lung tissue**

VASP expression was first analysed in normal lung tissue. No VASP expression was observed in pneumocytes (fig 1A) but alveolar macrophages showed moderate to strong immunoreactivity for VASP and the staining was mostly membranous (fig 1B). Vascular smooth muscle cells showed weak to moderate immunoreactivity for VASP and the staining model was in the cytosol and was homogenous (fig 1A, C).

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**Table 1 Characteristics of study patients**

<table>
<thead>
<tr>
<th>Patient no</th>
<th>Age/sex</th>
<th>pTNM*</th>
<th>Stage</th>
<th>Histological differentiation</th>
<th>Surgical intervention</th>
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<tr>
<td>1</td>
<td>53/F</td>
<td>T2N2M0</td>
<td>3a</td>
<td>Well</td>
<td>Right upper and middle lobectomies</td>
</tr>
<tr>
<td>2</td>
<td>55/M</td>
<td>T1N1M0</td>
<td>2b</td>
<td>Well</td>
<td>Right lower lobectomy</td>
</tr>
<tr>
<td>3</td>
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<td>T2N0M0</td>
<td>1b</td>
<td>Well</td>
<td>Left upper lobectomy</td>
</tr>
<tr>
<td>4</td>
<td>67/M</td>
<td>T2N0M0</td>
<td>1b</td>
<td>Well</td>
<td>Right upper lobectomy</td>
</tr>
<tr>
<td>5</td>
<td>79/M</td>
<td>T3N0M0</td>
<td>2b</td>
<td>Well</td>
<td>Left upper lobectomy</td>
</tr>
<tr>
<td>6</td>
<td>65/F</td>
<td>T1N0M0</td>
<td>1a</td>
<td>Well</td>
<td>Right lower lobectomy</td>
</tr>
<tr>
<td>7</td>
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<td>1a</td>
<td>Well</td>
<td>Right upper lobectomy</td>
</tr>
<tr>
<td>8</td>
<td>59/M</td>
<td>T2N1M0</td>
<td>2b</td>
<td>Well</td>
<td>Right upper and middle lobectomies</td>
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<tr>
<td>9</td>
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<td>Well</td>
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<tr>
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<td>Moderate</td>
<td>Right lower lobectomy</td>
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<tr>
<td>11</td>
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<td>Left lower lobectomy</td>
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<tr>
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<tr>
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<tr>
<td>18</td>
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<td>Poorly</td>
<td>Left lower lobectomy</td>
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<tr>
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<td>Right pneumonectomy</td>
</tr>
<tr>
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<td>3a</td>
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<td>Left lower lobectomy</td>
</tr>
<tr>
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<td>3a</td>
<td>Poorly</td>
<td>Right middle lobectomy</td>
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<tr>
<td>22</td>
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<td>2b</td>
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<td>Right upper lobectomy</td>
</tr>
<tr>
<td>23</td>
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<td>4**</td>
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<tr>
<td>24</td>
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<td>4**</td>
<td>Poorly</td>
<td>Right upper lobectomy</td>
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<tr>
<td>25</td>
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<td>T2N0M0</td>
<td>3a</td>
<td>Poorly</td>
<td>Left lower lobectomy</td>
</tr>
<tr>
<td>26</td>
<td>59/F</td>
<td>T2N2M0</td>
<td>3a</td>
<td>Poorly</td>
<td>Right middle lobectomy</td>
</tr>
</tbody>
</table>

*pTNM, pathological TNM.

**Patients with a solitary brain metastasis.**

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Reference:

1. Staining), 1+ (weak but detectable staining), 2+ (moderate or distinct staining). 3+ (intense staining). For each tissue a HSCORE value was derived by summing the percentages of cells stained at each intensity category and multiplying that value by the weighted intensity of the staining using the formula:

\[
\text{HSCORE} = \sum P_i \times (i + 1)
\]

where \(i\) represents the intensity scores and \(P_i\) is the corresponding percentage of the cells. Two slides were stained for each patient; in each slide three different areas and 500 cells in each area were evaluated under a microscope with 40x objective magnification. Two investigators (12% interobserver variability) who were blind to the slides determined the percentage of cells and the mean of their scores was used.

2. Immunohistochemistry was performed as described previously. The intensity of VASP immunoreactivity in lung tissues was semi-quantitatively evaluated as positively stained cells using the following intensity categories: – (no staining), 1+ (weak but detectable staining), 2+ (moderate or distinct staining), 3+ (intense staining). For each tissue a HSCORE value was derived by summing the percentages of cells stained at each intensity category and multiplying that value by the weighted intensity of the staining using the formula:

\[
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Bronchial gland cells and epithelial cells (surface epithelium, goblet cells) showed weak VASP immunoreactivity but some parabasal cells were moderately positive (fig 1D). All normal lung specimens had a similar immunohistochemical staining pattern for VASP.

**VASP expression in lung tissue with adenocarcinomas**

We first analysed the lung cells neighbouring the adenocarcinoma tissue on the same slide and found no difference in VASP expression from that of normal cells. Pneumocytes had no VASP expression while alveolar macrophages had moderate to strong immunoreactivity for VASP (fig 2A). No difference in VASP immunoreactivity was detected in alveolar macrophages or vascular smooth muscle cells between the groups (fig 2A).

The highest VASP immunoreactivity was detected in stage 3–4 adenocarcinoma tissue (fig 2B). Compared with normal bronchial epithelial cells, VASP expression was significantly higher in both stage 1–2 and stage 3–4 adenocarcinoma groups (fig 1D v fig 2B and C, p<0.001). Moreover, VASP expression in stage 3–4 adenocarcinomas was significantly higher than in stage 1–2 adenocarcinomas (fig 2B v fig 2C, p<0.001; fig 3).

Adenocarcinomas were also classified pathologically as well differentiated (grade I), moderately differentiated (grade II), and poorly differentiated (grade III). These differentiated cells generally showed moderate to strong immunoreactivity. Although there was a tendency for VASP immunoreactivity to increase from grade I to grade III, no statistically significant difference was found between the grades (p = 0.229, fig 4). However, the adenocarcinoma cells of each grade revealed a statistically significant increase in VASP expression compared with normal bronchial epithelial cells (p<0.001, fig 4).

**DISCUSSION**

The pathobiological characteristics of adenocarcinoma of the lung have not been completely delineated which makes it difficult to predict precisely the outcome after complete resection of the tumour. Many tumour related factors such as stage, histological type, pathological grade, and expression and activation status of p53, K-ras and c-erbB proteins have
Figure 2  Representative micrographs of human lung tissue with adenocarcinomas immunolabelled for VASP. (A) No VASP immunoreactivity is seen in pneumocytes (arrows) and very weak immunoreactivity was found in the epithelium (stars). The strongest immunoreactivity is seen in adenocarcinoma cells (arrowheads) where vascular smooth muscle cells (VMS) show weak to moderate immunoreactivity for VASP. (B) Considerably increased VASP immunoreactivity is seen in stage 3 adenocarcinomas (arrowheads) compared with normal bronchial epithelium (A) and stage 1 adenocarcinomas (C). Magnification: A ×50; B, C ×100.

Figure 3  Comparison of VASP immunoreactivity in normal bronchial epithelium, stage 1–2 adenocarcinomas, and stage 3–4 adenocarcinomas. Bronchial epithelium represents VASP immunoreactivity in normal lung tissue. All values are shown as mean (SD). *p<0.001.

Figure 4  Comparison of VASP immunoreactivity in normal bronchial epithelial cells and lung adenocarcinomas graded pathologically as grades I, II, and III. Bronchial epithelium represents VASP immunoreactivity in normal lung bronchial epithelial cells. No significant difference was found between the adenocarcinoma grades (p=0.229). All values are shown as mean (SD). *p<0.001.
been reported to be related to recurrence and survival of patients with adenocarcinoma. The cellular and molecular functions of VASP remain to be determined since it is a multifunctional protein which is associated with filamentous actin formation and is involved in the intracellular signalling pathway of the integrin-ECM interaction. It has therefore been suggested that it participates in cell adhesion and motility. As a membrane associated protein, VASP is widely distributed in the cytoplasm of migrating cells, especially at the leading edge of the cytoplasm. Moreover, this family of proteins is involved in focal adhesions.

Previous studies have suggested that the VASP protein family has a universal role in the control of cell motility and in the regulation of intracellular actin dynamics via a linear pathway from the receptor-ligand interaction in many systems. In this study, to our knowledge, we have shown for the first time that VASP has a spatial expression pattern in lung tissues depending on the cell type. In normal lung tissue VASP shows the strongest immunoreactivity in alveolar macrophages; these are migratory cells and it is possible that VASP may be involved in regulating the migration of these cells. The absence of VASP from pneumocytes and its very weak expression in normal bronchial epithelial cells—which are accepted as the main cell origin of adenocarcinomas—also support this suggestion since these cells are non-migratory cells of normal lung tissue. Furthermore, the presence of weak but membranous expression of VASP in the bronchial epithelium may suggest a role for VASP in cell-cell junctions where it modulates the formation of filamentous actin in cell-cell binding, since it has previously been found to bind to components of cadherin-catenin junctional complex and the transmission of signals at the cytoskeleton-membrane interface.

Localisation of VASP to the leading edge of a migrating cell can lead to the local accumulation of profilin which, in turn, can supply actin monomers to growing filament ends. VASP binds to the focal adhesion proteins vinculin and zyxin and this probably directs the phosphoprotein to focal adhesions and the leading edge of stimulated cells. Major therapeutic approaches in cancer have been based on inhibition of the ras signalling pathway with special emphasis on the MAPK arm. Transformation from benign to malignant cells can be affected by the expression of rho GTPases and ras effectors. In malignant epithelial cells inhibition of MAPK-rac signaling alters the expression and localisation of the actin regulating proteins vinculin and VASP which results in the loss of stable F-actin structures and the characteristics of actin based differentiation. The interaction between rac and VASP could therefore be an important target for cancer treatment.

Previous studies have suggested that VASP may have a role in trophoblast invasion, angiogenesis, cell proliferation, and tumorigenesis. In the present study we have shown that VASP expression is higher in adenocarcinomas than in normal bronchial epithelial cells and pneumocytes. Although we did not study the molecular mechanisms by which VASP could regulate tumorigenesis, we hypothesise that VASP may be involved in the transformation from normal bronchial epithelium and branchial gland cells to adenocarcinomas. Moreover, the fact that VASP immunoreactivity increases in parallel with tumour stage and that there is a statistically significant correlation between increasing VASP immunoreactivity and stage 3–4 adenocarcinomas suggests that the protein may assist in tissue invasion by adenocarcinomas. It has previously been shown that preventing VASP function or overproduction of VASP in naive NIH 3T3 fibroblasts results in neoplastic transformation, implying that normal cell growth may require the maintenance of VASP expression within a narrow range and suggesting a role for VASP in tumorigenesis and/or cancer progression.

There was no difference in VASP expression between adenocarcinomas of different grades, but a significant increase in VASP immunoreactivity was found with advancing adenocarcinoma stages from 1 to 4. This finding agrees with the results of previous studies implying that prognostic criteria in lung adenocarcinoma are related to the stage of the disease rather than the grade.

In conclusion, our results show for the first time the spatial and differential expression of VASP in normal lung tissue and in lung adenocarcinomas. A gradual increase in VASP expression in adenocarcinoma tissue in parallel to the increase in stage suggests that VASP may be involved in the invasive behaviour of lung adenocarcinomas, possibly by regulating intracellular F-actin formation, focal adhesion, and cell migration since adenocarcinoma invasion increases with increasing stage. Furthermore, we speculate that VASP may be involved in the differentiation of normal lung cells into lung adenocarcinomas, even though VASP expression is not related to the transformation from poorly to well differentiated adenocarcinoma tissue.

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This study was funded in part by Akdeniz University Scientific Research Project Unit.

REFERENCES
Clinical staging underestimates pathological stage in non-small cell lung cancer

Clinical staging of lung cancer should establish a cTNM reliably predictive of the pathological stage. This study analysed 2994 lung cancers, operated on with curative intent in Spain between 1993 and 1997, with the aim of determining the agreement between clinical and pathological staging.

93% of patients were male with a 57.5 pack-year smoking history. In 29% of the cases lung cancer was an incidental radiological finding. 98% of the cancers were non-small cell (59% squamous) and 80% underwent complete resection (55% lobectomy/bilobectomy, 32% pneumonectomy). Of the 2994 patients initially included in the series, 2606 had a clinical staging. 2710 were classified using the pathological staging, and a clinicopathological comparison was performed in 2377 cases (79%). The clinicopathological agreement for stages IA–IIIB was 47% (Kappa’s index 0.248), similar to that found in other studies (35–55%). The highest agreement (75%) was achieved for stages IA and IB (Kappa’s index 0.56) and the lowest for stages IIIB (22%) and IIA (8%). Clinical staging underestimated the pathological staging in 92% of stage IIA, 86% of IIB, 74% of IIB, and 15% of IB tumours.

Differences in staging protocols and in the characteristics of the population studied are likely to explain variability in the results between different studies. Future studies, using integrated positron emission tomography (PET)/CT scanning, are likely to result in a better agreement between clinical and pathological staging. This is important for treatment planning and the provision of accurate prognostic information.

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Thorax 2005 60: 576-581
doi: 10.1136/thx.2004.037622

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