Tumor-associated Antigen L6 and the Invasion of Human Lung Cancer Cells

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ABSTRACT

Metastasis is a coordinated process that depends on the interaction of cancer cells with the tumor microenvironment. Members of the transmembrane-4 superfamily (TM4SF) of surface proteins have been implicated in the regulation of cancer cell metastasis, and the expression of several TM4SF members on tumor cells is inversely correlated with patient prognosis. The tumor-associated antigen L6 (TAL6), a distant member of the TM4SF, is expressed on most epithelial cell carcinomas and is a target for antibody-mediated therapy. We examined whether TAL6 may play a role in cancer metastasis by using an established series of human lung carcinoma cell lines (CL1-0 to CL1-5) that exhibit increasing invasiveness in vitro and in vivo. We found that TAL6 expression correlated with the in vitro invasiveness of CL lung carcinoma cells ($r^2 = 0.98$) and human carcinoma cells ($r^2 = 0.69$). Forced expression of TAL6 on CL1-0 lung carcinoma cells significantly increased their in vitro invasiveness and decreased the survival of SCID mice in an experimental metastasis model. Specific antibody against TAL6 (monoclonal antibody L6) significantly reduced the migration and invasiveness of CL1-5 lung carcinoma cells. The effects of monoclonal antibody L6 on CL1-5 invasion required clustering of TAL6 on the cell surface. Real-time reverse transcription-PCR of lung cancer specimens showed that increased expression of TAL6 was significantly associated with early postoperative relapse ($P = 0.034$) and shorter survival ($P = 0.025$) in squamous cell lung cancer patients. Thus, TAL6 appears to be involved in cancer invasion and metastasis.

INTRODUCTION

Cancer metastasis is a complicated process involving a coordinated program of events that include changes in cell adhesion, polarized proteolysis and migration, intravasation into the circulation, subsequent adhesion to endothelial cells followed by extravasation, invasion, and induction of angiogenesis (1). Cell surface proteins and receptors are intimately involved in these processes. For example, loss of E-cadherin can reduce cell-cell adhesion and allow cancer cells to more readily escape tumors (2). Integrins play vital roles in regulating cell adhesion, motility, invasion, and angiogenesis (3–5), and metalloproteinases on tumor cells can degrade the ECM (6). Other surface proteins including receptor tyrosine kinases (7) and chemokine receptors (8) have been implicated in cancer metastasis. Although much has been learned about how surface proteins influence metastasis, the contribution of many receptors to the metastatic process remains poorly defined.

Tumor-associated antigen L6 is a distant member of the tetraspanin family (9) that was originally identified by mAb L6 (10). Members of the tetraspanin or TM4SF (5) of cell surface proteins possess four highly conserved hydrophobic transmembrane domains that form a small and large extracellular loop, a short cytoplasmic loop, and short NH₂ and COOH termini (11). TM4SF members such as CD9, CD63, CD82, and CD151 have been shown to regulate cancer cell motility and metastasis (12–15). The expression of tetraspanins has also been correlated with patient prognosis and survival (16–18).

TAL6 is expressed on most human lung, colon, breast, and ovarian tumors (10), and has generated interest as a target for antibody-mediated therapy (19–21). However, the biological function of TAL6 is largely unknown. TAL6 was found to be highly expressed on an invasive spermatogonia-derived cell line as compared with a noninvasive spermatocyte-derived cell line (22). We picked up TAL6 in a cDNA microarray screen for metastasis-associated genes (23), and TAL6 was reported recently to be involved in the migration of human keratinocytes (24). These findings and the strong association of several TM4SF proteins with cell motility suggested that TAL6 might also be involved in tumor cell metastasis. We used a human lung carcinoma model described previously that includes cell lines displaying different invasive and metastatic capabilities (CL1-0

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3 The abbreviations used are: mAb, monoclonal antibody; ECM, extracellular matrix; GAM, goat antimouse antibody; GAR, goat antirabbit antibody; MF, mean fluorescence; mAb L6, anti-TAL6 monoclonal antibody L6-20-4; RF, relative fluorescence; SCID, severe combined-immune deficiency; TM4SF, transmembrane-4 superfamily; RT-PCR, reverse transcription-PCR; TBP, TATA-box binding protein; $C_T$, threshold cycle.
and the sublines CL1-1 to CL1-5; Ref. 25) to study possible roles of TAL6 in lung cancer metastasis. We present data supporting the idea that TAL6 plays a role in cancer cell migration and invasion.

**MATERIALS AND METHODS**

**Cell Lines.** Human lung carcinoma cell lines with different invasive and metastatic capabilities (CL1-0 and its sublines CL1-1 to CL1-5) have been described (25). HT-1197 human bladder carcinoma (CRL-1473), Colo320 human colon carcinoma (CL-188), and NCI-H520 human lung carcinoma (HTB-182) cells were obtained from the American Type Culture Collection (Manassas, VA). H928 lung carcinoma cells were a gift from Dr. Ming-Yang Yeh, Cheng Hsin General Hospital, Taipei, Taiwan. NTU-B1 bladder carcinoma cells and SV40-immortalized normal human bronchial epithelial cells (BEAS-2B S-10; Ref. 26) were from Dr. Pan-Chyir Yang, Department of Internal Medicine, National Taiwan Medical School, Taipei, Taiwan. OVTW-59 ovarian carcinoma cells were generously provided by Dr. Chin-Tamg Lin, Department of Pathology, National Taiwan Medical School, and Dr. Pan-Chyr Yang, Department of Internal Medicine, National Taiwan Medical School, Taipei, Taiwan. OVTW-59 ovary carcinoma cells was obtained from the American Type Culture Collection (Bedford, MA) in PBS for invasion assays or with 100 µg/ml bovine collagen type I (invasion assay). In clustering experiments, CL1-5 cells were incubated with 0.5 mg/ml mAb OX8 or L6 for 1 h before 5 x 10⁵/mL of sample cells were added with the cells and after 24 h (invasion) or 6 h (migration) were counted as described previously (23).

**Antibodies.** Hybridomas secreting antibody against the rat CD8 α chain (OX8) was obtained from the European Collection of Cell Cultures (Wiltshire, United Kingdom). mAbs against CD63 (LP9), CD81 (JS64), CD9 (MM2/57), and integrin α2 (AK7) were purchased from Serotec Ltd. (Oxford, United Kingdom). A hybridoma-secreting antibody against a surface antigen on CL1-5 cells was a gift from Dr. M’iiss A. Hudson, Washington University School of Medicine, St. Louis, MO. All of the cells were cultured at 37°C, 5% CO₂ in RPMI 1640 supplemented with 10% bovine serum, 2.98 g/liter HEPES, 2 g/liter NaHCO₃, 100 units/ml penicillin, and 100 µg/ml streptomycin. All of the cells were free of Mycoplasma as determined by a PCR-based Mycoplasma detection kit (American Type Culture Collection). The effect of antibodies (20 µg/ml) on the growth of CL1-5 cells was determined by counting the cells in triplicate wells every 24 h. Doubling times were calculated by linear regression analysis.

**Flow Cytometric Assay.** Immunofluorescence staining of cells was performed as described (31). MF, measured on a FACScaliber flow cytometer (Becton Dickinson, Mountain View, CA), was calculated with FlowJo 3.2 (Tree Star, Inc., San Carlos, CA). RF was calculated as:

\[
RF(\%) = 100 * \frac{\text{MF}_{\text{S,L6}} - \text{MF}_{\text{S,C}}}{\text{MF}_{\text{CL1-5,L6}} - \text{MF}_{\text{CL1-5,C}}}
\]

where MF_{S,L6} and MF_{S,C} represent the MF of sample cells stained with mAb L6 or control antibody, respectively, and MF_{CL1-5,L6} or MF_{CL1-5,C} represent the MF of CL1-5 cells stained with mAb L6 and control mAb, respectively.

**In Vitro Migration and Invasion Assays.** Tumor cell migration and invasion were examined in a membrane invasion culture system. A polycarbonate membrane with 10-µm pores (Nucleopore Corp., Pleasanton, CA) was precoated with 100 µg/ml of human vitronectin (28). The reaction was terminated by addition of iodoacetamide to 2 mM. The flow-through from a protein-A affinity chromatography. L6 Fab fragments were produced by digesting mAb L6 (3 mg/ml) in 0.1 x Tris-HCl (pH 8.0), 2 mM EDTA, and 0.1 mM DTT with 0.03 mg/ml activated papain (Sigma Chemical Co., St. Louis, MO) for 3 h at 37°C. The reaction was terminated by addition of iodoacetamide to 2 mM. The flow-through from a protein-A affinity column was additionally purified on a HiLoad 16/60 Superdex 200 gelatin (Sigma Chemical Co.) in PBS for invasion assays or with 100 µg/ml gelatin (Sigma Chemical Co.) for migration assays. The cells that invaded through the coated membrane in 48 h (invasion) or 6 h (migration) were counted as described previously (23). In some experiments, the indicated concentrations of mAb L6 or control antibodies were added with the cells and after 24 h (invasion assay). In clustering experiments, CL1-5 cells were incubated with 0.5 µg/ml mAb OX8 or L6 for 1 h before 5 µg/ml second antibody (GAM or GAR) was added. The treatment was repeated after 24 h. Each experiment, performed in triplicate or quadruplicate, was repeated at least three times.

**Adhesion Assay.** EIA high protein binding flat-bottomed 96-well plates (Corning Inc., Corning, NY) were coated overnight at 4°C with 50 µl/well human fibronectin (60 µg/ml), human vitronectin (2 µg/ml), mouse laminin (16 µg/ml), bovine collagen type I (80 µg/ml), or human collagen type IV (80 µg/ml). All of the proteins were purchased from Sigma Chemical Co. Control plates were uncoated (0% binding) or coated with poly L-lysine (100% binding). The plates were washed and blocked (except for poly L-lysine-coated wells) with 5% BSA. CL1-0 (4 x 10^5/mL) or CL1-5 (6 x 10^5/mL) cells in RPMI 1640 containing 0.5% NuSerum were added to wells with 10 µg/ml human anti-TAL6 (L6-20-4), anti-influenza virus A nucleoprotein (HB65), and antihapten B virus surface antigen (H252B10) antibodies were obtained from the American Type Culture Collection. A hybridoma-secreting antibody against the rat CD8 α chain (OX8) was obtained from the American Cancer Tumor, which was free of poly L-lysine (100% binding). The plates were washed and antibody binding was measured at 405 nm after addition of horseradish peroxidase-conjugated streptavidin (1:1000; Jackson ImmunoResearch Laboratories, West Grove, PA) followed by 2,2'-azino-bis[3-ethylbenzthiazoline-6-sulfonic acid] substrate (28).

**TAL6 Cell Transfectants.** Human TAL6 cDNA was amplified by RT-PCR with primers P1, 5'-CCT AGG GAT CCA CCA TGT GCT ATG GGA AGT GTG CA-3' and P2, 5'-GGG TTG TCT AGA TTA GCA GTC ATA TTG CTG TTG GTG-3' from RNA isolated from CL1-5 cells as described (29). The PCR product was digested with BamHI and XbaI restriction enzymes (italicized) and subcloned into pdecf3 (generously provided by Dr. Jerome Langer, Robert Wood Johnson Medical School, Piscataway, NJ) under the control of the human polypeptide chain elongation factor 1α promoter (30). CL1-0 cells transfected with pdecf3 or pdecf3-L6 were selected in G418 (Calbiochem, San Diego, CA) to generate CL1-0/pdecf and CL1-0/L6 cells, respectively. CL1-0/L6 cells were sorted for high expression of TAL6 on a fluorescence-activated cell sorter to produce CL1-0/L6-H cells.

**MATERIALS AND METHODS**

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L6 or control antibody. The plates were briefly shaken, centrifuged at 500 × g, and then incubated at 37°C for 30 min. Adherent cells were fixated with 10% formalin and stained with 1% Toluidine blue in 10% formalin overnight. The cells were extensively washed, air dried, and then lysed with 2% SDS. The absorbance of the wells was measured at 600 nm in a microplate reader. Cell binding was calculated as:

\[
\text{Binding} \% = 100 \times \frac{A_S - A_B}{A_{PLL} - A_B}
\]

where \(A_S\) is the absorbance of sample wells, \(A_B\) is the absorbance of BSA-blocked wells, and \(A_{PLL}\) is the absorbance of poly L-lysine-coated wells. Each experiment, performed with 8–16 replicates, was repeated at least three times.

**Collagen Zymography.** CL1-0/pdef, CL1-0/L6, and CL1-5 cells were grown to confluence in six-well plates. The monolayers were washed three times with PBS and cultured overnight in 1 ml of serum-free RPMI 1640 supplemented with 5 μg/ml L6 or HB65. The culture supernatant was centrifuged at 1000 × g for 10 min to remove debris, and then 20 μl of the medium was mixed with 10 μl SDS PAGE sample buffer without reducing agent. The samples were electrophoresed in a 10.5% SDS-PAGE containing 1 mg/ml gelatin. The gel was washed in 50 mM Tris-HCl (pH 7.4) containing 2.5% Triton X-100 for 30 min. The gel was then incubated in activation buffer [50 mM Tris-HCl (pH 7.5), 150 mM NaCl, 10 mM CaCl₂, and 0.05% NaN₃] for 24 h at 37°C. The gel was subsequently stained with Coomassie blue and destained in water containing 10% methanol and 10% acetic acid.

**Patients and Specimens.** Fifty four patients who underwent surgery for non-small cell lung cancer at the National Taiwan University Hospital from September 1, 1994, through August 31, 1996, were included in the study. This investigation was performed after approval by the Institutional Review Board of the National Taiwan University Hospital. Written informed consent was obtained from all of the patients. None of the patients had received neoadjuvant chemotherapy or radiation therapy before surgery. Specimens of lung cancer tissue and adjacent normal lung tissue obtained at surgery were immediately snap-frozen in liquid nitrogen and stored at −80°C until use. WHO criteria (32) were used for histological classification. Tumor size, local invasion, and lymph node metastasis were determined at pathologic examination. The final disease stage was determined by a combination of surgical and pathologic findings, according to the current tumor-node-metastasis system for lung cancer staging (33). Among the 54 patients, 35 were men and 19 were women (mean age ± SD = 62.4 ± 10.6 years), 22 of whom had squamous cell carcinoma and 32 of whom had adenocarcinoma. The surgical-pathology stage of disease was stage I in 19 patients, stage II in 9, stage III in 20, and stage IV in 6. Tumor status was T₁ in 11 patients, T₂ in 28, T₃ in 5, and T₄ in 10. Twenty-five patients had no lymph node metastasis (N₀), and 29 had regional or mediastinal lymph node metastasis (N₁ in 8 patients and N₂ in 21 patients). Follow-up data were obtained from the patient medical charts and from our tumor registry service. Follow-up times ranged from 32 to 81.8 months. Relapse time was calculated from the date of surgery to the date of detection of local recurrence or systemic metastasis. Survival time was calculated from the date of surgery to the date of death. Patients who died of postoperative complications after surgery were excluded from the survival analysis.

**Real-Time PCR.** Total RNA was extracted from resected cancer tissue with an RNA extraction kit (RNeasy Mini Kit; Qiagen, Valencia, CA). The quality of RNA in samples was determined by electrophoresis through agarose gels and staining with ethidium bromide; 18S and 28S RNA bands were visualized with UV illumination. The samples used for the standard curve in the real-time quantitative RT-PCR were prepared by serial dilution to contain 250, 50, 10, and 2 ng of specific RNA. The serially diluted samples were distributed into aliquots and stored at −80°C until use. The primers used for L6 were as follows (34): forward primer = 5'-cgttgtgtctttcttgga-3'; reverse primer = 5'-ccagcccaagaagacat-3'. The sequence of the probe used to detect and quantify the RT-PCR product was 5'-ggtctgtagctctctctgc-3'. The probe was labeled at the 5' end with carboxyfluorescein and at the 3' end with N,N,N',N'-tetramethyl-6-carboxyrhodamine. The primers and probe used for quantitative RT-PCR of the TBP mRNA (internal control, GenBank accession no. X54993) were as described by Bieche et al. (35). The identities of PCR products were confirmed by DNA sequencing. Each amplification mixture (50 μl) contained 50 ng of sample RNA; 5× TaqMan EZ buffer (10 μl; PerkinElmer, Foster City, CA); 25 mM manganese acetate (6 μl); 300 μM AMPP deoxycytosine triphosphate, and dGTP; 600 μM dUTP; 5 units of Tth DNA polymerase; 0.5 units of AmpErase uracil-N-glycosylase; 200 nM forward and reverse primer; and 100 nM dual-labeled fluorogenic probe (Perkin-Elmer). The Tth DNA polymerase has both reverse transcriptase and Taq polymerase activity. For reverse transcription, the mixtures were incubated at 50°C for 2 min, 60°C for 30 min, and 95°C for 5 min for deactivation. The subsequent thermal cycling profile consisted of 40 cycles of denaturation at 94°C for 20 s, and primer annealing and extension at 62°C for 1 min. Each assay included a standard curve, a no-template control, and triplicate total RNA samples. The fluorescence emitted by the reporter dye was detected on-line in real time with the ABI prism 7700 sequence detection system (PE Applied Biosystems). The CT is the fractional cycle number at which the fluorescence generated by cleavage of the probe exceeds a fixed threshold above baseline. For a chosen threshold, a smaller starting copy number results in a higher CT value. In this study, we used TBP mRNA as an internal control (35). The relative amount of tissue L6 mRNA, standardized against the amount of TBP mRNA, was expressed as \(\Delta C_T = -[C_{\text{TLPd}} - C_{\text{TBP}}]\). The ratio of the number of L6 mRNA copies to the number of TBP mRNA copies was then calculated as \(2^{-\Delta C_T} \times K\), where K is a constant (36).

**Experimental Metastasis.** SCID mice obtained from the National Taiwan University were i.v. injected with 3 × 10⁶ CL1-0, CL1-5, or CL1-0/L6-H cells in PBS. Mice survival was followed for 80 days. All of the animal experiments were carried out with prior approval by the Animal Committee of the Institute of Biomedical Sciences. The ethical guidelines that were followed met the standards required by the Interdisciplinary Principles and Guidelines for the Use of Animals in Research.
Statistical Analysis. Statistical significance of differences between mean values was estimated with Excel (Microsoft, Redmond, WA) using the independent t test for unequal variances. Statistical analysis of clinical data was performed using SPSS for Windows software (version 10.0; SPSS Inc., Chicago, IL). Fisher’s exact tests and Student’s t tests were used to compare the clinicopathologic characteristics of tumors (and patients) with high and low expression of L6 mRNA. Survival curves were obtained by the Kaplan-Meier method, and the difference in survival and relapse times between groups with low and high expression of L6 antigen was analyzed with the log-rank test. P values less than 0.05 were considered to be statistically significant.

RESULTS

Correlation between TAL6 Expression and Cell Invasiveness. Fig. 1A shows that mAb L6 against human TAL6 specifically stained immortalized normal human bronchial epithelial cells (filled curve), CL1-0 (-----), and CL1-5 cells (----) were stained with mAb L6 or control mAb OX8 (shaded curves) and analyzed by flow cytometry. B, the invasion of CL lung carcinoma cells through membranes coated with Matrigel relative to the invasion of CL1-5 cells are shown. Bars, ±SE. C, the RF of CL cell lines after staining with mAb L6 or control mAb was calculated as described in “Materials and Methods.” D, linear regression analysis of the invasiveness of CL cell lines versus the RF of TAL6.

The correlation between TAL6 expression and malignancy. To additionally explore the relation between invasion and expression of TAL6, we quantified the invasiveness of five human lung carcinoma cell lines, CL1-0, CL1-1, CL1-2, CL1-3, and CL1-5 (Fig. 1B). The numbers of cells invading through a membrane coated with Matrigel were: CL1-0, 431 ± 144; CL1-1, 287 ± 79; CL1-2, 1350 ± 175; CL1-3, 1390 ± 360; and CL1-5, 2350 ± 300, showing a general trend of increasing invasiveness. The RF of cell lines after staining with mAb L6 also displayed a trend of increasing TAL6 expression (Fig. 1C). Linear regression analysis revealed a strong correlation (correlation coefficient r² = 0.98) between invasion and expression of TAL6 on CL lung carcinoma cells (Fig. 1D).

The correlation between TAL6 expression and invasiveness was additionally explored in a panel of human carcinoma cell lines that included lung (CL1-0, CL1-5 NCI-H520, and H928), bladder (HT1197, NTU-B1, and EJ), colon (Colo320), and ovarian (OVTW-59) carcinoma cells. Fig. 2A shows that TAL6 RF values ranged from 0.6% on Colo320 cells to 108% on H928 cells. HT119 (98 ± 35 cells) and CL1-0 (460 ± 57 cells) showed the lowest and highest RF, respectively. B, the invasion of cell lines through membranes coated with Matrigel relative to the invasion of CL1-5 cells is shown. Bars, ±SE. C, linear regression analysis of the invasiveness of cell lines versus the RF of TAL6.
cells) cell lines displayed the lowest invasion, whereas CL-1-5 (4000 ± 232), H928 (4040 ± 420), and NTU-B1 (3700 ± 1600) cells were most invasive (Fig. 2B). Linear regression analysis revealed a trend (r² = 0.69) of increasing invasion as the level of TAL6 on the cells increased (Fig. 2C).

In contrast to the trend observed for TAL6 expression, many TM4SF members display a trend of decreased expression with increasing malignancy. To determine whether CL lung carcinoma cells atypically express TM4SF proteins, we performed immunofluorescence analysis of CL-1-0 and CL-1-5 cells with mAbs against the TM4SF members CD9, CD63, and CD81. Fig. 3 shows that slightly more CD9 was expressed on CL-1-0 cells as compared with CL-5 cells. Similarly, both CD63 and CD81 were more highly expressed on CL-1-0 cells than on CL-1-5 cells. As before, TAL6 was more highly expressed on CL-1-5 cells than on CL-1-0 cells. The expression of a control protein (integrin α2) was similar on both CL-1-0 and CL-1-5 cells. These results demonstrate that CL lung carcinoma cells display the typical pattern of tetraspanin expression that is inversely correlated with malignancy.

**Forced Expression of TAL6 Increases the Invasiveness of Lung Carcinoma Cells.** A direct role for TAL6 in cancer cell invasion was investigated by engineering CL-1-0 cells to express high levels of TAL6. CL-1-0 cells were generated that expressed vector (CL-1-0/pdef) or TAL6 (CL-1-0/L6-H cells). Transfected cells were also sorted for enhanced TAL6 expression (CL-1-5/L6-H cells). Populations of cells were used in all of the experiments to eliminate possible artifacts associated with cell clones. Fig. 4A shows that TAL6 expression was highest on CL-1-5 cells followed by CL-1-0/L6-H and then CL-1-0/pdef cells. CL-1-0/L6 cells displayed TAL6 densities intermediate to CL-1-0/pdef and CL-1-0/L6-H (results not shown). The doubling times of CL-1-0/pdef, CL-1-0/L6-H, and CL-1-5 cells were similar (17.5, 17.9, and 18.0 h, respectively), demonstrating that forced expression of TAL6 did not affect cell viability or growth rate (Fig. 4B). Comparison of the adherence of CL-1-0/pdef, CL-1-0/L6-H, and CL-1-5 cells to purified ECM proteins revealed that overexpression of TAL6 significantly (P ≤ 0.05) increased CL-1-0/L6-H cell adhesion to collagen type IV (Fig. 4C). However, this minor effect is of questionable biological significance. CL-1-5 and CL-1-0 cells displayed similar binding to fibronectin, laminin, and vitronectin, but significantly more CL-1-5 cells adhered to collagen type I and collagen type IV. Interestingly, CL-1-5 cells displayed significantly lower adhesion than CL-1-0 cells to Matrigel, although collagen type IV is a major component of Matrigel. Fig. 4D illustrates that the invasiveness of CL-1-0/L6 (225% ± 18%) and CL-1-0/L6-H (257% ± 31%) cells was significantly (P ≤ 0.0005) higher than CL-1-0/pdef cells (100% ± 4.1%). Investigation of ECM degradation showed that culture supernatant from CL-1-5 cells possessed high collagenase activity as compared with CL-1-0 cells (Fig. 4E). However, expression of TAL6 did not appear to be related to collagenase activity; neither addition of L6 antibody to cells nor expression of TAL6 on CL-1-0 cells produced visible changes in collagenase activity. The in vivo malignancy of CL-1-0/L6-H cells was also examined in an experimental metastasis model. Previous studies have demonstrated that i.v. injection of CL-1-5 cells in SCID mice results in progressive growth of lung-tumor colonies with typical adenocarcinoma tumor morphology (25). Fig. 4F shows that mice i.v. injected with CL-1-0/L6-H cells exhibited a significant (P ≤ 0.05) reduction in mean survival time (43.3 ± 3.0 days) compared with mice that were injected with CL-1-0/pdef cells (57.8 ± 4.7 days). The mean survival time of mice injected with CL-1-5 cells was additionally reduced (36.6 ± 4.0 days). There was no significant difference in the mean survival times of mice injected with CL-1-0/L6-H or CL-1-5 cells. Taken together, these results suggest that high expression of TAL6 can increase both in vitro and in vivo cell invasiveness.

**Anti-TAL6 Antibody Decreases Cancer Cell Invasion.** Addition of mAb L6 to CL-1-5 cells in the upper chamber of a membrane invasion chamber significantly (P ≤ 0.0005) reduced their invasion by 41% as compared with cells treated with control mAb. mAb L6 also significantly (P < 0.05) reduced the migration of CL-1-5 cells through a gelatin-coated membrane by ~15% as compared with control antibody (Fig. 5B). Suppression of cell invasion and migration by mAb L6 was not caused by inhibition of cell growth, because the doubling time of CL-1-5 cells treated with L6 (20.9 h) was similar to cells treated with Ox8, a nonbinding control antibody (22.8 h), or mAb Ly-5D7, an antibody that binds to an unidentified surface antigen on CL-1-5 cells (23.8 h; Fig. 5C). In agreement with high TAL6 cell experiments, mAb L6 did not significantly alter the adhesion of CL-1-5 cells to fibronectin, laminin, collagen type I, collagen type IV, or vitronectin (Fig. 5D).

To additionally investigate the mechanism of antibody modulation of cell invasion, the effect of antibody dose on the invasion of CL-1-5 cells was examined. Fig. 6A shows that the invasion of CL-1-5 cells was inhibited by mAb L6 in a dose-dependent fashion. CL-1-5 invasion was inhibited significantly by concentrations of mAb L6 >2.5 μg/ml. However, an inter-
mediate concentration of mAb L6 (10 μg/ml) inhibited the invasion of CL1-5 cell significantly (P ≤ 0.005) more than did a high concentration (50 μg/ml) of antibody. This result suggested that clustering of TAL6 on cells was required to inhibit cell invasion, because monovalent binding is favored at high antibody concentrations. To directly test whether TAL6 clustering was required to suppress invasion, we prepared monovalent Fab fragments of mAb L6. Purification of the Fab fragments to remove bivalent antibodies resulted in a single band on SDS PAGE under reducing conditions corresponding to the light chain, and variable and CH1 regions of the heavy chain (Fig. 6B). The L6 Fab fragments retained antigen-binding activity as demonstrated by their competition of biotinylated mAb L6 binding to CL1-5 cells (Fig. 6C). L6 Fab fragments did not significantly (P > 0.35) reduce the invasiveness of CL1-5 cells, confirming that TAL6 bridging was required to reduce cancer cell invasiveness (Fig. 6D). We also examined whether clustering of TAL6 could enhance suboptimal antibody stimulation of cells. Fig. 6E shows that CL1-5 cells incubated with a low concentration of L6 (0.5 μg/ml) and a clustering second antibody (L6 + GAM) displayed significantly (P ≤ 0.05) reduced invasion compared with a nonclustering second antibody (L6 + GAR). The GAM second antibody did not directly inhibit CL1-5 invasion as shown by addition of control antibody with second antibody (OX8 + GAM). Taken together, these results demonstrate that clustering of TAL6 on the cell surface modulated cancer cell invasiveness.

Association of TAL6 mRNA Expression with Postoperative Relapse and Survival of Patients with Lung Cancer. Real-time quantitative RT-PCR was used to measure the number of TAL6 transcripts in lung cancer tissue from 54 patients with lung cancer. Tumor samples from lung cancer patients were arbitrarily classified as having either high or low TAL6 expression with a cutoff value taken as the mean TAL6 expression level from all of the patients. No relationship was seen between TAL6 expression and clinicopathologic characteristics, including age, sex, stage of disease, lymph node status, and tumor histology (Table 1). The median survival of low-expression patients (50.1 months) was longer than the high-expression patients (33.2 months), but the difference was not significant by the log-rank test (P = 0.234; Fig. 7A). Comparison of the limited number of squamous cell carcinoma patients revealed
enhanced malignancy in an experimental metastasis model in decreased invasiveness through Matrigel-coated membranes and high levels of TAL6. These cells displayed significantly demonstrated in CL1-0 cells that were engineered to express derived tumors. A direct role for TAL6 in cancer metastasis was TAL6 also correlated with the invasiveness of several epithelial-derived lung carcinoma cells correlated with the invasive capabilities of these cells. The expression of TAL6 on CL1-5 cells correlated with the notion that TAL6 plays an important role in cancer metastasis. Here we present several lines of evidence that implicate TAL6 in cancer metastasis. We found that expression of TAL6 on CL lung carcinoma cells correlated with the invasive capabilities of these cells. The expression of TAL6 also correlated with the invasiveness of several epithelial-derived tumors. A direct role for TAL6 in cancer metastasis was demonstrated in CL1-0 cells that were engineered to express high levels of TAL6. These cells displayed significantly increased invasiveness through Matrigel-coated membranes and enhanced malignancy in an experimental metastasis model in mice. Conversely, clustering TAL6 molecules on CL1-5 cells with mAb L6 modulated cell migration and invasion. Finally, increased expression of TAL6 was associated significantly with early postoperative relapse and shorter survival in lung squamous cell carcinoma patients. Taken together, our results support the notion that TAL6 plays an important role in cancer invasion and metastasis.

We used a panel of lung carcinoma cells to explore possible relationships between TAL6 expression and cancer cell metastasis. This cell model was developed from a poorly differentiated lung carcinoma cell line (CL1-0) by selecting subpopulations of cells that displayed enhanced invasiveness in a transwell invasion chamber (25). The five sublines (CL1-0 to CL1-5) not only displayed progressively increasing invasive-
ness in vitro, but also exhibited a similar trend of greater metastatic potential in SCID mice (25) and enhanced invasiveness in a tracheal graft assay (23). We also showed that CL1-5 cells were more malignant than CL1-0 cells in a SCID mouse experimental metastasis model (Fig. 4F). Therefore, the CL1-0/CL1-5 cell model captures relevant biological characteristics for studying the metastatic behavior of lung cancer cells. TAL6 is a distant relative of the TM4SF that includes at least 25 members (39). This superfamily is distinguished by the presence of four hydrophobic transmembrane domains oriented such that the NH\(_2\) end and COOH terminus are located intracellularly, and two loops are located extracellularly. TM4SF proteins exhibit promiscuous association with lineage-specific surface proteins, integrins, and other TM4SF members. In fact, many of the biological activities of TM4SF proteins are attributed to their association with integrins (40, 41). TM4SF proteins are thought to link integrins with cellular kinases and signaling molecules, thereby promoting lamellipodial extension and retraction (42). Tetraspanins have been nicknamed “molecular facilitators” to highlight their propensity to group specific cell-surface proteins, and enhance the formation and activity of signaling complexes (11).

TAL6 differs from most other TM4SF members in that it primarily influenced invasion, whereas most tetraspanins primarily act by modulating cell motility (39). The divergence of TAL6 function is also highlighted by the positive correlation of TAL6 expression with cancer cell invasion and poor patient prognosis. In contrast, the expression of most TM4SF members is inversely correlated with cancer cell metastasis and patient prognosis (16–18). Interestingly, PETA-3 (CD151) expression has also been positively correlated with cancer cell migration and metastasis (15). However, detailed analysis of conserved regions in TM4SF members indicates that TAL6 may belong to a new four-transmembrane superfamily comprising TAL6, IL-TMP, TM4SF5, and L6D (43).

Table 1 Clinicopathologic characteristics of tumors with low and high expression of tumor-associated L6 antigen mRNA

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>&lt;6.35</th>
<th>&gt;6.35</th>
<th>P</th>
</tr>
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<tbody>
<tr>
<td>Age, y, mean ± SD</td>
<td>61 ± 11</td>
<td>64 ± 10</td>
<td>0.286&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Sex, no. of patients</td>
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<td></td>
<td></td>
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<tr>
<td>Male</td>
<td>15</td>
<td>20</td>
<td>0.642</td>
</tr>
<tr>
<td>Female</td>
<td>10</td>
<td>9</td>
<td></td>
</tr>
<tr>
<td>Stage, no. of patients</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>I–II</td>
<td>15</td>
<td>13</td>
<td>0.290</td>
</tr>
<tr>
<td>III–IV</td>
<td>10</td>
<td>16</td>
<td></td>
</tr>
<tr>
<td>Tumor status, no. of patients</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>T&lt;sub&gt;1&lt;/sub&gt;–T&lt;sub&gt;2&lt;/sub&gt;</td>
<td>19</td>
<td>20</td>
<td>0.762</td>
</tr>
<tr>
<td>T&lt;sub&gt;3&lt;/sub&gt;–T&lt;sub&gt;4&lt;/sub&gt;</td>
<td>6</td>
<td>9</td>
<td></td>
</tr>
<tr>
<td>Lymph node status, no. of patients</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>N&lt;sub&gt;0&lt;/sub&gt;</td>
<td>12</td>
<td>13</td>
<td>1.00</td>
</tr>
<tr>
<td>N&lt;sub&gt;1&lt;/sub&gt;–N&lt;sub&gt;3&lt;/sub&gt;</td>
<td>13</td>
<td>16</td>
<td></td>
</tr>
<tr>
<td>Histology, no. of patients</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Squamous cell carcinoma</td>
<td>9</td>
<td>13</td>
<td>0.585</td>
</tr>
<tr>
<td>Adenocarcinoma</td>
<td>16</td>
<td>16</td>
<td></td>
</tr>
</tbody>
</table>

<sup>a</sup> Derived with Student’s t test; other P values were derived with Fisher’s exact test. All statistical tests were two sided.

<sup>b</sup> Tumor stage, tumor status, and lymph node status were classified according to the International System for Staging Lung Cancer (33).

Therefore, TAL6 may exert biological effects that are distinct from the “classical” tetraspanins.

One possible effector of TAL6 function is aminopeptidase N (CD13), which associates with TAL6 on the surface of human lung cancer cells. CD13 is a type II transmembrane Zn\(^{2+}\)-dependent metalloproteinase that catalyzes the cleavage of neutral amino acids from the NH\(_2\) terminus of peptides (44). CD13 expression has been correlated with the invasion of melanoma (45) and prostate carcinoma (46). Although CD13 has been implicated in the activation of collagenase IV (47), we did not find any evidence that TAL6 modulated the collagenase activity of cancer cells (Fig. 4E). Antibody-mediated clustering of TAL6 modulated the invasiveness of CL1-5 cells, suggesting that TAL6 can initiate cell signaling. The PDZ protein SITAC/syntenin 2α was shown recently to bind the peptide sequence Y-X-C-COOH at the COOH terminus of TAL6 (48). The syntenin-binding motif is conserved among tetraspanins.

Fig. 7 Kaplan-Meier survival plots for patients with non-small cell lung cancer, grouped according to tumor-associated antigen TAL6 mRNA expression. The relative amount of tissue L6 mRNA, standardized against the amount of TATA-box-binding protein mRNA, was expressed as \(-\Delta CT = -[CT(L6) - CT(TBP)]\). Patients were included in the high-expression group when the \(-\Delta CT\) value was \(≥6.35\). A, difference in overall survival between high (n = 26) and low (n = 19) TAL6 expression groups among lung cancer patients was not significant (P = 0.23). B, the difference in overall survival between the high (n = 11) and low (n = 5) TAL6 expression groups among the limited number of patients with squamous cell carcinoma is statistically significant (P = 0.025). All of the patients alive at their last follow-up are indicated by tick marks on the plot.
human, hamster, and murine TAL6, suggesting a possible role for this interaction in TAL6 function.

Taken together, our results suggest that TAL6 plays a role in cancer cell invasion. Given the broad distribution of TAL6 on most epithelial-derived cancer cells, dissecting the molecular mechanisms of TAL6 modulation of cell invasion may provide new molecular targets for intervention or diagnosis of tumor cell metastasis.

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