Differential expression and distribution of epithelial adhesion molecules in non-small cell lung cancer and normal bronchus

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Non-small cell lung cancer (NSCLC), which constitutes 80% of all lung cancers, encompasses several distinct subtypes, including the most common types of adenocarcinoma and squamous cell carcinoma (SCC). Gene expression profiles may help to better understand which pathogenetic mechanisms are associated with the development of lung cancer. Gene expression profiling has been applied to identify genes that are related to specific histological subtypes of lung cancer.^{1,2} In addition, several studies have focused on the relationship of gene expression profiles with clinical characteristics such as prediction of survival,^{1,3–5} chemosensitivity^{6,7} and metastatic behaviour,^{8,9} or to cellular processes such as apoptosis¹⁰ and cell cycle.¹¹

Our aim was to compare gene expression profiles in patients with adenocarcinoma and SCC in comparison with normal bronchus obtained from the same patient. We have focused on expression of epithelial cell–cell and cell–matrix adhesion molecules, as they are known to correlate with tumour progression. We validated the expression levels and investigated the distribution of protein and mRNA from most prominently expressed epithelium-related adhesion genes by immunohistochemistry (IHC) and RNA in situ hybridisation (ISH).

METHODS

Tissue specimens

Tumour and normal bronchus tissue were obtained from well-defined patients without chemotherapy and a smoking history of at least 45 packyears (table 1¹²). Tissues were obtained from four patients with adenocarcinoma and four patients with SCC during surgery. The tissue was partially snap frozen for cDNA array and IHC, and partially fixed and embedded in paraffin wax for ISH. The tissues were stained with haematoxylin and

Background: Changes in epithelial cell interactions have been implicated in carcinogenesis, tumour invasion and metastasis.

Aim: To screen for altered expression of epithelial adhesion genes in lung cancer development.

Methods: Gene expression profiles were assessed with cDNA expression arrays in eight non-small cell lung cancer (NSCLC) and eight normal bronchi obtained from the same patient. Immunohistochemistry (IHC) and RNA in situ hybridisation (ISH) were used to confirm the most prominently expressed adhesion molecules and to investigate their distribution at protein and mRNA levels.

Results: 43 differentially expressed cancer-related genes were identified in adenocarcinoma, squamous cell carcinoma (SCC) and normal bronchus. Five of these genes are related to epithelial adhesion—that is, integrin $\alpha 3$ (ITGA3), integrin $\beta 4$ (ITGB4), desmoplakin I and II (DSP), plakoglobin, and desmocollin 3 (DSC3). ITGA3 and ITGB4, showing predominantly cell–matrix staining, were up regulated in adenocarcinoma and SCC, respectively. ITGB4 also showed strong staining in SCC with IHC and ISH. Components of the desmosome adhesion complex DSP, plakoglobin and DSC3 were strongly up regulated in SCC and showed a distinct cell–cell staining pattern. DSP and plakoglobin were predominantly present at central, more differentiated tumour cells, whereas DSC3 showed a stronger staining in the peripheral basal cells of SCC tumour areas.

Conclusions: Lack of cellular adhesion may have an important role in the metastatic potency of a primary tumour. A possible association of strong presence and normal-distributed desmosomal molecules in SCC with the less frequent and late pattern of metastasis in SCC as compared with adenocarcinoma is suggested.

eosin for histological confirmation of the diagnosis. Patients were only included when the bronchus showed normal bronchus epithelium without metaplasia or dysplasia. The study was approved by the local medical ethics committee of the University Medical Center Groningen (Groningen, The Netherlands) and written informed consent was obtained from all patients.

cDNA expression array

Total RNA was isolated from frozen tissue sections using Strataprep total RNA miniprep kit (Stratagene, La Jolla, California, USA), according to the manufacturer's instructions, including DNase treatment.

Atlas Human Cancer 1.2 Arrays (Clontech, Palo Alto, California, USA) were hybridised according to the manufacturer's instructions. Briefly, 2–5 μ g of RNA was converted into cDNA and labelled with [α -³²P]dATP (Amersham Biosciences, 3000 Ci/mmol, 10 mCi/ml, Piscataway, New Jersey, USA), purified and hybridised overnight at 68°C. Filters were washed and exposed to a phosphorimaging screen for 1–4 days and scanned with Storm (Amersham Biosciences).

Data analysis

Images were analysed using Atlas Image software V.2.0 (Clontech). Natural logarithm (ln) was taken to improve a normal distribution of data. Data of the hybridisations were standardised to mean = 0 and standard deviation (SD) = 1. Subsequently, the principal components were calculated and

Abbreviations: DSC3, desmocollin 3; DSP, desmoplakin I and II; IHC, immunohistochemistry; ISH, in situ hybridisation; ITGA3, integrin $\alpha 3$; ITGB4, integrin $\beta 4$; NSCLC, non-small cell lung cancer; PCR, polymerase chain reaction; SCC, squamous cell carcinoma

Table 1 Description of clinical features of patients with adenocarcinoma (AC) and squamous cell carcinoma (SCC), without prior treatment

Patient no	Description (age, sex)	Location	Diagnosis	Packyears
1	70 years, M	LUL	AC, T1N1M0 Moderately differentiated	55
2	68 years, F	RLL	AC, T2N1M0 Moderately differentiated	53
3	71 years, M	RUL	AC, T2N2M0 Poorly differentiated	51
4	74 years, F	RUL	AC, T2N0M0 Poorly differentiated	56
5	75 years, M	LUL	SCC, T1N0M0 Moderately differentiated	62
6	72 years, M	LL	SCC, T2N1M0 Poorly differentiated	52
7	66 years, M	LUL	SCC, T3N0M0 Well differentiated	50
8	73 years, M	LUL	SCC, T2N1M0 Poorly differentiated	49

F, female; LL, left lung; LUL, left upper lobe; M, male; RLL, right lower lobe; RUL, right upper lobe.
Tumour-node-metastasis staging according to Mountain *et al.*¹²

the first principal component was deducted. Two independent selection criteria were applied to identify differentially expressed genes (fig 1).

Immunohistochemistry (IHC)

Mouse monoclonal antibodies directed against ITGA3 (1:1000, Telios, San Diego, California, USA), ITGB4 (1: 500 Telios), DSP (1:200, Chemicon, Temecula, California, USA), plakoglobin (γ -catenin, 1:1000, BD-Biosciences, San Jose, California, USA) and DSC3 (undiluted, Research Diagnostics, Flanders, New Jersey, USA) were used to stain the tissue sections according to standard laboratory procedures. Frozen sections were fixed in acetone for 10 min and incubated with the primary monoclonal antibodies for 1 h. Endogenous peroxidase activity was blocked with 0.075% hydrogen peroxide for 1 h. Positive staining was visualised after the second and third steps using peroxidase rabbit- α -mouse and goat- α -rabbit antibodies (Dako, Glostrup, Denmark) using 3-amino-9-ethyl-carbazole (AEC).

RNA in situ hybridisation

Total RNA of SCC was primed with oligo (dT) for cDNA synthesis according to the manufacturer's instructions (Invitrogen, Carlsbad, California, USA). Primers were selected from sequence files of the corresponding genes to obtain polymerase chain reaction (PCR) products of about 500 bp (table 2). After amplification, the PCR products were subcloned in the pCRII-TOPO vector (Invitrogen) according to the manufacturer's instructions. SP6 and T7 RNA polymerases were used to synthesise anti-sense and sense dioxigenin (DIG)-labelled RNA probes using the DIG RNA labelling kit (Sp6/T7; Roche, Basel, Switzerland). The sense probe was used as a negative control in the in situ hybridisation (ISH) experiments.

ISH was carried out on paraffin wax sections using standard laboratory protocols. Deparaffinised tissue sections were treated with 5, 10, 15 and 20 μ g/ml proteinase K (Roche) at 37°C for 1 h and washed with Tris-buffered saline. Slides were incubated with 1 ng/ μ l DIG-labelled probe in a hybridisation solution consisting of 5 $\times$ Denhardt's solution, 2 $\times$ SSC, 10% dextran, 30% formamide, 1 mg/ml t-RNA and 2 mg/ml fish sperm DNA overnight at 55°C. Slides were washed and treated for 30 min with 10 mg/ml RNase at 37°C. Anti-DIG-labelled alkaline phosphatase (Roche) in 0.1 M maleic acid buffer containing 0.15 M sodium chloride, 2% blocking buffer and 0.1% Triton X-100, was incubated on the slides for 1 h at room

temperature. Positive cells were visualised by overnight staining with nitroblue tetrazolium and BCIP (Roche) in TBS (pH 9.0) containing 50 mM magnesium chloride.

RESULTS

Gene expression in normal bronchus, adenocarcinoma and SCC

Comparison of tumour tissue with normal bronchus (c-1) and of adenocarcinoma with SCC (c-2) showed 43 differentially expressed genes, including 5 epithelium-related adhesion genes (fig 1). Two were integrins, ITGA3 being up regulated in adenocarcinoma in comparison with normal bronchus and SCC, and ITGB4 being up regulated in SCC compared with adenocarcinoma. The other three genes, all components of the desmosome adhesion complex, DSP, plakoglobin and DSC3, were up regulated in SCC in comparison with normal bronchus. Plakoglobin and DSC3 were also up regulated when compared with adenocarcinoma.

Distribution of integrins Immunohistochemistry

Strong cell-matrix and weak cell-cell staining of ITGA3-protein was observed in normal bronchial epithelium, and additionally in bronchial glands, muscular layer and small blood vessels (fig 2). In adenocarcinoma and SCC, strong cell-matrix and weak cell-cell staining of ITGA3 was observed at the borders of tumour areas. ITGB4 showed a staining pattern comparable with ITGA3 staining, except in the bronchial muscular layer, where no ITGB4 was detected. Additionally ITGB4 was more restricted to basal cell-matrix borders in the normal bronchus and in adenocarcinomas. In SCC tumour cells, ITGB4 staining was stronger compared with bronchial epithelium, whereas in adenocarcinoma tumour cells, staining was less prominent than in bronchial epithelium.

RNA-in situ hybridisation

ISH showed a positive staining for ITGA3 and ITGB4 in the basal cells of the normal bronchial epithelium (fig 3). Adenocarcinoma tumour cells and predominantly basal cells of SCC tumour areas showed a positive staining for ITGA3 and ITGB4. Overall, ISH showed a staining pattern comparable to IHC for ITGA3 and ITGB4.

Figure 1 Overview of differentially expressed genes in normal bronchus, adenocarcinoma (AC) and squamous cell carcinoma (SCC). Forty three gene names are listed at the right. The most differentially expressed genes are ranked per group from top to bottom. Selection criterion 1 (c-1) was based on identifying differentially expressed genes in tumour versus normal bronchus in each patient. Ratios were determined by subtracting the standardised ln values of normal bronchus tissue from those of the tumour tissue from the same patient (T:B ratio). A consistent T:B ratio of >absolute 1 (which means a 2.3-fold up regulation or down regulation) for each of the four individual samples per tumour type, was considered relevant. Selection criterion 2 (c-2) was based on identifying differentially expressed genes between AC and SCC by using the sum of four standardised ln values of all AC (AC-sum) or SCC (SCC-sum) samples. Differences were considered relevant when the AC-sum or SCC-sum is >absolute 3 and the difference between the sums is at least 4. The brightness of colour correlates with the degree of "normalised expression difference" as shown at the bottom of image, where green represents low and red represents high level of expression. Groups A–D represent the T:B ratios used for c-1, and groups E and F the tumour values used for c-2. (A) Genes up regulated in AC in comparison with normal bronchus (c-1) and SCC (c-2). (B) Genes down regulated in AC in comparison with normal bronchus (c-1). Two of the genes were also down regulated in SCC. (C) Genes up regulated in SCC in comparison with normal bronchus (c-1) and AC (c-2) except for two genes. (D) Genes down regulated in SCC in comparison with normal bronchus and in one case also in comparison with AC, whereas another case is down regulated in comparison with AC. (E) Genes with higher expression in AC in comparison with SCC. (F) Genes with higher expression in SCC in comparison with AC. c-1, selection criterion 1; c-2, selection criterion 2 (see Material and methods). ln, natural logarithm; ↑, up regulation; ↓, down regulation.

Distribution of desmosomal adhesion molecules Immunohistochemistry

In the normal bronchus, DSP showed weak staining in bronchial epithelial cells, at the basolateral borders of supra-basal cells and at the apical side of ciliated cells. Strong positive staining was observed in epithelial cells of bronchial glands (fig 2). In adenocarcinoma and SCC, positive cell–cell staining was clearly visible at the borders between tumour cells. The

strongest staining was shown in SCC, especially between tumour cells in the centre of SCC tumour areas.

Plakoglobin showed strong staining in the normal bronchus at apical and lateral borders of the basal bronchial epithelial cells and at cells of the bronchial glands. Adenocarcinoma and SCC showed strong tumour cell–cell staining. In SCC, cell–cell staining of tumour cells located in the centre of tumour areas was stronger than the surrounding basal tumour cells. Often,

Table 2 Primer sequences and Genbank accession number used for amplification of integrin $\alpha 3$, integrin $\beta 4$, desmoplakin I and II, plakoglobin and desmocollin 3

Gene	Forward primer 5'-3'	Reverse primer 5'-3'	Genbank reference	PCR product (bp)
Integrin $\alpha 3$	TCTGCCTAAGGTTGTCTACG	TTGGAAGCTAGCCTCAGCTTCTC	NM_002204	546
Integrin $\beta 4$	CTGTGAGATGGCCCAAGGAG	GGGTGCAAGGATGGAGTAGC	NM_000213	528
Desmoplakin I and II	CCTGCACAGGTGGCATCATC	CCGGAGCCGAAGACATGTTG	NM_004415	507
Plakoglobin	CAGGTGCGGTTCTCATCTG	AAGCACCTGGAGAGAGAAGC	NM_002230	531
Desmocollin 3	ACTGTGGAGCCTACCAAAAG	CCGGCAGGATTCCAAGGTCT	NM_024423	517

PCR, polymerase chain reaction.

plakoglobin also showed cytoplasmic staining in bronchial epithelial and tumour cells.

DSC3 showed weak staining in some normal bronchi, and if so, only at apical borders of basal bronchial epithelial cells. DSC3 was not detected in adenocarcinoma. By contrast, SCC showed strong cell–cell staining of tumour cells. The strongest staining was observed in the basal cells of SCC.

RNA-in situ hybridisation

ISH for DSP showed a strong staining in SCC especially in the central SCC tumour areas (fig 3). Plakoglobin showed a similar pattern in SCC. Plakoglobin staining in the normal bronchus and adenocarcinoma was less apparent compared with SCC. DSC3 staining was only visible in bronchial glands and not in bronchial epithelial cells of normal bronchus. Clear DSC3 staining was shown in SCC, with the strongest intensity in basal tumour cells, whereas no staining was observed in adenocarcinoma. Overall, ISH showed a pattern comparable to IHC for DSP, plakoglobin and DSC3.

DISCUSSION

Epithelial adhesion is a fundamental pathway in tissue architecture and epithelial differentiation. Changes in epithelial cell–cell and cell–matrix interactions have been implicated in carcinogenesis, tumour invasion and metastasis. Thus, there is a putative role of these adhesion genes in cancer susceptibility and metastasis. This study explored differential expression of cancer-related, inclusive adhesion-related, genes in NSCLC and normal bronchi from patients with a long smoking history. We identified 43 differentially expressed genes representing in part key regulators of biological functions including five epithelium-related adhesion genes, two subunits of integrin receptors, ITGA3 and ITGB4, and three genes involved in the desmosome complex, DSP, plakoglobin and DSC3. These genes were further investigated with IHC and RNA ISH and showed consistent results for all three techniques for three of the five genes (table 3).

The advantage of this study is that representative normal bronchus tissue were selected, containing the putative lung cancer precursor cells. We analysed both normal bronchus and NSCLC tissue from the same patient; moreover here, it is assumed that the greatest source of variation in gene expression profiles originates from interindividual biological variation.¹³ Another advantage of this approach is that genetic variance due to smoke exposure, which indeed can induce changes in the resulting gene expression profiles, will be excluded.^{14 15}

Integrins are a family of heterodimeric transmembrane glycoproteins, consisting of α and β subunits, which serve primarily as cell–matrix adhesion molecules but which can also be involved in cell–cell adhesion. Integrins have an important role in the regulation of cellular morphology, differentiation and proliferation. Of the seven existing integrins expressed on airway epithelial cells of healthy people, integrins $\alpha 3\beta 1$ and $\alpha 6\beta 4$ recognise components of the basement membrane

(laminins 5, 10 and 11) and function therefore as true cell–matrix adhesion molecules.¹⁶ Remarkably, integrin subunits $\alpha 3$ and $\beta 4$ were up regulated in our adenocarcinoma and SCC samples, respectively. On the basis of lower levels of integrin $\beta 1$ and $\alpha 6$, we could not access differential expression pattern for these two subunits. Differential expression for ITGA3 and ITGB4 were confirmed by IHC and ISH. This confirmation was most pronounced for ITGB4, which showed the strongest expression of both protein and mRNA in SCC, in contrast with the weakest expression in adenocarcinoma. ITGA3 and ITGB4 showed prominently cell–matrix staining in normal bronchus, adenocarcinoma and SCC. Previous immunohistochemical studies have shown comparable frequency, intensity and distribution of those integrins in normal bronchial epithelium and NSCLC.^{16–21}

Increasing evidence suggests modulation of desmosomes is an important step in the initiation and invasiveness of human malignancies. Desmosomes are adhesive intercellular junctions, made up of heterodimers of different isoforms of the transmembrane desmosomal cadherins, desmogleins and desmocollins, which are linked to cytokeratin intermediate filaments in epithelial cells through interactions with desmoplakins, plakoglobin and plakophilins (for a review, see Wheelock and Johnson²²). Little is known about the localisation of the desmosome components and their distribution in normal bronchial epithelium and NSCLC. So far, a tissue-dependent and differentiation-specific distribution of individual desmosomal components has only been described in squamous-like epithelial tissues, such as epidermis, hair follicles, mammary glands and oral mucosa,^{23–29} suggesting that desmosomal cadherins have a role in directing differentiation in epithelial tissues.^{26 30} This is the first study illustrating the distribution of differentially expressed desmosomal molecules in SCC.

We observed strong enhancement of RNA and protein expression in SCC for three desmosomal components: DSP, plakoglobin and DSC3. Normal bronchus and adenocarcinoma showed a relatively low or no expression of desmosome molecules. However, plakoglobin protein expression was observed in normal bronchus and adenocarcinoma at levels similar to SCC, which might be related to its role in adherens junction, attaching to actin filaments.³¹ Strongest expression of DSP and plakoglobin was observed in more differentiated cells centrally located in tumour areas, whereas DSC3 showed stronger expression in basal cells. Remarkably, this differentiation-specific distribution of the desmosomal molecules in SCC is very similar to the pattern in normal squamous epithelium of the skin. This strongly suggests a normal expression regulation of these molecules in malignant transformed SCC and may have implications for the biological behaviour of tumour cells. Lack of cellular adhesion has an important role in the metastatic potency of a primary tumour. Although additional studies are required to elucidate the exact relationship of desmosomal molecules with metastatic behaviour in lung cancer, the strong presence of these molecules in SCC and its

Figure 2 Immunohistochemistry on normal bronchus, adenocarcinoma (AC) and squamous cell carcinoma (SCC). Integrin $\alpha 3$ (ITGA3) and integrin $\beta 4$ (ITGB4) show predominantly cell-matrix staining. The desmosomal proteins desmoplakin I and II (DSP) and plakoglobin (PG) show cell-cell staining, with strongest staining between more central localised cells in SCC. Desmocollin 3 (DSC3) shows strong basal staining in SCC.

Figure 3 RNA in situ hybridisation staining of normal bronchus, adenocarcinoma (AC) and squamous cell carcinoma (SCC). Integrin $\alpha 3$ (ITGA3) and integrin $\beta 4$ (ITGB4) show a predominant staining in basal cells. The desmosomal genes desmoplakin I and II (DSP) and plakoglobin (PG) show evident staining of SCC tumour cells, most pronounced in central localised cells of the tumour areas in SCC. For desmocollin 3 (DSC3) strong basal staining was observed in SCC.

Table 3 Overview of the results of cDNA expression array (Array), immunohistochemistry (IHC) and RNA in situ hybridisation (ISH)

Gene	Tissue	Array	IHC	ISH
<i>Integrin $\alpha 3$</i>	Bronchus	w	s	s
	AC	s	s	s
	SCC	m	s	s
<i>Integrin $\beta 4^*$</i>	Bronchus	m	m	m
	AC	w	w	w
	SCC	s	s	s
<i>Desmoplakin I and II*</i>	Bronchus	w	w	w
	AC	m	m	w
	SCC	s	s	s
<i>Plakoglobin</i>	Bronchus	w	s	m
	AC	w	s	w
	SCC	s	s	s
<i>Desmocollin 3*</i>	Bronchus	w	w	w
	AC	w	w	w
	SCC	s	s	m

AC, adenocarcinoma; IHC, immunohistochemistry; ISH, in situ hybridisation; m, moderate expression or staining; s, strong expression or staining; SCC, squamous cell carcinoma; w, weak, or no expression or staining.

*Genes with consistent expression levels using the three different techniques.

apparently normal distribution may be associated with the less frequent and late metastasis pattern of SCC as compared with adenocarcinoma.

In summary, we reported a detailed description of distribution of differentially expressed adhesion molecules on protein and mRNA levels in NSCLC and normal bronchi. A remarkably high expression of desmosomal molecules was observed in SCC, with a specific central or peripheral localisation. This may signify a role of these molecules in tumour metastasis.

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