Downregulation of occludin affects the proliferation, apoptosis and metastatic properties of human lung carcinoma

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Abstract. Lung cancer is the most frequent and deadliest cancer in the world, especially in China. However, the molecular mechanisms involved in lung cancer remain unclear. Occludin (OCLN), one of the first identified tight junction proteins, has been revealed to be a necessary integral protein for TJ structure and function. In the present study, we investigated the role of occludin in lung tumorigenesis. We found that occludin protein expression was highly increased in human lung cancer patient samples. Western blotting results also revealed that occludin expression was different in several lung cancer cell lines, with the highest level in SPC-A1 cells. Moreover, occludin knockdown inhibited lung cancer cell proliferation in vitro and in vivo. In addition, occludin knockdown promoted the apoptosis of lung cancer cell lines and reduced the invasion ability. Mechanistically, the activity of key growth pathway AKT/PI3K was compromised after occludin knockdown. Expression of apoptosis-related proteins, BAX, caspase-3, caspase-9 and AIF, but not Bcl-2, were upregulated after silencing of occludin. Collectively, our findings for the first time identify the role of occludin as a tumor promoter and a pro-metastatic factor in lung cancer, demonstrating that occludin is a potential prognostic biomarker and therapeutic target in lung cancer.

Introduction

Lung cancer is the most common cancer worldwide (1). Moreover, it is the first leading cause of cancer-related deaths in China and is a major public health problem, due to limited therapeutic options in advanced stages (2). In addition, metastasis, which is formed by the spread of disseminated primary tumor cells to distant anatomic sites, leads to more than 90% of the deaths of cancer patients (3,4). Fundamentally, aberrant processing of genetic information, involving activation of oncogenes coupled with inactivation of tumor-suppressor genes, can lead to the carcinogenesis of lung cancer (5). In view of this, the discovery of novel molecular targets involved in lung cancer development and metastasis is critical to develop diagnostic markers and targeted therapies.

Since cancer cells must pass through a series of barriers in order to successfully metastasize to secondary foci, the tight junction (TJ) has become a crucial component in the process of cancer metastasis over the past decade (6,7). One of the first identified TJ proteins is occludin (OCLN), which has been revealed to be a necessary integral protein for TJ structure and function (8). Occludin, widely expressed in tissues and cells that have TJ, is a membrane protein with four trans-membrane domains forming two extracellular loops and a long cytoplasmic tail (9,10). Although occludin can bind directly to actin (11), cytoplasmic scaffolding protein zonaoccludens (ZO)-1 can also serve as a link between occludin and the actin cytoskeleton (10). However, accumulating evidence has revealed that the claudin family, containing more than 20 members, is the core component of TJ (12). Apart from the basic function, TJ have also been demonstrated to be associated with signal transduction such as cancer related mitogen activated protein kinase (MAPK) and Akt signaling pathways (13), which further implies its important role in cancer development.

As anticipated, previous studies revealed that dysregulation of TJ function is vital to the development and invasiveness of many cancers (14,15). Consistently, high expression of TJ-associated proteins has been revealed to promote apoptosis and inhibit the metastatic potencies of these cancer cells (16). These data revealed that the TJ-related proteins may function in a context-dependent manner. However, the role of occludin in lung cancer development and metastasis remains to be elucidated. In the present study, we silenced occludin in lung cancer cells successfully, and observed suppressed cell proliferation and increased apoptosis, as well as compromised invasion ability. Our findings indicate that occludin is a likely candidate of oncogenes in lung cancer.

Materials and methods

Clinical tissue specimens. Fresh lung cancer tissue samples and corresponding adjacent normal tissues were obtained from
patients with lung cancer who underwent surgical resections at Zhongnan Hospital of Wuhan University from March 2014 to December 2015. These samples were immediately frozen in liquid nitrogen and preserved at -80°C freezer. All the experiments were carried out in accordance with the approved guidelines of Hubei University of Medicine.

Cell culture. Lung cancer cell lines including A549, NCL-H1650, SPC-A1, HCC827, NCI-H1299, and MSTO-211H were obtained from the Cell Bank of the Chinese Academy of Sciences (Shanghai, China). All cells were cultured in Dulbecco's modified Eagle's medium (DMEM) with 10% fetal bovine serum (FBS; both from HyClone Laboratories; GE Healthcare Life Sciences, Logan, UT, USA). All cells were incubated at 37°C in 5% CO<sub>2</sub> humidified air.

siRNA transfection. For knockdown of occludin in lung cancer cell lines, siRNA was used. The sequences of the oligonucleotides were as follows: Control siRNA, CCUUUUAGGGGUGUUAAtt; occludin siRNA, GGGUCUGCUGUGACAUAAAtt. For transfection, cells were seeded in a 12-well plate with a density of 5x10<sup>4</sup> cells/well. After 24 h and 70-80% confluence, the cells were transfected with respective occludin siRNAs (50 nM) and control-siRNA (50 nM) in serum-free medium using Lipofectamine 2000™ (Thermo Fisher Scientific, Inc., Waltham, MA, USA) according to manufacturer's instructions. After incubation for 6 h at 37°C, the medium in each well was then replaced with DMEM, with 10% heat-inactivated FBS for the indicated time-points.

Real-time PCR analysis. Total RNA was extracted from lung cancer cell lines using TRIzol<sup>®</sup> Reagent (Invitrogen; Thermo Fisher Scientific, Inc.). Then RNase-free DNase was used to remove genome DNA. Reverse transcription was performed using AMV reverse transcriptase (Promega Corp., Madison, WI, USA) as well as a combination of random oligo(dT) primers according to the manufacturer's instructions. For the detection of the indicated gene expression in RNA levels, the real-time PCR cycling conditions were as follows: Initial denaturation at 95°C for 1 min; denaturation at 95°C for 5 sec; annealing extension at 60°C for 10 sec (a total of 40 cycles). All of the reactions were performed using the Bio-Rad Connect Real-Time PCR platform (Bio-Rad Laboratories, Hercules, CA, USA). Absorbance values were read at the extension stage and all data were analyzed using the 2<sup>ΔΔCt</sup> method.

Western blot analysis. Cells were lysed for total protein extraction in 2X SDS sample buffer [100 mM Tris-HCl (pH 6.8), 10 mM EDTA, 4% SDS, 10% glycerine]. A total of 30 µg denatured proteins, were run on 12% polyacrylamide gels containing SDS and electrophoresed onto nitrocellulose membranes. Membranes were then blocked with 5% non-fat dry milk and were immunoblotted with antibodies overnight. After extensive washing, the membranes were reacted with peroxidase-labeled corresponding secondary antibodies in PBS/T and again washed; finally, the immunoreactions were visualized by using an enhanced chemiluminescence (ECL) system (Thermo Fisher Scientific, Inc., Waltham, MA, USA). Primary antibodies to occludin (cat. no. ab67161), caspase-3 (cat. no. ab49976), caspase-9 (cat. no. ab2013) and AIF (cat. no. ab1998) were purchased from Abcam (Cambridge, UK) with 1:1,000 dilutions. Primary antibodies to BAX (cat. no. sc-493) and Bcl-2 (cat. no. sc-492) were purchased from Santa Cruz Biotechnology, Inc. (Dallas, TX, USA) with 1:200 dilutions. Primary antibodies to cytochrome c (cyt c; cat. no. 11940), p-AKT (cat. no. 4060), AKT (cat. no. 2920), p-P13K (cat. no. 4228), P13K (cat. no. 4257) and GAPDH (cat. no. 5174) were purchased from Cell Signaling Technology, Inc. (Danvers, MA, USA) with 1:1,000 dilutions. Secondary antibodies: Anti-rabbit HRP (cat. no. sc-2370) and anti-mouse HRP (cat. no. sc-2371) were obtained from Santa Cruz Biotechnology, Inc. with 1:5,000 dilutions.

CCK-8 viability assay. The growth of cells was assessed by the CCK-8 assay. An equal number of 2,000 cells were seeded in 96-well plate and cultured for various durations after siRNA transfection. At indicated time-points (24, 48, 72, 96 and 120 h), 10 µl of CCK-8 (Dojindo Molecular Technologies, Inc., Kumamoto, Japan) was added to the cell medium and incubated for another 1 h at 37°C. The absorbance was assessed quantitatively at 450 nm on an ELISA platereader (Model 680; Bio-Rad Laboratories).

Flow cytometric apoptosis analysis. SPC-A1 cells (250,000 cells/dish) were seeded in 10-cm dishes after siRNA transfection. Forty-eight hours later, the cells were also harvested, washed in PBS followed by staining with Annexin V and 7-AAD (Dojindo Molecular Technologies, Inc.) for 30 min when cell density reached 80%. The samples were then analyzed using a flow cytometer. The proliferation indices were analyzed using ModFit software (Verity Software House, Inc., Topsham, ME, USA) and were evaluated according to the formula (S+G2)/(G1+S+G2).

Flow cytometric cell cycle analysis. SPC-A1 cells (3x10<sup>5</sup>) were seeded in 10-cm dishes and transfected with siRNA for 48 h. Cells were harvested and washed twice with cold PBS and then fixed with ice-cold 70% ethanol for 1 h at 4°C. After centrifugation at 1,000 x g for 5 min, the cells were washed twice with PBS and resuspended with 0.5 ml of PBS containing PI (50 µg/ml) and RNase (100 µg/ml). The samples were then stained at room temperature in the dark for 15 min, and the cell cycle distribution was assessed by flow cytometry.

Transwell assays. To assess the invasiveness of lung cancer cells, we used Transwell chambers (Corning Inc., Corning, NY, USA) coated with Matrigel (BD Biosciences, Franklin Lakes, NJ, USA). Fully trypsinized transfected cells...
(1x10⁵ cells in 100 µl of serum-free DMEM) were plated into the upper chamber. DMEM (500 µl) supplemented with 20% FBS was added to the lower chamber. After incubation in a humidified atmosphere containing 5% CO₂ at 37˚C for 72 h, lung cancer cells that had invaded the lower chamber were fixed with methanol and stained with crystal violet. We used an inverted microscope to capture images of the invading cells manually. A Transwell assay was also performed to assess the cell migration in the same manner as the Transwell invasion assay with the exception of Matrigel coating.

Figure 1. Occludin is overexpressed in lung cancer tumor tissues. (A) qRT-PCR was performed to assess the mRNA level of occludin in lung cancer tumor tissues and paired adjacent normal tissues. GAPDH was used as a control. (B) Western blotting was used to assess the protein level of occludin in lung cancer tumor tissues and paired adjacent normal tissues. GAPDH was used as a control. (C) Quantification of the expression level of occludin in the two groups and normalized to GAPDH. (D) Quantification of the expression level of occludin in the two groups (by the grayscale value in A) and normalized to GAPDH. Error bars denote standard deviations. n=10, *P<0.05, **P<0.01, ***P<0.001. OCLN, occludin.
Mouse xenograft model. A mouse xenograft model was established as previously described (17). Briefly, Balb/c (nu/nu) (4-5 weeks old) nude mice were randomly divided into two groups and subcutaneously injected in the flank regions with 1.0x10^7 cells in 0.1 ml of PBS. The tumor size was assessed every 4 days with calipers and the tumor volume was calculated with the formula: (length x width^2)/2. Forty days following implantation, the mice were euthanized by asphyxiation in a CO2 chamber and tumors were excised immediately and images were captured. All procedures were conducted in accordance to the Animal Care and Use Committee Guidelines of Hubei University of Medicine.

Statistical analysis. All data were presented as the mean ± standard deviation (SD) and normalized relative to the negative control. Statistical analysis was performed using Prism 6 software. Differences among different groups were analyzed by one-way ANOVA. A P-value <0.05 was considered to indicate a statistically significant difference.

Results

Occludin is overexpressed in lung cancer tumor tissues. To determine whether occludin is involved in lung cancer development, we collected 71 pairs of lung cancer tumor tissues and their paired adjacent normal tissues and performed qRT-PCR to assess the relative expression levels of occludin. Notably, the relative expression level of occludin in the lung cancer tissues was significantly higher than that in the corresponding adjacent non-tumor tissues (Fig. 1A and C). Furthermore, in a small set of human lung cancer tumor and normal tissues from the same patients, all of the 10 tumor tissues exhibited increased occludin protein levels (Fig. 1B and D), which indicates that occludin may participate in the pathogenesis of lung cancer.

Figure 2. Occludin knockdown inhibits cell proliferation in vitro. (A) Western blotting was used to assess the protein level of occludin in various lung cancer cell lines. GAPDH was used as a control. (B) Quantification of the expression value of occludin in cell lines (by the grayscale value in A) and normalized to GAPDH. (C) Western blotting was used to assess the protein level of occludin in lung cancer cell line SPC-A1. GAPDH was used as a control. (D) Quantification of the expression level of occludin in SPC-A1 (by the grayscale value in C) and normalized to GAPDH. (E) CCK-8 growth assay of the control, siCtrl, and siOCLN of SPC-A1 cells. **P<0.01, ***P<0.001. OCLN, occludin.
Occludin knockdown inhibits cell proliferation. To analyze the expression pattern of occludin in lung cancer cells, six lung cancer cell lines, including A549, NCL-H1650, SPC-A1, HCC827, NCI-H1299 and MSTO-211H were examined. We determined that occludin was highly expressed in SPC-A1, moderately expressed in A549, HCC827, NCI-H1299 and MSTO-211H while relatively lowly expressed in NCL-H1650 cells (Fig. 2A and B). Therefore, we selected the SPC-A1 cell line to perform the following experiments. To investigate the functional role of occludin in lung carcinogenesis, siRNA-mediated loss-of-function experiments \textit{in vitro} were performed. The occludin expression levels of SPC-A1 cells transfected with occludin siRNA were significantly lower than that of SPC-A1 cells transfected with the scrambled sequence (Fig. 2C and D), indicating high knockdown efficiency.

Next, we evaluated the effect of occludin on lung cancer cell proliferation \textit{in vitro} by CCK-8 assay. Notably, we found that occludin knockdown resulted in significantly decreased cell proliferation in SPC-A1 cells at every time-point, indicating that occludin has a role in promoting the proliferation of SPC-A1 cells (Fig. 2E). These results demonstrated that overexpression of occludin promoted the proliferation of lung cancer cells.

\textbf{Occludin silencing induces cell cycle arrest and cell apoptosis.} We next investigated whether occludin promotes the proliferation of SPC-A1 cells by influencing cell cycle distribution \textit{in vitro}. The cell cycle analysis revealed that, compared with transfection with the scrambled siRNA, transfection with occludin siRNA resulted in a significant reduction of the SPC-A1 cells in the S and G2 phases and an increase in the G1 phase (Fig. 3A and C). Furthermore, we found that knockdown occludin significantly decreased the proliferation indices (Fig. 3D). Collectively, we demonstrated that the increase of occludin contributed to the proliferation of lung cancer cells by promoting cell cycle progression.

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**Figure 3.** Occludin silencing induces cell cycle arrest and cell apoptosis. (A) Flow cytometry was used to assess the cell cycle distribution of SPC-A1 cells stained with PI. Representative results are shown. (B) Flow cytometry was used to assess the percentage of apoptotic SPC-A1 cells after transfection with scramble or occludin siRNA. Representative scatter plots are shown. (C) Quantification of the percentage of G1, S, G2 cells assayed by flow cytometry. (D) Calculation of the proliferation indices according to flow cytometry. (E) Quantification of the percentage of apoptotic SPC-A1 cells. *\(P<0.05\), **\(P<0.01\), ***\(P<0.001\). OCLN, occludin.
We further examined the effect of occludin on apoptosis using flow cytometry. As anticipated, the early apoptotic cells (Annexin V⁺/7AAD⁻) were significantly increased after occludin knockdown in SPC-A1 cells (Fig. 3B and D), indicating that plays an anti-apoptotic role. In summary, we hypothesized that occludin supports lung cancer growth by exerting anti-apoptotic effects and promoting cell cycle progression in vitro.

Occludin silencing inhibits cell invasion and migration. Considering that TJ proteins are significantly correlated with the metastasis of various cancers, we further studied the role of occludin in the regulation of invasion ability by Transwell assays. Knockdown of occludin in SPC-A1 cells markedly attenuated their invasive ability (Fig. 4A and B), suggesting that occludin has a critical function in improving the invasion of lung cancer cells. Next, we explored the effect of occludin on
cell migration and found that occludin silencing significantly decreased the migration capacity as well (Fig. 4C and D). Therefore, we determined that occludin was a pro-metastatic factor in lung cancer cells.

**Occludin silencing inhibits lung cancer cell proliferation in a xenograft tumor model in vivo.** We further confirmed the aforementioned results in a xenograft tumor model in vivo. siCtrl and siOCLN SPC-A1 cells were injected subcutaneously into two groups of nude mice randomly. As anticipated, the tumor growth curve revealed that tumors derived from the siOCLN group grew more slowly than those from the siCtrl group (Fig. 5A). The tumor volume of the siCtrl group was 974±98.6 mm³, whereas the tumor volume of siOCLN group was 547±64.3 mm³ 40 days after implantation (Fig. 5B). These results demonstrated that overexpression of occludin promoted the proliferation of lung cancer cells in vivo.

**Occludin knockdown regulates apoptosis-related proteins and AKT/PI3K signaling.** To discover the mechanisms of occludin in lung cancer growth, we investigated the proliferation and apoptosis-related signaling pathways. We found that the AKT/PI3K pathway activation, which is essential for most cell growth, was compromised after occludin silencing (Fig. 6A and B). Accordingly, we detected the protein level of apoptosis-related molecules and found that the expression of BAX, caspase-3, caspase-9, AIF, and Bcl-2 were upregulated while anti-apoptotic Bcl-2 was decreased in SPC-A1 cells transfected with occludin siRNA (Fig. 6C and D). Thus, we surmised that occludin regulated lung cancer cell growth by affecting the AKT/PI3K pathway and apoptosis-associated factors.

**Discussion**

Tight junctions (TJs) the most apical intercellular structures in epithelial and endothelial cells are considered to play important roles in the formation of cell polarity and paracellular permeability (18,19). Additionally, accumulating evidence has revealed that the disruption of TJ structure has been linked to tumor development, which may be causally involved in malignant phenotypes, such as local tumor growth, invasion, and metastasis at distant sites (6,7,14,15). In the present study, we first deduced the critical role of occludin, a crucial component of TJs, in lung cancer. Occludin, which was overexpressed in
lung cancer tumor tissues, regulated the growth and metastasis of lung cancer cells in vitro and in vivo by controlling important proliferation pathways and a common set of cell apoptosis-related genes. Collectively, our study revealed the promotive role of occludin in lung cancer.

The two most fundamental traits of cancer cells involve their ability to sustain uncontrolled proliferation and resist cell death (5). An important finding of this study was the identification of occludin as a tumor promoter by altering the proliferation-associated pathways as well as the expression of apoptosis-related genes. The AKT/PI3K pathway, which is stimulated by insulin (20,21), participates in cell proliferation among various cancer types. In our study, we revealed that occludin may promote lung cancer cell proliferation by modulating this key pathway. However, whether lung cancer development is associated with insulin levels remains unclear. Apoptosis is an evolutionarily conserved process in the regulation of tissue homeostasis during normal development. However, several studies have recognized that inadequate apoptosis leads to the malignant phenotype, since it is widely believed that impaired susceptibility to respond to numerous apoptotic signals is associated with oncogenic transformation (22-24). In the present study, we determined that occludin knockdown could increase apoptosis, which may be due to the altered apoptosis-related gene profiles. This is consistent with a previous study that revealed that occludin also regulates Bax, Bcl-2 and caspase-3 expression in liver cancer cell lines (25). Occludin-deficient mice also displayed pronounced numbers of apoptotic primary hepatocytes and immortalized cells stimulated by serum-free conditions compared with the wild-type cells (13), in line with our data and the results in mammary gland cells (26). The modulation of the expression levels of these genes indicated that occludin may affect mitochondrial-mediated apoptosis (27). However, the association between occludin and mitochondria function warrants further study in the future. In contrast, overexpression of occludin enhanced the sensitivity to H2O2-induced cell death in HeLa cells (28). Thus, the function of occludin in apoptosis is complex and may depend on the cell type. In summary, overexpression of occludin contributed to limitless proliferation as well as acquisition of apoptotic resistance of lung cancer cells and participated in the carcinogenic process accordingly.

It is well accepted that the metastatic process involves a complicated interplay between altered cell adhesion, survival, migration, and homing on distant target organs (29). Notably, loss of occludin also displayed suppressive and progressive cancer phenotypes, such as invasive properties, which resulted in an important contribution to metastatic inhibition. Therefore, our data revealed that forced expression of occludin in tumors favors various steps affecting lung cancer development. We could not exclude the possibility that occludin may have some unidentified but more notable effects on tumor cells. Thus, considering the complexity of carcinogenesis, the in vivo function of occludin in lung cancer warrants further exploration in the future.

In summary, we revealed that high occludin expression confers malignant growth of lung cancer cells largely by promoting proliferation and blocking apoptosis in vitro and in vivo, and suggests that it may be a potential biomarker for lung cancer progression. Furthermore, our results revealed that occludin is a likely candidate as a tumor-promoter gene in certain types of cancer and may pave the way to new therapeutic modalities. Occludin depletion even antagonistically would be a potential therapeutic strategy for lung cancer.

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Availability of data and materials

The datasets used during the present study are available from the corresponding author upon reasonable request.

Authors' contributions

MW and JY conceived and designed the study. MW, YL, XQ, NW and YT performed the experiments. MW and JY wrote the paper. All authors read and approved the manuscript and agree to be accountable for all aspects of the research in ensuring that the accuracy or integrity of any part of the work are appropriately investigated and resolved.

Ethics approval and consent to participate

All experimental protocols were approved by the Ethics Committee of Zhongnan Hospital of Wuhan University.

Consent for publication

Not applicable.

Competing interests

The authors state that they have no competing interests.

References


