Possible involvement of the Hedgehog and PDPK1–Akt pathways in the growth and migration of small-cell lung cancer

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Abstract

Background: Small-cell lung cancer (SCLC) accounts for approximately 15% to 20% of all lung cancers, and it is the leading cause of tumor-related deaths globally. This study explored the molecular mechanisms underlying the development of SCLC.

Methods: The correlations of phosphoinositide-dependent kinase-1 (PDPK1), p-Akt, and Hedgehog expression with patient characteristics were analyzed using SCLC specimens, and their expression was measured in BEAS-2B cells (control) and the SCLC cell lines H82, H69, H446, H146, and H526. Transfection experiments were performed to inhibit or activate gene expression in cells. We then measured the proliferation and migration of H146 cells.

Results: PDPK1, p-Akt, and Hedgehog expression was significantly higher in SCLC tissues, and their expression was correlated with patient characteristics. p-Akt expression was significantly correlated with Hedgehog expression. In H146 cells, PDPK1 and p-Akt were significantly upregulated. Silencing of PDPK1 or Akt and inhibition of Hedgehog significantly inhibited the proliferation and migration of H146 cells. PDPK1 and Akt affected Hedgehog expression, but Hedgehog did not affect PDPK1 or p-Akt expression.

Conclusions: The interaction between the PDPK1–Akt pathway and the Hedgehog pathway influences the prognosis, growth, and migration of SCLC.

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Introduction

Lung cancer is currently the leading cause of tumor-related death worldwide. Small-cell lung cancer (SCLC) accounts for approximately 15% to 20% of lung cancer cases. Although SCLC is initially sensitive to chemotherapy, resistance is often acquired within a short period, and the disease progresses rapidly. Among patients receiving standard treatment, the median survival of those with non-metastatic lung cancer is approximately 17 months, versus 9 to 11 months for those with metastatic lung cancer. The prominent features of SCLC include frequent lymphatic and hematogenous metastasis and an extremely poor prognosis. Approximately 80% of patients with SCLC are diagnosed with clinical or subclinical lymphatic or hematogenous metastasis. Common metastatic sites include the lungs, brain, liver, and bones. Consequently, the long-term survival rate is extremely low. Therefore, to improve the diagnosis, prevention, and treatment of SCLC, it is important to study the pathogenesis and migration mechanism of SCLC, identify novel genes specific for SCLC, and determine the roles of these genes in SCLC pathogenesis.

Current studies have found that chemokines, cytokines, integrins, cadherins, and metalloproteinases are closely related to tumor metastasis. Akt is a serine- and threonine-specific kinase that is essential for cellular processes, including proliferation, apoptosis, and migration. After Akt binds to phosphatidylinositol triphosphate (PI3K) on the membrane in the correct manner, it can be phosphorylated by phosphoinositide-dependent kinase-1 (PDPK1) and mammalian target of rapamycin complex 2 at Thr308 and Ser473, respectively. This phosphorylation activates Akt, enabling it to regulate the phosphorylation of downstream proteins. Studies illustrated that inhibition of the PDPK1–Akt interaction significantly inhibits the proliferation of mouse hepatoma cells, indicating a role of the PDPK1–Akt signaling pathway in the proliferation of hepatoma cells.

The Hedgehog signaling pathway regulates the differentiation and growth of a variety of cells during embryonic development and plays an important role in the maintenance of environmental stability in mature organs and the maintenance of stem cell functions in certain mature organs. Numerous reports revealed that abnormally activated Hedgehog signaling is closely related to the occurrence and development of various human tumors, such as basal cell carcinoma, SCLC, pancreatic cancer, prostate cancer, and gastric cancer. However, the interaction of Hedgehog with the PDPK1–Akt signaling pathway and the role of this interaction in the development and migration of SCLC have seldom been studied. Hence, in the present study, we first examined the expression of Hedgehog, PDPK1, and Akt in patients with SCLC via immunohistochemical analysis to evaluate their relationships with clinical characteristics. Then, we
explored the roles of Hedgehog and the PDK1–Akt pathway in the proliferation and migration of SCLC cell lines using siRNA techniques.

Materials and methods

Patients and specimen collection

We studied specimens from patients who were diagnosed with SCLC and treated at Central Hospital of Xuhui District, Shanghai, China between July 2005 and January 2018. Specimens were fixed in 10% neutral-buffered formaldehyde for 24 hours at room temperature and processed for histopathologic and immunohistochemical evaluation. The control specimens were obtained from normal lung tissue located at least 5 cm from lung cancer tissue in patients with SCLC. Clinical factors, including sex, age, cancer location and differentiation, lymph node metastasis, TNM stage, and lesion size were obtained from patients’ medical records. The TNM stage was determined according to the Union for International Cancer Control guidelines. This study was approved by the ethics committee of Central Hospital of Xuhui District (approval No. SHXH2017-3265). Written informed consent was obtained from all patients. All procedures performed in studies involving human participants complied with the ethical standards of the institutional and/or national research committee and with the 1964 Declaration of Helsinki and its later amendments or comparable ethical standards.

Immunohistochemistry assay

First, 5-μm paraffin-embedded sections of specimens were deparaffinized and rehydrated. Endogenous peroxidase activity was blocked by incubating the sections with 3% H2O2 at 37°C. The sections were incubated with primary antibodies against PDK1 (ab52893, 1:100, Abcam, Cambridge, UK), p-Akt (ab38449, 1:100, Abcam), and Hedgehog (ab53281, 1/100; Abcam) overnight at 4°C. Afterward, the sections were washed and incubated with biotin-conjugated secondary antibodies (Sigma-Aldrich, St. Louis, MO, USA) and streptavidin-conjugated peroxidase (Abcam) for 30 minutes at room temperature. The expression of PDK1, p-Akt, and Hedgehog was measured using an image analysis system (Leica Q500 MC, Leica Instruments Ltd., Wetzlar, Germany). Ten fields of representative tissue cores of tissue microarray specimens were randomly selected and observed. One hundred positive cells were examined in each specimen. The number of positive units (PUs) of each positive cell was calculated as described by Wan et al., using the equation following equation: PU = (Ga – Gb)/Gmax x 100, where Ga means the average gray scale of positive staining, Gb means the average gray scale of background, and Gmax means the maximum gray scale of the image analysis system.

Survival curve

We analyzed the correlations between the expression of p-Akt and Hedgehog and the survival rate of patients with SCLC by dividing patients into two groups: the p-Akt– and/or Hedgehog-negative group and the p-Akt–positive and Hedgehog-positive group. Survival was assessed for 240 weeks in each group. The distribution for overall survival was estimated using the Kaplan–Meier method. The difference of survival between the groups was compared using the log-rank test.

Cell culture

BEAS-2B cells (normal lung epithelial cells, control) and the SCLC cell lines H82, H69, H446, H146, and H526 cells were purchased from American Type Culture
Collection (Manassas, VA, USA). The various cell lines were cultured in a humidified atmosphere of 5% CO₂ at 37°C in RPMI 1640 medium (Gibco, Thermo Fisher Scientific, Waltham, MA, USA) or Dulbecco’s modified Eagle’s medium (DMEM, Gibco). Both media were supplemented with 10% fetal bovine serum (Gibco). To confirm the effects of Hedgehog in normal lung epithelial and SCLC cell functioning (migration and proliferation), we treated BEAS-2B and H146 cells with the Hedgehog inhibitors cyclopamine (LC Laboratories, Woburn, MA, USA) and GDC-0449 (LC Laboratories) at a concentration of 1 μM for 72 hours, then measured cell viability and migration.

Western blot analysis
Cultured cells were first lysed in RIPA buffer (Thermo Fisher Scientific) and solubilized in SDS sample buffer. The protein concentrations in samples were determined using the Bradford assay (Bio-Rad, Hercules, CA, USA). The lysates were then separated via 12.5% sodium dodecyl sulfate–polyacrylamide gel electrophoresis (20 μg of total protein/lane). The separated proteins were transferred to a polyvinylidene difluoride membrane (MilliporeSigma, Burlington, MA, USA). After blocking in 5% (w/v) non-fat dry milk in 0.1% Tween 20 in TBS (Thermo Fisher Scientific) for 1 hour at room temperature, the membrane was incubated with primary antibodies against Hedgehog, PDPK1, p-Akt, and β-actin (#4967, 1:2000, Cell Signaling Technology, Danvers, MA, USA) overnight at 4°C. Next, membranes were incubated with horseradish peroxidase-conjugated goat antibodies (#7054; 1:10000, Cell Signaling Technology). Finally, the membranes were washed with saline and was examined using the ECL Plus Western Blotting Detection System (Amersham Life Science, Little Chalfont, UK) and visualized using the Image Lab V4.0 electrochemiluminescence system (Bio-Rad).

siRNAs
siRNAs targeting PDPK1 and Akt were purchased from Qiagen (Hilden, Germany). The transfection of siRNAs was performed using Hyperfect transfection reagent (Qiagen) according to the manufacturer’s instructions. siRNAs targeting PDPK1 and Akt were transfected into H146 cells. The targeted sequences of PDPK1 were 5'-CACGCCUAACAGGAGC GUAUUA-3' (siPDPK1-1), 5'-CAAGAGC GACCUCGUGGAGAA-3' (siPDPK1-2), and 5'-GCAAGACGACGAGACGUUAU UA-3' (siPDPK1-3). The targeted sequences of Akt were 5'-AAGGAGAGACGATG GACTTCCGAT-3' (siAkt-1), 5'-AAGG AC GGGGACATCAAGATA-3' (siAkt-2), and 5'-CCAUGAACGAGUUUGAGUAG CC-3' (siAkt-3). In total, 25 nM siRNA was used for each transfection. After 24 hours, the transfected cells were assayed. Only the results for siRNA-1 are presented in the figures.

Vector construction and transfection
H146 cells were seeded in six-well dishes (5 x 10⁵ cells/well) and grown to 70% to 80% confluence. For vector construction, the open-reading frame of PDPK1 cDNA was cloned into the pcDNA 3.1 vector (Ambion, Austin, TX, USA). The primers were as follows: 5'-ATGGGCCAGGACC ACCAGCCA GCTGTATGAGC-3' (forward) and 5'-AGTGACGTGTCGCCG CA GGCCACCGGA-3' (reverse). To express constitutively active Akt (CA-Akt), cells were transfected with either empty pcDNA 3.1 or pcDNA 3.1 vectors encoding CA-Akt kinase (Addgene Watertown, MA, USA). For transient transfection, cells were transfected with
pcDNA 3.1, pcDNA 3.1-PDPK1, or pcDNA 3.1 CA-Akt vectors (1–2 μg of total DNA per well) at approximately 80% confluence using Lipofectamine™ 2000 (Invitrogen, Thermo Fisher Scientific) according to the manufacturer’s instructions. The transfected H146 cells were incubated for 40 hours at 37°C before use in experiments.

**Cell proliferation assay**

Cell proliferation assays were performed using Cell Counting Kit-8 (CCK-8, Beyotime, Shanghai, China) according to the manufacturer’s protocol. In total, 100 mL of cell suspension was added to a 96-well plate and cultured in an incubator for 24 hours at 37°C and 5% CO2. Next, 10 mL of the CCK-8 solution was added to each well. Cell culture was continued for another 4 hours. The absorbance at 450 nm was then determined using a microplate reader (BioTek, Winooski, VT, USA) and was used to calculate the cell viability.

**Wound closure assay**

Cell invasion was measured using a wound closure assay. Cells were cultured in a 24-well plate in RPMI 1640 medium containing 10% FBS at 37°C until a monolayer formed. After the cells had reached 80% to 90% confluence, a 1-mm stainless steel probe was sterilized and used to draw a horizontal line on the bottom of each well. The cell debris was carefully removed with PBS. Next, cell culture was continued for 20 hours under the same conditions. An image of the width of the wound was taken and measured using ImageJ software (US National Institutes of Health, Bethesda, MD, USA). The extent of wound closure was calculated using the following formula: wound closure = ((wound width at 0 hours) – (wound width at 20 hours)/(wound width at 0 hours)) × 100%.

**Transwell assay**

Cell migration was assayed using Transwell assay. A hydrated Transwell chamber (Corning, Corning, NY, USA) was placed in a 24-well plate, and 500 μL of RPMI 1640 medium containing 15% FBS was added to the outside of the chamber. Next, 200 μL of the tumor cell suspension (containing 5 × 10^5 cells) was added to the chamber, followed by culture in RPMI 1640 medium containing 2% FBS. Twenty-four hours later, the Transwell chamber was removed and rinsed with PBS. The inserts were removed and dried. The cells in the upper chamber that had not migrated were removed using a cotton swab. Five hundred microliters of 0.1% crystal violet were added to stain the cells, and the number of cells that had migrated to the lower layer of the microporous membrane was counted under a fluorescence microscope (Keyence, Osaka, Japan). Ten fields were counted for each sample.

**Statistical analysis**

The paired t-test was used to compare differences between control and cancer tissues collected from the same patients. Survival curves were analyzed using the two-sided log-rank test. The data are presented as the mean ± standard error of the mean, and the data were analyzed using one-way analysis of variance followed by the Student–Newman–Keuls post hoc test. Statistical analysis was performed using IBM SPSS 17.0 (SPSS Inc., Chicago, IL, USA). *P < 0.05* was considered significant.

**Results**

**PDPK1, p-Akt, and Hedgehog expression in SCLC specimens**

PDPK1, p-Akt, and Hedgehog expression in normal and SCLC specimens from 65
patients was examined via immunohistochemical analysis, as presented in Figure 1. Quantitative analysis revealed that the PUs of PDPK1, p-Akt, and Hedgehog were all significantly higher in SCLC specimens than in control specimens (all \( P < 0.05 \)). SCLC specimens had significantly higher rates of positive PDPK1, p-Akt, and Hedgehog staining than control specimens (\( P < 0.05 \)).

**Correlations between the PUs of PDPK1, p-Akt, and Hedgehog and the clinicopathological parameters of SCLC**

We analyzed the correlations of the PUs of PDPK1, p-Akt, and Hedgehog with the clinicopathological parameters of lung carcinoma, including gender, age, cancer location, cancer differentiation, lymph node metastasis, TNM stage, and lesion size. The results are presented in Table 1.

The PUs of PDPK1, p-Akt, and Hedgehog were not significantly correlated with sex, age, and cancer location. The PU of Hedgehog was significantly correlated with “moderate to poor” cancer differentiation, N1–3 lymph node metastasis, a TNM stage of II to IV, and lesion size ≥3 cm (all \( P < 0.05 \)). The PUs of PDPK1 and p-Akt were significantly correlated with a TNM stage of II to IV and lesion size ≥3 cm (\( P < 0.05 \)).

**Correlation between the expression of p-Akt and Hedgehog**

The correlation between p-Akt and Hedgehog expression was analyzed using p-Akt and Hedgehog PUs and Pearson’s correlation analysis, as presented in Figure 2a. Pearson’s \( r \) was 0.5884 (95% confidence interval = 0.1089–0.8458, \( P = 0.021 \)), which indicated a significant correlation between p-Akt and Hedgehog expression.

![Figure 1](image-url)  
**Figure 1.** PDPK1, p-Akt, and Hedgehog expression in SCLC specimens. (a–c) PDPK1, p-Akt, and Hedgehog expression in normal tissues and SCLC specimens was examined via immunohistochemical analysis. (d–f) Quantitative analysis. (g–i) The rates of PDPK1, p-Akt, and Hedgehog positivity in normal (control) and SCLC tissues. #\( P < 0.05 \) versus control.  
PDPK1, phosphoinositide-dependent kinase-1; SCLC, small-cell lung cancer.
Table 1. Correlations of Hedgehog, PDPK1, and p-AKT PUs with patient clinicopathological parameters

|                  | N  | Hedgehog PUs | P  | PDPK1 PUs | P  | p-AKT PUs | P  |
|------------------|----|--------------|----|-----------|----|-----------|----|
| Sex              |    |              |    |           |    |           |    |
| Male             | 30 | 38.3 ± 10.6  | 0.86| 28.9 ± 7.8| 0.75| 35.6 ± 4.9| 0.82|
| Female           | 35 | 41.2 ± 11.4  |    | 33.6 ± 9.7|    | 33.9 ± 6.2|    |
| Age (years)      |    |              |    |           |    |           |    |
| <60              | 26 | 37.8 ± 9.8   | 0.76| 31.3 ± 7.6| 0.83| 33.8 ± 7.8| 0.92|
| ≥60              | 39 | 42.6 ± 11.7  |    | 28.9 ± 7.5|    | 36.1 ± 7.9|    |
| Cancer location  |    |              |    |           |    |           |    |
| Central          | 33 | 38.5 ± 12.6  | 0.92| 29.5 ± 6.9| 0.88| 32.8 ± 8.3| 0.81|
| Peripheral       | 32 | 43.8 ± 12.4  |    | 31.2 ± 7.4|    | 36.9 ± 6.8|    |
| Cancer differentiation | |   |    |           |    |           |    |
| Well             | 28 | 25.9 ± 8.9   | 0.01| 32.6 ± 5.2| 0.87| 36.4 ± 5.8| 0.86|
| Moderate to poor | 37 | 45.8 ± 11.5  |    | 30.9 ± 6.7|    | 33.2 ± 7.1|    |
| Lymph node metastasis | |   |    |           |    |           |    |
| N₀               | 34 | 19.9 ± 4.5   | 0.01| 30.9 ± 6.2| 0.83| 35.6 ± 5.2| 0.92|
| N₁–₃            | 31 | 60.3 ± 15.8  |    | 31.9 ± 10.1| 0.83| 33.5 ± 6.7|    |
| TNM stage        |    |              |    |           |    |           |    |
| Stage I          | 25 | 20.3 ± 5.1   | 0.01| 14.6 ± 5.2| 0.01| 16.8 ± 4.3| 0.01|
| Stage II–IV      | 40 | 51.2 ± 13.6  |    | 39.6 ± 9.4|    | 38.2 ± 4.5|    |
| Lesion size      |    |              |    |           |    |           |    |
| <3 cm            | 21 | 18.5 ± 3.8   | 0.01| 17.2 ± 6.3| 0.01| 19.2 ± 5.7| 0.01|
| ≥3 cm            | 44 | 55.6 ± 15.2  |    | 36.5 ± 7.8|    | 39.5 ± 6.6|    |

PDPK1, phosphoinositide-dependent kinase-1; PU, positive unit.

Figure 2. Correlations of p-Akt and Hedgehog expression with the survival of patients with SCLC. (a) The correlation between p-Akt and Hedgehog expression was analyzed using Pearson’s correlation analysis. (b) Correlations between p-Akt and Hedgehog expression and the survival of patients with SCLC. The survival of each patient was recorded for 240 weeks. PU, positive unit.
**Correlations between the expression of p-Akt and Hedgehog and the survival rates of patients with SCLC**

As presented in Figure 2b, the survival rate was significantly higher in patients with p-Akt-negative and/or Hedgehog-negative lesions than in those with p-Akt-positive and Hedgehog-positive lesions ($P = 0.0118$).

**PDPK1, p-Akt, and Hedgehog expression in SCLC cell lines**

As presented in Figure 3a, Hedgehog expression was significantly higher in H146 and H446 cells than in BEAS-2B cells (both $P < 0.05$). PDPK1 and p-Akt expression was significantly higher in H146 cells than in BEAS-2B cells ($P < 0.05$, Figure 3b–c). Based on this result, H146 cells were used in the subsequent experiments.

**Changes of Hedgehog and PDPK1 expression and Akt phosphorylation in H146 cells**

Figure 4a illustrates that PDPK1 expression was significantly inhibited by PDPK1 siRNA but was enhanced by CA-PDPK1 transfection (both $P < 0.05$). Similar findings were observed for p-Akt phosphorylation (both $P < 0.05$, Figure 4b). To analyze the correlations of PDPK1 and p-Akt expression with Hedgehog expression, we examined the effects of PDPK1 siRNA on p-Akt and Hedgehog expression in H146 cells. As presented in Figure 4c, PDPK1 siRNA significantly inhibited Hedgehog expression versus scrambled siRNA transfection ($P < 0.05$). In addition, PDPK1 siRNA transfection significantly inhibited the ratio of p-Akt/Akt ($P < 0.05$, Figure 4d). Next, we transfected H146 cells with CA-PDPK1 and CA-Akt, then measured the changes in Hedgehog expression. The results (Figure 4e) revealed that Hedgehog expression was significantly increased following PDPK1 and p-Akt overexpression (both $P < 0.05$). When we treated H146 cells with the Hedgehog inhibitors cyclopamine and GDC-0449, PDPK1 and p-Akt expression was not affected (Figure 4f,g). These results suggested that PDPK1 and Akt affected Hedgehog expression, but Hedgehog did not influence PDPK1 or p-Akt expression.

![Figure 3.](image-url)
Effects of PDPK1 or Akt siRNA transfection on the migration and proliferation of H146 cells

We treated H146 cells with scrambled siRNA, PDPK1 siRNA, or Akt siRNA, then measured the migration and proliferation of H146 cells. As highlighted in Figure 5a–d, H146 cell migration was inhibited by PDPK1 siRNA or Akt siRNA transfection (both \( P < 0.05 \)). As presented in Figure 5e, PDPK1 siRNA or Akt siRNA transfection also inhibited cell proliferation (both \( P < 0.05 \)).

Effects of Hedgehog inhibitors on the migration and proliferation of cells

We treated H146 cells with cyclopamine and GDC-0449, then measured the migration and proliferation of H146 cells. As presented in Figure 6, cyclopamine and GDC-0449 treatment significantly decreased the viability of both BEAS-2B and H146 cells (all \( P < 0.05 \)). As highlighted in Figure 7, cyclopamine and GDC-0449 exposure also significantly decreased the migration of H146 cells (both \( P < 0.05 \)).

Discussion

Previous studies revealed that PDPK1 is highly expressed in malignant tumors, including breast cancer, non-small-cell lung cancer (NSCLC), and pancreatic cancer. It is speculated that the gene acts as a proto-oncogene in the regulation of tumor progression.\(^8\)–\(^10\) Studies indicated that PDPK1 promotes the proliferation,
invasion, and migration of gallbladder cancer cells by upregulating JunB. It has also been reported that epidermal growth factor-induced PDPK1 promotes the metastasis of head and neck squamous cell carcinoma by upregulating fibronectin. Akt promotes or inhibits other survival pathways, such as the PKC, RTK–MAPK, and JAK–STAT3 pathways. Inhibiting the expression and activation of Akt both blocks the oncogenic activity of this gene and inhibits a variety of lung cancer-related genes, such as growth factors, growth factor receptors, and Ras. This suppression hinders the development of lung cancer. Recent studies found that Akt promotes the growth and proliferation of tumor cells, inhibits cell apoptosis, increases the tolerance of cells to hypoxia, induces angiogenesis, and stimulates the invasion and metastasis of tumor cells by phosphorylating various downstream substrates. In the present study, we found that PDPK1 and p-Akt expression was significantly enhanced in SCLC tissues. The rates of PDPK1, p-Akt, and Hedgehog positivity were significantly higher in SCLC tissues than in normal tissues. PDPK1 and p-Akt expression was positively correlated with a TNM stage of II to IV and lesion size.
Figure 6. Effects of Hedgehog inhibitors on the proliferation of (a) BEAS-2B and (b) H146 cells. BEAS-2B and H146 cells were treated with cyclopamine and GDC-0449, and proliferation and migration were evaluated using Cell Counting Kit-8. \#P < 0.05 versus the scrambled siRNA group. PDPK1, phosphoinositide-dependent kinase-1.

Figure 7. Effects of the Hedgehog inhibitors cyclopamine and GDC-0449 on the invasiveness and migration of H146 cells. Following treatment, (a and c) wound closure and (b and d) migration were evaluated. \#P < 0.05 versus the scrambled siRNA group. PDPK1, phosphoinositide-dependent kinase-1.
These results confirmed the close relationship between the PDPK1–p-Akt pathway and the clinical parameters of SCLC.

There is substantial evidence of the role of the Hedgehog pathway in the development of lung cancer. Watkins et al.\textsuperscript{13,14} discovered the activation of the Hedgehog pathway in SCLC. In the process of tracheal epithelial repair, the epithelial expression of Hedgehog ligand and Gli was significantly enhanced. Watkins and colleagues later discovered that this phenomenon also existed in the embryonic lung.\textsuperscript{14} The application of a Hedgehog monoclonal antibody inhibits the growth of SCLC cell lines but not that of NSCLC cell lines expressing Hedgehog, indicating that SCLC cells are dependent on the ligand-mediated activation of the Hedgehog pathway \textit{in vitro}.\textsuperscript{13} The results in the present study illustrated that Hedgehog expression was significantly enhanced in SCLC tissues. The rate of Hedgehog positivity was significantly higher in SCLC tissues than in normal tissues. Hedgehog expression was positively correlated with "moderate to poor" cancer differentiation, $N_{1-3}$ lymph node metastasis, a TNM stage of II to IV, and lesion size $\geq 3\text{ cm}$. To confirm the effects of Hedgehog on normal lung epithelial and SCLC cell functioning (proliferation and migration), we treated BEAS-2B and H146 cells with the Hedgehog inhibitors cyclopamine and GDC-0449, then measured cell viability and migration. The results revealed that Hedgehog inhibition significantly decreased the viability and migration of both BEAS-2B and H146 cells. Consistent with previous findings, these results suggested that Hedgehog plays an important role in the development and migration of SCLC.

To date, the relationship between the Hedgehog and Akt pathways has not been clarified. It has been reported that the Akt signaling pathway inhibitor LY294002 could reduce Hedgehog morphogen activity in cells.\textsuperscript{15} Ke et al.\textsuperscript{16} found that Hedgehog signaling promoted epithelial–mesenchymal transition in ovarian cancer, which required the incorporation of the Akt signaling pathway. Moreover, Yoo et al.\textsuperscript{17} reported that lymphatic vessel density, which was closely related to lymph node metastasis, was higher in Hedgehog-positive tissues than in Hedgehog-negative tissues, suggesting that Hedgehog signaling requires PI3K–Akt signaling to promote lymphangiogenesis. Recently, Williamson et al.\textsuperscript{18} revealed that after inhibiting the Hedgehog signaling pathway or silencing Gli1, the viability and invasiveness of thyroid cancer cell lines were significantly reduced. These results suggested that the promotion of tumor migration and metastasis by the Hedgehog signaling pathway required the participation of Akt and c-Met. To investigate the interaction of Akt and Hedgehog in the development of SCLC, we analyzed the correlation between p-Akt and Hedgehog expression. As highlighted in Figure 2, p-Akt expression was positively correlated with Hedgehog expression. Furthermore, we analyzed the correlations of p-Akt and Hedgehog expression with patient survival rates and found that patients with p-Akt- and/or Hedgehog-negative tumors had a significantly higher survival rate than those with p-Akt- and Hedgehog-positive tumors. These results revealed that the interaction of Akt and Hedgehog was critical for the prognosis and growth of SCLC.

To further study the roles of PDPK1, p-Akt, and Hedgehog in SCLC, we cultured several SCLC cell lines (H82, H69, H446, H146, and H526) and measured protein expression using western blotting. Hedgehog expression was significantly higher in H146 and H446 cells than in control cells, whereas PDPK1 and p-Akt expression significantly upregulated in H146 cells. Based on this result, H146 was
used for the subsequent experiments. We examined the effects of PDPK1 siRNA transfection on p-Akt and Hedgehog expression, in H146 cells, finding that PDPK1 silencing significantly inhibited Hedgehog expression and decreased the p-Akt/Akt ratio. Furthermore, after H146 cells were transfected with PDPK1 siRNA or Akt siRNA, their proliferation and migration were significantly inhibited. These results clarified that PDPK1 could activate the Akt pathway and Hedgehog, then promote the proliferation and migration of SCLC cells (H146 cells). It has been reported that activation of the Akt pathway enhanced the Hedgehog–Gli pathway through multiple mechanisms. Park et al. found that Hedgehog was independently activated in SCLC cells. The activation of Smoothened, a molecule involved in the Hedgehog signaling pathway, promoted the clonogenicity of human SCLC cells in vitro and the development of SCLC in mice in vivo. Pharmacological blockade of Hedgehog signaling inhibited the growth of mouse and human SCLC cells, suggesting a crucial cell-intrinsic role of Hedgehog signaling in the development of SCLC. To further investigate the interaction between the PDPK1–Akt pathway and Hedgehog signaling, we transfected H146 cells with CA-PDPK1 and CA-Akt, then measured the changes of Hedgehog expression. The results (Figure 4d) illustrated that Hedgehog expression was significantly increased by PDPK1 or p-Akt overexpression. However, Hedgehog inhibition did not affect PDPK1 and p-Akt expression (Figure 4e–f). These results suggested that PDPK1 and Akt affected Hedgehog expression, but Hedgehog did not affect PDPK1 or p-Akt expression.

**Conclusion**

Our study suggests that the interaction between the PDPK1–Akt pathway and Hedgehog promotes the poor prognosis and growth of SCLC. Inhibition of the PDPK1–Akt pathway or Hedgehog signaling might serve as a therapeutic strategy to slow the progression of SCLC and delay recurrence in individuals with SCLC. The limitations of the study included the lack of an animal study; we should confirm the role of the PDPK1–Akt pathway and Hedgehog signaling in animal SCLC models in the near future.

**Declaration of conflicting interest**

The authors declare that there is no conflict of interest.

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**Authors’ contributions**

NW performed the cell experiments. YZ performed the clinical data analyses. ZW performed the statistical analyses. BH revised the manuscript. RH designed and supervised the study. All authors have read and approved the manuscript.

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