DOK7V1 influences the malignant phenotype of lung cancer cells through PI3K/AKT/mTOR and FAK/paxillin signaling pathways

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Abstract. Downstream of tyrosine kinase 7 transcript variant 1 (DOK7V1) is a docking protein mediating signal transduction between receptors and intracellular downstream molecules. Our previous study indicated that DOK7V1 was decreased in lung cancer and its lower expression was associated with a decreased survival rate. The 5-year overall survival rate for patients with lung cancer was 20.2 and 18.6% for high and low DOK7 expression, respectively; the 5-year disease-free survival rate for patients with lung cancer was 14.3 and 16.9% for high and low DOK7 expression, respectively. DOK7V1 inhibited proliferation and migration, but enhanced adhesion, of lung cancer cells. In the present study, the effect of DOK7V1 and its domains [pleckstrin homology (PH) and phosphotyrosine-binding (PTB) domain] on the malignant phenotype and associated signaling pathway in lung cancer cells was investigated. The results indicated that truncation of DOK7V1 domains (DOK7V1Δ-PH and DOK7V1Δ-PTB) inhibited the proliferation and migration of lung cancer cells which exhibited the same trend as DOK7V1, whereas DOK7V1Δ-PH and DOK7V1Δ-PTB exhibited different functions from those of DOK7V1 in cell matrix adhesion. Consistently, DOK7V1 overexpression in lung cancer cells suppressed the phosphoinositide 3-kinase (PI3K)/protein kinase B (AKT)/mammalian target of rapamycin (mTOR) signaling pathways, but activated the focal adhesion kinase (FAK)/paxillin signaling pathway. Taken together, these results indicate that DOK7V1 may inhibit proliferation and migration via negatively regulating the PI3K/AKT/mTOR signaling pathway, and increase adhesion by upregulating the FAK/paxillin signaling pathway in lung cancer cells.

Introduction

Lung cancer remains the leading cause of cancer mortality and has become an increasingly serious public health burden globally (1). Lung cancer is categorized into small cell lung cancer (SCLC) and non-SCLC (NSCLC) following pathological pattern. NSCLC accounts for ~80% of all lung cancer cases, including squamous cell carcinoma, adenocarcinoma, large cell carcinoma and adenosquamous cell carcinoma (2). NSCLC is often diagnosed at an advanced stage and the 5-year survival rate remains ~15% (3). Therefore, there is an urgent requirement to identify and screen for tumor-associated genes, further investigate the molecular mechanisms of NSCLC tumorigenesis and development, and identify novel biomarkers for early diagnosis and therapeutic targets of NSCLC. Tumor-associated genes rely on diverse signaling pathways to influence the proliferation, migration and invasion of tumor cells, and subsequently disseminate from primary tumors to metastasize through lymphatic and blood vessels. Understanding the molecular mechanisms underlying tumor metastasis has a critical significance in formulating treatment and early diagnosis. Receptor tyrosine kinases (RTKs) serve a vital function in regulating cellular process, including cell migration, proliferation, differentiation, survival and cell cycle regulation (4,5). Deregulation of RTKs includes chromosomal translocations, autocrine activation, overexpression of RTKs or gain-of-function mutations which occurs in cancer (6). The downstream of tyrosine kinase (DOK) protein family consists of seven members, mediating intracellular signaling transduction downstream of RTKs (7-9). DOK proteins share a topology domain characterized by a central phosphotyrosine-binding (PTB) domain, an N-terminal pleckstrin homology (PH) domain and Src homology 2 target motifs in the C-terminus (7,10). On the basis of their location, DOK proteins are divided into three subgroups: DOKs 1-3 are mainly expressed in hematopoietic tissues (11), DOKs 4-6 are primarily present within the nervous system (12,13), and DOK7 is predominantly in skeletal muscle and the heart, which
exhibits a distinct function compared with other members (14). Aberrant expression of DOK proteins has been identified in multiple malignancies. DOKs 1-3 co-operatively suppress aggressive histiocytic sarcoma (11). DOK2 was regarded as a biomarker of patients with gastric cancer with poor prognosis (15,16). In a mouse model, deletion of DOKs 1-3 resulted in abnormal proliferation of bronchoalveolar stem cells and alveolar type II cells, which subsequently progressed to lung cancer (7). In addition, a decreased expression level of DOK2 was observed in human lung cancer tissues, and forced DOK2 overexpression repressed proliferation of lung cancer cells (7).

DOK7, as a non-catalytic adaptor protein, is essential for the activation of the tyrosine kinase muscle-specific kinase (MuSK) and acetylcholine receptor clustering, which are indispensable for neuromuscular junctions (17,18). The function of DOK7 in cancer was revealed gradually. Demethylation of DOK7 decreased the proliferation and invasion of esophageal squamous cell carcinoma cells (19). The PTB domain is the region through which DOK7 interacts with the juxtanemembrane region of MuSK, although the PTB and PH domains are required for the activation of MuSK (18,20,21). The PH domain consists of ~120 amino acids, mediating intracellular and extracellular signaling, also acting as important constituents of the cytoskeleton (22-24). PTB was also able to contact the cell membrane, and regulate intracellular and extracellular signaling. For example, PH and PTB domains of DOK4 are required for DOK4 localization at the membrane (10). The DOK1 PH domain appears to be required for tyrosine phosphorylation of the protein and its normal localization to a subcellular membrane component (10).

In our previous study, we identified that DOK7 transcripts were decreased in lung cancer, and the lower DOK7 level was associated with poorer overall survival and progression-free survival (25). Overexpression of the DOK 7 transcript variant 1 (DOK7V1) limited the proliferation and migration, but enhanced the adhesion to the extracellular matrix, of lung cancer cells (25). However, the underlying molecular mechanism of DOK7V1 and the functions of its domains in lung cancer cells remain unknown. Therefore, the aim of the present study was to investigate the functions of its domains and the molecular mechanism underlying its involvement in lung cancer cells.

Materials and methods

Cell culture. Lung cancer A549, SKMES-1 and H3122 cell lines were purchased from the American Type Culture Collection (Manassas, VA, USA). A549 and SKMES-1 cells were routinely cultured in Dulbecco's modified Eagle's medium (Thermo Fisher Scientific, Inc., Waltham, MA, USA) and H3122 cells were cultured in RPMI-1640 medium (Thermo Fisher Scientific, Inc., Waltham, MA, USA), containing 5% CO₂ supplemented with 10% fetal bovine serum (Shanghai ExCellBio, Inc., Shanghai, China), in an incubator at 37°C with a 95% humidified atmosphere containing 5% CO₂.

Construction of DOK7V1 plasmid and transfection. DOK7V1-Flag and pEnter (Vector) plasmids were purchased from Vigene Biosciences, Inc. (Rockville, MD, USA). DOK7V1Δ-PH and DOK7V1Δ-PTB were constructed using a Q5® Site-Directed Mutagenesis kit (Without Competent Cells) (New England BioLabs, Inc., Ipswich, MA, USA). Purified DOK7V1 and control plasmids were transfected into A549 and SKMES-1 cells using an EasyJet Plus electroporator (EquiBio Ltd., Altrincham, UK). H3122 cells were transfected with DOK7V1, DOK7V1Δ-PH and DOK7V1Δ-PTB, respectively, using Neonect™ DNA transfection reagent (Neofectbiotech Co., Ltd., Beijing, China).

RNA isolation and reverse transcription-polymerase chain reaction (RT-PCR). Total RNA was isolated from A549, SKMES-1 and H3122 cells using TRIzol® reagent (Invitrogen; Thermo Fisher Scientific, Inc.). cDNA was generated from each 1 µg RNA sample using a QuantiTect Reverse Transcription kit (Qiagen GmbH, Hilden, Germany), according to the manufacturer's protocol. PCR was performed as described previously (26) with the following primer sequences: DOK7, 5'-ACTGGGCTGCGTCTTCTTCC-3' (forward) and 5'-TCGGACGATGCTCAACAG-3' (reverse); and GAPDH, 5'-GGCTGCTTTTAACTCTGGTA-3' (forward) and 5'-GACTGGTGCTAGTCTCTT-3' (reverse).

Western blot analysis. Cells were collected and lysed using Radioimmunoprecipitation Assay Lysis and Extraction Buffer (Thermo Fisher Scientific, Inc.), followed by centrifugation at 13,000 x g for 15 min at 4°C. Total protein concentrations in the supernatant were determined using a DC Protein Assay kit (Bio-Rad Laboratories, Inc., Hercules, CA, USA) and an ELx800 spectrophotometer (Bio-Tek Instruments, Inc., Winooski, VT, USA). Equal amounts (30 µg) of protein samples were separated by SDS-PAGE (10% gel) and blotted onto a nitrocellulose membrane. The membrane was blocked using 5% skimmed milk in Tris-buffered saline for 1 h at room temperature. Proteins were then probed using anti-human FLAG (1:1,000; cat. no. F3040; Sigma; Merck KGaA, Darmstadt, Germany), anti-DOK7 antibody (1:1,000; cat. no. ab75049; Abcam, Cambridge, MA, USA), anti-protein kinase B (AKT) antibody (1:1,000; cat. no. 2920; Cell Signaling Technology, Inc.), anti-phospho-p-AKT (p)-AKT (1:1,000; cat. no. 4051; Cell Signaling Technology, Inc.), anti-phosphoinositide 3-kinase (PI3K) antibody (1:500; cat. no. sc-365290; Santa Cruz Biotechnology, Inc.), anti-mammalian target of rapamycin (mTOR) antibody (1:1,000; cat. no. ab2732; Abcam), anti-p-mTOR antibody (1:1,000; cat. no. 2971; Cell Signaling Technology, Inc.), anti-Rho-associated protein kinase (Rock) antibody (1:1,000; cat. no. sc-17794; Santa Cruz Biotechnology, Inc.), anti-p-Rock antibody (1:1,000; cat. no. ab2732; Abcam), anti-focal adhesion kinase (FAK) antibody (1:1,000; cat. no. sc-1688; Santa Cruz Biotechnology, Inc.), anti-p-FAK antibody (1:500; cat. no. sc-11766; Santa Cruz Biotechnology, Inc.), anti-paxillin antibody (1:1,000; cat. no. sc-635174; Santa Cruz Biotechnology, Inc.), anti-p-paxillin antibody (1:500; cat. no. sc-635020; Santa Cruz Biotechnology, Inc.), anti-human GAPDH antibody (1:500; cat. no. sc-47724; Santa Cruz Biotechnology, Inc.) and corresponding horseradish peroxidase (HRP)-conjugated goat anti-rabbit and goat anti-mouse immunoglobulin G secondary antibodies (1:3,000; cat. nos. ZB-2301M and ZB-2305; OriGene Technologies, Inc.).
Beijing, China), respectively. Protein bands were visualized and analyzed using Luminata Forte western blot HRP substrate (Merck KGaA) and a UV imager (Uvitec, Inc., Cambridge, UK).

**Immunofluorescence.** Cells on glass coverslips were rinsed with PBS three times, and then fixed with 4% paraformaldehyde for 20 min at room temperature. Anti-human FLAG was diluted 1:1,000 in blocking buffer (1% bovine serum albumin in PBS), prior to adding to the coverslips and incubating for 1 h at room temperature. Following three washes with PBS, the coverslips were incubated using Alexa Fluor® 488-conjugated secondary antibodies (1:100; cat. no. A‑21205; Thermo Fisher Scientific, Inc.) for 45 min at room temperature. Following three further washes with PBS, nuclei were stained with DAPI (1:500,000; cat. no. D‑1306; Thermo Fisher Scientific, Inc.) for 2 min at room temperature, followed by three further washes with PBS. The coverslips were mounted on glass slides and incubated at room temperature overnight. Immunofluorescence images were visualized using a confocal microscopy system (Leica TCS SP8; Leica Microsystems GmbH, Wetzlar, Germany) with a 63X oil-immersion objective.

**In vitro cell proliferation assays.** A total of 3,000 cells in 200 µl culture medium was added to each well of a 96-well plate. A total of five plates were used to obtain cell density readings following incubation at 37°C for up to 4 days. Following incubation, the medium was removed and 100 µl 10% Cell Counting Kit-8 (Dojindo Molecular Technologies, Inc., Kumamoto, Japan) was added for 1 h at 37°C. Subsequently, the absorbance was determined at a wavelength of 450 nm using a spectrophotometer (BioTek Instruments, Inc.).

**In vitro tumor cell migration assay.** Cells (1x10^6 cells/well) were seeded in a 6-well plate and cultured in an incubator at 37°C overnight. The cell monolayer was scratched with a 200-µl pipette tip to create an artificial wound prior to washing twice with PBS to remove floating cells. The migration of cells was monitored and recorded every 6 h using an inverted light microscope at x10 and x20 magnification for 24 h. The wound distance was determined and analyzed using ImageJ software (version 1.62; National Institutes of Health, Bethesda, MD, USA).

**In vitro tumor cell Matrigel adhesion assay.** A 96-well plate was coated with 5 µg Matrigel/100 µl/well and air-dried overnight at room temperature. Following rehydration, 20,000 cells/200 µl/well were seeded into each well for 40 min. Following incubation, medium was removed and the plate was washed three times with PBS to remove the non-adherent cells. Adherent cells were fixed with 4% formaldehyde for 30 min prior to being stained with 0.5% crystal violet solution for 10 min at room temperature. Subsequently, crystal violet was dissolved in 10% acetic acid and the absorbance at 570 nm was determined using a spectrophotometer (BioTek Instruments, Inc.).

**Gene set enrichment analysis (GSEA).** The association between biological processes/pathway, phenotypes and DOK7 expression level was analyzed using the GSEA program (version 2.2; www.broad.mit.edu/gsea). Samples from The Cancer Genome Atlas (TCGA) datasets were separated into high or low DOK7 expression groups using the median as the cut-off. GSEA was used to calculate a pathway Enrichment Score that assessed whether genes from a predefined gene set
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of PI3K/AKT/mTOR and FAK/paxillin signaling pathways were enriched among the high- (or low-) ranked genes or distributed randomly. Default settings were utilized when using software applications. Significance was determined by permutation analysis (1,000 permutations) and calculation of the false discovery rate (FDR). A gene set was considered to be significantly enriched when the FDR score was <0.05.

Statistical analysis. Data are presented as the mean ± standard deviation. Experimental procedures were repeated independently at least three times. Statistical analysis was performed using a two-sided Student's t-test for two-group comparisons and by one-way ANOVA, followed by a Bonferroni post hoc test, for multiple group comparisons. All statistical analyses were performed using Prism (version 5; GraphPad Software, Inc., La Jolla, CA, USA). P<0.05 was considered to indicate a statistically significant difference.

Results

Overexpression of DOK7V1 has an inhibitory effect on proliferation and migration, but a positive effect on adhesion in lung cancer cells. To select the optimal cell lines for further investigation, the expression of DOK7V1 in A549, SKMES-1 and H3122 cells was determined using RT-PCR analysis. The stable transfection of DOK7V1 in the lung cancer cell lines was verified by RT-PCR and western blot analysis (Fig. 1). To further confirm the effect of DOK7V1 on malignant phenotypes of lung cancer cells, the experiments were performed in A549 and SKMES-1 cells. The results indicated that overexpression of DOK7V1 significantly decreased the proliferation and migration, but significantly increased the adhesion, of A549 and SKMES-1 cells (Fig. 2).

PH domain and PTB domain determined the distribution of DOK7V1 in membrane and cytoplasm. DOK7V1 contains two functional domains: A PH domain and a PTB domain. To investigate the structural determinant regulating the intracellular distribution and expression of DOK7V1, as well as the biological function, two DOK7V1 truncated fragments were constructed based on the wild-type protein, DOK7V1Δ-PH and DOK7V1Δ-PTB (Fig. 3A). The immunofluorescence imaging results indicated that wild-type DOK7V1 was located and expressed in the membrane and cytoplasm. The truncated versions of DOK7V1 without the two functional domains led to a shift in the localization of DOK7V1 protein from the cytoplasm to be perinuclear (Fig. 3B). The results indicated that these two functional domains determined the distribution of DOK7V1 in the membrane and cytoplasm, which may influence the function of DOK7V1 in lung cancer cells.

Effect of the PH domain and PTB domain of DOK7V1 on the proliferation, migration and adhesion of lung cancer cells.
To further assess the biological function of DOK7V1Δ-PH and DOK7V1Δ-PTB, these truncated versions were stably transfected into H3122 cells and their expression was determined by western blotting. DOK7V1, DOK7V1Δ-PH and DOK7V1Δ-PTB were identified to exhibit increased expression in stably transfected cells compared with the corresponding control cells (Fig. 4A). It was identified that cell proliferation was significantly decreased from the third day, and migration was significantly decreased at 18 h, in cells singly overexpressing DOK7V1Δ-PH and DOK7V1Δ-PTB compared with vector control cells, which exhibited the same trend as DOK7V1 (Fig. 4B and 4C). The adhesion assay revealed an increased rate in DOK7V1-overexpressing cells compared with the control group; however, the DOK7V1Δ-PH and DOK7V1Δ-PTB truncated versions lost the tumor-promoting effects identified in DOK7V1 (Fig. 4D).
No differences exist between the two truncated versions and DOK7V1 in inhibiting PI3K/AKT/mTOR signal pathway phosphorylation. To further study the signal pathways in which DOK7V1 is involved, the protein levels and phosphorylation levels of AKT, PI3K, mTOR and Rock in A549 cells were determined. Decreased levels of AKT, PI3K and Rock phosphorylation were observed in A549 cells with DOK7V1 overexpression, whereas the total AKT, PI3K and Rock protein levels were similar in comparison with their corresponding controls (Fig. 5A). mTOR phosphorylation and the total mTOR protein levels were decreased in A549 cell lines with DOK7V1 overexpression (Fig. 5A). The AKT phosphorylation status was investigated in H3122 cells transfected with various DOK7V1 constructs. Wild-type DOK7V1 overexpression significantly arrested AKT activation, and this inhibition was similar in cells expressing the DOK7V1Δ-PH and DOK7V1Δ-PTB truncated versions (Fig. 5B). Taken together, these results indicated that the two truncated version of DOK7V1 did not eliminate the inhibitory function of DOK7V1 in activating AKT.

DOK7 overexpression is negatively associated with activation of the PI3K/AKT/mTOR signaling pathway and is positively associated with adherent ability. To further analyze the function of DOK7 expression in PI3K/AKT/mTOR signaling in lung cancer, GSEA was performed using a lung cancer dataset from TCGA. The lung cancer samples were separated according to the median DOK7 level in the specimens into high or low expression groups. Enrichment plots of GSEA demonstrated that the gene set of PI3K/AKT/mTOR signaling was primarily enriched in the low DOK7 expression group (Fig. 7A). The association between DOK7 expression and activation of FAK/paxillin signaling in lung cancer was identified using GSEA. As presented in Fig. 7B, the gene signatures of adhesion, which emphasized the importance of FAK in its description, were enriched in patients with high DOK7 expression. These results further suggested that, in lung cancer samples, DOK7 expression was negatively associated with PI3K/AKT/mTOR signaling activation and is positively associated with activation of the FAK/paxillin signaling pathway.

Discussion

In our previous study, we preliminarily studied the function of DOK7V1 in lung cancer cells and the association with the prognosis of patients with lung cancer (25). In the present study, the focus was on the mechanisms and functions of DOK7V1 and its domains in the lung cancer cells. The results of the present study suggested a hitherto unknown regulatory mechanism of DOK7V1 in lung cancer cells, which influences...
The functions of the DOK protein family in lung cancer have gradually been identified. DOKs 1-3 have been identified to be absent from or decreased in lung cancer (23). In mouse models, a lack of DOKs 1-3 promoted the proliferation of bronchoalveolar stem cells and alveolar type II cells, which resulted in the development of lung cancer, particularly in triple knockout mouse models (11). Decreased DOK2 levels were identified in primary lung adenocarcinomas and lymph node metastases, whereas no difference in expression of DOK1 and DOK3 was identified in lung cancer compared with normal tissues, although a decreased DOK3 level was observed in lymph node metastasis tissues when compared with its expression in primary tumor (7). DOK2 was identified as a tumor suppressor gene of lung cancer, since DOK2 was associated with epidermal growth factor mutation to promote lung cancer (27). DOK7 was the primary focus in the present study. In breast cancer, DOK7 is downregulated owing to the hypermethylation of its CpG islands (22). The results of this study indicated that DOK proteins may serve an inhibitory function in cancer development, and gene deletion or hypermethylation of DOK proteins may occur in cancer cells. In addition, our previous study identified a lower expression level of DOK7 in lung cancer and DOK7V1 overexpression, resulting in a decrease in proliferation and migration, and an increase in adhesion in lung cancer cells (25). In the present study, the experiments in which DOK7V1 inhibited proliferation and migration, but promoted adhesion in other lung cancer cell lines were repeated to confirm the effect of DOK7V1 on the malignant phenotype. Furthermore, the change of invasion following transfection of lung cancer cells with DOK7V1 was investigated, but no difference between the test group and control group was identified.

DOK7V1 is a 1,515-residue protein, containing a PH domain, a PTB domain and C-terminal sites of tyrosine...
phosphorylation. DOK7 is a substrate of MuSK and also an activator of its kinase activity, and is therefore an important protein for forming the vertebrate neuromuscular junction, and may facilitate trans-autophosphorylation of the kinase activation loop via a dimeric arrangement of its PH and PTB domains (28). A truncation of the domains may lead to changes in protein localization. A recent study revealed that DOK7 bound to membranes containing phosphatidylinositol phosphates (PIPs) through PH domain binding, facilitating local clustering of PIP molecules in the bilayer (29). In the present study, the PH and PTB domains were identified to determine the localization of DOK7V1 in the membrane and cytoplasm, as revealed by the nuclear membrane distribution of PH- and PTB-truncated DOK7V1 proteins (DOK7V1Δ-PH and DOK7V1Δ-PTB). Does this change in location have an effect on the function of the protein? In the present study, in vitro functional experiments indicated that DOK7V1 overexpression inhibited H3122 cell proliferation, motility and adhesion, and DOK7V1Δ-PH and DOK7V1Δ-PTB have the same effect on proliferation and motility as DOK7V1, but with the enhancement of adhesion eliminated.

Dysregulation of the PI3K/AKT/mTOR signaling pathway has been implicated in the cancerous migration, proliferation and poor prognosis of various types of cancer, including lung cancer (30-32). Tumor suppressor genes decreased the proliferation and migration of NSCLC by inhibiting the PI3K/AKT/mTOR signaling pathway (33). The in vitro results of the present study indicated that DOK7V1 inhibited the phosphorylation of AKT, PI3K, mTOR and ROCK in A549 cells. The two truncated versions of DOK7V1 also could not activate AKT in H3122 cells. PI3K/AKT signaling appears to rely on DOK7 in the cytoplasm or associated with the RTK. The weakened PI3K/AKT signaling may be caused by the retention of truncated DOK7V1 at the perinuclear area. Further analysis indicated that an increased level of DOK7V1 was negatively associated with the activation of PI3K/AKT/mTOR signaling. According to these results, we hypothesize that DOK7V1 overexpression may inhibit the proliferation and migration by suppressing the activation of PI3K/AKT/mTOR signaling pathway. However, a clear conclusion cannot be made. Investigation of whether DOK7V1 decreases the proliferation and migration of lung cancer cells following inhibition of the PI3K/AKT/mTOR signaling pathway is required in future studies.

Hyperactivation of the FAK/paxillin signaling pathway was identified to be significantly associated with cell adhesion (34-36). FAK and paxillin dynamics serve an essential function in regulating the adhesion of various cells (37). In the present study, an increased adhesion rate of cells overexpressing DOK7V1 compared with that of the control group was revealed; however, the DOK7V1Δ-PH and DOK7V1Δ-PTB truncated versions eliminated the tumor-promoting effects observed in DOK7V1. Conversely, FAK and paxillin were also weakened, which provides further evidence of the association between its location and function. Furthermore, it was confirmed that the gene signatures associated with FAK/paxillin signaling activation were enriched in patients with high DOK7 expression. These results further suggested that in lung cancer specimens, DOK7 expression was positively associated with activation of the FAK/paxillin signaling pathway. Certainly, further research is required to reach a clear conclusion.

In summary, it was identified that DOK7V1 downregulation is associated with poor prognosis of patients with lung cancer. The truncations of the DOK7V1 domains appeared to have effect on the localization of the protein. DOK7V1 overexpression reversed the malignant phenotypes of H3122 cells, including proliferation, migration and adhesion. DOK7V1Δ-PH and DOK7V1Δ-PTB were able to eliminate the function of DOK7 in adhesion, but not in proliferation and migration. The two truncated version retained the inhibitory effect of DOK7V1 on AKT activation, but inactivated the enhanced effect of DOK7V1 on FAK and paxillin. We hypothesize that DOK7V1 may inhibit proliferation and migration via negatively regulating the PI3K/AKT/mTOR signaling pathway and increasing adhesion by upregulating the FAK/paxillin signaling pathway in lung cancer cells. The results of the present study provide a novel basis to improve our understanding of the pathogenesis of lung cancer.

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Availability of data and materials

The datasets used and/or analyzed during the present study are available from the corresponding author on reasonable request.

Authors' contributions

WGJ, GC, HZ and LY designed the study. GC, HY, LY and SL prepared and revised the paper. All authors had final approval of the submitted and published versions of the paper.

Ethics approval and consent to participate

Not applicable.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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