

Thymidine kinase 1 silencing retards proliferative activity of pancreatic cancer cell via E2F1-TK1-P21 axis

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Abstract

Objectives: Thymidine kinase 1 (TK1) is one of the salvage enzymes engaged in the synthesis of DNA. Although a pro-carcinogenic role of TK1 has been reported in various types of cancers, its role in pancreatic ductal adenocarcinoma (PDAC) is still unknown. The study is aimed to elaborate the function of TK1 in PDAC and the potential mechanisms in the following study.

Materials and methods: TK1 expression was analysed by immunohistochemistry, real-time PCR and Western blot, and its relationship with clinicopathological characteristics of PDAC patients was further investigated. To verify the function of TK1 and potential mechanism, TK1 siRNA was used to transfect PDAC cells and performed a series of assays in cell and animal models.

Results: The level of TK1 expression was higher in cancerous tissues compared with matched adjacent tissues. TK1 overexpression was associated with progression of PDAC and poor prognosis. Knockdown of TK1 could suppress cell proliferation via inducing S phase arrest mediated by upregulation of P21. Further mechanism investigation suggested that transcription factor E2F-1 could directly regulate the TK1 and promote tumour proliferation.

Conclusions: The results suggested that TK1 might be involved in the development and progression of PDAC by regulating cell proliferation and show that TK1 may work as a promising therapeutic target in patients with PDAC.

1 | INTRODUCTION

Pancreatic ductal adenocarcinoma (PDAC) is one of the most aggressive digestive malignancies and is the fourth most common cause of cancer-related death in the United States. Patients diagnosed with PDAC have a poor prognosis with an overall 5 year survival rate of only 8% and a median survival of 6 months.¹ Although the morbidity of this malignant tumour has been increased over the last several years, there is no obvious improvement in the survival rate. Surgical resection may prolong survival and provide a unique curative option. Unfortunately, the surgical resection rate is <20% because of difficulty for diagnosis

at early stage.² Therefore, an increasing understanding of the complex molecular basis of PDAC and exploring novel therapeutic targets may contribute to the clinical treatment of this aggressive malignancy.

Thymidine kinase 1 (TK1), a cell cycle-dependent marker, is a pyrimidine salvage enzyme involved in the synthesis and repair of DNA. Thus, TK1 is closely correlated with cell proliferation and cell cycle activity.^{3,4} It is reported that wide distribution and overexpression of TK1 are found in most neoplastic cells.⁵ Progression and development of cancer are complex biological processes. Although the causes and exact mechanisms are still unknown, the anomaly of the cell cycle and abnormal regulation of cell proliferation are recognized to be elementary events in any malignant transformation. Numerous studies have documented the TK1 protein expression patterns could act as

XZ, CS and YP contributed equally to this work.

clinicopathological predictive markers in various types of cancers like breast cancer,⁶ kidney cancer,^{7,8} lung cancer,⁹ and gastrointestinal cancer;¹⁰ however, little has been explored about the expression and biological function of TK1 in PDAC. To further identify the role of TK1 in PDAC, we evaluated the expression patterns in PDAC cell lines and tissues and investigated the possible molecular mechanism.

2 | MATERIALS AND METHODS

2.1 | Tissue microarray

This study was approved by the Ethics Committee of the First Affiliated Hospital of Nanjing Medical University. TK1 mRNA expression of 165 patients with pancreatic cancer was downloaded from The Cancer Genome Atlas (TCGA, USA) data portal (<http://cancergenome.nih.gov/>). Seventy-two pancreatic cancer tissues and matched adjacent tissues microarray were obtained from Shanghai Zhuoli Biotechnology Co., Ltd (Zhuoli Biotechnology Co, Shanghai, China). All tissue samples applied in this study were fixed in formalin, embedded paraffin and further used for IHC.

2.2 | Immunohistochemical staining and analysis

Formalin-fixed paraffin-embedded tissues were sliced into 5 μm thickness, and then deparaffinized with xylene, and gradually rehydrated in graded ethanol solutions. Next, the slides were incubated with the anti-TK1 antibody (1:150; Abcam, Cambridge, MA, UK) at 4°C overnight, followed by incubation with biotinylated goat anti-rabbit serum streptavidin-peroxidase conjugate at room temperature for 30 minutes. Finally, the sections were developed with diaminobenzidine and counterstained with haematoxylin. Two independent pathologists who were blinded to assess the positivity and intensity of the tissue sections. The percentage of TK1-positive staining cells was scored as 0 (0%), 1 (1%), 2 (2%) ... 99 (99%), 100 (100%). The staining intensity was given 0 (none), 1 (low), 2 (medium) and 3 (high) scores, respectively. A combination of the two scores was calculated by the following formula for all cases: IHC score = positive rate score \times intensity score.¹¹

2.3 | RNA isolation and quantitative real-time PCR

Total RNA was isolated from cell pellets using TRIzol reagent (Life Technologies, Carlsbad, CA, USA) according to the manufacturer's instruction. The total RNA was then reverse-transcribed with PrimeScript RT Master Mix (Takara, Tokyo, Japan). For detection of the mRNA expression, qPCR was performed on the Step One Plus Real-Time PCR system (Applied Biosystems, Carlsbad, CA, USA) with FastStart Universal SYBR Green Master. Three independent experiments were performed with the following condition: 95°C for 10 minutes, 40 cycles of denaturation at 95°C for 15 seconds and 60°C for 1 minute. β -actin was used as an internal control. The specific primers of human TK1, E2F-1 and β -actin were designed by Primer Premier 5 and checked by Oligo 6. The $2^{-\Delta\Delta\text{Ct}}$ method was used to analyse the relative expression.

2.4 | Cell lines and cell culture

Human PDAC cell lines (MiaPaCa-2, BxPc-3, CFPAC-1, SW1990 and PANC-1) and the normal human pancreatic ductal cell line hTERT-HPNE were all purchased from American Type Culture Collection (ATCC). hTERT-HPNE cells were cultured in keratinocyte serum-free medium supplemented with epidermal growth factor and bovine pituitary extract according to the recommendation of ATCC. PDAC cell lines were cultured in Dulbecco's modified Eagle's medium (DMEM; Gibco, Thermo Fisher Scientific, New York, USA) supplemented with 10% foetal bovine serum (FBS), 100 U/mL penicillin and 100 $\mu\text{g}/\text{mL}$ streptomycin at 37°C in a humidified atmosphere containing 5% CO_2 .

2.5 | Transfection of small interfering RNA

The siRNA and the scrambled negative control of TK1 and E2F-1 used in this study were synthesized and purchased from GenePharma (Shanghai, China). The siRNA and the scrambled negative control sequences were as follows: TK1 siRNA#1, 5'-GGUGAUAAGUAUGCCAAATT-3' (sense) and 5' UUUUGGCAUACUUGAUCACCTT-3' (antisense); TK1 siRNA#2, 5'-CCU UCCAGAGGAAGCCAUUTT-3' (sense) and 5'-AAUGGCUUCCUCUG GAAGGTT-3' (antisense); TK1 siRNA#3, 5'-GGC CG GA CAACA AAGAGAATT-3' (sense) and 5'-UUCUCUUUGUUG UCCGG CCTT-3' (antisense); negative control, 5'-UUCUCCGAACGUGUCACGU TT-3' (sense) and 5' ACGUGACACGUUCGGAGAATT-3' (antisense). PANC-1 and CFPAC-1 cells were respectively transfected with siRNA (final concentration: 20 nmol/L) and the scrambled negative control using jetPRIME[®] transfection reagent (Polyplus, ILLKIRCH, France). The efficiency of all siRNAs was detected by qRT-PCR and Western blot.

2.6 | Western blot analysis

A Total Protein Extraction Kit (Keygen BioTECH, Nanjing, China) was used to extract protein from collected cells which contains PMSF, protease inhibitors and phosphatase inhibitors. Extracted protein was mixed with 5 \times SDS-PAGE and further boiled for 5 minutes. Standard methods of Western blot were used to analyse each protein expression,¹² β -actin protein was used as an endogenous reference.

2.7 | Proliferation assay

2.7.1 | Cell proliferation assay

The PDAC cells were seeded into 96 well plates at a density of 2.5×10^3 cells per well after co-incubation with siRNA for TK1. 10 μL of Cell Counting Kit-8 reagent (Dojindo, Tokyo, Japan) and 100 μL complete medium were mixed and added to each well to detect the cell viability. After incubation in dark at 37°C for 3 hours, each well was analysed at 450 nm absorbance.

2.7.2 | Clone formation assay

Human pancreatic cancer cell lines treated with or without siRNA of TK1 were plated at a density of 1×10^3 cells/well in 6 well plates. Cells were cultured for 10 days in DMEM supplemented with 10% FBS, penicillin (100 U/mL) and streptomycin (100 μ g/mL) in a 37°C humidified atmosphere of 5% CO₂. The colonies were stained with crystal violet solution, and then the number was counted.

2.7.3 | Cell cycle analysis

PANC-1 and CFPAC-1 cells were harvested after transfection of siRNA for TK1 and negative controls for 72 hours and fixed with 70% ethanol overnight at 4°C, and then incubated with 500 μ L PI/RNase staining buffer at 37°C for 15 minutes. Cell cycle was analysed by flow cytometer (Gallios, Beckman Coulter, Brea, CA, USA).

2.7.4 | Apoptosis detection

Pancreatic ductal adenocarcinoma cells were harvested after transfection for 48 hours and resuspended in 500 μ L Annexin V binding buffer. 5 μ L PE Annexin V and 7-AAD Viability Staining Solution were used to stain tumour cells. After incubation in dark at room temperature for 15 minutes, cells were analysed by flow cytometer (Gallios, Beckman Coulter).

2.7.5 | Transwell assay

Cell migration and invasion were respectively performed using Matrigel-coated and non-Matrigel-coated transwell chambers purchased from BD Biosciences (Franklin Lakes, NJ, USA). For both invasion and migration assays, 2.5×10^4 cells in 200 μ L serum-free DMEM after transfection with siRNA or negative controls were seeded in the upper chambers, the lower chambers were filled with 10% FBS DMEM. After cultured 24 and 48 hours, cells on the underside of the chamber were counted.

2.7.6 | Wound-healing assay

Wound-healing assays were performed in 6 well plates to assess cell migration. A scratch was made 24 hours after transfection. The cells were then cultured in serum-free DMEM for 24 hours. To assess migrated cells and wound area, pictures were taken at 0, 12 and 24 hours.

2.7.7 | Chromatin immunoprecipitation (ChIP)

Chromatin immunoprecipitation (ChIP) was carried out with the Magna Chromatin Immunoprecipitation kit (Millipore, Darmstadt, Germany). Immunoprecipitations were performed with anti-E2F-1 antibodies. ChIP products and genomic input DNA were analysed by qRT-PCR (E2F-1 primer sequences: forward, 5'-CCTGGAATTATCTATCTTGC

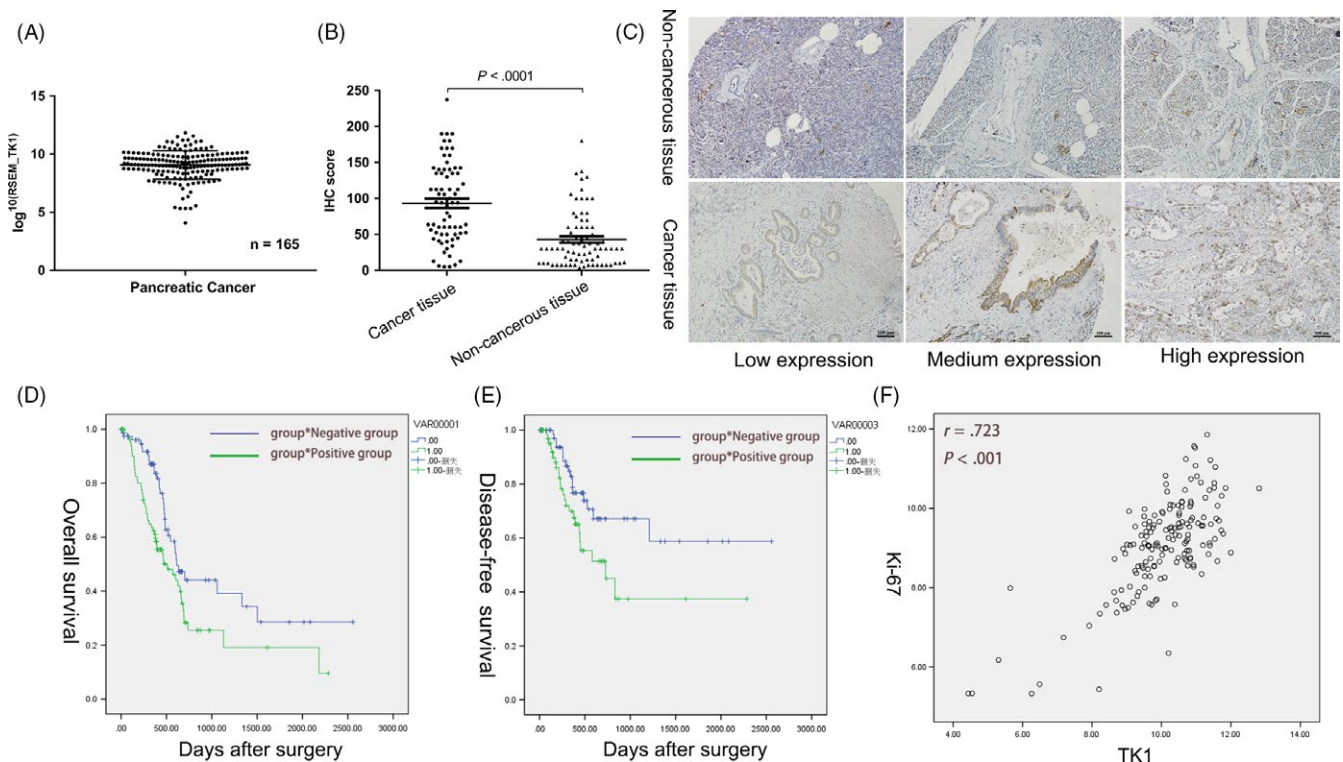


FIGURE 1 Expression and roles of thymidine kinase 1 (TK1) in pancreatic ductal adenocarcinoma (PDAC). A, TK1 mRNA expression in 165 PDAC samples obtained from TCGA was shown. B, C, Higher expression of TK1 was observed in pancreatic cancer than adjacent non-tumour tissues by IHC. D, E, Patients with relatively high expression of TK1 have a shorter overall survival and disease-free survival. F, The level of TK1 expression was positively correlated with Ki67

TABLE 1 Association of TK1 expression with clinicopathological features of PDAC

		TK1		
		Low	High	P
Histology stage	Well or moderate	61	55	.253
	Poor	21	28	
N stage	Absent	22	26	.608
	Present	60	57	
T stage	T1 or T2	18	9	.042*
	T3 or T4	64	74	
TNM stage	I-IIA	19	25	.313
	IIB-IV	63	58	

PDAC, pancreatic ductal adenocarcinoma; TK1, thymidine kinase 1; TMN, tumour-node-metastasis. *P<.05

AGAA-3'; reverse, 5'-AATGAATGAA ATGCAGCTTTTAAAC-3', TK-1 primer sequences: forward, 5'-GCAATCCACCTACTCTG-3'; reverse, 5'-CTCCAAGGCC GTCCCG CAGT-3'). Data of ChIP are shown as the percentage of input normalized to control purifications.

2.7.8 | Animal studies

Animal ethics was approved by Animal Core Facility of Nanjing medical university. Five-week-old male BALB/c nude mice were obtained from Model Animal Research Center of Nanjing University. To evaluate the proliferative function of TK1, each mouse was implanted subcutaneously with 1×10^6 PANC-1 cells. A total of 12 mice were randomly divided into two groups after 4 weeks of tumour injection. siRNA of TK1 or negative control injected into the tumour once every 48 hours. 4 weeks later, the nude mice were sacrificed, and the tumours were excised and photographed.

2.8 | Statistical analysis

Statistical analysis was performed by SPSS 20.0 software. A two-tailed Mann-Whitney *U* test and Student's *t* test were used to analysing the

TABLE 3 Relationship between TK1 expression and patient characteristics

Characteristics	TK1 IHC score	
	Standard deviation	P
Gender		
Male	48.44 ± 43.524	.640
Female	36.46 ± 29.277	
Age (year)		
≤65	44.33 ± 35.860	.631
>65	36.46 ± 29.277	
TNM stage		
I-IIA	66.44 ± 43.880	.034
IIB-IV	39.60 ± 37.200	
Diameter (cm)		
≤2	72.34 ± 48.937	.051
>2	40.96 ± 37.220	
Location		
Head	40.14 ± 35.841	.174
Body, tail	56.45 ± 47.449	
T stage		
T1 or T2	44.65 ± 40.563	.724
T3 or T4	42.50 ± 30.856	
N stage		
Absent	49.09 ± 41.213	.091
Present	33.27 ± 33.482	
M stage		
Absent	45.91 ± 40.858	.631
Present	28.33 ± 12.910	
Blood vessel invasion		
Absent	49.60 ± 45.346	.572
Present	37.62 ± 29.585	
Histological grade		
I/I-II/II	38.38 ± 39.396	.009
II-III/III	59.17 ± 36.754	

TABLE 2 Univariate and multivariate analyses

	Univariable analysis			Multivariable analysis		
	HR	95% CI	P	HR	95% CI	P
TK1	1.734	1.116-2.693	.014	1.588	1.005-2.510	.048
Age	1.456	0.942-2.250	.091			
Gender	0.934	0.608-1.434	.755			
HS	1.556	0.990-2.445	.055			
N stage	2.001	1.193-3.358	.009	1.113	0.352-3.5166	.856
T stage	2.396	1.186-4.838	.015	1.287	0.585-2.832	.53
TMN stage	2.323	1.333-4.046	.003	1.806	0.513-6.357	.357

CI, confidence interval.

statistical differences of proliferative activity and motility of the cells. Survival analysis was assessed by Kaplan–Meier plots. Cox proportional hazard regression model was used to evaluate the independent prognostic factors. Results were presented as the mean \pm SD. A significant difference was considered statistically with P -value $< .05$.

3 | RESULTS

3.1 | Expression of TK1 in PDAC

Generally, expression of TK1 mRNA in PDAC tissues ($n = 165$) was observed based on the data provided by TCGA (Figure 1A). The correlation between TK1 expression and the clinicopathologic characteristics in PDAC from all 165 patients in TCGA were shown in Table 1. It was shown that high level of TK1 was significantly correlated with higher T staging and positively correlated with Ki67 expression (Figure 1F). Furthermore, the Kaplan–Meier survival curves showed that PDAC patients with high level of TK1 have a shorter survival (Figure 1D,E). Cox proportional hazard regression model suggested that TK1 was an independent prognostic factor for PDAC patients (Table 2).

To further detect the TK1 protein expression in PDAC, 72 pancreatic cancer tissues and matched adjacent tissues were analysed, which revealed that protein levels of TK1 were significantly higher in cancerous tissues compared to that in adjacent non-cancerous tissues (Figure 1B,C). We also analysed the association between TK1 expression and clinicopathological characteristics in these patients, and TK1 overexpression was correlated with the TNM stage and histological grade (Table 3).

3.2 | Knockdown of TK1 expression by siRNA

TK1 expression was also measured in five PDAC cell lines (MiaPaCa-2, SW1990, BxPc-3, CFPAC-1 and PANC-1) and a normal human pancreatic ductal cell line (hTERT-HPNE) by qRT-PCR and Western blot. Compared to hTERT-HPNE, TK1 expression was higher in all cancer cell lines (Figure 2A,B). CFPAC-1 and PANC-1 were chosen to further study due to the relatively higher expression of TK1. CFPAC-1 and PANC-1 were used to conduct siRNA-related assays. The interference efficiency of three siRNAs (labelled as siRNA#1, siRNA#2 and siRNA#3) was detected at both mRNA and protein levels. siRNA#3 with the highest efficiency was chosen for further studies (Figure 2C,D).

3.3 | Effects of gene silencing of TK1 on cell proliferation, apoptosis and cell cycle

We supposed that TK1 played a vital role in the proliferation of PDAC cells because of its correlation with Ki67. Therefore, CCK-8 and clone formation assays were performed to investigate the effect of TK1 on PDAC cell proliferation. As shown in Figure 3, after transfection with siRNA of TK1, the proliferation of PDAC cells was remarkably decreased than the control group. Moreover, the cell clone formation assay showed the similar result. The number of cell colonies was decreased significantly in TK1-knockdown cells group than the control group.

To investigate the effect of TK1 knockdown on proliferation of PDAC cells in vivo, a xenograft model of subcutaneous tumour in

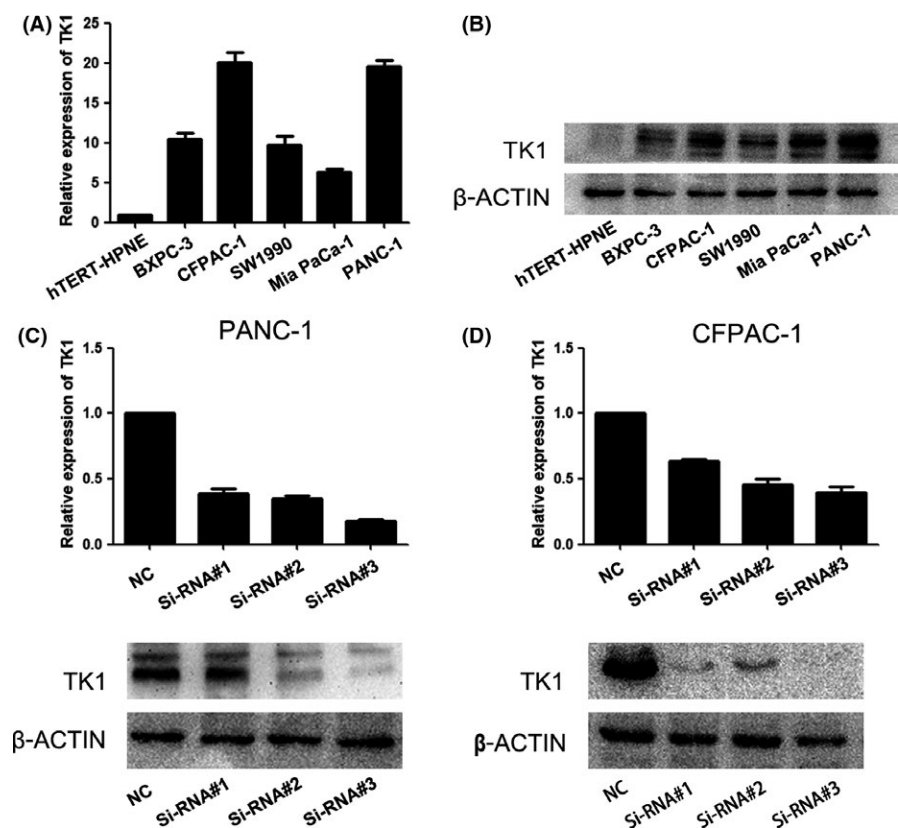


FIGURE 2 Knockdown of thymidine kinase 1 (TK1) by siRNA. A, B, Relative expression of TK1 in different PDAC cell lines was detected by Western blot. C, D, The different interference efficiency of three siRNAs for TK1 was evaluated by qRT-PCR and Western blot. Each error bar indicates the variation between the means of three independent experiments

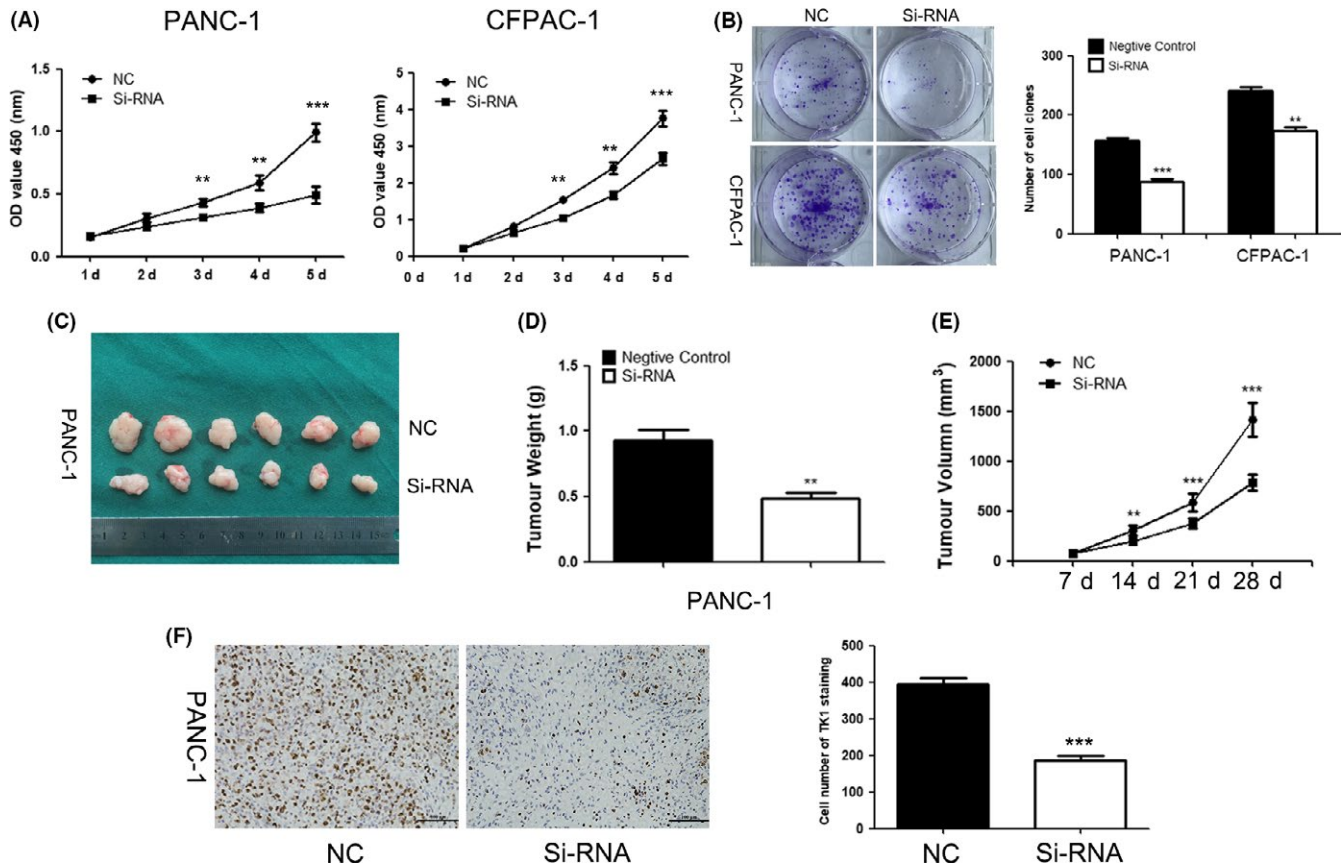


FIGURE 3 Effects of thymidine kinase 1 (TK1) downregulation on cell proliferation (A) CCK-8 assay showed that inhibition of TK1 expression suppressed PDAC cell proliferation. (B) The colon formation assay was performed to verify that gene silencing of TK1 suppressed cell proliferation. (C) Tumour formation after injecting siRNA of TK1 or negative control in nude mice. (D, E) The tumour volume was measured and calculated every week after injection, the weights of tumours obtained from the animal model were decreased after TK1 downregulation by siRNA. (F) Ki-67 index of tumours in TK1 downregulation group or negative control group ** represents $P < .01$, and *** represents $P < .001$

nude mice was applied. Subcutaneous tumours were harvested and taken images (Figure 3C). The volume and weight of tumours from the TK1-interfered group were lower than the negative control group (Figure 3D,E). We further detected the expression of Ki-67 by IHC, and decreased Ki-67 expression was observed in the TK1-knockdown group (Figure 3F). These results above revealed that gene interfering of TK1 might attenuate the proliferation of PDAC cells in vivo and in vitro.

No significant statistical difference of the percentage of apoptotic cells was observed between TK1 downregulation group and negative control group (Figure 4A). Meanwhile, there was little change in expression of the apoptosis-related protein, apoptosis may not be involved in proliferation change of TK1 downregulation. As shown in Figure 4C, downregulation of TK1 induced an increased cell percentage in S phase compared to control group. These results indicated that interference of TK1 could suppress PDAC cell proliferation by S phase arrest.

Western blot was used to further examine the effect of downregulation of TK1 on the expression of cell cycle and apoptotic related proteins. Several proteins (cyclin A, cyclin B1, cyclin D1, cyclin E, CDK2, CDK4, CDK1, P21, P27 which might play a fundamental role in S phase) were further assessed by Western blot. It has been

proven that Cyclin E and CDK2 could regulate normal S phase progression,^{13,14} and we found downregulation of TK1 decreased the expression levels of cyclin E and CDK2 and increased the expression of P21. The result indicated that downregulation of TK1 in PDAC cells may induce S phase arrest by upregulating P21.

3.4 | Downregulation of TK1 exhibit no effect on migration and invasion

We conducted migration and invasion assay in vitro to explore the effects of TK1 knockdown on migration and invasion of PDAC cells. As shown in Figure S1, downregulation of TK1 had no impact on migration and invasion of PDAC cells. The result was also confirmed by detecting the key proteins involved in migration and invasion of cancer cells with Western blot (Figure S1).

3.5 | Expression of TK1 was directly regulated by E2F-1

As an essential gene involved in DNA replication and synthesis, it is reported that TK1 was directly regulated by E2F-1 in several studies.¹⁵⁻¹⁸ To find out whether E2F-1 regulates the TK1 transcription in PDAC cells,

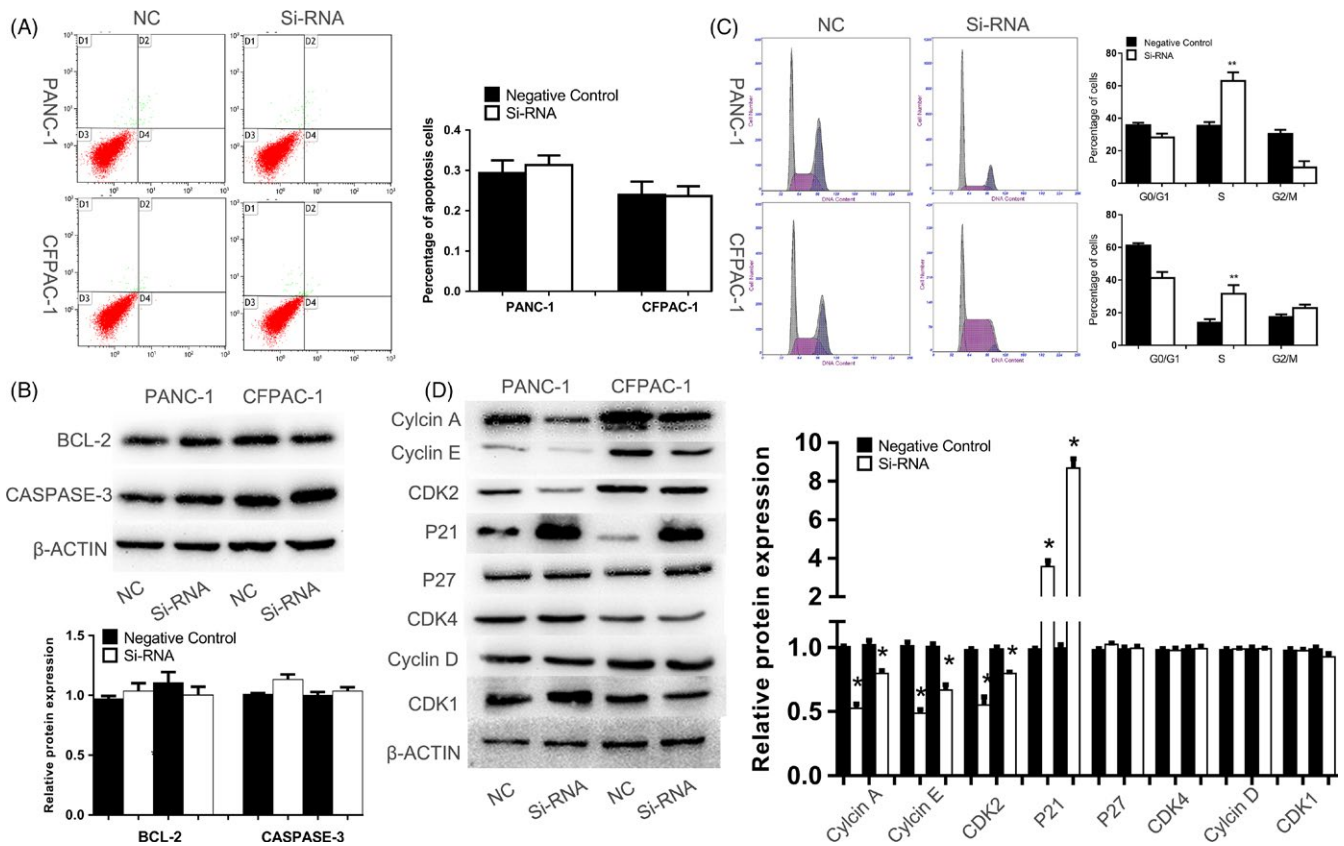


FIGURE 4 Effects of gene silencing of thymidine kinase 1 (TK1) on apoptosis and cell cycle. A, B, There was no statistical difference on the percentage of apoptotic cells and apoptosis-related proteins between two groups. C, D, Western blot indicated increased P21 was responsible for S phase arrest which was induced by TK1 knockdown. * represents $P < .05$, and ** represents $P < .01$

qRT-PCR, ChIP and Western blot assays were conducted. Firstly, the interference efficiency of three siRNAs (labelled as siRNA#1, siRNA#2 and siRNA#3) was detected by qRT-PCR and Western blot; siRNA#1 with the highest efficiency was chosen for further studies (Figure 5A). qRT-PCR and Western blot assay showed that the expression of TK1 was decreased in E2F-1 knockdown cells at mRNA and protein levels (Figure 5B). Besides, the results of ChIP assay indicated that E2F-1 could bind to the promoter of TK1 and the binding efficiency was decreased in E2F-1 knockdown cells (Figure 5C). Furthermore, the proliferation of PDAC cells was inhibited after E2F-1 knockdown (Figure 5D). These results suggested that E2F-1 might directly regulate the transcription of TK1 to further affect the cell proliferation in PDAC cells.

4 | DISCUSSION

In this study, we have identified high expression of TK1 was tightly correlated with the progression and prognosis of PDAC. In addition, E2F-1 could directly regulate TK1 transcription and promote PDAC cell proliferation. As far as we know, this study first identified aggressive characters of TK1 in PDAC, which suggested that TK1 could function as a prognostic and therapeutic target for PDAC.

It was generally known that plentiful oncogenes and complex mechanisms played a vital role in development and progression of

PDAC. TK1 was a type of TK which plays a significant role in mammalian cell cycle due to its vital function in DNA replication and repair. Increasing activity of TK1 during S phase could promote cell cycle from S phase to G2/M phase.¹⁹ TK1 was overexpressed in mitotic or malignant cells, but absent in quiescent cells.²⁰⁻²² Lots of studies have focused on the diagnostic and prognostic value of TK1 in many tumours.²³ Mao et al⁶ discovered the application of TK1 as a cell proliferation marker in human breast cancer. Kruck et al⁸ observed that high TK1 expression has been described to be associated with a higher stage, and grade in renal cell carcinoma. Moreover, TK1 has been assessed as a marker for prediction of biochemical recurrence in prostate cancer.²⁴ However, the expression pattern and function of TK1 in PDAC are not elucidated. In this study, relationship between TK1 expression and patient characteristics analysis indicated that high-TK1 expression correlated with Ki67, higher T stage (from TCGA data statistics), and advanced tumour stage (from IHC data statistics). We also observed that overexpression of TK1 in PDAC was associated with a shorter OS time and DFS time of PDAC patients. Further analysis identified that TK1 was an independent risk factor for poor prognosis.

Furthermore, we found that downregulation of TK1 could inhibit cell proliferation in vivo and in vitro, and the mechanisms involved in this phenomenon were still unclear. Cell cycle and apoptosis were confirmed to be involved in the process of proliferation.^{25,26} Therefore, we

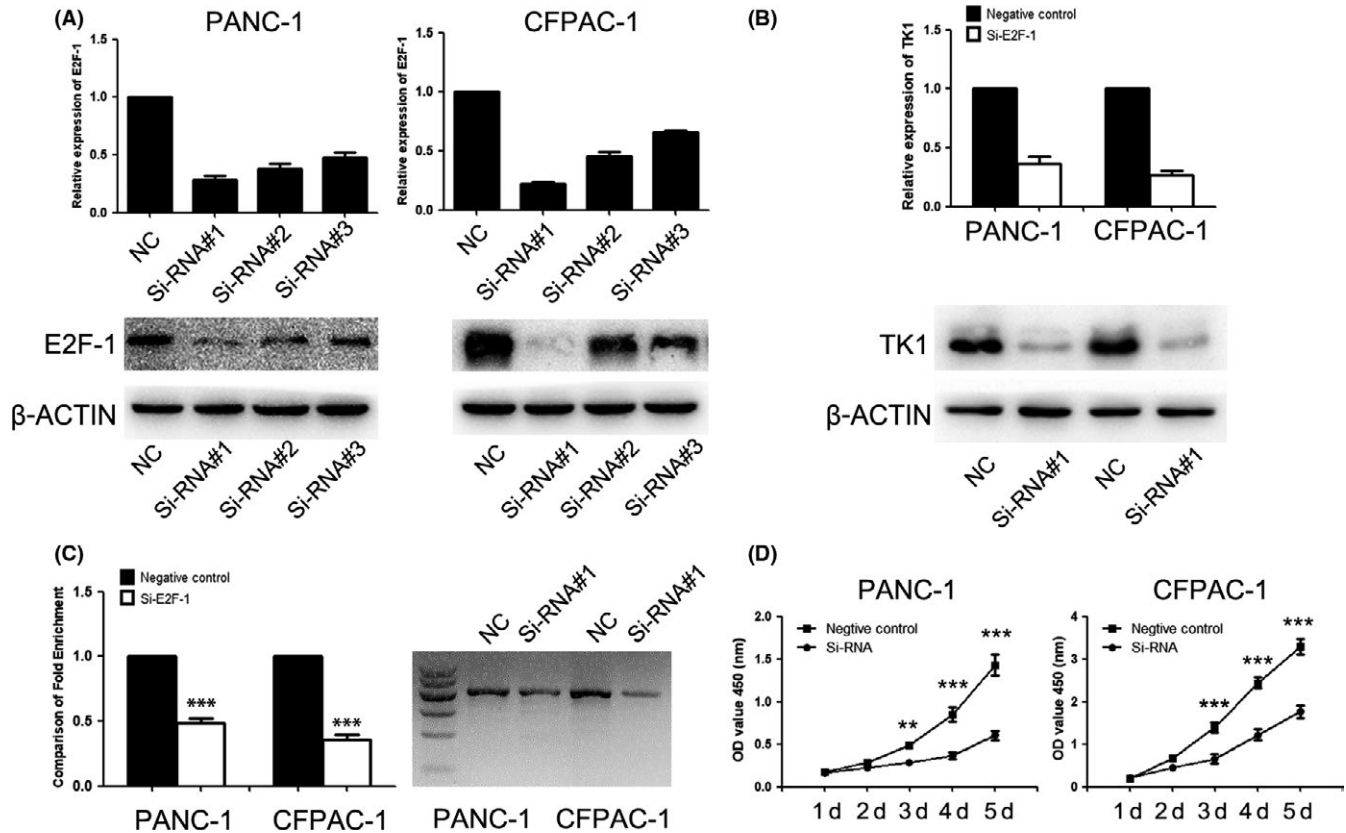


FIGURE 5 Expression of TK1 was directly regulated by E2F-1. A, Western blot and RT-PCR were used to validate the interference efficiency of three siRNAs for E2F-1. B, The expression of TK1 was downregulated by siRNA for E2F-1. C, The promoter of TK1 could be combined with E2F-1, and TK1 expression was decreased in E2F-1 knockdown group. D, The proliferation PDAC cells were affected after E2F-1 knockdown. ** represents $P < .01$, and *** represents $P < .001$

supposed that whether these processes contribute to the proliferation of PDAC cells. Although no obvious changes were observed in apoptosis, S phase arrest was observed in PDAC cells by TK1 knockdown. We found that the inhibition of S phase was concomitant with decreased CDK2, cyclin A and cyclin E expressions in TK1 knockdown PDAC cell, as well as increased P21 expression. P21, known as an inhibitor of the cyclin-dependent kinase, could inhibit the activation of CDK1, CDK2 and CDK4.²⁷⁻³⁰ TK1 might interact with P21 by combining with the C-terminal domain of P21.³¹ Our data showed that CDK2 was obviously downregulated after interfering TK1; however, CDK1 and CDK4 expressions might not be affected by the change of TK1. PDAC cell cycle arrest after TK1 downregulation might be mediated by P21/CDK2 interaction.

No evidence has shown that overexpression of TK1 was correlated with blood vessel invasion in clinicopathologic characteristics of PDAC patient. Accordingly, migration, invasion and Western blot assays further proved that there was no relation between TK1 overexpression and migration or invasion of PDAC cells.

We have identified the downstream genes regulated by TK1, and the upstream regulatory factors for TK1 were further investigated in our study. E2F-1, as an important transcription factor, is required for timely control of plentiful genes involved in DNA replication, repair and cell cycle.¹⁵ The previous study has reported that E2F-1 could directly bind to the promoter of TK1.¹⁵⁻¹⁸ We also

confirmed that E2F-1 was an upstream regulatory gene of TK1 in PDAC. Furthermore, there were some other factors that could regulate TK1, including C-MYC,³² FHIT,³³ SP1³⁴ and Prohibition,¹⁷ which might relate to potential pathways and involved in regulation of TK1. Further studies should be designed to explore the mechanism of TK1 in PDAC.

To the best of our current knowledge, this study first revealed the overexpression of TK1 in PDAC. Abnormal expression of TK1 is associated with progression of PDAC and a poor prognosis of PDAC patients. Downregulation of TK1 impedes PDAC proliferation both in vitro and in vivo experiments. As a transcription factor, E2F-1 could directly mediate expression of TK1. Therefore, TK1 could serve as a new anti-cancer target for treating PDAC.

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CONFLICT OF INTEREST

The authors confirm that there is no conflict of interest.

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SUPPORTING INFORMATION

Additional Supporting Information may be found online in the supporting information tab for this article.

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