Lentivirus-mediated short hairpin RNA targeting the APRIL gene suppresses the growth of pancreatic cancer cells in vitro and in vivo

FENG WANG1*, LIN CHEN2*, ZHEN-BIAO MAO1, JIAN-GUO SHAO2, CHANG TAN3 and WEI-DA HUANG3

1Department of Clinical Laboratory, Affiliated Hospital of Nantong University, Nantong, Jiangsu 226001; 2Department of Digestive Medicine, The Third People’s Hospital of Nantong City, Nantong, Jiangsu 226006; 3Department of Biochemistry, School of Life Science, Fudan University, Shanghai 200433, P.R. China

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Abstract. RNA interference (RNAi) is an evolutionarily conserved process of gene silencing in multiple organisms, which has become a powerful tool for investigating gene function by reverse genetics. Herein, we constructed a short hairpin RNA (shRNA) lentiviral expression vector targeting a proliferation-inducing ligand (APRIL) gene in the CFPAC-1 cell (a type of cell strain of human pancreatic cancer) in order to observe the inhibitory effect of APRIL gene’s shRNA on the growth of the CFPAC-1 cell in vitro and in vivo. The results showed that lentivirus-mediated RNAi effectively inhibited the expression of APRIL mRNA and protein in CFPAC-1 cells. Moreover, it can inhibit the growth of pancreatic cancer cells in vitro and in vivo. Our study indicates that lentivirus-mediated gene therapy is an attractive strategy in the treatment of pancreatic cancer and justifies the use of lentivirus in cancer gene therapy studies.

Introduction

Cancer cells are characterized by self-sufficiency in growth signals, insensitivity to anti-growth signals, sustained angiogenesis, tissue invasion and metastasis, limitless replicative potential and evasion of apoptosis (1). Despite the advances in surgery, radiotherapy and chemotherapy, the long-term survival rates of pancreatic cancer have not significantly improved. Gene therapy for pancreatic cancer is currently under investigation in clinical trials (2-4).

RNA interference (RNAi) has been widely used in the study of gene therapy for cancer (5-7). Many groups have reported the use of synthesized oligonucleotides or siRNA encoding plasmids to induce RNAi in mammalian cells by transfection, though this is still limited in its application, especially when it is necessary to generate long-term gene silencing in vivo (8,9). To circumvent this problem, lentivirus-delivered RNAi was developed. Viral vectors combined with RNAi provide useful tools in elucidating gene function by the analysis of loss-of-function phenotype and to explore the application of RNAi in gene therapy. Furthermore, the use of retroviral vectors can greatly expand the cell types available for RNAi analysis. Thus, the lentivirus vector is considered to be a promising gene delivery tool (10-12).

A proliferation-inducing ligand (APRIL) gene is a recently found new member of the tumor necrosis factor (TNF)’s superfamily. Studies have confirmed that the APRIL gene is overexpressed in many tumor tissues and tumor cell lines, particularly in the digestive system carcinomas, such as pancreatic cancer, gastric cancer, colon carcinoma, hepatoma and esophageal carcinoma etc, which suggests that APRIL plays an important role in the occurrence and development of these tumors (13-15).

Our previous experiments have further confirmed that the APRIL protein is overexpressed in pancreatic cancer tumors, yet, it is not expressed or has a weak expression in normal pancreatic gland tissues. Moreover, several pancreatic cancer cell lines have been screened and it has been found that the CFPAC-1 cell line is highly expressed in the APRIL gene. To study the effect mechanisms of the APRIL gene in the occurrence and development of pancreatic cancer, we constructed lentiviral vectors of RNAi targeting the human APRIL gene, then observed the inhibitory effect of APRIL gene’s short hairpin RNA (shRNA) on the growth of CFPAC-1 cells in vitro and in vivo, in order to probe the feasibility of gene therapy for pancreatic cancer.

Materials and methods

Cell culture. The 293T packaging cell line and human pancreatic cancer cell line CFPAC-1 (Academy of Life
Science, China) were maintained in Dulbecco’s modified Eagle’s medium (DMEM, Invitrogen, USA), supplemented with 10% fetal bovine serum (FBS, Hyclone, Logan, USA) and antibiotics (100 mg/ml streptomycin and 100 U/ml penicillin) at 37°C in a humidified incubator containing 5% CO₂.

**Construction of lentiviral vector.** Three self-complementary hairpin DNA oligos targeting APRIL mRNA were synthesized; the sequences were 5’-TAATCCAGGATGCTGGTTAAGGCTTTTTTTTC-3’ and 5’-CTGAGAAAAAAAAGCATCCTGGAAGGACGTAGGATGCTGGTTAAGGATAAGGCTTTTTTTTC-3’ and 5’-CTGAGAAAAAAAATTCCTGCACATCCGCAAAGAC-3’, named as APRILshRNA control. DNA oligos were annealed and inserted in the human U6 promoter site of the pGCL-GFP vector (Qiagen, The Netherlands). Then, the RNAi cassette was cloned into the latter vector and lentiviral vectors expressing shRNA were constructed. They were then confirmed by RT-PCR and DNA sequencing identification (3730xl DNA Analyzer, ABI, USA). The two primers targeting both sides of the pGCL-shRNA vector’s insert location were designed, forward primer, 5’-GTGTCACTAGGCGGGACGTGACACGTTCGGAGAATTACATCCGCAAAGAC-3’; reverse primer, 5’-AAAGGGTGTATCCCTGGCAGACGTGACACGTTCGGAGAATTACATCCGCAAAGAC-3’, named as APRILshRNA1, 5’-TAAGCCTTATCCTACGTCCTTCTCTCTTGAAACGTGACACGTTCGGAGAATTACATCCGCAAAGAC-3’ and 5’-CTGAGAAAAAAAAGCATCCTGGAAGGACGTAGGATGCTGGTTAAGGATAAGGCTTTTTTTTC-3’ and 5’-CTGAGAAAAAAAATTCCTGCACATCCGCAAAGAC-3’, named as APRILshRNA2, 5’-TCATACATCCAGAACAGCACCACATTCTCTTGAAACGTGACACGTTCGGAGAATTACATCCGCAAAGAC-3’; reverse primer, 5’-AAAGGGTGTATCCCTGGCAGACGTGACACGTTCGGAGAATTACATCCGCAAAGAC-3’.

**Construction of recombinant retrovirus and transfection into CFPAC-1 cells.** Lentiviral vector DNA packages and packaging vectors (pHelper1.0, pHelper2.0) (Qiagen) were then transfected into 293T cells. After transfection, the cells were incubated at 32°C to increase viral titer. Forty-eight hours later, the supernatant containing the retroviral particles was collected, filtered through the 0.45 μm low protein binding filter and the titer of lentiviruses was determined. CFPAC-1 cells were maintained in DMEM and were plated into 6-well plates at 3x10⁵ cells/well. Twenty-four hours later, the cells were infected with viral supernatants. After transfection, the cells were incubated at 32°C to increase viral titer. Forty-eight hours later, the supernatant containing the retroviral particles was collected, filtered through the 0.45 μm low protein binding filter and the titer of lentiviruses was determined. CFPAC-1 cells were transfected at 6 days post-transfection as indicated by the expression of GFP.

**Real-time RT-PCR analysis.** Total RNA was extracted using a RNA purification kit from Invitrogen following the manufacturer’s instructions. The real-time RT-PCR reactions were run on a real-time PCR system (iQ5 real-time PCR, Bio-Rad, USA) with the following cycle conditions: 95°C for 10 sec, 40 cycles at 95°C for 5 sec and at 60°C for 30 sec. A standard curve for APRIL was created using serially diluted total RNA from CFPAC-1 cells and used to quantify relative APRIL mRNA levels. The sequences of the APRIL gene primers were: forward primer, 5’-GGTATCCCTGGCAGACGTGACACGTTCGGAGAATTACATCCGCAAAGAC-3’; reverse primer, 5’-CTGTCACATCGGAGTCTATCCTGGAAGGACGTAGGATGCTGGTTAAGGATAAGGCTTTTTTTTC-3’; the product was 174 bp. β-actin was served as a control for normalization, the primers were: forward primer, 5’-GTGGAACGTGACACGTTCGGAGAATTACATCCGCAAAGAC-3’; reverse primer, 5’-AAAGGGTGTATCCCTGGCAGACGTGACACGTTCGGAGAATTACATCCGCAAAGAC-3’; the product was 302 bp.

**Western blot analysis.** The CFPAC-1 cells were lysed with a denaturing SDS-PAGE sample buffer using standard methods. Protein lysates were separated by 10% SDS-PAGE and transferred onto nitrocellulose membranes. The membranes were blocked with TBS containing 0.1% Triton X-100 and 5% nonfat milk overnight at 4°C, then incubated with anti-human APRIL antibody (dilution at 1:1000, Prosci, CA) at 4°C overnight. After washing, the membranes were incubated with HRP-conjugated mouse Ig at room temperature for 1 h. Signal detection was carried out with an ECL system (Amersham Biosciences, USA).

**Flow cytometry analysis of apoptosis.** The CFPAC-1 cells were harvested and fixed with ice-cold 70% ethanol. Fixed samples were centrifuged at 250 x g for 5 min, treated with RNase (0.25 mg/ml) and re-suspended in propidium iodide (50 g/ml) at room temperature. Propidium iodide-stained cells were analyzed by flow cytometer (Coulter Epics XL, Beckman, USA).

**MTT assay cell growth viability.** The cell growth following transfection was evaluated by MTT assay. Cells at a concentration of 5x10⁵ per well were seeded in a 96-well plate and incubated for 24 h. Seventy-two hours after transfection, the cells were incubated at 32°C to increase viral titer. Forty-eight hours later, the supernatant containing the retroviral particles was collected, filtered through the 0.45 μm low protein binding syringe filter and the titer of lentiviruses was determined. CFPAC-1 cells were maintained in DMEM and were plated into 6-well plates at 3x10⁵ cells/well. Twenty-four hours later, the cells were infected with viral supernatants in the presence of polybrene (6 μg/ml final concentration) for 12 h at a multiplicity of infection (MOI) of 5, then added to a fresh medium with fresh viral supernatants. After 24 h, the cells were incubated with a fresh viral supernatant for an additional 12 h. The transfected CFPAC-1 cells were subcultured at an appropriate density in fresh DMEM and 90% of the cells were transfected at 6 days post-transfection as indicated by the expression of GFP.

**Tumorigenicity experiments and the treatment of established tumors.** To determine whether APRILshRNA silence APRIL gene could inhibit tumor development in vivo, non-transfected control cells, APRILshRNA control, APRILshRNA1, APRILshRNA2 and APRILshRNA3 transfected CFPAC-1 cells (1x10⁶ cells in 100 μl) were injected subcutaneously into the axilla of each BALB/c nude mouse. The tumor growth was monitored weekly in two dimensions and tumor size was calculated according to the formula v = a²b/2, where a and b are the shortest and longest diameters, respectively. To determine the effect of APRIL inhibition on established tumors, CFPAC-1 cells (1x10⁶ cells in 100 μl) were injected subcutaneously into the axilla of each mouse and allowed to develop measurable tumors. Then, the mice were treated with
the APRILshRNA control, APRILshRNA1, APRILshRNA2 and APRILshRNA3 (1x10^7 transducing units/tumor) every 5 days; the mice treated with 0.9% sodium chloride were used as controls.

Statistical analysis. Average values were expressed as mean ± standard deviation (SD). Statistical significance between the different groups was determined by the Student’s t-test. P-values <0.05 were considered significant.

Results

Lentivirus-mediated RNAi inhibits APRIL gene expression in CFPAC-1 cells. In order to exclude an off-target silencing effect mediated by specific shRNA, we employed 3 different sequences of shAPRIL in our present study. To evaluate the inhibition of APRIL mRNA expression, real-time RT-PCR was performed 72 h after transfection, the APRIL mRNA expression in APRILshRNA1 and APRILshRNA2 transfected CFPAC-1 cells were reduced by 75 and 71%, respectively, as compared with the non-transfected control, APRILshRNA control or APRILshRNA3, reduces APRIL mRNA in CFPAC-1 cells (P<0.05).

Lentivirus-mediated RNAi inhibits APRIL protein expression in CFPAC-1 cells. Western blot analysis was performed 72 h after transfection. The APRIL protein expression demonstrated a significant reduction in APRILshRNA1 (3.83±1.4% of β-actin) and APRILshRNA2 (4.27±1.9% of β-actin) transfected CFPAC-1 cells respectively, as compared with a non-transfected control (23.5±3.2% of β-actin), APRILshRNA control (22.1±3.5% of β-actin) and APRILshRNA3 (19.3±4.0% of β-actin) transfected ones (P<0.05) (Fig. 2), suggesting that APRILshRNA1 and APRILshRNA2 strongly blocked APRIL expression, whereas no obvious inhibition of APRIL protein was observed in non-transfected control cells and APRILshRNA control and APRILshRNA3 transfected cells.

APRIL RNAi enhanced apoptosis and growth inhibition in CFPAC-1 cells. To further evaluate whether silencing the APRIL gene in CFPAC-1 cells may promote cell apoptosis, flow cytometry analysis was performed and the apoptosis index was calculated. In the cells transfected with the non-transfected control, APRILshRNA control, APRILshRNA1, APRIL shRNA2 and APRILshRNA3, the apoptosis index was (4.8±0.5)% (4.0±0.8)% (15.3±0.7)% (14.1±1.2)% and (5.5±0.9)%, respectively. Apoptosis indexes were significantly higher in APRILshRNA1 and APRILshRNA2 transfected cells as compared with the non-transfected control, APRIL shRNA control and APRILshRNA3 transfected ones (P<0.05).

MTT analysis revealed a cell growth inhibition consistent with the results of an apoptosis analysis. As shown in Fig. 3, cell growth was significantly inhibited in APRILshRNA1 and APRILshRNA2 transfected cells as compared with the non-transfected control, APRILshRNA control and APRIL shRNA3 transfected ones (P<0.05).

APRIL RNAi inhibits tumor growth in vivo. To determine whether APRILshRNA silences the APRIL gene could inhibit tumor development in vivo, non-transfected control cells, APRILshRNA control, APRILshRNA1, APRILshRNA2 and APRILshRNA3 infected CFPAC-1 cells were injected into the nude mice and the tumor growth was monitored weekly. When compared with non-transfected control cells, APRILshRNA control and APRILshRNA3 infected CFPAC-1 cells, the APRIL shRNA1 and APRILshRNA2 transfected cells developed much smaller tumors in the nude mice (P<0.05), (Fig. 4A).
and tissue-specific RNAi delivery to patients (16). RNAi, gene transfer strategies, especially viral vectors for efficient development. Particularly attractive from a medical standpoint, holds substantial promise for basic research and drug gene expression mediated by small double-stranded RNAs—

**Discussion**

The phylogenetically conserved cellular phenomenon of RNAi, the sequence-specific posttranscriptional silencing of gene expression mediated by small double-stranded RNAs—holds substantial promise for basic research and drug development. Particularly attractive from a medical standpoint, is the juxtaposition of new RNAi methodology with established gene transfer strategies, especially viral vectors for efficient and tissue-specific RNAi delivery to patients (16). RNAi, combined with the versatility and robustness of lentiviral vector-mediated gene delivery to a wide range of cell types offers the possibility of a long-term downregulation of specific target genes *in vitro* and *in vivo*. The use of silencing lentivectors allows for a rapid and convenient way of establishing cell lines (or transgenic mice) that stably express shRNAs for analysis of the effects produced by the knockdown of a specific gene (17).

Pancreatic cancer is the fourth leading cause of cancer-related death in the USA. The disease has a high mortality rate and the 5-year survival rate is estimated to be 4%. Currently, surgical resection is only possible in 20% of patients; even then, the overall 5-year survival rate is only 25% (18). Despite scientific efforts and significant progress in understanding the basic cellular event in pancreatic cancer, survival rates have not changed much during the last 20 years. Prognosis in pancreatic cancer remains unsatisfactory due to its late clinical presentation, low surgical resectability rates and resistance to chemotherapy. Novel therapeutic strategies are needed urgently in order to improve the prognosis of patients with pancreatic cancer (19). During the course of cancer development, gene abnormality such as mutation or amplification can be detected frequently, which may be related with tumor survival, progression and metastasis. In human pancreatic cancer, a series of specific gene (oncogene and tumor suppressor gene) mutations are observed, including the APRIL gene.

APRIL, also known as TALL-2 and TNFSF13, is a new found member in the TNF superfamily. It is closely related to the B cell activation factor from the TNF family (BAFF). These two TNF ligands share the receptors, B cell maturation antigen (BCMA) and transmembrane activator, calcium modulator and cyclophilin ligand interactor (TACI) (20). Recently, APRIL binding to the heparan sulfate side-chains of proteoglycans (HSPG) was demonstrated (21,22). In animal models, APRIL shows a tumor-promoting activity, as their overexpression induces the development of B cell neoplasia (23,24). A direct binding onto tumor cells via HSPG was observed *in vitro* experiments and confirmed *in situ* and indicates a potential role for HSPG/APRIL interactions in the development of tumors (25).

APRIL tumor-promoting activity is not restricted to B cell lymphomas. Furthermore, APRIL was shown to provide a proliferative/survival signal to tumor cells. This activity has been observed significantly *in vivo* by the overexpression of APRIL in tumor cells or by blocking endogenous APRIL (26-28).

In this study, we constructed the shRNA lentiviral expression vector targeting APRIL gene in CFPAC-1 cells, then observed an inhibitory effect of APRIL gene's shRNA on the growth of CFPAC-1 cells *in vitro* and *in vivo*. The results showed that APRILshRNA1 and APRILshRNA2, rather than the non-transfected control, APRILshRNA control or APRILshRNA3, effectively inhibited the expression of APRIL mRNA and protein in CFPAC-1 cells. Moreover, APRILshRNA1 and APRILshRNA2, rather than a non-transfected control, APRILshRNA control or APRIL shRNA3, inhibited the growth of the pancreatic cancer cells *in vitro* and *in vivo*. Our present study suggests that APRIL is a feasible RNAi target gene for pancreatic cancer. Thus,
stable lentivirus-mediated APRILshRNA holds great promise as a novel approach for APRIL-positive pancreatic cancer treatment.

Furthermore, our study indicates that lentivirus-mediated gene therapy is an attractive strategy in the treatment of pancreatic cancer and justifies the use of lentivirus in cancer gene therapy studies. However, further promote the technique into a gene therapeutic approach, an effective and safe protocol should be developed. In summary, further studies should focus on the lentivirus vector delivery strategies that can carry APRILshRNA specifically into APRIL-positive cancer cells with low toxicity and high efficiency.

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References