p53-inducible long non-coding RNA PICART1 mediates cancer cell proliferation and migration

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Abstract. Long non-coding RNAs (lncRNAs) function in the development and progression of cancer, but only a small portion of lncRNAs have been characterized to date. A novel lncRNA transcript, 2.53 kb in length, was identified by transcriptome sequencing analysis, and was named p53-inducible cancer-associated RNA transcript 1 (PICART1). PICART1 was found to be upregulated by p53 through a p53-binding site at -1808 to -1783 bp. In breast and colorectal cancer cells and tissues, PICART1 expression was found to be decreased. Ectopic expression of PICART1 suppressed the growth, proliferation, migration, and invasion of MCF7, MDA-MB-231 and HCT116 cells whereas silencing of PICART1 stimulated cell growth and migration. In these cells, the expression of PICART1 suppressed levels of p-AKT (Thr308 and Ser473) and p-GSK3β (Ser9), and accordingly, β-catenin, cyclin D1 and c-Myc expression were decreased, while p21Waf/cip1 expression was increased. Together these data suggest that PICART1 is a novel p53-inducible tumor-suppressor lncRNA, functioning through the AKT/GSK3β/β-catenin signaling cascade.

Introduction

Based on the transcript size, non-coding RNAs (ncRNAs) are divided into two groups: small non-coding RNAs and long non-coding RNAs (lncRNAs) (1,2). Biologically, lncRNAs are specifically characterized as RNA transcripts that are longer than 200 nt in length and do not have an open reading frame (ORF). To date, several biological functions of lncRNAs have been described, including i) signal molecules in gene transcription, ii) decoys to titrate transcription factors, iii) directors for chromatin, and iv) scaffolds for proteins to form ribonucleoprotein complexes (1,2). Differential expression of lncRNAs is a well-known event and regarded as a hallmark of cancer (3). Many lncRNAs have been characterized in cancers and they function as tumor suppressors or oncogenes. For example, imprinted H19 is an lncRNA first reported in mammalian cells, encoded by a gene located on chromosome 11p15.5; H19 is involved in fetal and placental growth (4). H19 has been reported as a marker in breast and cervical cancers (5). In gastric cancer, the lncRNA H19 promotes carcinogenesis and metastasis by interacting with miR-675 (4). Other characterized lncRNA oncogenes and tumor suppressors include ANRIL, HOTTIP, HOTAIR, BCAR4, lncRNA-p21 and XIST (6).

In cancer, lncRNAs have emerged as critical players involved in signaling transduction, DNA damage repair, cell cycle control, epigenetic modifications, and chromosome modeling (6). For example, the lncRNA HOTAIR regulates chromatin dynamics and promotes metastasis in multiple types of cancers, including breast cancer, colorectal cancer, non-small cell lung carcinoma and hepatocellular carcinoma (7). The aberrant activation of cellular signal transduction is well documented in cancer, promoting unlimited growth and proliferation of cancer cells. lncRNAs function in cellular signaling pathways and modulate cancer development and progression. For instance, the lncRNA ATB stabilizes IL-11 mRNA and activates the STAT3 signaling cascade which drives cell colonization at distant metastatic sites (8). BCAR4 (breast cancer anti-estrogen resistance 4) increases migration and invasion of breast cancer cells through activation of a non-canonical Hedgehog signaling pathway.
pathway (9). CCAT2 (colon cancer-associated transcript 2) and MALAT1 (metastasis-associated lung adenocarcinoma transcript 1) promote cancer progression and metastasis through activation of the Wnt signaling pathway (10,11). The present study characterized a new tumor suppressor lncRNA, named p53-inducible cancer-associated RNA transcript 1 (PICART1). This PICART1 lncRNA consists of 3 exons, 2533 bp in length (gene ID: 284080) and is encoded by a gene located at 17q21.33. Our data demonstrated that PICART1 inhibited cell proliferation and migration in breast and colorectal cancer cells through suppression of the AKT/GSK3β/β-catenin signaling pathway, functioning as a tumor suppressor.

Materials and methods

Ethics statement. IRB protocol was approved by Springfield Committee for Research Involving Human Subjects.

Human sample procurement. Frozen breast (n=50 pairs) and colon (n=50 pairs) cancer and matched normal adjacent tissues were procured following approved IRB protocol from the Tissue Bank of Simmons Cancer Institute at the Southern Illinois University. These frozen specimens were used for RNA extract and real-time RT-PCR analyses of PICART1 expression.

Cell culture. Human breast cancer cells (MCF7 and MDA-MB-231) were cultured in Dulbecco’s modified Eagle’s medium (DMEM; Thermo Fisher Scientific, Waltham, MA, USA) supplemented with 10% FBS (Sigma-Aldrich, St. Louis, MO, USA), 1% penicillin/streptomycin at 37˚C, 5% CO₂. The basal medium for HEK 293T cells was the same as that for supplemented with 10% FBS (Sigma-Aldrich), 1% penicillin/streptomycin at 37˚C, 5% CO₂. MDA-MB-231 cells were cultured in Dulbecco’s modified Eagle’s medium (DMEM; Thermo Fisher Scientific, Waltham, MA, USA) supplemented with 10% FBS (Sigma-Aldrich), 1% penicillin/streptomycin at 37˚C, 5% CO₂. Human breast cancer cells (HCT116 and HCT8) were grown in RPMI-1640 medium (Thermo Fisher Scientific) supplemented with 10% FBS (Sigma-Aldrich), 1% penicillin/streptomycin at 37˚C, and 5% CO₂. Human colorectal cancer cells through suppression of the AKT/GSK3β/β-catenin signaling pathway, functioning as a tumor suppressor.

Cell transfection and infection. For packaging of lentiviruses, 293T cells were seeded in 10-cm dishes. Cells were transfected with expression plasmids pCDH, PICART1, pGreenpuro, shRNA-2 or shRNA-3 with packaging plasmids REV, VSVG and GAG for 24 h. Then the medium was changed and cells were continued to culture for 48 h to collect viruses, 293T cells were seeded in 10-cm dishes. Cells were transfected with expression plasmids pCDH, PICART1, pGreenpuro, shRNA-2 or shRNA-3 with packaging plasmids REV, VSVG and GAG for 24 h. Then the medium was changed and cells were continued to culture for 48 h to collect viral particles (15). For infection, cells were spread at 60% of confluence and exposed to lentviruses (MOI=1) with 1:2000 diluted Polybrene (Millipore, Billerica, MA, USA). Western blotting. Cells were lysed in cell lysis buffer (Roche, Indianapolis, IN, USA) with a protease inhibitor cocktail followed by centrifugation at 14,000 rpm for 5 min to collect supernatant proteins. Protein separation, membrane blotting and antibody probing were conducted as previously described (16). Antibodies used were anti-p53 (dilution 1:1000; Santa Cruz Biotechnology, Santa Cruz, CA, USA), anti-pAKT (Th3208) (dilution 1:1000; Cell Signaling Technology, Danvers, MA, USA), anti-pAKT (Ser473) (dilution 1:1000; Cell Signaling Technology), anti-AKT (dilution 1:1000; Cell Signaling Technology), anti-GSK3β (S9) (dilution 1:1000; Cell Signaling Technology), anti-GSK3β (dilution 1:1000; Cell Signaling Technology), anti-GSK3β (dilution 1:1000; Cell Signaling Technology).
Cell Signaling Technology), anti-β-catenin (dilution 1:1000; Cell Signaling Technology), anti-cyclin D1 (dilution 1:1000; Cell Signaling Technology), anti-c-Myc (dilution 1:500; Santa Cruz Biotechnology) and anti-p21Waf/cip1 (dilution 1:1000; Cell Signaling Technology). β-actin was used as an internal control.

Luciferase activity. The dual luciferase assay kit (Promega, Madison, WI, USA) was used following the manufacturer's protocol. Cells were collected and lysed for luciferase assay 48 h after transfection. Renilla luciferase was used as an internal control for normalization (17).

Transwell migration and invasion assays. Transwell migration assays were performed using 8.0-µm pore inserts (BD Biosciences, San Jose, CA, USA). A total of 2-5×10^4 cells were suspended in 500 µl serum-free medium and loaded into upper wells; lower chambers were filled with 750 µl complete medium with 10% FBS. For the invasion assays, inserts were coated with 1 mg/ml Matrigel and pre-incubated at 37˚C for 2 h. A total of 3-7×10^4 cells were suspended in 500 µl serum-free medium and loaded into the coated inserts; lower chambers were filled with 750 µl complete medium. Migration and invasion chambers were incubated in a 5% CO₂ incubator at 37˚C for 36-48 h. Cells were stained and counted under a microscope for 5 random fields (18).

Wound healing assays. Cells were seeded in 6-well plates at 90% confluence and cultured in medium supplemented with 2% FBS. Scratches were made using a 200-µl tip. Migration distance was estimated from images (4 fields) taken at each time point (18).

Cell proliferation. For the cell proliferation assays, cells were split into 96-well plates at 2000-3000 cells/per well. Viable cells were measured by tetrazolium salt, 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assays over 5 days (19). For cell counting assays, cells were split into 12-plates. At the indicated time points, cells were collected and subjected to trypan blue staining. Viable cells were counted in a Vi-cell counter (Beckman Coulter, Brea, CA, USA).

Statistical analysis. Data were analyzed by SPSS Statistical software (SPSS Inc., Chicago, IL, USA). Values are shown as mean ± SD. The Student's t-test was used for the analyses. A P-value of ≤0.05 was considered to indicate a statistically significant difference.

Results

Identification of a novel long non-coding RNA transcript PICART1. p53 is a master transcription factor regulating the expression of a wide range of coding and non-coding genes. HDM2 (MDM2 in mice) binds to the p53 protein and inhibits its transactivation activity in target gene expression and triggers ubiquitination-proteasome degradation (20). A lead compound, RITA (reactivation of p53 and induction of tumor cell apoptosis), binds to p53 and thus leads to p53 conformational changes and dissociation from HDM2, reactivating p53 (21). Using transcriptome RNA sequencing analysis in MCF7 cells (p53wt) treated with RITA (100 nM), we recognized 10 new non-coding RNA species with pivotal expression changes in response to RITA treatment. Table I summarizes the basic information of these new RNA transcripts induced by RITA. LOC284080 (PICART1), p53 inducible cancer-associated RNA transcript 1.
**PICART1 is downregulated in breast and colon cancer cell lines and tissues.** As a potential p53 target, PICART1 was first examined in cancer cell lines and tissues to see whether this lncRNA species is involved in tumor development and progression. As shown in Fig. 1A, in 50 paired breast cancer and adjacent normal tissues, PICART1 expression was decreased by >2-fold in 32 (64.0%) breast cancer cases compared to that in the adjacent normal breast tissues, with a maximal decrease of >30-fold. Similarly, in 50 paired colon cancer and adjacent normal tissues, PICART1 expression was decreased by >2-fold in 23 (46%) malignant tissues with a maximal decrease of >25-fold (Fig. 1B). PICART1 expression was also downregulated in cancer cell lines. As shown in Fig. 1C, PICART1 expression was high in non-transformed human mammary epithelial cells MCF10A, but decreased by ~7.0-, 2.0-, 3.0- and 2.5-fold in the MCF7, MDA-MB-231, HCT116 and HCT8 cells, respectively, compared to the MCF10A cells. These data suggest that PICART1 is downregulated in breast and colon cancer cell lines and tissues, functioning as a potential tumor suppressor.

**p53 upregulates PICART1 expression through a p53 response element in its promoter.** As a transcription factor, p53 may stimulate PICART1 expression through a promoter-mediated mechanism. With a p53 scan program MatInspector software (https://www.genomatix.de/online), we analyzed the promoter region of PICART1 and identified 3 putative p53 response elements ~3000 bp upstream of PICART1 transcripts (Fig. 3A). Specifically, these 3 putative p53 response elements were located at -1808 to -1783 bp, -1524 to -1499 bp, and -1131 to -1106 bp. We then cloned this 3000-bp promoter fragment into a pGL4 basic luciferase reporter vector and constructed four promoter-luciferase reporter plasmids that contained -3000, -1592, -1227 and -681 bp, respectively. The -3000 bp hosted all three putative p53 response elements, the -1592 bp contained putative p53 response elements 2 and 3, and the -1227 bp had only putative p53 regulatory element 3 (Fig. 3B). These promoter fragments all demonstrated promoter activity driving luciferase reporter expression, but the promoter activity of -1227 and -681 bp vectors was lower compared to the longer -1592 and -3000 bp fragments, indicating existence of enhancer(s) in the longer region (Fig. 3B).
We further tested the response of these promoter fragments to RITA. Our results showed that the -3000-bp fragment was activated by RITA at 0.05 to 0.5 µM in a dose-dependent manner in both MCF7 and HCT116 cells (Fig. 3C), but the
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-1592 bp fragment was not (data not shown). These data suggested that the putative p53 element 1 is functional, but the putative p53 elements 2 and 3 are not. We then conducted co-transfections of the PICART1 promoter (-3000 bp) with p53 gene or shRNA in MCF7 and HCT116 cells. Our results demonstrated that luciferase activity was increased by p53 expression but decreased by p53 silencing compared to the vector control (Fig. 3D). We examined binding of p53 to this putative element 1 in the PICART1 promoter using a ChIP assay. The results demonstrated that p53 binds to the element 1 (Fig. 3E). We further characterized this putative p53 element 1 by targeted point mutations (Fig. 3A, right). As a result, mutations at the putative p53 responsive element 1 lowered down basal activity of the PICART1 promoter (Fig. 3B) and
eradicated its response to p53 (Fig. 3F). These data suggested that p53 upregulates the PICART1 expression through the p53 responsive element 1 in its promoter.

**PICART1 inhibits cell proliferation, migration, and invasion.** As a target of p53 tumor suppressor, we further estimated the biological function of PICART1 in cancer cells, using the ectopic expression and silencing cell models. Full length PICART1 cDNA was cloned for ectopic expression, and 3 PICART1-specific shRNA constructs were made for silencing. Fig. 4A shows the targeted ectopic expression (left) and silencing (right) of PICART1 in MCF7 and HCT116 cells.
The data demonstrated that shRNA-2 and shRNA-3 knocked down PICART1 expression to ~40-60% compared to the vector control. We estimated the cell growth and proliferation of these cell lines with targeted expression and silencing of PICART1, and the results showed that the ectopic expression of PICART1 notably inhibited the proliferation of MCF7 and HCT116 cells as tested by MTT assays (Fig. 4B) and viable cell counting method (data not shown). In contrast, silencing of PICART1 by shRNAs led to an increase in cell growth and proliferation in the MCF7, HCT116 and MDA-MB-231 cells (Fig. 4C). Furthermore, FACScan analysis showed that PICART1 induced cell cycle arrest at the G2/M phase (Fig. 4D).
We further investigated the effects of PICART1 expression on migration and invasion of cancer cells. As shown in Fig. 5A, ectopic expression of PICART1 in MCF7 and MDA-MB-231 cells led to a notable decrease in the cell migration through micro-holes in the Transwells. In contrast, PICART1 silencing enhanced cell migration (Fig. 5B). Similar results were obtained in the invasion assays, and the ectopic expression of PICART1 suppressed the invasion of MCF7 and HCT116 cells (Fig. 5C). Altogether, these data suggest that PICART1 inhibits the proliferation, migration, and invasion of breast and colon cancer cells.

**Figure 6.** PICART1 regulates cell proliferation through the AKT/GSK3β/β-catenin cascade. (A) Inhibition of the AKT/GSK3β/β-catenin pathway effectors by PICART1 expression. (B) Activation of the AKT/GSK3β/β-catenin pathway effectors by PICART1 silencing. (C) AKT inhibitor. MCF7 cells with PICART1 silencing were treated with 5.0 µM of MK-2206, an AKT inhibitor, for 24 h. Protein expression and cell proliferation were estimated. *P<0.05 and **P<0.01 compared to the controls. (D) A hypothetical functional model of PICART1.

**PICART1 regulates cell proliferation through the AKT/GSK3β/β-catenin signaling pathway.** We further investigated the underlying mechanisms through which PICART1 functions. To address this question, we explored the effects of PICART1 expression on main oncogenic signaling pathways, such as AKT and ERK signaling. PICART1 did not have notable effects on ERK signaling (data not shown). However, as shown in Fig. 6A, ectopic expression of PICART1 led to a decrease in pAKT (Thr308 and Ser473), suggesting that PICART1 may function through the AKT signaling cascade.
We then explored the downstream targets/effectors of AKT. Our results showed that pGSK3β (Ser9) and β-catenin expression was decreased by PICART1. β-catenin functions as a key transcription factor after translocation into the nucleus (22). As a result, the expression of cyclin D1 and c-Myc was decreased by PICART1 (Fig. 6A). c-Myc is a suppressor of p21Waf1/cip1 expression (23). Therefore, in the tested cells, we observed an increase in p21Waf1/cip1 expression due to downregulation of c-Myc by PICART1 (Fig. 6A). In contrast, silencing of PICART1 led to an increase in pAKT (Thr308 and Ser473), pGSK3β (Ser9), β-catenin, cyclin D1 and c-Myc expression but a decrease in p21Waf1/cip1 expression (Fig. 6B). To further confirm that PICART1 regulates cell growth and proliferation through the AKT/β-catenin signaling pathway, we treated PICART1-silenced cells (where AKT signaling was activating) with 5.0 µM MK-2206, an AKT1/2 inhibitor. Our result showed that exposure to the AKT inhibitor abolished the stimulation of PICART1 silencing on cell proliferation (Fig. 6C). Taken together, our data suggest that PICART1 inhibits proliferation of MCF7 and HCT116 cells through control of the AKT/GSK3β/β-catenin pathway (Fig. 6D).

Discussion

Long non-coding RNAs (lncRNAs) are RNA transcripts in the genome which function as oncogenes or tumor suppressors, playing a critical role in cancer progression and development (24). This study characterized a new lncRNA named PICART1. PICART1 is p53-inducible and functions as a tumor suppressor via regulating the AKT/GSK3β/β-catenin signaling pathway. This study also characterized a p53-response element in the promoter of the PICART1 gene.

p53 is a critical transcription factor that regulates expression of a wide range of genes, including protein-coding and non-coding genes. This makes p53 an important protein in multiple cellular processes, such as DNA repair and genome stability, cell cycle control, apoptosis, and senescence (25). As a transcription factor, p53 functions as a tetramer, binding to a sequence-specific response element in the promoter of target genes and controlling gene expression (26). This study revealed PICART1 as a new lncRNA target of p53, which is supported by several lines of evidence. The most direct evidence lies in the effects of targeted expression or silencing of p53 on the expression of endogenous PICART1 and on its promoter activity, and in the characterization of an active p53 response element in the PICART1 promoter. Targeted mutations of this element decreased the promoter activity at baseline and response to p53 expression or silencing. In addition, RITA is considered a p53 activator, restoring the function of p53 from its negative regulator HDM2 (MDM2 in mice); doxorubicin induces DNA damage and thus activates p53. Our results showed that both RITA and doxorubicin that activate p53 through different mechanisms triggered PICART1 expression, providing an additional line of evidence that p53 regulates PICART1 expression. However, p53 may not be the sole signaling pathway that regulates PICART1 expression. It is noted that in the tested cells, PICART1 expression was notstringently correlated with the p53 status. The PICART1 level was high in the non-transformed MCF10A cells that harbors a wild-type p53, but decreased in MCF7, HCT116, and HCT8 cells that also have a wild-type p53. It would be interested to further characterize the other factors that may affect PICART1 expression.

Our data showed that PICART1 expression was higher in response to RITA exposure than to ectopic p53 expression. This may be ascribed to the nature of different treatments. Unlike drug treatment, the efficiency of p53 transfection, i.e., the percentage of the cells transfected, rarely reaches 100%. Thus, PICART1 expression was affected by p53 delivery only in a portion of cells. More importantly, p53 is a well-known cell apoptotic tumor suppressor (27). Gene transfection of p53 often leads to high expression and resultant cell death and RNA degradation, thus resulting in lower PICART1 induction observed in p53 delivery than in RITA exposure. In fact, PICART1 expression was downregulated much more efficiently in the p53-silenced cells due to the increased survival of these cells.

Promoter motif analysis recognized 3 putative p53 response elements in the PICART1 promoter, but only one was functional. The luciferase reporter activity assays showed that the full length of the promoter fragment (-3000 bp) that contains all three elements presented a dose-dependent response to RITA treatment, while the promoter fragment -1592 bp that harbors the elements 2 and 3 was not responsive to RITA. This excluded the possibility that the elements 2 and 3 are functional p53 binding motifs. Therefore, our targeted mutation analyses were focused on element 1 located at -1808 to -1783 bp, and our study results confirmed its activity responding to p53, thus strengthening the conclusion of p53 regulation on the PICART1 expression.

Cancer cells are characterized by unlimited growth and proliferation and increased migration and invasion, leading to development of tumors (28). p53-induced expression, coupled together with the decreased expression in cancer cells and tissues, suggests that PICART1 may function as a tumor suppressor. To test this hypothesis, we investigated the effects of targeted expression or silencing of PICART1 on proliferation, migration, and invasion of breast and colon cancer cells. Our result showed that ectopic expression of PICART1 suppressed cell proliferation and induced cell cycle arrest in G2/M phase while silencing of PICART1 enhanced cell proliferation. In addition, PICART1 expression markedly inhibited cell migration and invasion while silencing of PICART1 promoted cell migration and invasion. These results, along with the downregulation in breast and colon cancer, suggest that PICART1 acts as a tumor suppressor.

lncRNAs function as tumor suppressors through modulating cellular signaling networks (29). PI3K/AKT and Wnt/β-catenin are two master oncogenic signals regulating cell proliferation, survival, migration, and invasion (30,31). AKT is a serine-threonine kinase activated by phosphorylation at Thr308 and then Ser473 (32). In the canonical Wnt pathway, Wnt protein binds to Frizzled receptors, suppressing the activity of glycogen synthase kinase-3β (GSK3β). Without Wnt signaling, β-catenin associates with Axin, adenomatosis polyposis (APC), protein phosphatase 2A (PP2A), glycogen synthase kinase 3β (GSK3β), and casein kinase 1α (CK1α), forming a destruction complex, degrading β-catenin by targeting for ubiquitination (33-35). An activation of Wnt signaling drives the destruction complex to translocation and...
disruption, allowing β-catenin accumulation and translocation into the nucleus. Once in the nucleus, β-catenin associates with Tcf/Lef transcription factors and drives expression of target genes, such as cyclin D1 and c-Myc, promoting cell proliferation and migration (22,36-38). AKT cross-talks with Wnt/β-catenin signaling through phosphorylation at Ser9 and inactivation of GSK3β leading to increase of β-catenin level (39).

In this study, we found that PICART1 suppressed AKT activity with decreased phosphorylation levels of Thr308 and Ser473, which in turn lowered the pGSK3β (Ser9) and β-catenin levels, suppressing cyclin D1 and c-Myc expression. The latter is a repression transcription factor of p21Waf1/cip1 by sequestering SPI (23). Thus we observed an increase in p21Waf1/cip1 expression by PICART1. It is currently unknown how PICART1 suppresses phosphorylation activation of AKT. Considering the multiple functional possibilities of lncRNAs, PICART1 may function as an adaptor to couple AKT with phosphatases, leading to dephosphorylation. PICART1 may also bind to AKT and block the access of AKT kinases for phosphorylation. Further study is warranted. It is also noteworthy that p21Waf1/cip1 regulates both G1/S and G2/M transition (40,41). In this study, targeted PICART1 expression induced cell cycle arrest at G2/M phase, but not at G1. It is well known that cell cycle regulation is a complicated event with involvement of multiple proteins, which may lead to differential phenotypes (42).

In summary, this study characterized a new lncRNA, named PICART1. PICART1 was upregulated by p53 through a sequence-specific response element in the promoter. PICART1 was downregulated in breast and colon cancer cells and tissues and functioned as a tumor suppressor, inhibiting cell proliferation, migration, and invasion through modulation of the AKT/ GSK3β/β-catenin signalling cascade.

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