Long non-coding RNA MEG3 inhibits the proliferation of cervical carcinoma cells through the induction of cell cycle arrest and apoptosis

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Cervical cancer remains an important public health problem worldwide. New and effective therapeutic strategies targeting cervical cancer are urgently needed. Long non-coding RNAs (lncRNAs) are newly identified regulators in tumorigenesis and tumor progression. To investigate the role of lncRNA MEG3 in the development of cervical cancer, we examined MEG3 expression in 18 pairs of cervical cancer and matched adjacent non-neoplastic tissues. Real-time quantitative RT-PCR (qRT-PCR) results showed high expression levels of MEG3 in non-neoplastic tissues, but markedly lower levels in cancer tissues. We further investigated whether the restoration of MEG3 expression might affect the proliferation of cervical carcinoma cells. Ectopic expression of MEG3 inhibited the proliferation of human cervical carcinoma cells HeLa and C-33A in vitro. On the other hand, knockdown of MEG3 promoted the growth of well-differentiated cervical carcinoma HCC94 cells. Further investigation into the mechanisms responsible for the growth inhibitory effects revealed that overexpression of MEG3 resulted in the induction of G2/M cell cycle arrest and apoptosis. These results identified an important role of MEG3 in the molecular etiology of cervical cancer and implicated the potential application of MEG3 in cervical cancer therapy.

Key words: cervical cancer, long non-coding RNAs; MEG3, apoptosis

Although the incidence of cervical cancer has been greatly reduced with the application of pap smear screening, cervical cancer is still the third most commonly diagnosed cancer in women worldwide, with estimated 529,800 new cases diagnosed annually [1]. Therefore, new and effective therapeutic strategies for cervical cancer are urgently needed.

Noncoding RNAs (ncRNAs) are newly discovered regulators of diverse biological functions. In cancer, ncRNAs have been shown to have both supportive and inhibitory effects in tumor development [2, 3]. ncRNAs can be grouped into two major classes based on the sizes of their transcripts: small ncRNAs (< 200 bp) and long ncRNAs (lncRNAs) (≥ 200 bp) [4, 5]. Small ncRNAs comprise a broad range of RNA species, and many of them are associated with 5′ or 3′ regions of genes. microRNA is one of the most important small ncRNAs that have been documented to play important roles in cancer development [6-8].

Recent studies identified an important role for lncRNAs in tumor progression [9]. LncRNAs, ranging from 200 nucleotides to over 10 kb, are abundantly transcribed by the mammalian genome [10, 11]. Increasing evidence supports the association of aberrant lncRNA expression with human cancers. As lncRNAs are emerging as critical components of the cancer transcriptome, it is reasonable to anticipate that lncRNAs may contribute to the tumorigenesis and progression of cervical cancer. Gibb EA et al [12] provided the first lncRNA expression profile in the human cervical intraepithelial neoplasia (CIN) transcriptome and demonstrated aberrant lncRNA expression in early-stage neoplasia.

Maternally expressed gene 3 (MEG3), an imprinted gene, is a tumor suppressor gene located in chromosome 14q32 [13]. The potential functions of MEG3 have been studied in a number of cancer types. MEG3 is down-regulated in several cancer types. In addition, overexpression of MEG3 inhibits pituitary tumor cell proliferation, suggesting that MEG3 is a potential tumor suppressor gene [14]. Likewise, re-expression of MEG3 inhibits the proliferation of tumor cells in vitro in glioma [15] and meningioma [16]. Further,
MEG3 is reported to be dysregulated in certain tumors, such as hepatocellular carcinoma [17], Wilms tumors [18], and multiple myeloma [19]. However, the role of MEG3 in tumorigenesis and progression of cervical cancer remains unknown.

In this study, we examined the expression of lncRNA MEG3 in 18 pairs of cervical cancer and matched non-neoplastic tissues using real-time quantitative RT-PCR (qRT-PCR). Moreover, we investigated the impact of ectopic expression of MEG3 in proliferation, cell cycle progression and apoptosis in vitro in human cervical carcinoma cells HeLa and C-33A.

Materials and methods

Tissue samples. Eighteen pairs of primary cervical cancer and matched adjacent non-neoplastic tissues were obtained from the Department of Gynecology and Obstetrics in the First Affiliated Hospital of Xinxiang Medical University. The study protocol was approved by the Institutional Review Board of Xinxiang Medical University. Written informed consent was obtained from all patients enrolled in the study. Tissues were cut into pieces smaller than 0.5 cm in any single dimension and immediately immersed into 2 ml RNA later solution (Ambion, Foster City, CA). Tissue samples were stored in liquid nitrogen until use. Tissue sections from each sample were reviewed and classified by a pathologist.

Quantitative reverse transcriptase PCR (qRT-PCR). Total RNA was isolated from cervical cancer and matched non-tumor tissues using the RNeasy kit (Qiagen, Grand Island, NY) according to manufacturer’s instructions. Reverse transcription reactions were carried out with 1 mg total RNA using the PrimeScript RT reagent kit (TaKaRa BIO, Shiga, Japan). Random hexamer primers were used in the RT reactions. Real-time PCR was performed on a Bio-Rad CFX-96 real-time PCR system (Bio-Rad, Hercules, CA) using SYBR® Premix DimerEraser kit (TaKaRa, Shiga, Japan) following the manufacturer’s instructions. MEG3 primer sequences were obtained from previous publication [15] as following: Sense, CTGCCCATCTACACCTCAG; Antisense, CTCTCCGCGCTGCTGCGCTAGGGGCT. GAPDH was used as a housekeeper gene for the qRT-PCR reactions. Each test was done in triple replication and the 2-ΔΔCt method was used to calculate the expression of lncRNA MEG3 in tissue samples.

Cell culture. Human cervical carcinoma cells lines HeLa, C-33A and HCC94 (well-differentiated) were obtained from the Cell Bank of the Chinese Academy of Sciences and maintained in modified Eagle’s media (MEM) containing 10% fetal bovine serum (FBS) at 37°C in a 5% CO2 atmosphere.

MTS cell growth assay. Plasmid pEGFP-MEG3 was constructed by subcloning MEG3 cDNA in an EcoRI BamHI fragment into the pEGFP vector. HeLa or C-33A cells were transiently transfected with pEGFP-MEG3 using Lipofectamine 2000 (Life technologies, Carlsbad, CA). These cells were reported to have high transfection efficiencies [20]. Cell proliferation was evaluated using colorimetric MTS assay (Promega, Madison, WI) following the manufacturer’s protocol. MTS assay measures the restoration of 3-(4,5-dimethylthiazol-2-yl)-5-(3-carboxymethoxyphenyl)-2-(4-sulphophenyl)-2H-tetrazolium (MTS) to formazan by metabolically active cells.

siRNA transfection and MTS cell growth assay. Well-differentiated cervical carcinoma HCC94 cells were grown in 24-well culture plates to 80-90% confluence overnight and then transfected with 25 nM non-targeting siRNA control, or synthetic siRNAs against MEG3 (Guangzhou RiboBio Co., Ltd.) for 48 h using LipofectamineTM transfection reagent (Invitrogen) following the manufacturer’s protocol. Following transfection, the cells were harvested and MEG3 knockdown was confirmed by qRT-PCR. HCC94 cells were transiently transfected with siRNA-MEG3 or control non-targeting siRNA for 24 to 72 h, and cell proliferation was evaluated by MTS cell growth assay as described above.

Cell cycle analysis. HeLa or C-33A cells (5×10^5) were plated in 60 mm dishes overnight. The cells were then transiently transfected with pEGFP-MEG3 or control vector for 48 h. Adherent cells were trypsinized and resuspended in cold PBS. Three volumes of cold 70% ethanol were added in a dropwise manner while vortexing. Cell suspensions were fixed for 24 h at -20°C, washed twice with cold PBS. Cells were resuspended in 100 µg/ml RNase A in 100 µl PBS. Five min later, 50 µg/ml propidium iodide in 400 µl PBS was added. Cell cycle progression was analyzed using a BD FACScan flow cytometer. Cell cycle analysis was performed using the FCS Express 4 software (De Novo Software, Los Angeles, CA).

Apoptosis assay. HeLa or C-33A cells (5×10^3) were plated in 60 mm dishes overnight. The cells were then transiently transfected with pEGFP-MEG3 or control vector for 48 h. Cells were collected by trypsinization, washed twice with cold PBS and then resuspended in 1 × Binding buffer at a concentration of 1 × 10^6 cells/ml. 100 µl cell suspension was transferred to a 5 ml tube and stained with annexin V and 7-aminoactinomycin D (7-AAD) (BD Biosciences, San Diego, CA) according to the manufacturer’s protocol. Stained cells were immediately analyzed using a BD FACSScan flow cytometer. Data analysis were performed using the FCS Express 4 software.

Western blot analysis. The cells were transfected with pEGFP-MEG3 or control vector for 48 h, and immediately lysed in RIPA buffer (10 mm Tris, pH 7.4, 150 mm NaCl, 5 mm EDTA, 8% glycerol, 1% Triton X-100, 0.1% SDS, 0.5% sodium deoxycholate, 10 mm Na3VO4, 1 mm sodium fluoride, and Complete Protease Inhibitor Mixture (Roche Diagnostics)). 40 µg of total protein per sample was separated by SDS-PAGE, blotted onto polyvinylidene fluoride membranes that were blocked for 1 h at room temperature with 5 % nonfat dry milk in TBS containing 0.05 % Tween 20. The membranes
were then incubated overnight at 4°C in nonfat dry milk in TBS containing 0.05% Tween 20 with the indicated primary antibodies. Membranes were washed in nonfat dry milk in TBS containing 0.05% Tween 20 and incubated with anti-rabbit (1:1,000) or anti-mouse (1:1,000) secondary antibody conjugated to horseradish peroxidase (Amersham Pharmacia Biotech, Piscataway, NJ). Visualization of the protein bands was done using the enhanced chemiluminescence plus kit as recommended by the manufacturer (Perkin-Elmer, Boston, MA). The following primary antibodies were obtained from Santa Cruz Biotechnology: p53 (sc-6243), p21 (sc-397), cyclin B1 (sc-245), Cdk1/Cdk2 Antibody (sc-53219), caspase 3 (sc-7148), caspase 8 (sc-56070), and actin (sc-8432).

Statistical analysis. Statistical significances between groups were determined by two-tailed Student’s t-test. A p-value of < 0.05 was considered significant.

Results

Down-regulation of IncRNA MEG3 expression in cervical cancer. Obtained data revealed that IncRNA MEG3 was markedly down-regulated in moderate (CIN2) and severe (CIN3) histopathologic grades of CIN compare to non-neoplastic cervical tissues [12] (Fig. 1A, upper panel). MEG3 expression was reversely correlated with the severity of the tumor.

To further evaluate the clinical relevance of MEG3, expression of MEG3 in 18 pairs of cervical cancer and matched non-neoplastic tissues were analyzed by qRT-PCR. High expression levels of MEG3 were detected in non-neoplastic tissues, but markedly lower levels were identified in cancer tissues (P < 0.01, Fig. 1B).

To further confirm our finding that MEG3 is down-regulated in cervical cancer in comparison to non-tumor tissues, we analyzed published data on Oncomine Database and found increased expression of MEG3 in non-tumor cervical tissues compared with cervical cancers (Fig. 1A, lower panel). These results strongly indicate that MEG3 may contribute to tumorigenesis and malignant progression of cervical cancer.

MEG3 inhibits cell proliferation in human cervical carcinoma cell lines in vitro. Based on the observation that MEG3 expression level is high in normal cervical tissues but is low or undetectable in cervical cancer, we hypothesized that MEG3 is involved in the regulation of cervical cell proliferation. We tested this hypothesis in human cervical cancer cell lines HeLa and C-33A, which express very low levels of MEG3 [21]. HeLa or C-33A cells were transiently transfected with pEGFP-MEG3 or control vector, and cell viability was examined using the MTS cell growth assay. HeLa and C-33A cells were efficiently transfected with the pEGFP vector, as indicated by the green fluorescence introduced to the cells (data not shown). Over-expression of MEG3, as confirmed by qRT-PCR (Fig. 2A, left two panels), significantly (P < 0.05) inhibited the growth of HeLa or C-33A cells compared with control (Fig. 2B). To study whether down-regulation of MEG3 promotes cervical cancer cell proliferation, MEG3 was knocked down by siRNA in the well-differentiated HCC94 cells which have high levels of endogenous MEG3 (Fig. 2A, right panel). siRNA-MEG3 transfection increased the growth of HCC94 cells (P < 0.05, Fig. 2B). These results indicate that MEG3 inhibits the proliferation of human cervical cancer cells in vitro.
MEG3 induces G2/M cell cycle arrest and apoptosis. Induction of cell cycle arrest is an important anti-proliferation mechanism. Next we investigated whether the growth inhibitory activity of MEG3 involved the control of cell cycle progression. Over-expression of MEG3 resulted in cell cycle arrest at G2/M phase in HeLa and C-33A cells (Fig. 3A). Microscopic observations revealed significant amount of cell death in addition to growth arrest following MEG3 transfection (data not shown). Therefore, we investigated whether MEG3 could also promote apoptotic cell death. HeLa and C-33A cells were transiently transfected with MEG3 for 48 h and apoptosis was analyzed by annexin V/7-AAD staining. Over-expression of MEG3 induced both early (annexin V+/7-AAD-) and total (annexin V+) apoptosis in HeLa and C-33A cells (Fig. 3B).

The p53 protein is a potent tumor suppressor and one of the important players in apoptosis signaling. LncRNAs have been shown to physically associate with p53, suggesting that lncRNAs may function through p53 pathway [22]. Therefore, we examined whether MEG3 regulates the expression of p53. Overexpression of MEG3 resulted in increased expression of p53 in Hela and C33A cells, as assessed by Western blot analysis (Fig. 4). Additionally, MEG3 increased the cleavage of p53 downstream target caspase 3 (Fig. 4). To elucidate the mechanisms for the G2/M phase arrest in MEG3-transfected cells, we investigated the expression of pivotal protein (cyclin B1 and CDK1) for G2/M transition [23]. Compared with control transfected cells, transfection with MEG3 resulted in decrease in cyclin B1 and CDK1 expression (Fig. 4). p21 is one of the p53 targets and acts as an inhibitor of G2/M transition [24]. As shown in Fig. 4, elevated p21 level was detected following MEG3 transfection. These results suggest that MEG3 may induce apoptosis through p53-mediated pathway.
Figure 3. Over-expression of MEG3 induced G2/M arrest and apoptosis in cervical cancer cells. HeLa and C-33A cells were transiently transfected with pEGFP-MEG3 or control vector for 48 h. A. Cell cycle analysis was performed by PI staining using flow cytometry. Representative histograms were presented (right panel). Results shown are representative of two independent experiments. *, p < 0.05. (left panel). B. Apoptosis was assessed by annexin V-FITC/7-AAD staining using flow cytometry. Representative histograms were presented (right panel). Results shown are representative of two independent experiments. *, p < 0.01.
LncRNAs belong to a newly identified class of RNA molecules which do not have protein-coding capacity [25]. LncRNAs control transcription of target genes, which indicates that the difference in IncRNA profiling between normal and cancer cells may not merely be the secondary effect of cancer transformation. Further, IncRNAs may be associated with cancer progression [26]. Thus, IncRNA expression profiles may be of important value in cancer diagnosis, prognosis and therapy.

In the present study, we evaluated the expression of IncRNA MEG3 in cervical cancer tissues and matched non-tumor tissues. We demonstrated that MEG3 expression was significantly (P < 0.01) decreased in cervical cancer tissues compared with adjacent non-tumor tissues. In addition, the search of Oncomine database and published literature confirmed our findings that MEG3 is significantly down-regulated in cervical cancer compared with normal tissues. Further, ectopic expression of MEG3 inhibited the proliferation of human cervical carcinoma cells HeLa and C-33A in vitro. Knockdown of MEG3 promoted cell proliferation in well-differentiated cervical cancer HCC94 cells.

Braconi et al.[17] reported that ectopic expression of MEG3 induced apoptosis in hepatocellular cancer PRC/PRF/5 cells. MEG3-induced apoptosis was also observed in U251 and U87 glioma cells [15]. We further demonstrated that over-expression of MEG3 promoted apoptotic cell death and induced G2/M cell cycle arrest in HeLa and C-33A cells. We investigated the molecular mechanisms responsible for apoptosis and G2/M cell cycle arrest induced by MEG3 in HeLa and C-33A cells. Western blot analysis results showed that CDK1 and cyclinB1 were down-regulated by the overexpression of MEG3, indicating that CDK1 and cyclinB1 may be involved in the induction of G2/M cell cycle arrest by MEG3. In addition, MEG3 activates p53 and its downstream targets caspase 3 and p21. These results suggest that MEG3-mediated inhibition of tumor cell proliferation is due, at least in part, to the induction of G2/M cell cycle arrest and apoptosis through p53 and cyclin-CDK pathways.

The detailed mechanisms by which MEG3 inhibits tumor growth remain to be determined. Gordon FE et al [27] discovered increased expression of angiogenic genes in the brains of mouse MEG3-null embryos, suggesting that deletion of MEG3 promotes angiogenesis. It is well known that tumor growth and metastasis both require angiogenesis [28]. Therefore, enhanced angiogenesis may be another mechanism by which inactivation of MEG3 contributes to tumor development.

In conclusion, we demonstrate that MEG3 is significantly down-regulated in human cervical cancer tissues compared with matched non-neoplastic tissues. Over-expression of MEG3 inhibits human cervical cancer cell growth, which is associated with the induction of G2/M cell cycle arrest and apoptosis. These findings, together with findings by others, indicate that MEG3 is a tumor suppressor and may be a potential target for cancer therapy.

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References

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