Long non-coding RNA MIR22HG suppresses cell proliferation and promotes apoptosis in prostate cancer cells by sponging microRNA-9-3p

Wei Zhang, Chunmei Shi, Qian Xu, Xinfeng Chen, Hua Zhu, and Bing Zheng

Department of Urology, The Second Affiliated Hospital of Nantong University, Nantong, Jiangsu, China

ABSTRACT

The present study was designed to discuss long non-coding RNA (lncRNA) MIR22HG expression in prostate cancer and to address its effect on prostate cancer cells. MIR22HG and microRNA (miR)-9-3p expressions in prostate cancer cells were examined with the use of quantitative real-time PCR (qRT-PCR). Cell counting kit (CCK)-8, colony formation, and TUNEL were conducted to determine cell viability and apoptosis. Immunofluorescence was employed for the detection of Ki67 expression, and western blotting was applied for the examination of apoptosis-related proteins. The relationship of MIR22HG and miR-9-3p was verified employing luciferase reporter assay. Indeed, low MIR22HG expression was discovered in prostate cancer cells. Subsequently, in vitro loss-of-function studies revealed that MIR22HG overexpression suppressed cell proliferation but promoted cell apoptosis, accompanied with a reduction in Ki67 and Bcl-2 expressions, as well as an elevation in Bax and cleaved caspase 3 expressions. In addition, MIR22HG was identified as a sponge of miR-9-3p and the impacts of MIR22HG overexpression on cell proliferation and apoptosis were partly hindered by miR-9-3p overexpression. In summary, MIR22HG acts as an anticancer gene in prostate cancer via inhibiting cell proliferation and promoting apoptosis by sponging miR-9-3p. This article may provide a novel insight into the treatment of prostate cancer.

ARTICLE HISTORY

Received 24 January 2022
Revised 11 May 2022
Accepted 13 May 2022

KEYWORDS

MIR22HG; miR-9-3p; prostate cancer; proliferation; apoptosis

Highlights

- LncRNA MIR22HG suppresses cell proliferation and promotes apoptosis in prostate cancer cells
- MIR22HG acts as an anti-cancer gene in prostate cancer by sponging miR-9-3p.
- MIR22HG may be a candidate of the targeted therapy of prostate cancer treatment.

Introduction

Prostate cancer is a disease most prevalent among American men and ranks as the second major contributor to cancer-associated death in older men [1]. Recently, with the changes of aging, lifestyle, and environment, the incidence and mortality of prostate cancer in China have shown a significant increase in the past decade [2]. Patients, who receive close monitoring and endocrine therapy at a local early stage, can maintain a relatively satisfactory prognosis within 5 years; however, due to the hidden symptoms of prostate cancer, many patients are diagnosed at middle-advanced stages [3–5]. At present, androgen deprivation therapy (ADT) has become widespread and is regarded as the standard care for patients. Despite the fact that ADT initially has a high response rate and significantly delays the recurrence of prostate cancer, most prostate cancers relapse over time and become resistant to androgen-free therapy, thus evolving from androgen-dependent to castration-resistant, leading to the metastasis and recurrence of prostate cancer [6–8]. The identification of oncogenes and tumor suppressor genes indicates that genetic factors play an important role in the pathogenesis of
prostate cancer [9]. In recent years, long non-coding RNAs (lncRNAs) have been identified as key determinants of cancer biology owing to their function in regulating cancer cell behavior [10,11]. LncRNA NR_028502.1, which is located at 17p13.3 and defined as a lncRNA by Encyclopedia of DNA Elements (ENCODE), was named miR22HG (human miR-22 host gene) for its role as a host gene of miR-22. Emerging literature have found that lncRNA MIR22HG has a dual role in a variety of cancers. Being an oncogene, lncRNA MIR22HG was upregulated in esophageal cancer [12]; Meanwhile, lncRNA MIR22HG was expressed at a low level and was closely related to tumor microvascular invasion in hepatocellular carcinoma [13,14]. However, no studies have reported the specific involvement of MIR22HG in prostate cancer. Clinical investigations revealed that MIR22HG expression is decreased in prostate cancer, and the depressed expression of MIR22HG is markedly related to higher Gleason score and T stage, indicating that MIR22HG may be related to the occurrence and development of prostate cancer [15].

Therefore, the study intended to provide a detailed examination of the specific function of MIR22HG in prostate cancer, offering a novel insight into the targeted therapy of prostate cancer.

**Methods and materials**

**Cell culture**

American Type Culture Collection (ATCC; Manassas, VA, USA) was the supplier of normal prostate epithelial cell line RWPE-2 (cat. no. CRL-11610) and prostate cancer lines including 22Rv1 (cat. no. CRL-2505), DU145 (cat. no. HTB-81), LNCaP (cat. no. CRL-1740) and PC3 (cat. no. CRL-1435). RPMI1640 medium supplied by Thermo Fisher Scientific was chosen as the medium for the cultivation of above cell lines, to which 10% fetal bovine serum (FBS) that also provided by Thermo Fisher Scientific, 100 mg/ml streptomycin and 100 U/ml penicillin were added as required. The culture conditions were at 37°C with 5% CO₂.

**Quantitative real-time PCR (qRT-PCR)**

Total RNA extraction processed with Trizol reagent (Invitrogen) was applied for concentration assays adopting Nanodrop 2000 (Thermo Fisher Scientific, Inc.). The synthesis of complementary DNA (cDNA) was conducted through reverse transcription with the aid of a PrimeScript RT reagent kit (Takara Bio, Shiga, Japan) in compliance with the standard procedures of supplier. qRT-PCR was undertaken in accordance with the instructions of the SYBR Premix Ex Taq™ II kit (Takara Bio). The used primers were listed as follows: MIR22HG, forward, 5’-CGGACGCAGTGTCTTGC-3’, reverse, 5’-GCTTTAGCTGGTGCTAGACCA-3’; miR-9-3p, forward, 5’-TCTTTGTATCTAGCTGTAT-3’, reverse, 5’-GAACATGTCTGCGTATCTC-3’; β-actin, forward, 5’-GAAGATCAAGATCATTGCTC-3’, reverse, 5’-TACTCCGTCTGCTAGTCAA-3’; U6, forward, 5’-GCTTCGGCGACACATATCATATAAA-3’, reverse, 5’-CGCTTACGAATTTGCGTGTCAT-3’. The 2^−ΔΔCt method was applied for the calculation of gene expressions [16]. Relative expression level of MIR22HG was normalized to the internal control β-actin, and the relative expression level of miR-9-3p was normalized to U6 [17].

**Cell transfection**

Cells were specifically subjected to transfection with the application of using Lipofectamine 3000 (Invitrogen, Carlsbad, CA, USA) [17]. The pcDNA3.1 vector targeting MIR22HG (pcDNA-MIR22HG) and empty vector (pcDNA-NC), as well as miR-9-3p mimic and mimic NC were provided by GenePharma (Shanghai, China). Subsequently, DU145 cells were severally subjected to transfection with the above plasmids. Gene expression was decided utilizing qRT-PCR 48 h post transfection.

**Cell counting kit-8 (CCK-8) assay**

Cells that inoculated in 96-well plates (1 × 10^4 cells/well) were grown for the indicated times (24, 48 and 72 h). Then, further incubation for
2 h was carried out at 37°C following the addition of 10 μl of CCK-8 solution (cat. no. C0037; Beyotime Institute of Biotechnology, Shanghai, China). An absorbance per well was tested at 450 nm with the employment of a microplate reader (Thermo Fisher Scientific, Inc.) for 1 h at room temperature. Subsequently, the cells were treated with TUNEL reaction solution for 1 h at 37°C before being processed with DAPI staining. Eventually, TUNEL-positive cells were photographed with the help of a fluorescence confocal microscope (magnification, x200; Leica Microsystems GmbH).

**Cell colony formation**

The cells were inoculated in 6-well plates (500 cells/well) and incubated in an incubator containing 5% CO₂ at 37°C for 2 weeks. The medium was kept fresh every 2–3 days during the period. The PBS-washed cells were then subjected to 4% formaldehyde fixation for 15 min and 0.1% crystal violet staining for 20 min at indoor temperature, followed by an observation of colony formation under an inverted light microscope (low magnification; Olympus Corporation) [18].

**Immunofluorescence**

Cells that inoculated in 6-well plates underwent 4% formaldehyde fixation for 10 min and 0.1% Triton X-100 permeabilization at room temperature for 5 min. Subsequently, cells were blocked with 5% BSA (Thermo Fisher Scientific, Inc.) at room temperature for 1 h, followed by the probe with anti-Ki67 antibody (1:200; cat. no. ab16667; Abcam) at 4°C overnight and further incubation with Alexa Fluor® 488-conjugated secondary antibody (1:400; cat. no. A11008; Molecular Probes; Thermo Fisher Scientific, Inc.) for 1 h. Meanwhile, DAPI staining was applied to process the nucleus away from light. Eventually, image capture was undertaken with the application of a fluorescence confocal microscope (magnification, x200; Leica Microsystems GmbH) [19].

**TUNEL assay**

The terminal deoxynucleotidyl transferase-mediated terminal deoxynucleotidyl transferase-mediated dUTP-digoxigenin nick end labeling (TUNEL) system (Promega, Madison, WI, USA) was applied to assess cell apoptosis [18]. Cells seeded on glass coverslips underwent 4% formaldehyde fixation for 10 min, protease K (10 mg/mL, Sigma, USA) pretreatment for 20 min, as well as blockade of 5% BSA (Thermo Fisher Scientific, Inc.) for 1 h at room temperature. Subsequently, the cells were treated with TUNEL reaction solution for 1 h at 37°C before being processed with DAPI staining. Eventually, TUNEL-positive cells were photographed with the help of a fluorescence confocal microscope (magnification, x200; Leica Microsystems GmbH).

**Western blotting**

RIPA lysis buffer with the addition of protease inhibitor Cocktail (Beyotime Institute of Biotechnology) was applied to extract total proteins, followed by the protein concentration determination adopting BCA kit. Proteins (40 μg) was subjected to SDS-PAGE and electrophoresis was performed under a voltage of 60 V-90 V. The isolated protein was then electrically moved to PVDF membrane and sealed with 5% skimmed milk at room temperature for 1 h. The primary antibodies against Bcl-2 (1:1,000; cat. no. ab196495; Abcam), Bax (1:1, 000; cat. no. ab182733, Abcam), cleaved caspase 3 (1:1, 000; cat. no. #9661, Cell Signaling Technology), caspase 3 (1:1, 000; cat. no. #9662, Cell Signaling Technology) and GAPDH (1: 2, 500; cat. no. ab9485, Abcam) were added for overnight incubation at 4°C. On the next day, PVDF membrane was washed with Tris buffered saline with Tween®20 (TBST), and then incubated with horseradish peroxidase-labeled secondary antibody (1:5, 000; cat. no. sc-2004; Santa Cruz Biotechnology, Inc.) at 37°C for 2 h in a shaker. Electrochemiluminescence kit (EMD Millipore, Billerica, MA, USA) was used to visualize protein signals. Finally, Image J software version 1.52 (National Institutes of Health, Bethesda, MD, USA) was used to quantitatively measure the gray level of protein bands.

**Dual luciferase-reporter assay**

The target fragments of wide type (WT) and mutant (MUT) of MIR22HG were subcloned into the pmirGLO dual-luciferase vector (Promega) to generate pmirGLO-MIR22HG-WT/MUT reporter vectors. DU145 cells were co-transfected with WT- or MUT-reporter vectors with miR-9-3p mimic or mimic NC using Lipofectamine 3000 (Invitrogen), respectively. After 48 h, Dual Luciferase Reporter Assay System
(Promega) was applied in accordance with its requirements. The relative luciferase activity was compared to Renilla luciferase activity to reduce the effect of intrinsic variability on experimental accuracy [17].

**Bioinformatics analysis**

MIR22HG expression data in patients with multiple cancers were obtained from Gene Expression Profiling Interactive Analysis database (GEPIA; http://gepia.cancer-pku.cn/).

**Statistical analysis**

The data analysis was done by means of GraphPad Prism version 6.0 (GraphPad software). The presentation method of findings was in the form of mean ± standard deviation (SD) from at least three independent experiments. Assessment of difference was performed by Student’s t-test or one-way ANOVA. A probability level of <0.05 meant that these experimental figures exhibited statistically significance.

**Results**

**MIR22HG was downregulated in prostate cancer**

First, we aimed to understand the expression level of MIR22HG in prostate cancer. MIR22HG expression data in patients with multiple cancers were obtained from GEPIA. The expression level of MIR22HG was downregulated in the majority of cancer tumor samples by contrast to its paired normal samples.

![Figure 1](image_url)

**Figure 1.** MIR22HG was downregulated in prostate cancer. (a) The gene expression profile of MIR22HG across all tumor samples and paired normal tissues. (b) The gene expression profile of MIR22HG in prostate cancer tumor samples and normal samples. *p < 0.05. (c) MIR22HG expression in normal prostate epithelial cell line and prostate cancer lines. ***p < 0.001 vs RWPE-2.
Expectedly, MIR22HG was expressed at a higher level in prostate cancer tumor tissues than that in normal tissues (Figure 1b). Based on this, the expression level of MIR22HG in normal prostate epithelial cell line RWPE-2 and prostate cancer lines (22Rv1, DU145, LNCaP and PC3) was also detected. The results revealed that compared to RWPE-2, MIR22HG was greatly downregulated in prostate cancer cell lines, particularly in DU145 cells (Figure 1c), further demonstrating the downregulation of MIR22HG expression in prostate cancer.

**MIR22HG inhibited cell proliferation and promoted apoptosis in DU145 cells**

To explore the biological role of MIR22HG in prostate cancer, DU145 cells were transfected with pcDNA-MIR22HG to overexpress MIR22HG (Figure 2a). A series of cellular biological experiments revealed that MIR22HG greatly inhibited cell proliferation ability, evidenced by the reduction of cell viability, cell colony formation and Ki67-positive cells upon MIR22HG overexpression (Figure 2b-d). In addition, the apoptotic cells were increased when MIR22HG was overexpressed in DU145 cells (Figure 3a). Meanwhile, reduced protein level of Bcl-2 as well as elevated protein levels of Bax and Cleaved caspase3 were also observed in cells with MIR22HG overexpression (Figure 3b). Taken together, these findings suggested that MIR22HG overexpression greatly suppressed cell proliferation and promoted apoptosis in DU145 cells.

**miR-9-3p is a direct target of MIR22HG**

Next, the potential mechanism of MIR22HG underlying its regulatory function in prostate cancer was explored. By searching on Starbase website...
there may be a binding site between MIR22HG and miR-9-3p, indicating a potential targeting interaction between MIR22HG and miR-9-3p (Figure 4a). Then, dual luciferase-reporter assay was performed and verified this targeting relationship (Figure 4b), implying that miR-9-3p was likely to be a direct target of MIR22HG. Moreover, compared to RWPE-2 cells, a marked upregulation of miR-9-3p was discovered in prostate cancer cell lines (Figure 4c), and overexpression of MIR22HG greatly reduced the expression level of miR-9-3p (Figure 4d). These results suggested that MIR22HG directly targeted miR-9-3p and negatively regulated miR-9-3p expression in DU145 cells.

**miR-9-3p partly abolished the biological functions of MIR22HG in DU145 cells**

Finally, to verify the regulatory relationship between MIR22HG and miR-9-3p, DU145 cells were subjected to transfection with MIR22HG or cotransfection with miR-9-3p mimic or mimic-NC. First, the results in Figure 5a demonstrated a successful transfection upon miR-9-3p mimic. Then, a series of cellular biological experiments were conducted as aforementioned, and the results...
exhibited that the inhibitory effects of MIR22HG on cell viability, cell colony formation, and cell apoptosis were partly abolished by miR-9-3p (Figure 5b-e), revealing an important role of MIR22HG/miR-9-3p axis during the progression of prostate cancer.

Discussion
In this study, we demonstrated that MIR22HG displayed antioncogenic features in contrast to miR-9-3p. It was demonstrated that miR-9-3p was a direct target of MIR22HG, and could weaken the inhibitory effects on cell proliferation and promotive effects on cell apoptosis by MIR22HG. In this way, data in this study suggested that MIR22HG suppressed the progression of prostate cancer through sponging miR-9-3p. Our findings indicated that MIR22HG could be used as a novel marker of prostate cancer and serve as a therapeutic target for prostate treatment.

Currently, more and more lncRNAs have been characterized as functional transcripts that play crucial roles in a variety of biological processes, as well as the pathological status of prostate cancer [20]. lncRNAs are frequently dysregulated in cancers. Of note, based on transcriptomic studies, hundreds of prostate cancer-associated lncRNAs have been discovered to exert multiple biological functions on the initiation and development of prostate cancer [21]. For instance, upregulation of lncRNA PCAT1 was reported to promote prostate cancer cell growth and tumor growth by upregulating late androgen response genes [22]; lncRNA LINC00844, a novel co-regulator of androgen receptor that plays a central role in the androgen transcriptional network, exerted anti-migratory and anti-invasive properties in prostate cancer cells, thereby suppressing the development and progression of prostate cancer [23]. LncRNA MIR22HG is one of the important lncRNAs in various tumor carcinogenesis and tightly linked to cancer cell proliferation, invasion, and tumor growth [12,24]. For example, MIR22HG silencing prevented the loss of viability, repressed migration, and invasion, and induced apoptosis in esophageal adenocarcinoma cells [12]. Interfering with MIR22HG reduced cell proliferation rate prevented tumor growth and invasion in vivo, thus inhibiting glioblastoma progression [24]; Upregulation of MIR22HG inhibited endometrial

Figure 4. miR-9-3p partly abolished the biological functions of MIR22HG in DU145 cells. (a) The binding sites between MIR22HG and miR-9-3p from Starbase website (http://starbase.sysu.edu.cn/) prediction. (b) Luciferase reporter assay was applied for the verification of binding interaction. ***p < 0.001 vs mimic NC. (c) miR-9-3p expression in normal prostate epithelial cell line and prostate cancer lines. ***p < 0.001 vs RWPE-2. (D) DU145 cells were subjected to transfection with pcDNA-NC or pcDNA-MIR22HG, and miR-9-3p expression was tested employing qRT-PCR. **p < 0.01 vs pcDNA-NC.
carcinoma cell proliferation and promoted cell apoptosis [25]; MIR22HG exerted its tumor suppressive activity by inhibiting cell survival, proliferation, and tumor metastasis in colorectal cancer [26]. Accordingly, in the present study, we demonstrated that MIR22HG acted as an antioncogene in prostate cancer by suppressing cell proliferation and promoting cell apoptosis in prostate cancer cells.

It is noted that MIR22HG acts as an antitumor player through regulating targeted miRNAs, which helps to elucidate the molecular mechanism.
of MIR22HG in prostate cancer, and studies have found that MIR22HG can play an antitumor role through regulating targeted miRNAs. miRNAs are short (18–25 nt) and highly stable RNAs, which are involved in post-transcriptional regulation of gene expression [27]. For example, MIR22HG could target miR-5000-3p and regulate the expression of miR-5000-3p to inhibit the proliferation and migration of laryngeal cancer cells, thereby inhibiting the occurrence and development of laryngeal cancer [28]; MIR22HG could also regulate its target gene miR-10a-5p and subsequently influence downstream NCOR2 expression, thereby inhibiting the ability of hepatocellular carcinoma cells to proliferate, migrate and invade [14]. We also verified a binding relationship between MIR22HG and miR-9-3p. miR-9-3p has been reported to be associated with the prognosis of patients with esophageal squamous cell carcinoma (ESCC), and its overexpression has close relation with the poor survival of patients with ESCC [29]. The expression of miR-9-3p is upregulated in human medullary thyroid cancer tissues and cells, while miR-9-3p inhibitor can significantly inhibit the cell cycle progression of medullary thyroid cancer cells and promote cell apoptosis [30]. In addition, there was a remarkable rise in miR-9-3p expression in the serum of prostate cancer patients by contrast to that of healthy control, indicating that miR-9-3p may be involved in the pathogenesis of prostate cancer [31]. As expected, we demonstrated that miR-9-3p has a higher expression in prostate cancer cell lines in comparison with that in normal prostate epithelial cell line. MIR22HG could direct target miR-9-3p and negatively regulate miR-9-3p. Overexpression of miR-9-3p could weaken the biological functions of MIR22HG in DU145 cells, evidencing that MIR22HG-induced apoptosis and -suppressed proliferation ability may be through the inhibition of miR-9-3p.

Conclusions

To sum up, prostate cancer cells showed low level of lncRNA MIR22HG. MIR22HG acted as a molecular sponge of miR-9-3p and hinder the progression of prostate cancer by inhibiting cell proliferation and promoting apoptosis. Thus, MIR22HG may be a prospective biological marker for the treatment of prostate cancer.

Disclosure statement

No potential conflict of interest was reported by the authors.

Funding

The study was supported by Nantong City Scientific Project (JCZ21079) and Jiangsu Commission of Health (BJ21010).

Ethical approval

This article does not contain any studies with human participants performed by any of the authors.

Availability of data and materials

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

Author contribution

Concept design: Bing Zheng;
Experiments conduction and statistical analysis: Wei Zhang, Chunmei Shi, Qian Xu, and Xinfeng Chen;
Data analysis and interpretation: Wei Zhang, Chunmei Shi, and Hua Zhu;
Manuscript preparation and editing: Bing Zheng. Wei Zhang;
All authors reviewed the approved the final version of the manuscript.

ORCID

Bing Zheng http://orcid.org/0000-0002-4967-8408

References

[1] Siegel RL, Miller KD, Jemal A. Cancer statistics, 2018. CA Cancer J Clin. 2018;68(1):7–30.
[2] Du Z, Sun T, Hacisuleyman E, et al. Integrative analyses reveal a long noncoding RNA-mediated sponge regulatory network in prostate cancer. Nat Commun. 2016;7:10982.
[3] Wilt TJ, Jones KM, Barry MJ, et al. Follow-up of prostatectomy versus observation for early prostate cancer. N Engl J Med. 2017;377(2):132–142.
[4] Loeb S, Folkvaljon Y, Makarov DV, et al. Five-year nationwide follow-up study of active surveillance for prostate cancer. Eur Urol. 2015;67(2):233–238.
[5] Klotz L, Vesprini D, Sethukavalan P, et al. Long-term follow-up of a large active surveillance cohort of patients with prostate cancer. J Clin Oncol. 2015;33:272-277.

[6] Petrylak DP. Current state of castration-resistant prostate cancer. Am J Manag Care. 2013;19:358-65.

[7] Wang Q, Li W, Liu XS, et al. A hierarchical network of transcription factors governs androgen receptor-dependent prostate cancer growth. Mol Cell. 2007;27(3):380-392.

[8] Tran C, Ouk S, Clegg NJ, et al. Development of a second-generation antiandrogen for treatment of advanced prostate cancer. Science. 2009;324(5928):787-790.

[9] Shen MM, Abate-Shen C. Molecular genetics of prostate cancer: new prospects for old challenges. Genes Dev. 2010;24(18):1967-2000.

[10] Spizzo R, Almeida MI, Colombatti A, et al. Long non-coding RNAs and cancer: a new frontier of translational research? Oncogene. 2012;31(43):4577-4587.

[11] Gutschner T, Diederichs S. The hallmarks of cancer: a long non-coding RNA point of view. RNA Biol. 2012;9(6):703-719.

[12] Su W, Guo C, Wang L, et al. LncRNA MIR22HG abrogation inhibits proliferation and induces apoptosis in esophageal adenocarcinoma cells via activation of the STAT3/c-Myc/FAK signaling. Aging (Albany NY). 2019;11(13):4587-4596.

[13] Dong Y, Yan W, Zhang SL, et al. Prognostic values of long non-coding RNA MIR22HG for patients with hepatocellular carcinoma after hepatectomy. Oncotarget. 2017;8:114041-114049.

[14] Wu Y, Zhou Y, Huan L, et al. LncRNA MIR22HG inhibits growth, migration and invasion through regulating the miR-10a-5p/NCOR2 axis in hepatocellular carcinoma cells. Cancer Sci. 2019;110:973-984.

[15] Shen H, Weng XD, Yang D, et al. Long noncoding RNA MIR22HG is down-regulated in prostate cancer. Math Biosci Eng. 2019;17:1776-1786.

[16] Livak KJ, Schmittgen TD. Analysis of relative gene expression data using real-time quantitative PCR and the 2 (-Delta Delta C(T)) Method. Methods. 2001;25:402-408.

[17] Long H, Li Q, Xiao Z, et al. LncRNA MIR22HG promotes osteoarthritis progression via regulating miR-9-3p/ADAMTS5 pathway. Bioengineered. 2021;12:3148-3158.

[18] Zhang Y, Zhang Y, Wang S, et al. SP1-induced lncRNA ZFPM2 antisense RNA 1 (ZFPM2-AS1) aggravates glioma progression via the miR-515-5p/Superoxide dismutase 2 (SOD2) axis. Bioengineered. 2021;12:2299-2310.

[19] Gong H. Pinocembrin suppresses proliferation and enhances apoptosis in lung cancer cells in vitro by restraining autophagy. Bioengineered. 2021;12:6035-6044.

[20] Mitobe Y, Takayama KI, Horie-Inoue K, et al. Prostate cancer-associated lncRNAs. Cancer Lett. 2018;418:159-166.

[21] Hua JT, Chen S, He HH. Landscape of Noncoding RNA in Prostate Cancer. Trends Genet. 2019;35:840–851.

[22] Guo H, Ahmed M, Zhang F, et al. Modulation of long noncoding RNAs by risk SNPs underlying genetic predispositions to prostate cancer. Nat Genet. 2016;48:1142–1150.

[23] Lingadahalli S, Jadhao S, Sung YY, et al. Novel lncRNA LINC00844 Regulates Prostate Cancer Cell Migration and Invasion through AR Signaling. Mol Cancer Res. 2018;16:1865–1878.

[24] Han M, Wang S, Fritah S, et al. Interfering with long non-coding RNA MIR22HG processing inhibits glioblastoma progression through suppression of Wnt/beta-catenin signalling. Brain. 2020;143:512–530.

[25] Cui Z, An X, Li J, et al. LncRNA MIR22HG negatively regulates miR-141-3p to enhance DAPK1 expression and inhibits endometrial carcinoma cells proliferation. Biomed Pharmacother. 2018;104:223–228.

[26] Xu J, Shao T, Song M, et al. MIR22HG acts as a tumor suppressor via TGFbeta/SMAD signaling and facilitates immunotherapy in colorectal cancer. Mol Cancer. 2020;19:51.

[27] Bartel DP. MicroRNAs: genomics, biogenesis, mechanism, and function. Cell. 2004;116(2):281–297.

[28] Chen H, Ali M, Ruben A, et al. E2F6-mediated down-regulation of MIR22HG facilitates the progression of laryngocarcinoma by targeting the miR-5000-3p/FBXW7 axis. Mol Cell Biol. 2020;40(10):e00496–19.

[29] Zhong X, Huang G, Ma Q, et al. Identification of crucial miRNAs and genes in esophageal squamous cell carcinoma by miRNA-mRNA integrated analysis. Medicine (Baltimore). 2019;98(27):e16269.

[30] Chen Y, Zhang S, Zhao R, et al. Upregulated miR-9-3p promotes cell growth and inhibits apoptosis in medullary thyroid carcinoma by targeting BLCAP. Oncol Res. 2017;25(8):1215–1222.

[31] Tinay I, Tan M, Gui B, et al. Functional roles and potential clinical application of miRNA-345-5p in prostate cancer. Prostate. 2018;78(12):927–937.