Micro-ribonucleic acid 29b inhibits cell proliferation and invasion and enhances cell apoptosis and chemotherapy effects of cisplatin via targeting of DNMT3b and AKT3 in prostate cancer

Background: Micro-ribonucleic acids (miRNAs) are crucial regulators in malignant tumors. miRNA-29b (miR-29b) has been identified as a tumor suppressor in prostate cancer (PCa). However, very few studies have investigated the effects of miR-29b in PCa, especially the mechanism and its association with chemotherapy. Our study aimed to explore the role and mechanism of miR-29b in PCa.

Materials and methods: The expression levels of miR-29b were detected in ten clinical PCa specimens and four different PCa cell lines through quantitative real-time polymerase chain reaction. After miR-29b mimics and inhibitors were successfully transfected into LNCaP, the 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide assay was then used to investigate cell proliferation and cisplatin sensitivity of PCa cells. Cell cycle, cell apoptosis, and cell invasion were detected via flow cytometry, annexin V–fluorescein isothiocyanate labeling, and transwell assay, respectively. Based on bioinformatic methods, Western blot analysis, and dual-luciferase reporter assay, novel target genes of miR-29b were identified.

Results: miR-29b was downregulated in PCa tissues compared with matched adjacent nontumor tissues. In the androgen-independent PCa cell line (LNCaP-AI), the expression of miR-29b was much lower than the androgen-dependent PCa cell line (LNCaP). Subsequent studies showed that forced expression of miR-29b inhibited cell proliferation and cell invasion and induced cell apoptosis in PCa. Upregulation of miR-29b also enhanced the chemosensitivity of PCa cells to cisplatin. Moreover, we identified DNMT3b and AKT3 as novel target genes of miR-29b in PCa.

Conclusion: Taken together, the results showed that miR-29b plays a tumor-suppressive role in PCa. It inhibits cell biological behavior and enhances the chemotherapy effects of cisplatin through its involvement in epigenetic regulation and PI3K/AKT pathway.

Keywords: miRNA-29b, prostate cancer, cell biological behavior, chemosensitivity, DNMT3b, AKT3
Micro-ribonucleic acids (miRNAs) are endogenous small noncoding RNAs. They work as post-transcriptional regulators of gene expression by binding to the 3′-untranslated region (UTR) of target messenger RNAs (mRNAs). One miRNA can bind to a number of mRNA transcripts, and in turn one mRNA can be targeted by a widespread panel of miRNA species. Evidence has shown that miRNAs are crucial regulators of cancer by targeting different signaling pathways and multiple transcripts.

The miRNA (miR)-29 family consists of three members – miR-29a, miR-29b, and miR-29c – differing only in two or three bases. As a member of miR-29 family, miR-29b is generally recognized as a fundamental regulator of the epithelial–mesenchymal transition (EMT) involved in cancer metastasis and chemoresistance. Several reports have revealed that miR-29b is a critical regulator of cancer, mostly through targeting oncogenes and exerting epigenetic regulation. Studies have reported that miR-29b is downregulated in PCa and acts as a tumor suppressor. However, there are very few studies focusing on the role and underlying mechanisms of miR-29b in PCa. Based on these facts, we undertook this study to investigate the specific roles of miR-29b in PCa. Its effects on cell biological properties, chemoresistance, and target genes in PCa were explored.

**Materials and methods**

**Clinical prostate specimens**

Clinical specimens were obtained from patients at the Second Xiangya Hospital of Central South University, Changsha. All the patients had been diagnosed with PCa and had undergone radical prostatectomy during April 2012 to October 2013. Cancer tissues and matched noncancerous tissues were obtained immediately after the prostatectomies were resected. The specimens were then maintained at −80°C. The protocol was approved by the Institutional Review Board of Central South University.

**Cell lines and cell culture**

The human PCa cell lines PC3, DU145, and LNCaP were procured from the cell bank of the Chinese Academy of Sciences in Beijing, People’s Republic of China (PRC) and were cultured from the cell bank of the Chinese Academy of Sciences in Beijing, People’s Republic of China (PRC) and were cultured in Roswell Park Memorial Institute (RPMI) 1640 medium containing 10% fetal bovine serum (FBS), 100 U/mL penicillin, and 100 μg/mL streptomycin (Sigma-Aldrich, St Louis, MO, USA) in a humidified incubator at 37°C containing 5% CO₂. The androgen-independent cell line LNCaP-AI was derived from LNCaP after culturing in androgen-removed culture medium. The LNCaP-AI cell line was cultured in RPMI 1640 medium with FBS after processing with 10% activated carbon and glucose. A miR-29b mimic sequence was synthesized by GenePharma Technology (Shanghai, PRC). The miR-29b mimic sequence was transfected at working concentrations using Lipofectamine 2000 reagent (Thermo Fisher Scientific, Waltham, MA, USA).

**Quantitative real-time PCR**

Total RNA was extracted using Trizol reagent from the cells (Trizol RNA-extraction kit; Qiagen, Venlo, the Netherlands) according to the manufacturer’s protocol. For miRNA-expression analysis, complementary deoxyribonucleic acid (DNA) was synthesized from total RNA (10 ng) using a TaqMan miRNA reverse-transcription kit (Thermo Fisher Scientific) with miRNA-specific primers. RNU6 was used as an internal control. The primer of miRNA amplification was miR-29b-forward 5′-GGGTAGCACCATATTGAAATC-3′, miR-29b-reverse 5′-TTGGCAGCTCATTGAAATC-3′, U6-forward 5′-CTGCCTTCGCGAGCGACG-3′, and U6-reverse 5′-AACGCTTACGAGGTTAATC-3′. Quantitative real-time polymerase chain reaction (qRT-PCR) was carried out using All-in-One™ q-PCR Mix (GeneCopoeia, Rockville, MD, USA) on an ABI Prism 7900HT system (Thermo Fisher Scientific). Template-negative and reverse transcription-negative conditions were used as controls. All experiments were repeated three times, and all assays contained a negative-control reaction group. All samples were normalized to internal controls. The fold change in expression was then obtained using the ddCt method.

**Transfection with miR-29b mimics**

The hsa-miR-29b mimics and negative-control miRNA were respectively used (hsa-miR-29b mimics sense 5′-UAGCACCACUUUAAUUGGUGIU3′, antisense 5′-AACACUGAUUUCUAAUGUGCUA-3′; negative-control miRNA sense 5′-UUUCGCACGUGUACGUTT-3′, antisense 5′-AGGUGUACACGGUGGAGAATT-3′). LNCaP cells were seeded in six-well plates and cultured with serum-free medium for 24 hours. Lipofectamine 2000 was used to transfect cells with miR-29b mimics, inhibitors, or negative-control miRNA according to the instructions. Concentrations for miR-29b mimics and inhibitors were 2 μM. RPMI 1640 containing 10% FBS was used to culture transfected cells. miR-29b mimics and negative-control miRNA were synthesized by GenePharma Technology. The transfection efficiencies of miRNA in LNCaP cells were confirmed based on the expression level of miR-29b by quantitative real-time (qRT)-PCR and immunofluorescent assay. Transfection efficiency more than 80% was defined as...
successful transfection. Fluorescent cells and normal cells were observed under light microscopy and fluoroscopy, respectively. Transfection efficiency in our experiment was more than 80%, indicating a successful transfection (Figure 1A). Relative miR-29b expression levels in the transfection group were significantly higher than the negative-control group (*P<0.05, Figure 1B).

Cell proliferation and cisplatin sensitivity (MTT assay)
Based on the level of miR-29b expression, the LNCaP cell line was selected for the research. Cisplatin was obtained from Selleck Chemicals (Shanghai, PRC). According to the results of our preliminary experiment, the optimum cisplatin concentration was 5.0 μg/mL. LNCaP cells of each group were plated in 96-well plates at 1x10⁴ per well in a final volume of 100 μL. 3-(4,5-Dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) was added after incubation for 24 hours in a humidified incubator at 37°C with 5% CO₂. The plates were incubated for another 4 hours. Dimethyl sulfoxide (Sigma-Aldrich) was used at 37°C for 10 minutes following removal of culture medium. The absorbance was measured at 570 nm by a microplate reader (Molecular Devices, Sunnyvale, CA, USA).

Cell-cycle, cell-apoptosis, and cell-invasion assays
To evaluate the effect of miR-29b on the cell cycle and cell apoptosis, flow cytometry and annexin V–fluorescein isothiocyanate (FITC) labeling was used. The cells transfected with miR-29b mimics or negative-control miRNA were harvested and resuspended in 100 mL of flow-cytometry staining buffer. The buffer contained 1× phosphate-buffered saline (PBS) and 2% FBS. The cells were then stained with propidium iodide (PI; Sigma-Aldrich) staining solution. PI fluorescence was measured, and PI fluorescence-positive cells were counted as dead cells. The collected cells were resuspended in 0.5 mL of PBS. Then, they were fixed with 4.5 mL of 70% ethanol for 4 hours, washed with PBS, and stained with PI staining solution. The cell-cycle distribution was determined by using a FACSCalibur flow cytometer (FL-2 channel). Cell apoptosis was detected via an annexin V-FITC labeling kit purchased from Kaiji Biotech (Nanjing, PRC), and was carried out.
according to the supplier’s instructions. FITC-labeled cells were counted and analyzed by a FACS Aria™ flow cytometer (BD Biosciences, San Jose, CA, USA).

Matrigel (BD Biosciences) was utilized according to the instructions. Diluted with FBS-free RPMI 1640 with a ratio of 1:8, the diluted Matrigel was then added to the transwell bottom chamber. LNCaP cells were treated with 0.25% trypsin and added to RPMI 1640 to prepare a 1×106/mL single-cell suspension. A transwell chamber was then put into a 24-well plate. RPMI 1640 (600 μL) containing 10% FBS and 200 μL of the prepared single-cell suspension were added. After 48 hours’ incubation at 37°C in an incubator with 5% CO₂, all of the liquid was removed from the transwell chamber and the bottom chamber. Nitrocellulose membranes were used for Western blot analysis. The membrane was then washed with PBS three times, immersed in methanol, and kept for 20 minutes at room temperature, followed by hematoxylin staining for 10 minutes. The cells that migrated through the pores to the lower surface of the membrane were counted under a microscope (×400).

Bioinformatics and Western blot analysis
Identification of potential miR-29b target genes was performed using the TargetScan and PicTar databases. The putative miRNA targets included the most recent TargetScan predictions and the 3′-UTR targets reported by Ru et al. Protein lysates were separated by 10% sodium dodecyl sulfate polyacrylamide gel electrophoresis (Sigma-Aldrich) and transferred to nitrocellulose blotting membranes (Pierce, Rockford, IL, USA). The membranes were then incubated with fluorescence-labeled secondary antibody IgG (1:4000 dilution). GAPDH was used as the control. The company information and catalog numbers for antibodies SOCS-1, Mcl-1, DNMT3b, MMP2, IFN-γ, SMAD3, SOCS5, AKT3, and GAPDH were as follows: ABCAM/ab9870, Cst/5453, Abcam/ab2851, Abcam/ab124294, Cst/3159, Cst/9523, Abcam/ab3695, Cst/8018, and SANTA/SC-365062, respectively. After exposure and development, protein expression was analyzed using a gel-imaging analysis system (Gel-Pro 4.0). Western blot assays were performed in triplicate.

Dual-luciferase reporter assay
The Dual-Luciferase® Reporter Assay System (Promega, Fitchburg, WI, USA) was used according to the protocol. Dual-luciferase reporter constructs containing the putative miR-29b binding sites from DNMT3b and AKT3 were used as described previously. Briefly, partial sequences of the 3′-UTR of AKT3 (454 nt) and sequences of the 3′-UTR of DNMT3b (443 nt) were inserted in the pmirGLO vector (Promega). Constructs were cotransfected into LNCaP cells cultured in 12-well plates using Lipofectamine 2000. Briefly, 30 nM miRNA mimics, inhibitors, or control miRNA was transfected with 0.6 μg of the pMiR 3′-UTR clones. Luciferase activity was measured at 48 hours posttransfection.

Statistical analysis
SPSS 16.0 software was used for statistical analysis. All data are presented as means ± standard deviation of three independent experiments. Student’s two-tailed t-test was used for comparisons of two independent groups. P-values <0.05 were considered statistically significant.

Results
Expression levels of miR-29b in prostate cancer specimens and cancer cell lines
Ten patients receiving radical prostatectomy were included in our study, with approval of the Institutional Review Board of Central South University. All the patients provided written informed consent. The age range of these ten patients was 56–74 years, with an average age of 68.1 years. There were seven T1–T2 patients of and three T3 patients according to TNM (tumor, node, metastasis) staging. There were four G1–G2 patients and six G3–G4 patients of according to pathologic grading (Table 1). miR-29b expression was significantly lower in cancer tissues than in noncancerous tissues (P<0.05, Figure 1C). However, no statistical significance and no correlation between miR-29b levels and PSA, prostate-specific antigen; TNM, tumor, node, metastasis. (Promega). Constructs were cotransfected into LNCaP cells cultured in 12-well plates using Lipofectamine 2000. Briefly, 30 nM miRNA mimics, inhibitors, or control miRNA was transfected with 0.6 μg of the pMiR 3′-UTR clones. Luciferase activity was measured at 48 hours posttransfection.

Table 1 Patient characteristics

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Abbreviations: PSA, prostate-specific antigen; TNM, tumor, node, metastasis.
represented the androgen-independent cell line, was significantly lower than miR-29b expression in LNCaP cells (Figure 1D). Since LNCaP cells showed relatively low miR-29b expression level and had been mostly used in our previous studies, they were selected for our next experiments.

Upregulation of miR-29b expression inhibits cell growth and invasion and induces apoptosis of PCa cells

Cell-cycle changes were examined by flow cytometry in the miR-29b mimic-transfected group (miR-29b group) compared with the negative-control group (Cont-miR). In the miR-29b group, the proportion of sub-G0/G1-phase cells increased significantly, and the proportion of S-phase and G2/M-phase cells decreased significantly compared with the Cont-miR group (P<0.05, Figure 2A), indicating that overexpression of miR-29b inhibited cell growth in PCa.

Cell apoptosis was then determined by annexin V-FITC labeling. The total apoptosis rate in the miR-29b group was significantly higher than in the Cont-miR group (P<0.05, Figure 2B), suggesting that overexpression of miR-29b induced cell apoptosis in PCa.

Similar results were revealed in transwell chambers for invasion assay between the miR-29b group and Cont-miR group. The invasion cells were more significantly reduced in the miR-29b-transfected group compared to the Cont-miR group (P<0.05, Figure 2C), demonstrating that increased miR-29b expression repressed cell invasion in PCa.

**Figure 2** Upregulation of miR-29b expression inhibited growth and invasion and induced apoptosis of prostate cancer cells. **Notes:** (A) In the miR-29b mimic-transfected group (miR-29b), the proportion of sub-G0/G1-phase cells increased significantly, and the proportion of S-phase and G2/M-phase cells decreased significantly compared with the negative-control group (Cont-miR). (B) The percentage of apoptotic cells in the miR-29b group was significantly higher than in the Cont-miR group (P<0.05). (C) The invasion cells were significantly reduced in the miR-29b-transfected group compared to the negative-control group (P<0.05). **Abbreviations:** miR, micro-ribonucleic acid; FITC, fluorescein isothiocyanate.
Overexpression of miR-29b enhances chemosensitivity of PCa cells to cisplatin

There were five groups: the blank control group (Blank), LNCaP transfected with negative-control miRNA (Cont-mir), LNCaP transfected with negative-control miRNA plus cisplatin (Cont-mir + cisplatin), LNCaP transfected with miR-29b mimics (miR-29b), and LNCaP transfected with miR-29b mimics plus cisplatin (miR-29b + cisplatin). The LNCaP cell-growth rate in miR-29b was significantly lower than in Blank and Cont-mir (P<0.05), indicating that miR-29b inhibited cell proliferation of PCa. The cell-growth rate in miR-29b + cisplatin was significantly lower than Cont-mir + cisplatin and miR-29b (P<0.05), suggesting that overexpression of miR-29b enhanced the chemotherapy of cisplatin on PCa (Figure 3A).

miR-29b directly targets DNMT3b and AKT3

Potential miR-29b target genes were searched using the TargetScan and PicTar databases. Eight potential candidate target genes were screened, including SOCS-1, Mcl-1, DNMT3b, MMP2, IFNγ, SMAD3, SOCS3, and AKT3. In parallel, GAPDH was also included as a control. Western blot analysis was then done to detect the protein-expression levels of these nine genes (Figure 3B). Among these, the protein expression of Mcl-1, DNMT3b, and AKT3 was significantly decreased in the miR-29b mimic-transfection group and increased in the miR-29b inhibitor-transfection group (Figure 3C). Since previous research by Steele et al has demonstrated that Mcl-1 is a target gene in PCa, we focused on the genes of DNMT3b and AKT3.12 We found putative miR-29b target sites in the DNMT3b 3′-UTR and AKT3 3′-UTR (Figure 4A). Dual-luciferase reporter assays showed reduced luciferase activities in miR-29b mimic-transfected cells and increased luciferase activities in miR-29b inhibitor-transfected cells in both the DNMT3b (Figure 4B) and AKT3 groups (Figure 4C). The results demonstrated that miR-29b targets DNMT3b and AKT3 directly.

Discussion

miRNAs have been known to affect cellular biological behaviors, including the cell cycle and cell proliferation, apoptosis, and invasion.5 There are huge associations of miRNAs with tumor etiology, progression, and prognosis in multiple human cancers.9 Various studies have demonstrated that miR-29b acts as a tumor-suppressive miRNA in many different cancers. However, relatively little research has focused on the effects of miR-29b on PCa. Ru et al demonstrated that miR-29b can inhibit cell growth by targeting antiapoptotic and prometastatic molecules in PCa. miR-29b also acts as an antimetastatic miRNA in PCa cells by regulating EMT signaling.10,12

In our study, we have illustrated that miR-29b expression is decreased in PCa tissues compared with noncancerous tissues, in accordance with previous research results.10 However, no statistical significance and no correlation between levels of miR-29b and Gleason grade or TNM stage.

Figure 3 Overexpression of miR-29b enhances chemosensitivity of cisplatin on prostate cancer cells. miR-29b targets DNMT3b and AKT3.

Notes: (A) Compared with the blank-control (blank) and negative-control miR (Cont-mir) groups, the relative cell-growth rate in the miR-29b group was significantly lower along with action time, indicating that miR-29b inhibited cell proliferation of prostate cancer. Cell viability in the miR-29b group plus cisplatin was significantly lower than the miR plus cisplatin-control group and the miR-29b group, suggesting that overexpression of miR-29b enhanced the chemotherapy of cisplatin for prostate cancer. (B) Western blotting of eight potential genes’ expression at the protein level in the blank, negative-control mirna, mir-29b inhibitor-transfection, and miR-29b mimic-transfection groups. (C) The quantitative protein expression of DNMT3b, AKT3, and Mcl-1 were significantly increased in the miR-29b mimic-transfection group (Mimics) and decreased in the miR-29b inhibitor-transfection group (Inhibitors). *P<0.05.

Abbreviation: miR, micro-ribonucleic acid.
was found. In PCa cells, miR-29b expression was lower in androgen-independent cells than in androgen-dependent cells. This might indicate the association of miR-29b with castration-resistant PCa (CRPC) progress. To assess miR-29b's effects on cellular biological behavior, miR-29b mimics were transfected into PCa cells, and the cell cycle, proliferation, apoptosis, and invasion were then detected. Our findings showed that forced expression of miR-29b inhibited cell proliferation and cell invasion and induced cell apoptosis in PCa. Moreover, miR-29b upregulation enhanced the chemosensitivity of cisplatin on PCa, implying a potential role in solving the issue of chemoresistance in PCa.

With regard to how miR-29b potentiates the effects of enhancing chemosensitivity of PCa cells, there are some possible mechanisms. In acute myeloid leukemia, the epigenetic regulation of miR-29b can enhance the chemotherapy effects.\(^\text{11}\) Probably, miR-29b could also increase the chemosensitivity of PCa through epigenetic regulation. Moreover, EMT is involved in the resistance of cancer cells to chemotherapy, and miR-29b is generally recognized as a fundamental regulator of EMT involved in cancer metastasis and chemoresistance.\(^\text{12,14}\) miR-29b could also increase the chemosensitivity of PCa cells to cisplatin through targeting EMT-related factors. What is more, studies have shown that the PI3K/AKT pathway is involved in the mechanism of cisplatin effects in various cancers.\(^\text{15,16}\) miR-29b has been identified as a negative regulator of the PI3K/AKT pathway.\(^\text{17}\) Our findings demonstrated that AKT3 is a target gene of miR-29b in PCa. As a result, miR-29b may potentiate its effects of PCa chemosensitivity to cisplatin through regulating the PI3K/AKT pathway. In summary, miR-29b could regulate cancer chemosensitivity, possibly through exerting epigenetic effects and regulating the EMT process and PI3K/AKT pathway.

Previous studies on PCa have identified Mcl-1, COL1A1, COL4A1, and MMP-2 as target genes of miR-29b.\(^\text{12}\) Subsequent research indicated that miR-29b directly targets Snail, a key gene involved in the metastatic phenotype.\(^\text{18}\) These results partly shed light on how miR-29b exerts tumor-suppressor effects on PCa. To further investigate possible functional mechanisms of miR-29b in PCa and to identify more target genes of miR-29b, we used bioinformatic methods and Western blot to screen potential candidate target genes. Dual-luciferase reporter assays were then performed to identify target genes, since Mcl-1 has been identified as a target gene of miR-29b multiple myeloma as well as PCa.\(^\text{12,18}\) Therefore,
we focused on \(DNMT3b\) and \(AKT3\) to detect if these two genes were novel target genes of miR-29b in PCa. In this article, the results of dual-luciferase reporter assays identified that miR-29b targeted \(DNMT3b\) and \(AKT3\) directly.

Epigenetic regulations are alterations in gene expression, including DNA methylation, histone modification, and RNA-associated silencing, which are very important pathways in modulating malignant cells. DNA methylation can affect specific gene expression and result in tumor occurrence or antitumor activity.\(^{24}\) \(DNMT3b\) is a member of the DNA methyltransferase family, which generally exerts DNA-methylation effects, accounting for inactivation of tumor-suppressor genes in many cancer cells. Various studies have shown that miR-29b can target DNA methyltransferases and downregulate the global DNA methylation in malignant cells, through which miR-29b exerts its tumor-suppressive role. Increased miR-29b expression could lead to marked downregulation of the expression of DNA methyltransferases, including DNMT1, DNMT3a, and DNMT3b, by direct targeting or indirect interaction in several cancers.\(^{25-27}\) In PCa, hypermethylation of tumor-suppressor genes has been found to be associated with advanced PCa. DNMT activity and DNMT levels were higher in PCa cell lines compared to their nonneoplastic cells, and DNMT3b activity was significantly higher in more aggressive PCa cell lines.\(^{28}\) Moreover, DNMT inhibitors could be potential therapeutic agents for advanced PCa.\(^{29}\) In this study, we observed that forced expression of miR-29b inhibited DNMT3b protein levels by directly targeting DNMT3b, consistent with previous studies in other tumors. This result indicates that epigenic regulation of miR-29b is a very important pathway of tumor suppression in PCa.

Activation of the PI3K pathway, resulting from loss of the \(PTEN\) tumor-suppressor gene, is one of the predominant pathways in PCa.\(^{30}\) The serine/threonine-kinase AKT family is the primary downstream mediator of PI3K signaling. The AKT family consists of three members – AKT1, AKT2, and AKT3 – that are highly homologous isoforms. Although sharing a high degree of homology, the AKT family shows distinct roles in human cancers.\(^{31,32}\) In triple-negative breast cancer, downregulation of AKT3 significantly inhibits the growth of cancer cells.\(^{33}\) AKT3 is also involved in modulating drug sensitivity.\(^{34}\) In PCa, knockdown of the AKT family inhibits the growth of cancer cells in vitro and in vivo, suggesting that targeting the AKT family could be an effective treatment for PCa.\(^{35}\) The regulation of the miR-29 family on cell proliferation and apoptosis is also dependent on the PI3K/AKT pathway in acute myeloid leukemia.\(^{36}\) In multiple myeloma, the PI3K/AKT pathway is a negative regulator of miR-29b by reducing AKT phosphorylation.\(^{37}\) All these previous studies have implicated that AKT3, possibly the AKT family, is a crucial protein in the cellular processes of malignant tumors, including cell growth and drug sensitivity. AKT3 is also involved in the regulation of miR-29b in several cancers. In PCa, knockdown of the AKT family inhibits the growth of cancer cells in vitro and in vivo, suggesting that targeting the AKT family could be an effective treatment for PCa.

In summary, our findings demonstrate that miR-29b was downregulated in PCa and was also downregulated in an androgen-independent PCa cell line compared to an androgen-dependent cell line, suggesting its potential role in the CRPC process. miR-29b can inhibit cell proliferation and invasion and enhance cell apoptosis and chemosensitivity of PCa. Furthermore, we have shown for the first time that miR-29b exerts its tumor-suppressive role through involvement in epigenetic regulation and the PI3K/AKT pathway. Further studies with larger samples are needed to elucidate the effects and mechanism of miR-29b in the CRPC process and chemoresistant cancer.

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Author contributions

All authors contributed toward data analysis, drafting and revising the paper and agree to be accountable for all aspects of the work.
Disclosure

The authors report no conflicts of interest in this work.

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