Mangiferin inhibition of proliferation and induction of apoptosis in human prostate cancer cells is correlated with downregulation of B-cell lymphoma-2 and upregulation of microRNA-182

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Abstract. Mangiferin, a flavonoid extracted from the mango tree, possesses anti-inflammatory, antibacterial, anti-herpes simplex and antitumor activity, and is able to affect immune function. The present study investigated the anticancer effects of mangiferin treatment on PC3 human prostate cancer cells, and the potential underlying mechanisms. In the present study, an MTT assay was used to analyze the proliferation of PC3 cells. Subsequently, flow cytometry and colorimetric assay kits were utilized to measure the PC3 cell apoptotic rate. The expression levels of B-cell lymphoma-2 (Bcl-2) and microRNA-182 (miR-182) were detected using western blot analysis and quantitative reverse transcription-polymerase chain reaction, respectively. Finally, miR-182 and anti-miR-182 were transfected into PC3 cells, which were used to investigate the effects of mangiferin. Mangiferin treatment reduced the proliferation of PC3 human prostate cancer cells in a concentration- and time-dependent manner. In addition, mangiferin was able to promote apoptosis and induce the caspase-3 activity of PC3 human prostate cancer cells. Mangiferin treatment was also able to significantly reduce Bcl-2 expression levels and enhance miR-182 expression in PC3 cells. Finally, it was observed that mangiferin inhibited proliferation and induced apoptosis in PC3 human prostate cancer cells, and this effect was correlated with downregulation of Bcl-2 and upregulation of miR-182.

Introduction

Prostate cancer is a disease that affects older males, and is one of the most common malignant tumors amongst males in Europe (1). Previously, prostate cancer was observed relatively rarely in China; however, in recent years the incidence of prostate cancer and associated mortality rates have been increasing (1). In 2008, the worldwide standardized incidence rate of prostate cancer was 28.5/100,000 and the standardized mortality rate was 7.5/100,000 (2-4). Therefore, prostate cancer has become a significant area of research.

Cell apoptosis may be inseparable from the development of prostate cancer. Apoptosis, or programmed cell death, differs from necrosis, the natural cell death process (5). Reduced apoptosis and increased proliferation are thought to be the primary mechanisms underlying the formation of the majority of tumors. Inhibition of apoptosis may also possess a significant role in the progression of prostate cancer, and B-cell lymphoma-2 (Bcl-2) is thought to be one of the most significant genes required for apoptosis (6).

MicroRNAs (miRs) are a class of endogenously expressed non-protein encoding RNAs (2). Mature miRs are single-stranded and consist of ~22 nucleotides (7). Previous studies have indicated that miRs may be involved in human tumorigenesis and development. Numerous tumors exhibit alterations in miR expression levels, and these miRs may have a role in tumorigenesis, functioning as cancer genes or tumor suppressor genes (8). Casanova-Salas et al (9) demonstrated that miR-182 was a potential biomarker for prostate cancer diagnosis, prognosis predictions and for improvement of the predictive capability of existing biomarkers. Siva et al (10) reported that molecular assays may be used to detect miR-182 expression in prostate cancer.

Mangiferin is a flavonoid, extracted from the mango tree; and studies have revealed that it is capable of inhibiting the growth of liver and colon cancer cells, K562 leukemia cells and other tumor cells (11-14). Pharmacological experiments indicate that mangiferin is capable of effectively treating chronic bronchitis (3). Mangiferin is the primary active ingredient in the Tibetan capillary treatment of hepatitis and is the antiviral active ingredient of rhizoma anemarrhenae, a Traditional Chinese Medicine (4). In the present study, the anticancer effects of mangiferin on prostate cancer cells were investigated and its apoptosis-inducing effects were studied at the cellular level, with the aim of identifying the underlying mechanisms.

Materials and methods

Experimental materials. Mangiferin (purity, >98%) was obtained from Sigma-Aldrich (St. Louis, MO, USA). The chemical structure of mangiferin is illustrated in Fig. 1.

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RPMMI-1640, fetal bovine serum (FBS) and penicillin/streptomycin were obtained from Gibco Life Technologies (Carlsbad, CA, USA). MTT and chemiluminescence reagent kits were purchased from Sangon Biotech Co., Ltd. (Shanghai, China). Cleaved caspase-3 and caspase-9 colorimetric assay, bicinchoninic acid (BCA) Protein assay and UNIQ-10 Column Trizol Total RNA Isolation kit were purchased from Sangon Biotech Co., Ltd. (Shanghai, China). TRizol reagent and oligo (dT) primers were purchased from Invitrogen Life Technologies (Carlsbad, CA, USA). SYBR chemistry and Annexin V-fluorescein isothiocyanate (FITC)/propidium iodide (PI) Apoptosis Detection kits were obtained from Shanghai BestBio Co. (Shanghai, China).

Cell culture. PC3 human prostate cancer cells were purchased from Shanghai Cell Bank of the Chinese Academy of Sciences (Shanghai, China) and cultured in RPMI-1640 medium containing 10% (v/v) FBS with penicillin (100 U/ml)/streptomycin (100 U/ml) in 5% CO2 at 37°C. Approximately every 2-3 days, the culture medium was removed and replaced. PC3 cells were treated with mangiferin in complete RPMI-1640 medium.

Cell proliferation assay. PC3 cells were cultured with mangiferin (10, 20 and 40 µM) for 0, 24, 48 and 72 h. Proliferation of PC3 cells was quantified using an MTT assay. PC3 cells (5x10^4/ml) were seeded into flat-bottomed 96-well culture plates and ~10 µl MTT (5 mg/ml) was added to each well. Cells were incubated for 4 h in 5% CO2 at 37°C. Following this, the culture medium was removed, and 200 µl dimethyl sulfoxide (Invitrogen; Thermo Fisher Scientific, Inc., Waltham, MA, USA) was added to each well and incubated for 10-20 min. The absorbance was measured at 540 nm using a Labsystems Multiskan MS Plate Reader (Thermo Fisher Scientific, Inc.).

Flow cytometry. PC3 cells were cultured with mangiferin (10, 20 and 40 µM) for 72 h. Cells were collected using centrifugation at 3,000 x g for 10 min at 4°C and washed twice with ice-cold phosphate buffered saline (PBS). In accordance with the manufacturer's protocol, cells were incubated with 5 µl Annexin V-FITC on ice for 30 min in the dark. PI (10 µl) was then added to the cells and they were subsequently observed using flow cytometry, using a LSRII (BD Biosciences, San Jose, CA, USA).

Caspase-3 activity assays. PC3 cells were cultured with mangiferin (10, 20 and 40 µM) for 72 h. Caspase-3 activity was detected using the cleaved caspase-3 colorimetric assay kit (Sangon Biotech Co., Ltd.) according to the manufacturer's instructions. The fluorescence of caspase-3 activity was detected at a wavelength of 405 nm, using a Labsystems Multiskan MS Plate Reader.

Western blot analysis of Bcl-2 expression. PC3 cells were treated with various concentrations of mangiferin (10, 20 and 40 µM) for 72 h. Cells were collected using centrifugation at 3,000 x g for 10 min and washed twice with ice-cold PBS. Cells were resuspended with 200 µl protein lysate and centrifuged at 12,300 x g for 15 min at 4°C for 15 min at 4°C. Protein concentration was determined using the BCA Protein Assay kit (Sangon Biotech Co., Ltd.). Approximately 50 µg protein was electrophoresed using a 10% polyacrylamide gel (Beyotime Institute of Biotechnology, Nanjing, China) containing SDS. Proteins were transferred onto nitrocellulose membranes (Millipore Corporation, Bedford, MA, USA) at 4°C for 2-4 h. The membranes were blocked in 5% non-fat milk for 1-2 h and subsequently incubated with rabbit anti-Bcl-2 (15071S; 1:1,000; Cell Signaling Technology, Boston, MA, USA) and rabbit anti-human β-actin (D110001-0025; 1:500; Sangon Biotech Co., Ltd.) antibodies overnight at 4°C with agitation. Following washing with Tris-buffered saline and Tween-20 PBST (with 0.1% Tween 20; Beyotime Institute of Biotechnology) monoclonal anti-rabbit immunoglobulin IgG conjugates (C2162; 1:1,000, Beijing Applygen Technologies, Inc., Beijing, China) for 2 h. Subsequently, membranes were washed with TBST and bands were observed using an enhanced chemiluminescence kit Stripping Buffer for Western Blot (PA114, Tiangen Biotech Co., Ltd.).

Reverse transcription-quantitative polymerase chain reaction (RT-qPCR). PC3 cells were cultured with mangiferin (10, 20 and 40 µM) for 72 h. TRizol reagent (Invitrogen Life Technologies) was used to extract total RNA, and total RNA was purified using an RNA kit (Sangon Biotech Co., Ltd.). Complementary DNA was reverse transcribed using oligo (dT) primers (Invitrogen Life Technologies). SYBR chemistry (BestBio, Co.) was used for PCR. All primers were purchased from Sangon Biotech Co., Ltd. The following primer sequences were used: miR-182 forward, 5'-GGGGAAGGCTGTTTTC-3' and miR-182 reverse, 5'-GGGATTCTTACATGTGAGGCAGGTTCTCAC-3'; U6 forward, 5'-CGGCCTCAGCAGCAGTACTA-3' and U6 reverse, 5'-CGCTTCAAGATTTGCAGTCA-3'.

miRNA transfection. miR-182 mimic and miR-182 inhibitor were purchased from Sangon Biotech Co., Ltd. miR-182 mimic (100 nmol/l) and miR-182 inhibitor (100 nmol/l) transfections were conducted in PC3 cells using Lipofectamine™ 2000 (Invitrogen Life Technologies) according to the manufacturer's instructions. Following transfection for 6 h, culture medium was removed and replaced with complete medium. Subsequently, PC3 cells were incubated for 18 h in 5% CO2 at 37°C, prior to treatment with various concentrations of mangiferin (50 µM) for 72 h.
Figure 2. Treatment with mangiferin has an antiproliferative effect on PC3 cells. Values are expressed as the mean ± standard deviation. *P<0.01 compared with 0 µM mangiferin treatment control group.

Figure 3. Treatment with mangiferin enhances the apoptosis of PC3 cells. (A) Statistical analysis of cellular apoptosis levels and (B) caspase-3 activity. Values are expressed as the mean ± standard deviation. *P<0.01 compared with 0 µM mangiferin treatment control group. FCM, flow cytometry.

Figure 4. Bcl-2 expression is downregulated following treatment with mangiferin. (A) Effects of mangiferin on Bcl-2 and (B) statistical analysis of Bcl-2 protein expression levels. Values are expressed as the mean ± standard deviation. *P<0.01 compared with 0 µM mangiferin treatment control group. Bcl-2, B-cell lymphoma-2.

Figure 5. miR-182 expression is increased following treatment with mangiferin. Values are expressed as the mean ± standard deviation. *P<0.01 compared with 0 µM mangiferin treatment control group. miR-182, microRNA-182.
Statistical analysis. Data analysis was performed using SPSS version 17.0 (SPSS Inc., Chicago, IL, USA). Data were expressed as the mean ± standard deviation. Differences were assessed using the two-tailed Student’s t-test. \( P<0.05 \) was considered to indicate a statistically significant difference.

Results

Mangiferin exerts an antiproliferative effect on PC3 cells. To investigate the potential effects of mangiferin (0, 10, 20 and 40 \( \mu M \)) on the proliferation of PC3 cells, the present study used an MTT assay to examine antiproliferative activity. Mangiferin inhibited the proliferation of PC3 cells in a concentration-dependent manner. When PC3 cells were treated with mangiferin at a concentration of 20 \( \mu M \) for 72 h, or with mangiferin at a concentration of 40 \( \mu M \) for 48 or 72 h, the proliferation of PC3 cells was significantly reduced compared with that of the control group treated with 0 \( \mu M \) mangiferin (Fig. 2).

Mangiferin enhances apoptosis of PC3 cells. To assess the effect of mangiferin (0, 10, 20 and 40 \( \mu M \)) on the apoptosis of PC3 cells, apoptosis and caspase-3 activity were analyzed. Mangiferin accelerated apoptosis and enhanced the caspase-3 activity of PC3 cells in a concentration-dependent manner. In addition, when treated with mangiferin (20 and 40 \( \mu M \)) for 72 h, the apoptosis of PC3 cells was significantly increased compared with that of the control group treated with 0 \( \mu M \) mangiferin (Fig. 3A). Furthermore, the caspase-3 activity of PC3 cells treated with 20 and 40 \( \mu M \) mangiferin was significantly enhanced compared with that of the control group treated with 0 \( \mu M \) mangiferin (Fig. 3B).

Bcl-2 expression is downregulated following mangiferin treatment. In order to elucidate the effect of mangiferin on the expression of Bcl-2, PC3 cells were exposed to 0, 10, 20 and 40 \( \mu M \) mangiferin for 72 h, and the expression of Bcl-2 was analyzed. Bcl-2 protein expression was downregulated...
Following exposure to mangiferin at all concentrations. Furthermore, mangiferin (20 and 40 µM) was able to significantly reduce Bcl-2 expression levels in PC3 cells compared with those of the control group treated with 0 µM mangiferin (Fig. 4A and B).

miR-182 expression is increased following mangiferin treatment. To examine the effect of mangiferin treatment on the expression of miR-182, PC3 cells were exposed to 0, 10, 20 and 40 µM mangiferin for 72 h, and the expression of miR-182 was detected. Following treatment with mangiferin (20 and 40 µM) for 72 h, miR-182 expression was significantly increased in PC3 cells (Fig. 5).

Overexpression of miR-182 inhibits Bcl-2 expression in PC3 cells. In order to further understand miR-182 expression in PC3 cells, an miR-182 mimic was transfected into PC3 cells. It was observed that miR-182 overexpression significantly inhibited Bcl-2 expression in PC3 cells (Fig. 6A and B).

miR-182 inhibitor abrogates the effect of mangiferin. To further evaluate the correlation between miR-182 expression levels and the mechanism of antiproliferative activity of mangiferin, miR-182 inhibitor was transfected into PC3 cells. The results of the present study demonstrated that transfection with miR-182 inhibitor significantly reduced miR-182 expression levels in PC3 cells (Fig. 7A). However, miR-182 inhibitor was also able to significantly reduce the antiproliferative and apoptotic effects of mangiferin (50 µM) on PC3 cells (Fig. 7B and C). Therefore, miR-182 inhibitor may be capable of rescuing Bcl-2 expression in PC3 cells (Fig. 7D).

Discussion

Prostate cancer is one of the most common types of cancer in the male reproductive system (5). Due to socioeconomic development, as well as changes in lifestyle and environment, the incidence and mortality rates of male primary prostate cancer tumors are changing (15). A recent study revealed that the incidence and development of prostate cancer may be associated with the dysregulation of apoptosis regulatory genes (16). Proliferation of the cells that lose their response to apoptosis, may contribute to the invasion and metastasis of prostate tumors. In the present study, it was revealed that mangiferin was capable of inhibiting the proliferation of PC3 cells in a concentration- and time-dependent manner. Pan et al (11) demonstrated that mangiferin was able to inhibit the proliferation of human nasopharyngeal carcinoma cells through G2/M arrest and induction of early apoptosis. In the present study, mangiferin accelerated the apoptotic rate and enhanced the caspase-3 activity of PC3 human prostate cancer cells in a concentration-dependent manner. Taken together, these results indicate that mangiferin may promote apoptosis and increase caspase-3 activity in cancer cell lines (17).

The Bcl-2 protein family contains significant genes involved in the regulation of apoptosis (18). The primary anti-apoptotic and pro-apoptotic members of the Bcl-2 family regulate apoptosis by controlling mitochondrial function (19). Bcl-2 inhibits apoptosis by preventing an increase in cytoplasmic calcium concentration (6). In previous studies, Bcl-2 expression has been observed to be a notable feature in certain human malignant tumors; and Bcl-2 is now believed to be an anti-apoptotic gene that promotes tumor occurrence through inhibiting apoptosis and prolonging the survival of cells (20). In the present study, mangiferin was able to significantly reduce the Bcl-2 expression levels in PC3 cells. Pan et al (11) reported that mangiferin induced cell apoptosis through Bcl-2 and Bcl-2-associated X protein (Bax) expression. Pal et al (21) hypothesized that mangiferin attenuates diabetic nephropathy by inhibiting Bcl-2 and Bax expression.

miRs are a class of small molecule RNA, involved in the regulation of gene expression, which have a significant role in cell growth, differentiation, apoptosis and tumorigenesis (7). Greater than half of human miR genes are located in a genomic region of a tumor-associated or fragile site. Expression profiling has been utilized to reveal that the expression of specific miRs may be indicative of the tumor stage and differentiation state (22). In the present study, treatment with mangiferin significantly increased miR-182 expression levels in PC3 cells. Notably, upregulation of miR-182 suppressed Bcl-2 expression in PC3 cells. In addition, suppression of miR-182 expression reduced the effect of mangiferin on PC3 cells and increased the expression levels of Bcl-2 in PC3 cells. Peng et al (8) reported inhibition of proliferation in PC3 human prostate cancer cells following atorvastatin treatment, due to downregulation of Bcl-2 and upregulation of miR-182. Overexpression of miR-182 has been observed to suppress the growth of uveal melanoma cells via a reduction in Bcl-2 expression levels (23).

Mangiferin is double-benzene pyrone flavonoid that exhibits antioxidant, hypoglycemic and renal protective effects. Mangiferin possesses anti-inflammatory, antibacterial and anti-herpes simplex virus effects; it affects gallbladder and immune function, as well as a protective effect on the simulated high altitude hypoxia liver injury (24-26). In conclusion, the results of the present study indicated that mangiferin is capable of exerting an antitumor effect on PC3 human prostate cancer cells, potentially via downregulation of Bcl-2 and upregulation of miR-182. In addition, the effects of mangiferin may be able to contribute to the characterization of prostate cancer and the development of novel therapies.

References


