Bergamottin exerts anticancer effects on human colon cancer cells via induction of apoptosis, G2/M cell cycle arrest and deactivation of the Ras/Raf/ERK signalling pathway

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Abstract

Introduction: Accumulating evidence has shown the potential of bergamottin as an anticancer agent. The present study was undertaken to evaluate the anticancer effects of bergamottin (µM) against colon cancer cells.

Material and methods: Antiproliferative effects were evaluated by WST-1 cell viability assay. Apoptotic effects were studied by DAPI and Annexin V/PI staining. Cell cycle analysis was carried out by flow cytometry. Transwell assay was used to study the effects on cell invasion. Protein expression was estimated by the western blot method.

Results: The results showed that bergamottin suppresses the proliferation of all the human colon cancer cell lines. Nonetheless, the growth inhibitory effects of bergamottin on the HT-29 and RKO cells were more significant (IC50, 12.5 µM). The anticancer effects of bergamottin on the HT-29 and RKO cells were mainly due to apoptosis. Bergamottin could considerably increase the expression of Bax and reduce the expression Bcl-2. The cleavage of caspase-3, 8 and 9 was also enhanced upon bergamottin treatment of the colon cancer cells. Flow cytometric analysis showed that bergamottin also induced G2/M cell cycle arrest of the HT-29 and RKO cells. Additionally, bergamottin could also suppress the invasion of HT-29 and RKO cells. The Raf/MEK/ERK pathway is regarded as one of the essential pathways involved in the development and progression of cancers. Herein, it was observed that bergamottin could concentration dependently block this pathway in colon cancer cells. In vivo study revealed that bergamottin could also suppress the growth of tumours in xenografted mice models.

Conclusions: Taken together, bergamottin suppresses the proliferation of colon cancer cells and may be utilised in the development of chemotherapy for colon cancer.

Key words: bergamottin, apoptosis, colon cancer, invasion, cell cycle arrest.

Introduction

For millennia, plants have been an amazing pool of therapeutically important metabolites [1]. Although many currently used drugs are of plant origin, plants continue to serve as a source of more drugs or drug leads [2]. The use of plant-derived drugs dates back to the 19th century and since then many phytochemicals have been used in cancer treatment [3].
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Material and methods

Cell culture conditions

The colon cancer cell lines HT-29, SW-948, RKO and SW480, and normal CDD-18Co cells were obtained from the Cancer Research Institute of Beijing (Beijing, China) and maintained in Dulbecco’s modified Eagle’s medium (Invitrogen Life Technologies, Massachusetts, USA), 100 µg/ml streptomycin and 100 U/ml penicillin G (Himedia, Pennsylvania, USA) in an incubator at 37°C with 5% CO₂.

Cell proliferation assay

The viability of the human colon cancer cells and normal astrocytes was monitored by WST-1 assay. In brief, HT-29 and RKO cells were cultured in 96-well plates at the density of 2 × 10⁴ cells/well and treated with 0 to 200 µM concentrations of bergamottin for 24 h at 37°C. This was followed by the incubation of the cells with WST-1 at 37°C for 4 h. The absorbance was then measured at 450 nm using a Victor3 microplate reader to determine the proliferation.

Apoptosis assay

The HT-29 and RKO cells (0.6 × 10⁶) were cultured in 6-well plates and treated with bergamottin at the concentrations of 0, 3.12, 6.2 and 12.5 µM for 24 at 37°C. The cells were then subjected to phosphate buffered saline (PBS) washing and subsequently stained with DAPI. Finally, the cells were examined under a microscope to detect the induction of apoptosis. For determination of the percentage of apoptosis the transfected cells were stained with annexin V/PI and subsequently examined by a flow cytometer.

Cell cycle analysis

The cultured human colon cancer HT-29 and RKO cells were firstly treated with varied concentrations of bergamottin for 24 at 37°C. The cells were then washed with PBS. Afterwards, the HT-29 and RKO cells were stained with propidium iodide (PI) and the distributed of the cells in cell cycle phases was assessed by a FAC5 flow cytometer.

Wound-healing assay

The bergamottin treated colon cancer cells were cultured at 37°C until confluence (80%). This was followed by removal of the media and subsequent washing of the cells with PBS. A scratch was made with a pipette tip and a photograph was taken. The plates were then kept at 37°C for 24 h and a graph was taken again under a microscope.

Cell invasion assay

The effects of bergamottin on the invasion ability of HT-29 and RKO cells was determined by transwell chambers with Matrigel. Around 200 ml cell cultures were placed onto the upper chambers and only medium was placed in the bottom wells. After 24 h of incubation, the cells were removed from the upper chamber and the cells that invaded...
ed via the chambers were subjected to fixation with methyl alcohol and subsequently stained with crystal violet. An inverted microscope was used to count the number of invaded cells at 200X magnification.

**Western blot analysis**

The HT-29 and RKO cells were cultured for 24 h at 37°C. Next the cultures were transferred to 50 ml centrifuge tubes and centrifuged at 12 000 rpm for 10 min. The pellets were washed with PBS and then resuspended in lysis buffer (20 mM HEPES, 350 mM NaCl, 20% glycerol, 1% Nonidet P 40, 1 mM MgCl₂, 0.5 mM EDTA, 0.1 mM EGTA, 1 mM DTT, 1 mM PMSF, protease inhibitor cocktail, and phosphatase inhibitor cocktail). BCA assay was used to determine the concentration of the proteins in each sample. Equal concentrations of the proteins from each sample were loaded and separated on the SDS-PAGE and then transferred to polyvinylidene fluoride membrane. These membranes were then treated with TBS and then exposed to primary antibody (Santa Cruz Biotechnology, Inc., Dallas, TX, USA) at 4°C for 24 h. Afterwards, the cells were treated with HRP-conjugated anti-rabbit secondary antibody and subsequently chemiluminescence reagent was used for the visualisation of the proteins.

**In vivo study**

National Institutes of Health standards for the care and use of laboratory animals and approved by Cancer Hospital of China Medical University under approval number CMU67/34/2018. The mice (4 weeks old, weighing around 24 ±2.2 g) were injected with 5 × 10⁶ HT-29 cells subcutaneously at the left flank. As the tumours were apparent, the mice (n = 5) for each group were injected intraperitoneally with DMSO (0.1%) dissolved bergamottin and diluted with 100 µl of normal saline at 0, 15, 30 and 60 mg/kg body weight and taken as the first day of the experiment. Bergamottin was given to the rats thrice a week while the con-

| No. | Cell line | IC₅₀ [µM] |
|-----|-----------|----------|
| 1   | HT-29     | 12.5     |
| 2   | SW-948    | 25       |
| 3   | RKO       | 12.5     |
| 4   | SW480     | 30       |
| 5   | CDD-18Co  | > 100    |

Table I. Antiproliferative effects of bergamottin against human colon cancer cells as examined by WST-1 assay and expressed as IC₅₀ (µM)

![Chemical structure of bergamottin](image)

**Figure 1.** A – Chemical structure of bergamottin. WST-1 cell viability showing the effects of bergamottin on the proliferation rate of HT-29 (B), RKO (C) and normal CDD-18Co (D) cells. The experiments were repeated thrice and shown as mean ± SD (*p < 0.05)
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Statistical analysis

The experiments were done three times. The values presented are means of three repeats ± SD. *P < 0.05, **P < 0.01 and ***P < 0.001 were considered statistically significant. Student’s t test using GraphPad prism 7 software was employed for statistical analysis.

Results

Bergamottin inhibits the proliferation of human colon cancer cells

The antiproliferative effects of bergamottin were evaluated by MTT assay on different human colon cancer cell lines. Bergamottin treatment resulted in a significant decrease in the viability of all the colon cancer cell lines and the IC_{50} varied from 12.5 to 30 µM (Table I). Nonetheless, more profound growth inhibitory effects were observed against the HT-29 and RKO cells as evident from the IC_{50} of 12.5 µM for bergamottin against these two cell lines (Figure 1 B). Next, the cytotoxic effects of bergamottin were also investigated on the normal CDD-18Co cells and interestingly the toxic effects were comparatively much lower on the normal cells as evident from the IC_{50} of > 100 µM (Figure 1 B). Microscopic analysis showed that bergamottin induced significant changes in the morphology of both HT-29 and RKO cells such as shrinkage, membrane blebbing and rounding (Figure 2).

Bergamottin triggers apoptosis in RKO and HT-29 cells

To reveal the reasons behind the anticancer effects of bergamottin, DAPI and annexin V/PI staining of the bergamottin treated HT-29 and RKO cells were performed. DAPI staining showed clear nuclear fragmentation indicative of apoptosis (Figure 3 A). Annexin V/PI staining revealed that the apoptotic HT-29 and RKO cell percentage increased upon bergamottin treatment. The percentage of HT-29 apoptotic cells was 2.05, 7.20, 12.62 and 28.9% and that of RKO apoptotic cells was 2.99, 6.5513, 79 and 26.01% at 0, 3.12, 6.25, 12.5 and 25 µM concentration of bergamottin (Figure 3 B). Western blotting was further used to confirm the apoptosis and the results showed that bergamottin significantly increased Bax expression, which was concomitant with depletion of Bcl-2 expression (Figure 4). Moreover, bergamottin also caused an increase in the cleavage of caspase-3, 8 and 9 in a concentration-dependent manner. These results suggest that bergamottin suppresses the growth of colon cancer cells at least partially via promotion of apoptosis.

Bergamottin induces G2/M cell cycle arrest of HT-29 and RKO cells

Cell cycle analysis of the bergamottin treated HT-29 and RKO cells was also performed to assess whether bergamottin triggers cell cycle arrest. The results showed that bergamottin caused an increase in the percentage of the G2/M phase cancer cells. The HT-29 and RKO increased from 1.21% and 0.75% in control to 87% and 61% at 12.5 µM con...
centration (Figure 5). Next, western blot analysis was performed to determine the expression of the regulatory proteins of cell cycle and it was revealed that bergamottin caused significant reduction in Cdc2, cyclin A, and cyclin B. Nonetheless, the levels of negative regulatory proteins of cell cycle progression (p21Cip1, and p27Kip1) were remarkably and dose-dependently depleted by bergamottin (Figure 5).

**Bergamottin blocks Raf/MEK/ERK signal transduction**

The effect of bergamottin was also assessed on the Raf/MEK/ERK signalling cascade to ascertain whether bergamottin induced G2/M arrest is via blocking of this pathway. It was found that bergamottin prompted concentration dependently suppression of the phosphorylation of Raf-1, MEK1/2 and ERK1/2 with no visible impact on Raf-1, MEK1/2 and ERK1/2 expression (Figure 6).

**Bergamottin inhibits invasion of HT-29 and RKO cells**

Transwell assays were performed to determine the effects of bergamottin on the invasion of the HT-29 and RKO cells. The results showed that bergamottin could decrease the invasion of the RKO and HT-29 colon cancer cells considerably (Figure 7 A). Additionally, the decrease of cell invasiveness was also associated with depletion of metalloproteinase 2 and 9 expression (Figure 7 B).
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**Figure 4.** Effect of bergamottin on expression of apoptosis-related proteins in HT-29 and RKO colon cancer cells as depicted by western blot analysis. The experiments were repeated thrice.

**Figure 5.** Flow cytometry showing the effects of bergamottin on the distribution of HT-29 and RKO cells in different cell cycle phases. The experiments were performed in triplicate.

**Bergamottin suppresses xenografted tumour growth in vivo**

To assess the in vivo anticancer effects of bergamottin, a xenografted mouse model was used. The results showed that bergamottin caused reduction in the xenografted tumour growth (Figure 8 A). The tumour weight was significantly decreased and the tumour volume showed a constant reduction over the period of 6 weeks (Figure 8 B). Additionally, no apparent toxicity was observed on the mice during the in vivo study (Figure 9).

**Discussion**

Colon cancer is a devastating cancer and its incidence is expected to escalate significantly in the next few decades [15]. The unsatisfactory clinical outcome, flawed treatment strategies and adverse effects of the existing drugs form the major hurdles in colon cancer treatment. Moreover, emergence of chemoresistance and frequent relapse make it more difficult to treat colon cancer [11]. Herein, the anticancer effects of a naturally occurring furanocoumarin, bergamottin, were evaluated against a panel of hu-
man colon cancer cell lines. Although bergamottin inhibited the growth of all the colon cancer cell lines, the IC\textsubscript{50} values were found to be lowest against the HT-29 and RKO cells. Additionally, the cytotoxicity of bergamottin was found to be very low against the normal CDD-18Co cells. These findings are consistent with previous studies wherein bergamottin was found to suppress the growth of cancer cells by blocking the STAT3 signalling pathway [16]. Similarly, another related coumarin, auraptene, has been shown to curb the growth of gastric cancer cells via induction
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of apoptosis [17]. Apoptosis is a systematic and orchestrated process in living cells that enables the elimination of harmful or defective cells from the body. Apoptosis has gained tremendous attention for its therapeutic implications in cancer [18]. Many anticancer agents halt the growth of cancer cells by promoting apoptosis [19]. In this study, we found that bergamottin induces apoptotic cell death of the HT-29 and RKO colon cancer cells. The expression of the important apoptotic biomarker proteins Bax and Bcl-2 [20] was also altered upon bergamottin treatment. Additionally, bergamottin enhances the cleavage of caspase-3, 8 and 9 as well as that of PARP, favouring apoptosis. In addition to apoptosis several plant-derived molecules have also been shown to inhibit cell division by arresting the cells at different checkpoints [21]. Herein we also observed that bergamottin caused the arrest of the HT-29 and RKO cells at the G2/M phase of the cell cycle. This was also associated with depletion of Cdc2, cyclin A, cyclin B1 and upsurge of the negative regulators of cell cycle progression (p21Cip1 and p27Kip1). Studies have shown that inhibitors of the Raf/MEK/ERK signalling pathway may provide effective anticancer agents for the management of diverse cancer types [22, 23]. However, the currently used regimen of drugs is disappointingly ineffective to block this pathway. Therefore the effects of bergamottin on this pathway were examined and it was found that bergamottin could block this pathway. Bergamottin also suppressed the migration and invasion of the colon cancer cells, which is agreement with previous studies wherein bergamottin has been reported to inhibit the invasion of glioma cells [9]. Given the promising results of the in vitro anticancer effects of bergamottin, xenograft-ed tumour mouse models were used to evaluate the anticancer effects of bergamottin in vivo. The results showed that bergamottin could inhibit the tumour growth in vivo, indicative of the anticancer potential of bergamottin.

In conclusion, the findings of the present study indicate that bergamottin exerts anticancer effects on human colon cancer cells via induction of apoptosis and cell cycle arrest. Bergamottin could also suppress invasion of colon cancer cells via blocking the Ras/Raf/ERK signalling cascade. All considered, bergamottin may prove beneficial in colon cancer treatment and warrants further research endeavours.
Conflict of interest

The authors declare no conflict of interest.

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Figure 9. A – Images of tumours from different treatment groups. Effect of bergamottin on tumour weight (B) and tumour volume at indicated concentrations and time intervals (C). The values are shown as mean ± SD (*p < 0.05)
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