Evodiamine induces apoptosis and inhibits metastasis in MDA-MB-231 human breast cancer cells in vitro and in vivo

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Abstract. Breast cancer remains the leading cause of cancer-related deaths among women. Owing to high efficiency and low toxic effects, further exploration of natural compounds from Chinese herbal medicine may be an efficient approach for breast cancer drug discovery. In this study, we investigated the effects of evodiamine on the growth and metastasis of MDA-MB-231 human breast cancer cells in vitro and in vivo. In vitro, evodiamine inhibited cell migration and invasion abilities through downregulation of MMP-9, urokinase-type plasminogen activator (uPA) and uPAR expression. Evodiamine-induced G0/G1 arrest and apoptosis were associated with a decrease in Bcl-2, cyclin D1 and cyclin-dependent kinase 6 (CDK6) expression and an increase in Bax and p27<sup>Kip1</sup> expression. Moreover, evodiamine regulated p-ERK and p-p38 MAPK expression. Evodiamine-induced apoptosis was enhanced by its combination with the extracellular signal-regulated kinase (ERK) inhibitor PD98059 or the p38 mitogen-activated protein kinase (p38 MAPK) inhibitor SB203580. Evodiamine-inhibited metastasis was partly blocked by combination with PD98059 or SB203580. In vivo, the administration of evodiamine (10 mg/kg) significantly reduced tumor growth and pulmonary metastasis. These results demonstrate that evodiamine possesses antitumor activities via inhibition of cell migration and invasion, arrest of the cell cycle and induction of cell apoptosis in MDA-MB-231 cells.

Introduction

Breast cancer is the most frequently diagnosed cancer and the leading cause of cancer-related death among females worldwide, accounting for 23% of the total new cancer cases and 14% of the total cancer deaths (1). The main causes of death of patients with breast cancer are due to the excessive proliferation and metastasis of cancer cells (2,3). To date, various anticancer agents which are natural products derived from plants, have been used clinically as first-line option drugs, such as paclitaxel, camptothecins and vinorelbine (4-6). However, there is still the need for more effective drugs with low toxic effects for breast cancer therapy. Hence, it is a challenge to discover an agent that can efficiently suppress the abnormal growth and metastatic progression of breast cancer cells.

Evodiamine, a quinolone alkaloid, is a bioactive component isolated from the fruit of Evodia rutaecarpa, which is extracted from medicinal plants such as Evodia rutaecarpa (Juss.) Benth, Evodia rutaecarpa (Juss.) Benth. var. officinalis (Dode) Huang and Evodia rutaecarpa (Juss.) Benth. var. bodinieri (Dode) Huang. Evodiamine has been shown to exhibit comprehensive pharmacological activities including antiobesity, anti-IAV, protection against myocardial ischemia-reperfusion injury and regulation of testosterone secretion (7-10); and particularly, its anticancer bioactivity is indispensable.

Evodiamine arrests cancer cell cycle distributions and induces cell apoptosis to play a role in antitumor growth. Previous studies have indicated that evodiamine-induced S phase arrest in LoVo cells is associated with a marked decrease in the protein expression of cyclin A, cyclin A-dependent kinase 2 and cdc25c (11). Evodiamine caused blockage of cells in the G2/M phase, accompanied by an increase in the protein expression of cyclin B1 and the phosphorylated form of p34(cdc2) (Thr 161) in prostate cancer LNCaP cells (12). Evodiamine inhibited the growth of thyroid cancer ARO cells, arrested the cells at the M phase, and induced apoptosis through inducing the activation of caspase-3, -8 and -9, and the cleavage of polyADP-ribose polymerase (13). Moreover, evodiamine was found to increase the expression of Bax and p53, decreased the expression of Bcl-2, lowered the mitochondrial transmembrane potential and induced the activation of caspase-3 in colorectal carcinoma COLO-205 cells (14).

Previous studies have revealed that evodiamine and paclitaxel may be regarded as leading compounds for use as antimetastatic agents acting through the inhibition of colon cancer cell migration without cytotoxicity (15). Evodiamine concentration-dependently inhibited the invasion of Lewis lung carcinoma (LLC) and B16-F10 melanoma in addition...
Materials and methods

Drugs and reagents. The compound evodiamine was purchased from Shanghai Research and Development Centre for Standardization of Traditional Chinese Medicine (Shanghai, China), and dissolved in dimethyl sulfoxide (DMSO). Dulbecco’s modified Eagle’s medium (DMEM), trypsin, penicillin and streptomycin were purchased from Gibco-BRL (Grand Island, NY, USA). 3-(4, 5-Dimethylthiazol-2-yl)-2, 5-diphenyltetrazolium bromide (MTT), DMSO, propidium iodide (PI), acridine orange (AO) and Matrigel were purchased from Sigma-Aldrich (St. Louis, MO, USA). Antibodies against Bcl-2, cyclin D1, p27Kip1, CDK6, p38 MAPK, p-p38 MAPK, p-ERK, p-SAPK/JNK, SAPK/JNK and GAPDH were purchased from Cell Signaling Technology Inc. (Danvers, MA, USA). Bax and ERK were purchased from Abgent (San Diego, CA, USA). Urokinase-type plasminogen activator (uPA) and uPAR were purchased from Santa Cruz Biotechnology Inc. (Santa Cruz, CA, USA). MMP-2 and -9 were purchased from Abcam (Cambridge, UK). IRDye™ fluorescence antibodies were obtained from LI-COR Biosciences (Lincoln, NE, USA). SB203580 and PD98059 were obtained from Biomol (Philadelphia, PA, USA).

Cell lines and culture conditions. The human breast cancer cell line MDA-MB-231 was purchased from the Cell Bank of the Type Culture Collection of the Chinese Academy of Sciences (Shanghai, China) and cultured in DMEM supplemented with 10% fetal bovine serum (FBS), 100 U/ml penicillin and 100 µg/ml streptomycin at 37°C in a humidified atmosphere of 5% CO2.

MTT assay. Cell viability was measured by the MTT assay. MDA-MB-231 cells (5x10⁴/ml) were plated in 96-well plates overnight, and then treated with various concentrations of evodiamine. After 24, 48, 72 h of drug exposure, 20 µl of MTT (5 mg/ml) was added to incubate for 4 h. Then 150 µl DMSO was added to dissolve the formazan crystals. The optical density (OD) was detected at 490 nm by an ELISA plate reader (BioTek Instruments, Inc., Winooski, VT, USA). The percentage of cell survival was calculated as follows: Cell survival rate (%) = (OD sample/OD control) x 100%.

Wound healing assay. MDA-MB-231 cells (2x10⁴/ml) were seeded in a 6-well culture plate to form a confluent monolayer. The monolayer of cells was then scrape-wounded with a sterile micropipette tip to create a wound of constant (~1 mm) width. After removal of the cellular debris with phosphate-buffered saline (PBS) twice, cells were treated with medium containing the indicated concentration of evodiamine at 37°C. After incubation for 24 h, MDA-MB-231 cells that had migrated to the wounded region were observed (x100 magnification).

Migration and invasion assays. Migration and invasion assays were performed in 8-µm-diameter pore size Transwell chambers in 24-well plates (Corning Incorporated, Corning, NY, USA). For the cell invasion assay, the internal surface of each polycarbonate membrane was coated with Matrigel (30 µg) for 30 min at 37°C for gel formation and then blocked with serum-free DMEM containing 0.2% bovine serum albumin (BSA). In brief, MDA-MB-231 cells (5x10⁴/ml) were seeded onto the upper chamber in 200 µl of serum-free medium containing evodiamine at concentrations of 0, 15, 30, 60 µM, respectively; the lower compartment of the chamber was filled with 600 µl DMEM supplemented with 0.2% BSA and 10% FBS. After incubation for 24 h at 37°C, the cells in the upper surface of the membrane were carefully removed with a cotton swab, and cells that had migrated or invaded to the lower surface of the membrane were fixed with methanol and stained with 0.1% crystal violet for 30 min. The migrated or invaded cells were then visualized and counted from 6 randomly selected fields (x200 magnification).

Cell cycle analysis. Cell cycle phase analysis was performed by PI staining. MDA-MB-231 cells were treated with different concentrations of evodiamine for 24 h. The cells were harvested, washed with PBS, and fixed in ice-cold 70% ethanol at 4°C overnight. The cells were washed with ice-cold PBS and stained with PI (50 µg/ml) in the dark for 30 min. Cell cycle distribution was analyzed using Mycle software (Beckman Coulter, Fullerton, CA, USA).

Annexin V-FITC binding assay. The quantification of apoptotic cells was performed by flow cytometric analysis using an Annexin V-FITC/PI Apoptosis Kit (Beckman Coulter, Miami, FL, USA). MDA-MB-231 cells were treated with different concentrations of evodiamine for 48 h. Annexin V and PI staining was performed according to the manufacturer’s instructions. After staining, the quantification of apoptotic cells was measured with a flow cytometer (Beckman Coulter). The percentage of early apoptosis was detected by Annexin V-positivity and PI-negativity, while the percentage of advanced apoptosis was detected by Annexin V-positivity and PI-positivity.

Acridine orange (AO) fluorescence staining. Apoptotic morphological analysis was performed using AO fluorescence staining. MDA-MB-231 cells were seeded on coverslips in 6-well plates and treated with evodiamine for 48 h. The coverslips were fixed with 95% ethanol for 15 min, acidified with 1% acetic acid for 30 sec, dyed with AO (0.1 mg/ml) for 10 min, differentiated with CaCl2 (0.1 mol/l) for 2 min, and observed under a fluorescence microscope (x400 magnification).
Terminal deoxynucleotidyl transferase-mediated dUTP nick end labeling (TUNEL) assay. Apoptotic cells in sections of tumor tissues were detected using a KeyGene TUNEL apoptosis detection kit (Kai-ji, Nanjing, China) according to the manufacturer's protocol. Briefly, tumor histological sections were permeabilized with 20 µg/ml protease K. Apoptosis was detected by labeling the 3'-OH ends of the fragmented DNA with Biotin-11-dUTP under TdT enzyme at 37°C for 60 min. The tumor slides were then incubated with streptavidin horseradish peroxidase (streptavidin-HRP) conjugate at 37°C for 30 min, followed by incubation with DAB and after-stain with hematoxylin. The TUNEL-positive cells (dark brown nuclei) were counted in 6 randomly selected fields under a microscope (x200 magnification). The apoptotic index (AI) was calculated as follows: AI (%) = number of apoptotic cells/total number of cells x 100%.

In-cell western analysis. In-cell western assay (19) was used to determine the level of protein expression in cells. MDA-MB-231 cells (1.5x10^5/ml) were plated in 96-well plates. The next day, the cells were treated with 60 µM evodiamine for 24 or 48 h. The cells were fixed with 4% formaldehyde for 20 min and washed with 0.1% Triton, and finally blocked with 10% non-fat milk overnight at 4°C. The cells were then incubated with the primary antibodies: Bcl-2, Bax, cyclin D1, p27Kip1, CDK6, MMP-2, MMP-9, uPA, uPAR, p38 MAPK, p-p38 MAPK, ERK, p-ERK, SAPK/JNK and p-SAPK/JNK overnight at 4°C, respectively. The housekeeping protein, GAPDH, was added to each well at the same time as the control. The wells were then incubated with the corresponding secondary IRDye™ 680 DX (red fluorescence) or IRDye™ 800 DX (green fluorescence) fluorescence antibody recommended by the manufacturer (Rockland, Gilbertsville, PA, USA) in the dark. Images were obtained using the Odyssey Infrared Imaging System (LI-COR Biosciences). The levels of protein expression were calculated as the ratio of the intensity of the target protein to that of GAPDH. The experiments were carried out in triplicate.

Animal tumor xenograft model and treatment with drugs. Six-week-old female BALB/c athymic mice were obtained from the Laboratory Animal Center at Shanghai University of Traditional Chinese Medicine. The mice were housed in barrier facilities with rodent chow and water under a specified-pathogen-free (SPF) condition throughout the experimental duration. MDA-MB-231 cancer cells (3x10^5) suspended in 100 µl of serum-free DMEM were injected into the mammary fat pad of the mice. One day after tumor cell transplantation, 10 mg/kg evodiamine (17) was intraperitoneally (i.p.) injected once every 2 days. The body weight of each mouse was measured each week after treatment. All mice were sacrificed after drug administration for 2 months, and the tumors were removed and weighed. The lung and tumor tissues were fixed with formalin. Thin sections were stained with hematoxylin and eosin (H&E) and metastatic nodules present in the lungs were counted. Representative fields (x100 magnification) for each group were photographed. All procedures conformed to the consideration of animal welfare and were approved by the Ethics Committee of Shanghai University of Traditional Chinese Medicine.

Statistical analysis. Experimental results are presented as the means ± standard deviation (SD). Differences were evaluated by the Student's t-test or one-way analysis of variance (ANOVA). P<0.05 was considered to indicate a statistically significant result.

Results

Anti-proliferative effects of evodiamine in MDA-MB-231 cells. The growth of MDA-MB-231 cells was inhibited by evodiamine in a dose- and time-dependent manner. Treatment at doses of 15, 30 and 60 µM did not cause cytotoxicity in MDA-MB-231 cells at 24 h (P>0.05); however, the survival rates of MDA-MB-231 cells were significantly decreased to 75.06 and 61.76% following treatment with 90 and 120 µM, respectively (P<0.01). Following treatment at doses of 15, 30, 60 and 120 µM for 48 h, the survival rates were decreased to 90.50, 84.40, 71.70, 51.44 and 42.47%, respectively (P<0.01). The half maximal inhibitory concentration (IC_{50}) at 48 h was ~90 µM. Following treatment at doses of 15, 30, 60, 90 and 120 µM for 72 h, the survival rates were decreased to 85.44, 77.45, 63.69, 47.49 and 38.77%, respectively (P<0.01) (Fig. 1).

Evodiamine inhibits the migration and invasion ability of MDA-MB-231 cells. In the wound healing assay, the gaps in the cells treated with 15, 30 and 60 µM evodiamine were wider than that in the untreated group (Fig. 2A). In the Transwell migration assay, the cells that had migrated to the lower chamber were reduced by evodiamine in a concentration-dependent manner. The number of migrated cells following treatment with 15, 30 and 60 µM evodiamine were reduced to ~90.41, 77.31, 50.14 and 33.89%, respectively (P<0.01) (Fig. 2B and D). In the Transwell invasion assay, the number of cells invaded through the Matrigel-coated filter was reduced by evodiamine dose-dependently. Compared with the untreated cells, the number of invaded cells following treatment 15, 30 and 60 µM evodiamine were reduced to ~90.41, 77.31, 50.14 and 33.89%, respectively (P<0.05 or P<0.01) (Fig. 2C and D).

Evodiamine induces G0/G1 cell cycle arrest of MDA-MB-231 cells. Exposure to different concentrations of evodiamine (0, 15, 30, 60, 90 and 120 µM) caused G0/G1 arrest of MDA-MB-231 cells. The growth of MDA-MB-231 cells was inhibited by evodiamine in a dose- and time-dependent manner. The half maximal inhibitory concentration (IC_{50}) at 48 h was ~90 µM. Following treatment at doses of 15, 30, 60, 90 and 120 µM for 72 h, the survival rates were decreased to 85.44, 77.45, 63.69, 47.49 and 38.77%, respectively (P<0.01) (Fig. 1).
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Evodiamine induces apoptosis in MDA-MB-231 cells. To confirm whether evodiamine inhibits MDA-MB-231 cell viability due to apoptosis, morphological observation and Annexin V-FITC binding assay of MDA-MB-231 cells were conducted. Chromatin condensation, nuclear fragmentation and apoptotic bodies were found in the evodiamine-treated cells (Fig. 4A). This result indicated that MDA-MB-231 cells underwent the typical morphological changes of apoptosis when exposed to 60 µM evodiamine for 48 h. To quantify apoptosis, the Annexin V-FITC apoptosis assay was further carried out by FACS. Treatment of MDA-MB-231 cells with different concentrations of evodiamine resulted in a dose-dependent increase in apoptotic cells (Fig. 4B). The detailed percentages of early apoptosis, late apoptosis and total apoptosis in MDA-MB-231 cells were calculated (Table II).

Evodiamine regulates the expression of proteins related to apoptosis, cell cycle, cell migration and invasion in the MDA-MB-231 cells. To investigate the underlying mechanism of the induction of apoptosis and cell cycle arrest and inhibition of cell migration and invasion by evodiamine, expression of proteins related to apoptosis, cell cycle and cell migration and invasion was determined by in-cell western analysis. Compared to untreated cells, evodiamine significantly decreased the level of Bcl-2 (P<0.01) and increased the level of Bax (P<0.05), resulting in upregulation of the Bax/Bcl-2 ratio. Evodiamine treatment decreased the levels of cyclin D1 (P<0.05) and CDK6 (P<0.01) and increased the level of p27^{kip1} (P<0.01); as well as decreased the levels of MMP-9, uPA and uPAR (P<0.05), whereas evodiamine treatment did not affect MMP-2 expression (P>0.05) (Fig. 5).
Evodiamine regulates the levels of p-ERK and p-p38 MAPK in MDA-MB-231 cells. The expression levels of p-ERK, p-p38 MAPK, p-JNK and each non-phosphorylated form were detected by in-cell western analysis. The levels of p-ERK, p-p38 MAPK and p-JNK expression were normalized to non-phosphorylated ERK, p38 MAPK and JNK, respectively. As shown in Fig. 6A, compared to the untreated cells, the levels of p-ERK and p-p38 MAPK were significantly decreased following exposure to evodiamine for 24 h (P<0.05). Moreover, compared to the untreated cells, the level of p-ERK was significantly decreased (P<0.05), while the level of p-p38 MAPK was increased following exposure to evodiamine for 48 h (P<0.05). The level of p-JNK remained constant following evodiamine treatment for 24 or 48 h (P>0.05). These results indicate that evodiamine regulates the activities of the ERK and p38 MAPK pathways.

The role of ERK and p38 MAPK in the induction of apoptosis and inhibition of migration. To address the role of ERK and p38 MAPK in evodiamine-induced apoptosis, MDA-MB-231 cells were treated with 20 µM PD98059 (ERK inhibitor), or 30 µM SB203580 (p38 MAPK inhibitor) alone and co-treated with 60 µM evodiamine for 48 h. Co-treatment of 20 µM PD98059 or 30 µM SB203580 with 60 µM evodiamine reduced the cell viability to 37.52% and 38.50%, respectively. Cell survival rate was decreased ~10%, compared to that following treatment with 60 µM evodiamine alone (P<0.05). FACS revealed that the percentage of apoptotic cells was significantly higher

Table I. Cell cycle distribution in MDA-MB-231 cells exposed to different concentrations of evodiamine for 24 h.

<table>
<thead>
<tr>
<th>Evodiamine (µM)</th>
<th>Sub-G1 (%)</th>
<th>G0/G1 (%)</th>
<th>S (%)</th>
<th>G2/M (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>0.13±0.03</td>
<td>53.29±2.73</td>
<td>15.50±0.96</td>
<td>27.01±0.56</td>
</tr>
<tr>
<td>15</td>
<td>7.79±1.88b</td>
<td>74.92±2.35b</td>
<td>6.48±1.30</td>
<td>8.04±2.75</td>
</tr>
<tr>
<td>30</td>
<td>10.69±3.89b</td>
<td>74.20±2.02b</td>
<td>5.27±2.29</td>
<td>6.13±3.46</td>
</tr>
<tr>
<td>60</td>
<td>15.93±1.89b</td>
<td>72.94±0.29b</td>
<td>3.38±0.68</td>
<td>3.85±1.46</td>
</tr>
<tr>
<td>90</td>
<td>24.28±3.46b</td>
<td>65.30±2.69b</td>
<td>3.99±1.47</td>
<td>3.90±2.82</td>
</tr>
<tr>
<td>120</td>
<td>32.19±1.84b</td>
<td>60.12±1.26b</td>
<td>3.71±0.13</td>
<td>2.46±1.17</td>
</tr>
</tbody>
</table>

Values are represented as the means ± SD. *P<0.05, **P<0.01, vs. untreated cells.

Table II. Percentage of apoptosis in the MDA-MB-231 cells exposed to different concentrations of evodiamine for 48 h.

<table>
<thead>
<tr>
<th>Evodiamine (µM)</th>
<th>Early apoptosis (%)</th>
<th>Late apoptosis (%)</th>
<th>Total apoptosis (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Annexin V+, PI −</td>
<td>Annexin V+, PI +</td>
<td></td>
</tr>
<tr>
<td>0</td>
<td>1.24±0.52</td>
<td>0.00±0.01</td>
<td>1.24±0.52</td>
</tr>
<tr>
<td>15</td>
<td>11.89±0.73</td>
<td>0.04±0.01</td>
<td>11.93±0.74</td>
</tr>
<tr>
<td>30</td>
<td>18.98±6.06</td>
<td>0.02±0.01</td>
<td>19.00±6.05</td>
</tr>
<tr>
<td>60</td>
<td>31.44±1.78</td>
<td>0.05±0.06</td>
<td>31.49±1.83</td>
</tr>
<tr>
<td>90</td>
<td>50.55±1.98</td>
<td>0.08±0.06</td>
<td>50.63±1.96</td>
</tr>
<tr>
<td>120</td>
<td>57.98±2.60</td>
<td>0.17±0.07</td>
<td>58.15±2.55</td>
</tr>
</tbody>
</table>

Values are represented as the means ± SD. *P<0.01 vs. untreated cells.

Figure 3. Effect of evodiamine on the cell cycle distribution of MDA-MB-231 cells. Cells were cultured in 0, 15, 30, 60, 90 and 120 µM of evodiamine for 24 h. Representative flow cytometry profiles are shown. Distribution of the cell cycle in MDA-MB-231 cells at different phases. The peak located left of the dashed line represents the sub-G1 apoptotic peak.
Figure 4. Evodiamine induces apoptosis in MDA-MB-231 cells. (A) MDA-MB-231 cells were treated with 0 and 60 µM evodiamine for 48 h, respectively, stained with acridine orange (AO), and photographed using fluorescence microscope. (B) MDA-MB-231 cells were treated with 0, 15, 30, 60, 90 and 120 µM evodiamine for 48 h and analyzed by FACS. Representative flow cytometry profiles are shown. The bottom-right quadrant indicates early apoptosis, the top-right quadrant indicates late apoptosis.

Figure 5. Effect of evodiamine on the expression of Bcl-2, Bax, p27Kip1, cyclin-dependent kinase 6 (CDK6), cyclin D1, MMP-2, MMP-9, urokinase-type plasminogen activator (uPA) and uPAR in MDA-MB-231 cells. The expression of p27Kip1, CDK6, cyclin D1, MMP-2, MMP-9, uPA and uPAR was determined after treatment with 60 µM evodiamine for 24 h. The expression of Bcl-2 and Bax was determined after treatment with 60 µM evodiamine for 48 h. (A) The fluorescence intensity ratio of proteins to GAPDH is shown as relative expression. Values are represented as the means ± SD. *P<0.05, **P<0.01, vs. untreated cells. (B) A representative image is shown. Red fluorescence indicates the expression of GAPDH in 680 channel and green fluorescence indicates the expression of MMP-9 in 800 channel. Experiments were repeated with similar results.
following treatment with the two inhibitors in combination with 60 µM evodiamine, compared to the cultures exposed to 60 µM evodiamine alone (P<0.05) (Fig. 6B).

Furthermore, to address the role of ERK and p38 MAPK in evodiamine-inhibited migration, MDA-MB-231 cells were treated with 20 µM PD98059 or 10 µM SB203580 alone or co-treated with 60 µM evodiamine for 24 h. As shown in Fig. 6C, co-treatment of 20 µM PD98059 or 10 µM SB203580 with 60 µM evodiamine increased the cell migration rate ~20%, compared to that following treatment with 60 µM evodiamine alone (P<0.05) (Fig. 6C). These results suggest that the ERK and p38 MAPK pathways may play an important role in evodiamine-induced apoptosis and inhibition of migration in MDA-MB-231 cells.

Evodiamine administration retards tumor growth, induces apoptosis and inhibits the pulmonary metastasis of MDA-MB-231 cells in nude mice. The average body weights of the control (saline-treated) and evodiamine-treated mice did not differ significantly throughout the study (Fig. 7A). Movement, digestion and swelling seemed normal in the evodiamine-treated mice. Moreover, the tumor weight in the evodiamine-treated group was lower than that of the control group (P<0.01) (Fig. 7B). H&E staining revealed a relatively higher nuclear to cytoplasmic ratio in the tumors from the control group when compared with the ratio in the evodiamine-treated group. TUNEL staining demonstrated that the number of apoptotic-positive cells in the evodiamine-treated group was significantly higher than that in the control group (Fig. 7C). The AI was subsequently calculated. The average AI was 13% in the evodiamine-treated group (Fig. 7D). Furthermore, the lungs in the evodiamine-treated mice contained less metastasized MDA-MB-231 cells morphologically, when compared to the control group (Fig. 7C). The average number of tumor nodules in the control group was 22, while that in the evodiamine-treated group was 13, indicating evodiamine treatment significantly decreased lung metastasis (P<0.01) (Fig. 7E). Collectively, these results suggest that evodiamine administration significantly retards tumor growth, induces apoptosis and inhibits pulmonary metastasis, without an abnormal response in the MDA-MB-231 xenograft model.

Discussion

Evodiamine was found to possess antitumor potential in different cancer cell lines, however, there is no available information on the effects of evodiamine on highly metastasis human breast cancer MDA-MB-231 cells. Moreover, the mechanism involved in the inhibition of growth and metastasis of MDA-MB-231 cells by evodiamine requires further investigation. In the present study, for the first time, we investigated the effects of evodiamine on the growth, migration and invasion of MDA-MB-231 cells and primarily focused on its molecular mechanism.

We found that evodiamine inhibited cell motility, migration and invasion of MDA-MB-231 cells (Fig. 2) and exerts an antimetastasis effect by inhibiting pulmonary metastatic nodules in xenograft BALB/c nude mice (Fig. 7). During the progression of cancer cell metastasis, matrix metalloproteinases (MMPs) play a crucial role in tumor invasion and metastasis formation due to their ability to degrade extracellular matrix proteins (20). In our study, the level of MMP-9 was downregulated by evodiamine (Fig. 5), and this result
was consistent with the evidence of evodiamine blocking the expression of TNF-induced NF-κB-regulated gene products including MMP-9 in KBM-5 cells (18). uPA and its receptor uPAR are usually overexpressed in breast cancer cells. uPA binding to uPAR directly contributes to the degradation of the extracellular matrix, and also mediates activation of MMPs, thereby promoting cancer cell invasion and migration (21,22). The levels of uPA and uPAR were downregulated by evodiamine (Fig. 5). Therefore, inhibition of the levels of MMP-9, uPA and uPAR may be, at least in part, responsible for the antimetastatic potential of evodiamine against MDA-MB-231 cells.

Treatment of MDA-MB-231 cells with evodiamine resulted in significant G0/G1 phase arrest (Fig. 3) and apoptosis induction (Figs. 4 and 7). Control of cell cycle progression in cancer cells is regarded to be an effective strategy for inhibition of tumor cell proliferation. Cyclin D1 and cyclin-dependent kinase 6 (CDK6) are overexpressed in breast cancer and play a role in G1 progression and oncogenesis. One of the CDK inhibitors, p27Kip1, inhibits a wide variety of cyclin-CDK complexes in vitro, and its overexpression blocks the progression of cells through the G1 phase (23,24). In our study, evodiamine treatment caused G0/G1 phase arrest accompanied by downregulation of the expression of cyclin D1 and CDK6 and upregulation of the expression of p27Kip1 (Fig. 5). Previous studies have reported that evodiamine blocks G0/G1 phase in L929 cells (25), induces S phase arrest in LoVo cells (11) and causes blockage of the G2/M phase in LNCaP cells (12). These studies showed that evodiamine arrests the cancer cell cycle at different phases. This may provide explanations...
concerning the types of cell lines and their specific response to evodiamine.

In our study, we found that evodiamine plays an anti-proliferative role via inducing MDA-MB-231 cell apoptosis. Apoptosis is tightly regulated by anti-apoptotic and pro-apoptotic molecules, including proteins of the Bcl-2 family. Among them, Bcl-2 and its dominant inhibitor Bax are key regulators of several survival and apoptosis pathways. Overexpression of Bcl-2 enhances cell survival by suppressing apoptosis, but overexpression of Bax accelerates cell death, and the ratio of Bax/Bcl-2 plays a critical role in determining whether cells will undergo apoptosis (26). Here, evodiamine increased pro-apoptotic Bax expression and decreased anti-apoptotic Bcl-2 expression, leading to upregulation of the ratio of Bax/Bcl-2 (Fig. 5). Thus, this finding confirmed that MDA-MB-231 cells underwent apoptosis induced by evodiamine through the Bax/Bcl-2 pathway.

Recent studies have demonstrated that mitogen-activated protein kinases (MAPKs), including Jun N-terminus kinase (JNK), p38 mitogen-activated protein kinase (p38 MAPK) and extracellular signal-regulated kinase (ERK), play crucial roles in cell survival, apoptosis and migration in tumor development and progression (27-29). The aberrant activation of ERK and p38 MAPK is associated with the promotion of migration. Activated ERK1/2 was found to result in high uPA expression, rapid cell proliferation and the migration of breast cancer cells (30,31). Daintain/AIF-1 activated p38 MAPK signaling pathway contributed to the upregulation of TNF-α and led to enhanced migration of MDA-MB-231 and MCF-7 cells (32). In our study, the expression levels of p-ERK and p-p38 MAPK were both downregulated following exposure to evodiamine for 24 h in MDA-MB-231 cells (Fig. 6A). These findings may account for the inhibition of MDA-MB-231 cell migration and invasion by evodiamine, and such anti-metastatic activities of evodiamine may partly involve the suppression of ERK and p38 MAPK signaling pathways.

In general, the activation of JNK/SAPK and p38 MAPK is associated with promotion of apoptosis, while ERK activity inhibits apoptosis (33). In the present study, the expression of p-ERK was downregulated while the expression of p-p38 MAPK was upregulated following exposure to evodiamine for 48 h in MDA-MB-231 cells (Fig. 6B). This result was consistent with the inhibitory effect of evodiamine on proliferation of A375-S2 cells. Evodiamine blocked the protective role of ERK in A375-S2 cells through the downregulation of p-ERK expression (34). Thus, evodiamine induces cell apoptosis through ERK inactivation and p38 MAPK activation.

According to the above findings, indicating that evodiamine regulates p-ERK and p-p38 MAPK, it was hypothesized that ERK and p38 MAPK may have some effects on evodiamine-mediated cell metastasis and apoptosis. In the present study, the anti-metastatic effect of evodiamine was partly blocked by PD98059 and SB203580 (Fig. 6C). Therefore, we may conclude that evodiamine-inhibited migration of MDA-MB-231 cells may partly occur through suppression of the ERK or p38 MAPK pathway. When ERK or p38 MAPK was blocked, the anti-migration potential of evodiamine was reduced.

Zelivianski et al (35) found that the ERK inhibitor PD98059 enhanced docetaxel-induced apoptosis of androgen-independent human prostate cancer cells. In the present study, PD98059 or SB203580 enhanced the potential of evodiamine-induced growth suppression in breast cancer MDA-MB-231 cells (Fig. 6B). The mechanism of PD98059 involved in evodiamine-mediated cell apoptosis was previously associated with decreased expression of c-Myc and cyclin D1 which are key targets of the MAPK pathway and activate the caspase cascade (36). Although activation of p38, to some extent, accounted for the growth inhibition by evodiamine, p38 inhibition also synergized evodiamine-induced cell apoptosis. p38 had dual effects during the regulation of cell growth. It phosphorylated p53, was involved in Fas/Fasl-mediated apoptosis pathway and induced Bax translocation to induce apoptosis, otherwise, it initiated the cell survival pathway, and inhibited apoptosis signal generation (37,38). In addition, many of the signals from other pathways may converge with that of the MAPK cascade and downstream factors may be involved in this response. For these reasons, the detailed mechanisms involved in the synergistic induction of apoptosis generating by co-treatment of PD98059 and SB203580 with evodiamine require further investigation.

In conclusion, the results of the present study indicate that evodiamine not only inhibits proliferation, induces G0/G1 phase arrest and apoptosis, but also inhibits migration and invasion of human breast cancer MDA-MB-231 cells. Moreover, evodiamine regulated ERK and p38 MAPK activities, and its pro-apoptotic and anti-metastatic effects were enhanced or weakened by an ERK inhibitor and a p38 MAPK inhibitor, respectively. Although further investigations are needed to verify the anticancer mechanism of evodiamine, this study emphasizes the promising application of evodiamine in breast cancer treatment.

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References


