**Hedyotis Diffusa Willd** extract induces apoptosis via activation of the mitochondrion-dependent pathway in human colon carcinoma cells

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Received June 9, 2010; Accepted August 2, 2010

DOI: 10.3892/ijo_00000785

Abstract. Hedyotis Diffusa Willd has been used as a major component in several Chinese medicine formulations for the clinical treatment of colorectal cancer. However, the molecular mechanism of the anti-cancer activity of Hedyotis Diffusa Willd remains unclear. In the present study, we investigated the cellular effects of the ethanol extract of Hedyotis Diffusa Willd (EEHDW) in the HT-29 human colon carcinoma cell line. We found that EEHDW inhibited the growth of HT-29 cells demonstrating EEHDW-induced cell morphological changes and reduced cell viability in a dose- and time-dependent manner. Furthermore, we observed that EEHDW treatment resulted in DNA fragmentation, loss of plasma membrane asymmetry, collapse of mitochondrial membrane potential, activation of caspase-9 and caspase-3, and increase of the ratio of pro-apoptotic Bax to anti-apoptotic Bcl-2, suggesting that the HT-29 cell growth inhibitory activity of EEHDW was due to mitochondrion-mediated apoptosis, which may partly explain the anti-cancer activity of Hedyotis Diffusa Willd.

Introduction

Colorectal cancer (CRC) is one of the leading causes of death in Western countries, and the incidence of this disease is also increasing in Asian countries (1). To date, chemotherapy is the main therapeutic approach for patients with advanced CRC; and 5-fluorouracil (5-FU)-based regimens are the standard treatment for these patients. However, due to drug resistance, systemic chemotherapy using 5-FU-based regimens produces objective response rates of <40% (2-4). Moreover, many currently used chemotherapeutic agents for cancer therapy have potent cytotoxic effects in normal cells and may induce DNA mutations that probably lead to secondary cancers (5). Both drug resistance and toxicity against normal cells limit the effectiveness of current CRC chemotherapy, thus increasing the necessity for the development of new therapeutic approaches (6,7). Natural products that have relatively fewer side-effects have been used clinically in China for thousands of years as important alternative remedies for a variety of diseases. Recently many researchers have investigated these and other natural products to discover novel anti-cancer agents (8-11). Hedyotis Diffusa Willd, belonging to the Rubiaceae family, is a medicinal herb widely distributed in Northeast Asia. As a well-known traditional Chinese folk-medicine, it is used for heat-clearing, detoxification, promotion of blood circulation and removal of blood stasis (12). Hedyotis Diffusa Willd has long been used as an important component in formulated prescriptions of Chinese traditional medicine to treat various types of cancer, including stomach and colon cancers (12-14). However, the precise mechanism of the potential tumorcidal activity of Hedyotis Diffusa Willd remains to be elucidated.

Cell death by apoptosis eliminates excess, redundant, abnormal cells in animals and hence is crucial for animal development and tissue homeostasis. Disturbed regulation of this vital process represents a major causative factor in the pathogenesis of cancers including CRC (15-17). Bcl-2 family proteins are key regulators of apoptosis (15,16), functioning as either suppressors such as Bcl-2, or promoters such as Bax.
One possible mechanism by which Bcl-2 family proteins regulate apoptosis is through their influence on the permeability of mitochondrial outer membrane (MOM) following homo- or hetero-association (18). It has been demonstrated that after activation, the pro-apoptotic Bax or Bak is sufficient to induce mitochondrial outer membrane permeabilization (MOMP) (19-22), releasing apoptogenic proteins such as cytochrome c and Diablo/Smac that trigger apoptosis by activating caspases and nucleases (22-26). The anti-apoptotic Bcl-2 protein may bind to active Bax to prevent it from damaging the MOM (19,26-28). The ratio of active anti- and pro-apoptotic Bcl-2 family members determines the fate of cells, and alteration of the ratio by aberrant expression of these proteins impairs the normal apoptotic program contributing to various apoptosis-related diseases (29,30). For instance, overexpression of Bcl-2 is commonly found in various cancers (31), which not only confers a survival advantage to the cancer cells but also causes resistance to conventional chemo- and radio-therapies. Therefore, promoting cell apoptosis via regulating the Bcl-2 family proteins has been the main focus in the development of anti-cancer therapies.

In order to extend the clinical observations of the potential anti-cancer effect of *Hedyotis Diffusa* Willd and help to establish a scientific foundation for further research, in this study, we evaluated the effect of the ethanol extract of *Hedyotis Diffusa* Willd (EEHDW) on the growth and apoptosis of HT-29 human colon carcinoma cells, and investigated the possible molecular mechanisms mediating its biological effect. We found that EEHDW inhibited the growth and induced apoptosis of HT-29 cells. EEHDW-induced apoptosis was accompanied by loss of mitochondrial membrane potential (∆Ψm), caspase-9 and caspase-3 activation and up-regulation of Bax to Bcl-2 ratio. Our finding suggests that promotion of cancer cell apoptosis via activation of the mitochondrion-dependent pathway probably is one of the mechanisms by which *Hedyotis Diffusa* Willd can be effective in the treatment of cancer.

**Materials and methods**

**Materials and reagents.** Dulbecco’s modified Eagle’s medium (DMEM), fetal bovine serum (FBS), penicillin-streptomycin, trypsin-EDTA, TRizol reagent and iBlot Western detection stack/iBlot dry blotting system, 5,5′,6,6′-tetraethylbenzimidol-carbocyanine iodide (JC-1), DNA ladder detection kit, caspase-3 and -9 colorimetric protease assay kits were purchased from Invitrogen (Grand Island, NY, USA). SuperScript II reverse transcriptase was provided by Promega (Madison, WI, USA). Bcl-2 and Bax antibodies, horseradish peroxidase (HRP)-conjugated secondary antibodies were obtained from Cell Signaling (Beverly, MA, USA). A fluorescein isothiocyanate (FITC)-conjugated Annexin V apoptosis detection kit was provided by Becton-Dickinson (San Jose, CA, USA). All the other chemicals used, unless otherwise stated, were obtained from Sigma Chemicals (St. Louis, MO, USA).

**Preparation of ethanol extract from Hedyotis Diffusa Willd.** *Hedyotis Diffusa* Willd (500 g) were extracted with 5000 ml of 85% ethanol using refluxing method and filtered. The ethanol solvent was then evaporated on a rotary evaporator (Shanghai Yarong, Model RE-2000, China). The resultant solution was concentrated to a relative density of 1.05, and the dried powder of ethanol extract of *Hedyotis Diffusa* Willd (named EEHDW) was obtained by spraying desiccation method using a spray dryer (Buchi, Model B-290, Switzerland). Stock solutions of EEHDW were prepared by dissolving the EEHDW powder in 40% DMSO to a concentration of 400 mg/ml, and stored at -20°C. The working concentrations of EEHDW were made by diluting the stock solution in the culture medium. The final concentration of DMSO in the medium was <0.5%.

**Cell culture.** Human colon carcinoma HT-29 cells were obtained from American Type Culture Collection (ATCC, Manassas, VA, USA). The cells were grown in DMEM containing 10% (v/v) FBS and 100 U/ml penicillin and 100 μg/ml streptomycin in a 37°C humidified incubator with 5% CO₂. The cells were subcultured at 80-90% confluency. Cells used in this study were subjected to no more than 20 cell passages.

**Evaluation of cell viability by MTT assay.** Cell viability was assessed by the 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl-tetrazolium bromide (MTT) colorimetric assay. HT-29 cells were seeded into 96-well plates at a density of 1.0×10⁴ cells/well in 0.1 ml medium. The cells were treated with various concentrations of EEHDW for 24 h or with 4 mg/ml of EEHDW for different periods of time. Treatment with 0.5% DMSO was included as vehicle control. At the end of the treatment, 10 μl MTT (5 mg/ml in phosphate buffered saline, PBS) were added to each well, and the samples were incubated for an additional 4 h at 37°C. The purple-blue MTT formazan precipitate was dissolved in 100 μl DMSO. The absorbance was measured at 570 nm using an ELISA reader (BioTek, Model EXL800, USA).

**Observation of morphologic changes.** HT-29 cells were seeded into 6-well plates at a density of 2.0×10⁵ cells/well in 2 ml medium. The cells were treated with various concentrations of EEHDW for 24 h. Cell morphology was observed using a phase-contrast microscope (Olympus, Japan). The photographs were taken at a magnification of x200.

**Detection of apoptosis by flow cytometry analysis with Annexin V/PI staining.** After incubated with various concentrations of EEHDW, apoptosis of HT-29 cells was determined by flow cyometry analysis using a fluorescence-activated cell sorting (FACS) caliber (Becton-Dickinson, CA, USA) and Annexin V-fluorescein isothiocyanate (FITC)/propidium iodide (PI) kit (Becton-Dickinson). Staining was performed according to the manufacturer’s instructions. The percentage of cells in early apoptosis was calculated by Annexin V-positivity and PI-negativity, while the percentage of cells in late apoptosis was calculated by Annexin V-positivity and PI-positivity.

**Detection of DNA fragmentation by gel electrophoresis.** DNA fragmentation was determined using DNA ladder detection kit (Invitrogen), following the manufacturer’s instructions. Briefly, HT-29 cells were detached from the plates by scraping
and washed in PBS. Cells (2x10^6) were resuspended in 70 μl TE lysis buffer and the enzyme A solution provided in the kit was added to the crude lysate and incubated at 37°C for 10 min. The resultant sample was then treated with enzyme B solution and incubated at 50°C for 30 min. DNA was precipitated with ammonium acetate solution and pre-cooled absolute ethanol followed by centrifugation at 16,000 x g for 10 min. The DNA pellet was washed with 70% ethanol and re-centrifuged at 16,000 x g for 10 min. Precipitated DNA was resuspended in the provided DNA suspension buffer and analyzed after separation by gel electrophoresis (1.5% agarose). The DNA bands were examined using a Gel Documentation System (BioRad, Model Gel Doc 2000, USA).

**Measurement of mitochondrial membrane potential (Δψm) by flow cytometry analysis with JC-1 staining.** JC-1 is a cationic dye that exhibits potential mitochondria-dependent accumulation, indicated by a fluorescence emission shift from green to red, which thus can be used as an indicator of mitochondrial potential. In this experiment, 1x10^6 treated HT-29 cells were resuspended after trypsinization in 1 ml of medium and incubated with 10 μg/ml of JC-1 (Invitrogen) at 37°C, 5% CO₂ for 30 min. Both red and green fluorescence emissions were analyzed by flow cytometry after JC-1 staining.

**Analysis of caspase activation.** The activities of caspase-3 and -9 were determined by a colorimetric assay using the caspase-3 and -9 activation kits (Invitrogen), following the manufacturer’s instructions. Briefly, after treated with various concentrations of EEHDW for 24 h, HT-29 cells were lysed with provided lysis buffer for 30 min on ice. The lysed cells were centrifuged at 16,000 x g for 10 min, and 100 μg of the protein were incubated with 50 μl of the colorimetric tetrapeptides, Asp-Glu-Val-Asp (DEVD)-p-nitroaniline (pNA) (specific substrate of caspase-3) or Leu-Glu-His-Asp (LEHD)-pNA (specific substrate of caspase-9) at 37°C in the dark for 2 h. Samples were read at 405 nm in an ELISA reader (BioTek, Model E XL800, USA). The data were normalized to the activity of the caspases in control cells (treated with 0.5% DMSO vehicle) and presented as fold of control.

**RNA extraction and RT-PCR analysis.** HT-29 cells (2x10^6) were seeded into 6-well plates in 2 ml medium and treated with various concentrations of EEHDW for 24 h. Total RNA from HT-29 cells was isolated with TRIzol reagent (Invitrogen). Oligo(dT)-primed RNA (1 μg) was reverse-transcribed with SuperScript II reverse transcriptase (Promega) according to the manufacturer’s instructions. The obtained cDNA was used to determine the mRNA amount of Bcl-2 or Bax by PCR with Taq DNA polymerase (Fermentas). GAPDH was used as an internal control for protein loading. After membranes were blocked in TBS with 0.25% Tween-20 (TBST) and incubated at 50˚C for 30 min with agitation at RT in SuperBlock T20 (TBS) blocking buffer (Thermo Scientific, Rockford, IL, USA), and the lysates were separated by Nusep 12% LongLife Tris Glycine iGels (Nusep Ltd., Australia) under a reducing condition using 200 V for 1 h. The proteins were then electrophoretically transferred onto nitrocellulose membranes using the iBlot Western detection stack/iBlot dry blotting system (Invitrogen). Membranes were blocked for 30 min with agitation at RT in SuperBlock T20 (TBS) blocking buffer (Thermo Scientific, Rockford, IL). Membranes were washed in TBS with 0.25% Tween-20 (TBST) and exposed to primary antibodies against Bcl-2 or Bax (1:1000, Cell Signaling Technology) overnight at 4°C with rocking. β-actin (1:1000, Cell Signaling Technology) was also measured as an internal control for protein loading. After membranes were washed in TBST, secondary horseradish peroxidase (HRP)-conjugated antibodies (anti-rabbit, Cell Signaling Technology) were added at 1:250 dilution for 1 h at room temperature and the membranes were washed again in TBST. Blots were developed using Super Signal Pico Substrate (Thermo Scientific, Rockford, IL, USA), and images were taken using a Kodak image station 400R (Kodak, Rochester, NY, USA).

**Statistical analysis.** All data are the means of three determinations and data were analyzed using the SPSS package for

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**Figure 1.** Effect of EEHDW on cell viability of HT-29 cells. (A) HT-29 cells were treated with the indicated concentrations of EEHDW for 24 h. (B) Cells were treated with 4 mg/ml of EEHDW for the indicated time periods. Cell viability was determined by the MTT assay. The data were normalized to the viability of control cells (100%, treated with 0.5% DMSO vehicle). Data are averages with S.D. (error bars) from at least three independent experiments. *P<0.05, **P<0.01, significant versus control cells.
EEHDW inhibits the growth of HT-29 cells. The effect of EEHDW on the viability of HT-29 cells was determined by MTT assay. As shown in Fig. 1A, treatment with 0.5-5 mg/ml of EEHDW for 24 h dose-dependently reduced cell viability by 13-63% compared to untreated control cells (P<0.01), with an estimated half-maximal inhibitory concentration (IC50) value of 4 mg/ml. The cell viability was decreased to 37% at the highest concentration of EEHDW (5 mg/ml) in this study. We also evaluated the effect of 4 mg/ml of EEHDW (IC50 value) on cell viability with incubation for different periods of time. As shown in Fig. 1B, treatment with 4 mg/ml of EEHDW led to a gradual decrease in cell viability with the increase of exposure time. These results suggest that EEHDW inhibits HT-29 cell growth or viability in a dose- and time-dependent manner. To further verify these results, we evaluated the effect of EEHDW on HT-29 cell morphology via phase-contrast microscopy, since cell morphology in culture is indicative of the healthy status of the cells. As shown in Fig. 2, untreated HT-29 cells appeared as densely packed and disorganized multilayers, whereas after incubation with various concentrations of EEHDW for 24 h many of the cells became rounded and shrunken, and detached from each other or floated in the medium. In addition, with 24 h exposure to EEHDW, the cells became less confluent. Taken together, these data demonstrate that EEHDW inhibits the growth of HT-29 cells.

EEHDW induces apoptosis of HT-29 cells. To determine whether the cell-growth suppressive effect of EEHDW is due to apoptosis, we examined EEHDW’s pro-apoptotic activity in HT-29 cells via Annexin-V/PI staining followed by FACS analysis (Fig. 3A). In this assay, Annexin V/PI double-negative population (labeled as LL in the FACS diagram) indicates viable cells; Annexin V-positive/PI-negative or Annexin V/PI double-positive population (labeled as LR or UR in the FACS diagram) represents cells undergoing early or late apoptosis, respectively. As shown in Fig. 3A and B, the percent of cells undergoing apoptosis following treatment with 0, 1, 3 and 5 mg/ml of EEHDW (including the early and late apoptotic cells) was 12, 22, 28 and 40%, respectively (P<0.05, versus untreated control cells). This indicates that EEHDW treatment induces HT-29 cell apoptosis in a dose-dependent manner. To confirm the pro-apoptotic function of EEHDW, we investigated the effect of EEHDW on cellular DNA fragmentation, a typical feature of apoptosis. As shown in Fig. 3C, DNA extracted from HT-29 cells treated with 4 mg/ml of EEHDW (IC50 value) for 24 h displayed a characteristic ladder pattern of discontinuous DNA fragments.

EEHDW induces the loss of mitochondrial potential (Δψm) and the activation of caspases-9 and -3. The mitochondrion-dependent pathway is the most common apoptotic pathway in vertebrate animal cells. The mitochondrial membrane permeabilization, accompanied by the collapse of electrochemical gradient across the mitochondrial membrane, is one of the key events during cellular apoptosis (32,33). This results in the release of numerous apoptogenic proteins, such as cytochrome c, from the mitochondria triggering the activation of caspases-9 and -3, and eventually inducing apoptosis. To investigate the mechanism of how EEHDW induces HT-29 cell apoptosis, we used FACS analysis with JC-1 staining to examine the change in mitochondrial membrane potential after EEHDW treatment. JC-1 is a lipophilic, cationic dye that selectively enters into mitochondria. In
healthy cells with high mitochondrial potential, JC-1 forms J-aggregates with intense red fluorescence (590 nm), whereas under apoptotic condition, the mitochondrial membrane potential collapses, so that JC-1 does not accumulate within the mitochondria but remains in the cytoplasm in monomeric form showing green fluorescence (525 nm). These fluorescence differences can be detected by FACS analysis using JC-1 green and red channels. As shown in Fig. 4, JC-1 fluorescence was shifted from a JC-1-green-bright/JC-1-red-bright signal in untreated HT-29 cells to a JC-1-green-bright/JC-1-red-dim signal in cells treated with EEHDW in a dose-dependent fashion, indicating EEHDW-induced loss of...
mitochondrial membrane potential in HT-29 cells. To identify the downstream effectors in the apoptotic signaling pathway, the activation of caspases-9 and -3 was examined by a colorimetric assay using specific chromophores, DEVD-pNA (specific substrate of caspase-3) and LEHD-pNA (specific substrate of caspase-9). As showed in Fig. 5A and B, EEHDW treatment significantly and dose-dependently induced activation of both caspases-9 and -3 in HT-29 cells (P<0.01 or 0.05, versus untreated control cells). These data suggest that EEHDW promotes HT-29 cell apoptosis via the mitochondrion-dependent pathway.

EEHDW regulated the expression of anti-apoptotic Bcl-2 and pro-apoptotic Bax. Bcl-2 family proteins are key regulators of mitochondrion-mediated apoptosis, including anti-apoptotic members such as Bcl-2 and pro-apoptotic members such as Bax. Tissue homeostasis is maintained by controlling the ratio of active anti- and pro-apoptotic Bcl-2 family proteins. Higher Bcl-2-to-Bax ratio by aberrant expression of the proteins is found commonly in various cancers. To further study the mechanism of EEHDW’s anti-cancer activity, we performed RT-PCR and Western blotting to examine the mRNA and protein expression of Bcl-2 and Bax in EEHDW-treated HT-29 cells. The results of the RT-PCR assay showed that EEHDW treatment profoundly increased Bax and reduced Bcl-2 mRNA expression in HT-29 cells (Fig. 6A); and the pattern of protein expression of Bax and Bcl-2 was similar to their respective mRNA levels (Fig. 6B). This suggests that EEHDW induces mitochondrion-dependent apoptosis in HT-29 cells through the regulation of expression of Bcl-2 family proteins.

Discussion
Cancer cells are characterized by an unregulated increase in cell proliferation and/or a reduction in cell apoptosis (15). In addition, disrupted apoptosis contributes to drug resistance of tumor cells, which has become a significant obstacle for the successful management of patients with malignant tumors including colorectal cancer (CRC) (2-4). Moreover, many currently used anti-cancer agents contain intrinsic and potent cytotoxicity to normal cells, which limits their long-term use and thereby their therapeutic effectiveness (5). These problems highlight the urgent need for the development of novel cancer chemotherapies. Since natural products, such as traditional Chinese herbal medicines (TCM), have relatively fewer side-effects as compared to modern chemotherapeutics and have long been used clinically to treat various types of diseases including cancer (8-11), discovering naturally occurring agents with pro-apoptotic activities is a promising approach for anti-cancer treatment. *Hedyotis Diffusa Willd* is an important traditional heat-clearing and detoxifying Chinese herb used in TCM formulations, with many reported pharmacological applications. Recently, *Hedyotis Diffusa Willd* has been demonstrated to be clinically effective in treating various cancers including CRC (12-14). However, the mode of action for its anti-tumor
is still largely unknown. Therefore, before *Hedyotis Diffusa Willd* can be further developed as an anti-cancer agent, its anti-tumor activity and underlying molecular mechanism should first be elucidated.

Here we reported for the first time that the ethanol extract of *Hedyotis Diffusa Willd* (EEHDW) reduces the viability and inhibits growth of human colon carcinoma HT-29 cells in a dose- and time-dependent fashion. Furthermore, we demonstrated that these effects on HT-29 cells result from the induction of apoptosis by EEHDW.

Apoptosis manifests in two major pathways. For the intrinsic pathway, death signals are integrated at the level of the mitochondria (therefore, also this pathway is referred to as mitochondrial-dependent). For the extrinsic pathway, death signals are mediated through cell surface receptors. Both pathways eventually lead to the activation of caspases and nucleases, resulting in the destruction of the cell (15-17). Mitochondrial outer membrane permeabilization (MOMP) is a key commitment step in the induction of cellular apoptosis, since it is the point of convergence for a large variety of intracellular apoptotic signaling pathways leading to the release of many apoptogenic proteins from the mitochondrial intermembrane space. During the process of MOMP, the electrochemical gradient across the mitochondrial membrane collapses. Therefore, the loss of mitochondrial membrane potential is a hallmark for apoptosis. Our data clearly showed that treatment with EEHDW leads to a collapse of mitochondrial membrane potential.

Caspases, represented by a family of cysteine proteases, are the key proteins that modulate the apoptotic response. Caspase-3 is a key executor of apoptosis, which is activated by an initiator caspase such as caspase-9 during mitochondrial-mediated apoptosis. In this study, we found that EEHDW induces the activation of both caspase-9 and -3 in HT-29 cells in a dose-dependent manner. Thus, EEHDW-induced HT-29 cell death is accompanied by an increase in the activities of caspases-9 and -3, which then stimulates the molecular cascades for apoptosis.

Mitochondrion-dependent apoptosis is mainly regulated by Bcl-2 family proteins. MOMP is thought to occur through the formation of pores in the mitochondria by pro-apoptotic Bax-like proteins, which can be inhibited by anti-apoptotic Bcl-2-like members. Therefore, the ratio of Bax to Bcl-2 is a critical determinant of the fate of cells. In this study, we demonstrated that EEHDW treatment dose-dependently enhances Bax mRNA expression and reduces Bcl-2 mRNA expression in HT-29 cells. This indicates that EEHDW induces apoptosis by affecting the ratio of Bax:Bcl-2 at transcriptional level. We further studied the role of EEHDW on the expression of proteins involved in the mitochondrial pathway. The results showed that EEHDW treatment up-regulates Bax protein expression and down-regulates Bcl-2 protein expression, which is in accordance with the pattern of their mRNA expression after EEHDW treatment.

In conclusion, our data for the first time demonstrate that EEHDW inhibits the growth of and induces HT-29 cell apoptosis via the mitochondrial-dependent pathway. These results suggest that *Hedyotis Diffusa Willd* may be a potential novel therapeutic agent for the treatment of colorectal and other cancers.

Acknowledgments

This work was supported by Open Fund of Fujian Key Laboratory of Integrative Medicine on Geriatrics (2008J1004) and Developmental Fund of Chen Keji Integrative Medicine (CKJ 2008056).

References


