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Underlying mechanism for the modulation of apoptosis induced by a new benzoindole derivative on HT-29 colon cancer cells

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Colorectal cancer is the third most common form of cancer affecting both men and women around the world. The chemical and biological studies of heterocyclic compounds have been an interesting area in pharmaceutical and medicinal chemistry. A new synthetic compound namely 2-(1,1-dimethyl-1*H*-benzo[e]indol-2-yl)-3-((2-hydroxyphenyl)amino)acrylaldehyde, abbreviated as DBID was screened for the antiproliferation effects against the colorectal cancer cell line, HT-29 and its possible mechanism of action was elucidated. To determine the IC₅₀ value, MTT assay was employed and further verified by the LDH release assay and apoptosis-inducing effect. DBID inhibited the proliferation of HT-29 cells and significantly increased the levels of caspase -8, -9 and -3/7 in the treated cells compared to untreated cells. Apoptosis features in HT-29 cells were detected in treated cells by using the AO/PI staining and flow cytometric analysis of Annexin V. The changes in expression of some apoptotic genes were confirmed by gene quantification using RT-PCR. The current study showed that the DBID compound exhibited chemotherapeutic activity, which was evident by significant increases in the expression and activation of caspase, up-regulation of the expression of specific apoptotic genes and exploitation of the apoptotic signaling pathways to trigger cancer cell death.

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Introduction

The American Cancer Society reported colorectal cancer as the third most common form of cancer in both genders around the world. Their report indicates that, for 2017, colorectal cancer would be the third leading cause of cancer deaths in women, and the second leading cause of death in men estimating about 50 260 deaths as a result from the disease. Recently, there has been a drastic increase in the number of colon cancer cases. This surge has highlighted this cancer as a health issue of serious concern, with 135 430 cases of colorectal cancer being estimated and expected to be reported in 2017.¹

Age is the one of the risk factors for colorectal cancer as 91% of patients are diagnosed at 50 years of age or older. The other risk factors are personal or family history of colorectal cancer and/or polyps, known to have chronic inflammatory bowel

disease, and some inherited genetic manifestations. Modifiable risks factors include alcohol consumption, long-term smoking, obesity, less physical activity and high amounts of red or processed meat in diet. The standard treatment for colon and rectal cancer is surgical removal, followed by adjuvant chemotherapy or radiotherapy. For localized cancers, surgical resection may be curative. However, this therapy is just moderately successful especially for late stage cancers; therefore, new approaches to the treatment of colorectal cancer are required.^{1–3}

The chemistry and biological study of heterocyclic compounds have been an interesting area for a long time in pharmaceutical and medicinal chemistry. Medicinally many indole derivatives are acknowledged to have a wide range of medicinal properties.^{4–6}

In the present study, new Schiff base 2-(1,1-dimethyl-1*H*-benzo[e]indol-2-yl)-3-((2-hydroxyphenyl)amino)acrylaldehyde (DBID) has been synthesized and the potential chemotherapeutic properties of DBID, as a novel synthetic compound and its possible mechanism of action was investigated on colorectal cancer cell line, HT-29.

Experimental

Synthetic compound

The new compound 2-(1,1-dimethyl-1*H*-benzo[e]indol-2-yl)-3-((2-hydroxyphenyl)amino)acrylaldehyde (DBID) was synthesized,

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characterized and confirmed by spectroscopic techniques. These data have already been published by the same authors.⁷ The compound was provided by the Department of Chemistry, University of Malaya.

Cell culture

To evaluate the antiproliferative activity of DBID, a human colonic carcinoma cell line, HT-29 (ATCC® HTB-38™) was purchased from the American Type Culture Collection (ATCC, VA, USA). Normal human colon cell CCD 841 CoN (ATCC® CRL-1790™) was provided from American Type Culture Collection, VA, USA. The HT-29 and CCD 841 CoN cells were cultured in ATCC-formulated McCoy's 5A Medium (ATCC® 30-2007™) and Eagle's Minimum Essential Medium (ATCC® 30-2003™), respectively. The cells were supplemented with 10% FBS (fetal bovine serum), and 1% penicillin–streptomycin (iDNA, South America), incubated at 37 °C in a 5% CO₂ incubator.

Antiproliferative effects of DBID

The inhibitory effect of DBID was investigated using a human colorectal carcinoma cell, HT-29 while the antiproliferative effect of DBID was evaluated by using the MTT (3-(4,5-dimethyl thiazol-2-yl)-2,5-diphenyl tetrazolium bromide) assay.⁸ Normal colon cell line, CCD 841 CoN was used to identify any cytotoxic effect of the compound. The cells were treated with various concentrations of the compound (0–50 µg ml⁻¹) for 48 h. All the experiments were performed in triplicate. In this study 5-fluorouracil (5-FU) was used as a standard drug. The IC₅₀ known as the concentration of the compound that caused 50% growth inhibition was determined based on the following formula;

$$\text{Inhibition (\%)} = \frac{[(\text{OD untreated} - \text{OD treated})/(\text{OD untreated})] \times 100}{\text{OD untreated}}$$

Determination of lactate dehydrogenase (LDH) release

The release of lactate dehydrogenase (LDH) into the culture medium was assessed using a CytoTox-ONE™ Homogeneous Membrane Integrity Assay kit (Promega, Madison, WI, USA) according to the manufacturer protocol. In this experiment, the cells were treated with different concentration of DBID for 24 h. The lysis solution (9% weight/volume solution of Triton® X-100 in water) was used to determine the maximum amount of LDH present. The percentage of cytotoxicity was calculated by using the following formula:

$$\text{Cytotoxicity (\%)} = 100 \times \frac{[(\text{sample LDH release} - \text{culture medium background})/(\text{maximum LDH release} - \text{culture medium background})]}{\text{maximum LDH release} - \text{culture medium background}}$$

Reactive oxygen species generation

The intracellular reactive oxygen species (ROS) generation was measured using Cellular Reactive Oxygen Species Detection Assay Kit (Abcam, Cambridge, UK; ab113851) according to the manufacturer protocol. The procedure was based on oxidation

of DCFH-DA to form the fluorescent compound, 2',7'-dichloro-fluorescein (DCF) by intracellular ROS and other peroxides. The cells were treated with DBID at the different concentrations for 24 h.

Fluorescence microscopic examination

To investigate the morphological characterization of HT-29 cells, propidium iodide and acridine orange double staining was performed where the cells were then observed under a fluorescence microscope according to the method described previously.^{9,10} The HT-29 cells were treated with two different concentrations of DBID for 24 h. The harvested cells were stained with an equal volume of fluorescent dye (AOPI) and observed under a fluorescence microscope.

Flow cytometric Annexin V

The Annexin V assay was carried out using FITC Annexin V Apoptosis Detection Kit supplied from BD Pharmingen™ (ApoAlert Annexin V, Clontech, California, USA). The cultured cells in a 25 cm flask were treated with the IC₅₀ concentration of DBID compound determined by MTT assay for 24 h. The cells were then subjected with the FITC Annexin V staining protocol to measure apoptosis.

Caspase activity assays

The caspase -3/7, -8 and -9 activities was evaluated by using commercial kits purchased from Promega Company (Madison, WI, USA) according to the manufacturer's protocol. The cells were treated with different concentrations of DBID for 24 h. Cells without any treatment, cell treated with respected caspase inhibitor and media alone without cells were used as the controls and blank.

Gene expression analysis by RT-PCR

HT-29 cells were treated with the IC₅₀ concentration of DBID and incubated for 24 h in 25 cm² culture flasks. Total RNA was extracted with RNeasy® plus mini kit (Cat. no. 74134, QIAGEN, Germany), and assessed for quality and quantity using absorption measurements (Thermo Scientific Nanodrop 2000 Spectrophotometer, Massachusetts, USA). Reverse transcription into cDNA was performed by using Two-Step qRT-PCR kit, high capacity RNA to cDNA by Applied Biosystems, USA. The relative quantification of the genes was performed using the TaqMan® Gene Expression Master Mix by following the manufacturer's protocol. The expression of two endogenous controls ACTB (Hs99999903_m1) and GAPDH (Hs03929097_g1) were used in this study. Apoptosis-related cysteine peptidase specific primers; caspase 3 (CASP3, Hs00234387_m1), caspase 8 (CASP8, Hs01018151_m1), caspase 9 (CASP9, Hs00609647_m1), apoptosis regulator; BCL2 (BCL2, Hs00608023_m1), BCL2 associated X (BAX, Hs00180269_m1), tumor protein p53 (TP53, Hs01034249_m1) and BH3 interacting domain death agonist (BID Hs00609632_m1) were purchased from TaqMan® (MGB probes, FAM™ dye-labeled). All experimental samples were performed in biological triplicates. The ΔΔCt method was used



to determine the relative expression and comparative Ct values analysis *via* differences between the treated and untreated groups were performed.

Statistical analysis

All experiments were carried out in triplicate, and the results were analyzed and presented as means \pm standard deviation (SD). Statistical analysis was done by using SPSS V.18.0 statistical software, Student's *t*-test. *P*-value less than 0.05 was considered statistically significant.

Results and discussion

In vitro antiproliferative effect

The colonic carcinoma cell line HT-29 was selected to determine the antiproliferative activity of the compound. In addition, CCD 841 CoN cells was included to investigate the cytotoxicity of the compound on normal colon cells. As shown in Table 1, DBID exhibited the potent antiproliferative effect against HT-29 cells ($IC_{50} = 11.85 \pm 2.7$) whereas the IC_{50} of the compound on normal colon cells ($IC_{50} > 150 \mu\text{g ml}^{-1}$) was significantly higher than HT-29 ($p < 0.05$). The 5-fluorouracil (5-FU) acting as the reference control showed high anti-proliferative activities against tumorigenic HT-29 cells. The findings of this study showed strong inhibitory effect of the DBID compound against the tumorigenic cell line; HT-29 without no cytotoxic effect on normal cells. The MTT assay showed that DBID at concentration of $11.86 \mu\text{g ml}^{-1}$ can inhibit 50% of HT-29 proliferation *in vitro*.

The leakage of LDH

The cytotoxic effect of DBID compound against HT-29 cells was further verified by the LDH release assay. The measurement of the LDH release is commonly used for testing cytotoxicity based on the integrity of the cell membrane. The loss of integrity of the cell membrane leads to the leakage of LDH from the cytoplasm into the surrounding culture medium in non-viable cells.¹¹ DBID compound induced LDH leakage in HT-29 cells when compared to untreated cells (Fig. 1). The LDH leakage from HT-29 cells treated with DBID compound was significantly different at concentration as low as $10 \mu\text{g ml}^{-1}$ compared to the untreated cells ($p < 0.05$).

In determining the release of lactate dehydrogenase (LDH), the significant percent of cytotoxicity of DBID against HT-29 cells was shown at concentration $10 \mu\text{g ml}^{-1}$, which was compatible

Table 1 Antiproliferative effects and cytotoxicity activities of a new benzindole derivative (DBID) on colon cancer cell line and normal human colon cell line^a

Compound	HT-29	CCD 841 CoN
DBID	11.85 ± 2.7	189.73 ± 4.1
5-Fluorouracil	3.9 ± 1.26	43.8 ± 6.3

^a Data are presented as IC_{50} . IC_{50} is the effective concentration of the compound which was used to prevent the 50% of the cells' growth ($\mu\text{g ml}^{-1}$). Results were expressed as means \pm SD ($n = 3$).

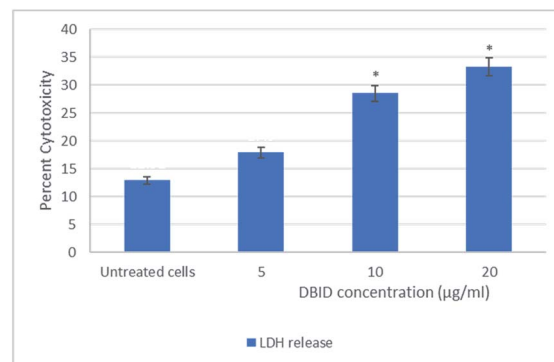


Fig. 1 Lactate dehydrogenase (LDH) release in HT-29 cells treated with different concentration of DBID compound for 24 h. Results were expressed at means \pm SD ($n = 3$). *Indicates significant differences compared to the control (untreated cells) verified by the Student's *t*-test ($p < 0.05$).

with the IC_{50} value of DBID ($11.85 \pm 2.7 \mu\text{g ml}^{-1}$) determined in the MTT assay. Based on the presented results, DBID compound was toxic to HT-29 cells, which resulted in a loss of membrane integrity, release of LDH into the culture medium, and generated a greater fluorescent signal in comparison with untreated cells. The anticancer effect of heterocyclic compounds has been proposed in many studies.^{12–16}

Intracellular ROS formation

The reactive oxygen species are produced continuously in living cells, as a result of metabolic and other biochemical reactions. Under oxidative stress conditions, excessive ROS leads to various biochemical and physiological impairments and promotes cell death. Fig. 2 illustrates the intracellular reactive oxygen species (ROS) generation in treated cells with different concentration of DBID for 24 h compared with untreated cells. The formation of ROS in HT-29 cells treated with the DBID compound increased in a dose-dependent manner and the strong significant production of ROS in treated cells was

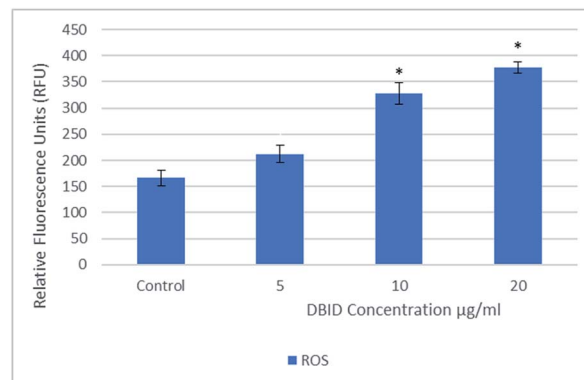


Fig. 2 Intracellular ROS generated in HT-29 cells treated with different concentration of DBID compound for 24 h. Results are expressed as mean \pm standard deviation. * $p < 0.05$ versus control (untreated cells) by Student's *t*-test.



generated at concentration of 10 and 20 $\mu\text{g ml}^{-1}$ upon treatment. At 5 $\mu\text{g ml}^{-1}$ of DBID concentration treatment, a slight increase of ROS was observed.

DBID was able to elevate the production of ROS in the treated cells significantly compared with untreated cells. The ROS concentration is vital in cells and tissue. Under oxidative stress conditions, excessive ROS leads to a various biochemical and physiological impairments and promotes cell death. The important role of intracellular ROS in apoptosis induction has been reviewed previously.¹⁷ Previous studies have suggested that the reactive oxygen species are responsible as part of the signaling process for activating an important mechanism for eliminating cancer cells such as apoptosis. It has been proposed that the disproportional increase in intracellular ROS leads to release cytochrome c from mitochondria which trigger caspase activation.^{18–20} There are several anticancer agents depend on this form of cell death for their efficacy.^{19,21} The present results propose the apoptosis inductions through the production of intracellular ROS formation in cells treated with DBID compound.

Fluorescence microscopic examination

The occurrence of apoptosis was qualitatively investigated. This involved the use of the two dyes of AO/PI and fluorescence microscopy. AO penetrates the plasma membrane of viable and early apoptotic cells where stained nucleated cells would generate a green-coloured fluorescence while PI would only enter the cells with compromised membranes and later produce an orange/red fluorescence intensity in dying, dead, and necrotic nucleated cells stained with PI.^{22,23}

As shown in Fig. 3, a normal morphology with green nuclei and also an intact structure was observed in untreated cells while the cells treated with DBID showed bright green nuclei indicating the condensation of nucleic chromatin and interpolated propidium iodide (orange fluorescence) amongst the fragmented DNA and nuclear fragmentation. Apoptosis characteristics *e.g.* cell shrinkage, membrane blebbing and chromatin condensation were observed in DBID treated cell. Fig. 3 shows remarkable morphological changes of the HT-29 cells treated with DBID in both concentrations (Fig. 3B and C) in apoptotic bodies. The results of this study proved that the activation of the caspase pathway may trigger apoptosis in HT-29 cells that had been treated with DBID.

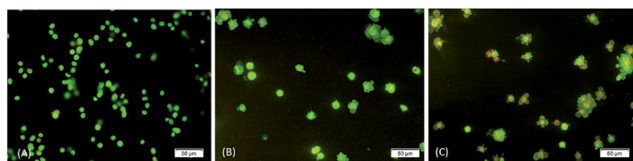


Fig. 3 Morphological changes of untreated and DBID-treated HT-29 cells. Fluorescence microscopic examination by using double fluorescent dye (AO/PI) staining method on cells treated with DBID after 24 h incubation. From left to right; (A) untreated HT-29 cells, (B) 10 $\mu\text{g ml}^{-1}$ DBID-treated, (C) 20 $\mu\text{g ml}^{-1}$ DBID-treated.

Annexin V assay

Annexin V/propidium iodide (PI) flow cytometry was performed to confirm the presence of apoptosis. Annexin V is used to detect apoptosis by targeting the loss of plasma membrane asymmetry by specific interaction with phosphatidylserine (PS) on the cell surface. The intact cell membrane of viable cells is not permeable to PI and represent AV–/PI– staining. The loss of plasma membrane asymmetry and strong affinity of AV-FITC with PS leads to AV+/PI– staining in the early apoptotic cells, whereas AV+/PI+ represents late apoptotic. Necrotic stage is representing by AV–/PI+ as a result of PI penetration through the membranes and intercalate into nucleic acid due to loss of plasma and nuclear membrane integrity. The flow cytometry of Annexin V results and analysis are presented in Fig. 4(A and B). The cells treated with DBID significantly underwent apoptosis after 24 h post treatment compared with untreated cells ($p < 0.05$). More than 40% of treated cells were at the early and late stages of apoptosis after 24 h (Fig. 4(B)). In addition, treatment with DBID showed no concurrent increase in necrotic cells. Consistent with AOPI staining results, flow cytometric Annexin V analysis of DBID treated cells clearly demonstrated that anti-proliferation and apoptosis in HT-29 treated cells are closely related.

Cellular caspase activities (caspase -9, -8 and -3/7)

The activation of caspases -9, -8 and -3/7 in HT-29 cells treated with different concentration of DBID compound were determined. Each of these assays were linear with respect to the number of cells undergoing apoptosis in the experiment. As shown in Fig. 5(A–C), the compound (DBID) significantly increased the activities of caspases -9, -8 and -3/7 in HT-29 cells,

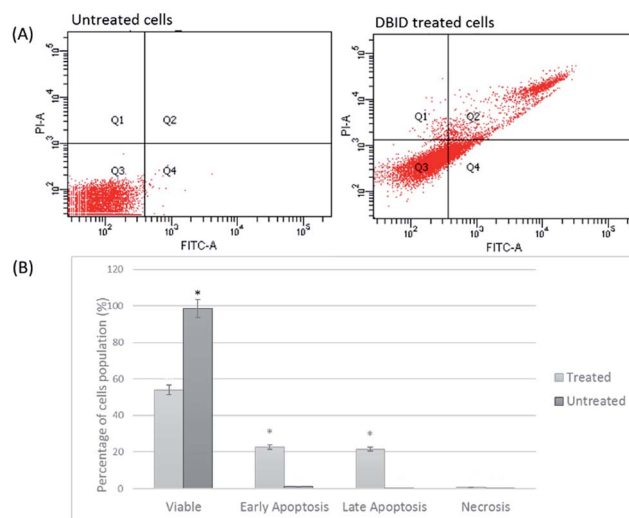


Fig. 4 Flow cytometric Annexin V/PI. (A) HT-29 cells (untreated and DBID treated) were stained with Annexin V/PI (B) flow cytometric analysis of the percentage level of the four quadrants representing viable cells (Annexin V–PI–), early apoptosis (Annexin V+PI–), late apoptosis (Annexin V+PI+) or necrotic (Annexin V–PI+) stages in treated and untreated cells. Values shown are percentages of each quadrant * $p < 0.05$.



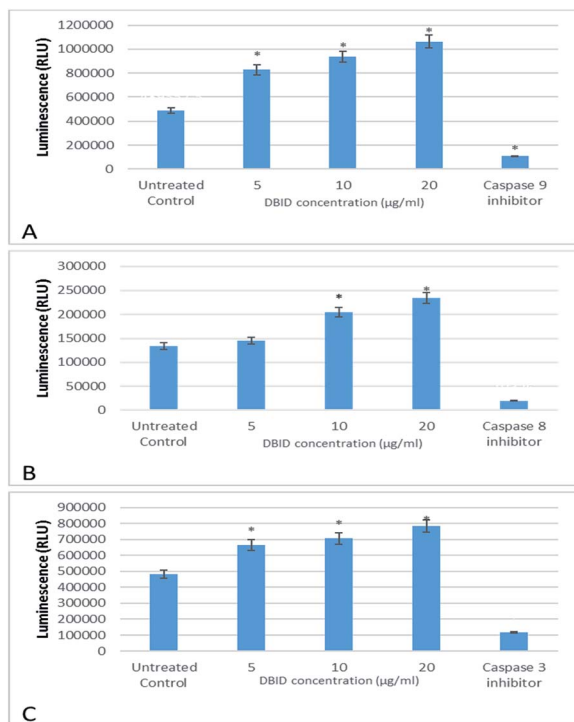


Fig. 5 The effect of DBID on caspase activity in HT-29 colon cancer cells. The activity of (A) caspase-9 (B) caspase-8 and (C) caspase-3/7. Data shown are the mean values of the triplicates. *Indicates significant differences from untreated cells ($p < 0.05$).

compared with the control in a dose-dependent manner ($p < 0.05$). Fig. 6 represents the caspase activity fold changes over control cells. The caspase-8 activity was significant only at the concentration of 10 and 20 $\mu\text{g ml}^{-1}$ of the compound ($p < 0.05$). The activity of caspase -3/7 and -9 in treated cells was steadily increased in a dose-dependent manner and statistically significant compared to the untreated cells ($p < 0.05$). Apoptosis is typically accompanied by the activation of caspase proteins. Caspases are cysteine–aspartic proteases or cysteine-dependent aspartate-directed proteases, a family of cysteine proteases that play essential roles in apoptosis. Caspase-9 is known as an indicator for the activation of a branch caspase cascade, namely, the intrinsic (mitochondria mediated) apoptotic pathway and caspase-8 is a marker for the extrinsic (receptor) pathway of apoptosis. Caspase-3 is activated in apoptotic cells by both the extrinsic and intrinsic pathways.²⁴ The results of caspase activity suggested that the intrinsic or mitochondrial pathway were mainly mediated by the activation of caspase-9, and also the extrinsic or death receptor pathway mediated by the activation of caspase-8, triggered the apoptosis induction in treated cells. These results revealed that the caspase -9 and -8 as initiator caspases and caspase-3/7 as executioner caspases were involved in the mechanism of apoptosis induced by the compound on HT-29 cells. Consistent with this result, the cytotoxic effect and antiproliferative activity of heterocyclic compounds synthetic compounds, against colon cancer cells by activating caspases and inducing apoptosis have also been shown in a previous study.^{25–27}

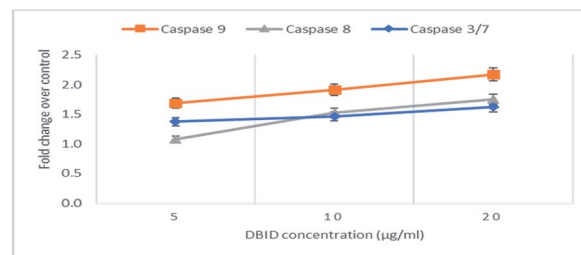


Fig. 6 The fold change in caspase activity of HT-29 cells treated with DBID at different concentrations.

Gene expression

The HT-29 cells were treated with the IC_{50} concentration of the DBID compound to further investigate the possible pathways involved in antiproliferative effect of the compound at gene expression level using real time-PCR technique.

The gene expression study of some of these genes involved in apoptosis showed the overexpression of TP53, BID and BAX in HT-29 cells. Furthermore, the genes encode the cysteine–aspartic acid protease (caspase) family *e.g.* CASP9, CASP8 and CASP3 were also up-regulated in this study. Fig. 7 shows that CASP3, CASP9, CASP8, TP53, BAX and BID were significantly overexpressed in treated HT-29 cells by a mean factor of 2.48, 2.79, 1.67, 3.7, 4.1 and 1.47, respectively. The apoptosis regulator gene, BCL2 was slightly downregulated in DBID treated cells by a mean factor of 0.89.

The activation of caspases is known as a hallmark of apoptosis which proposals new therapeutic agents for the treatment of cancer cells.^{28–31} The p53 tumor suppressor gene (TP53) encodes the tumor protein p53, a transcriptional factor that binds to DNA and activates the expression of downstream genes, which mediates cellular stress responses *i.e.* cell-cycle arrest, senescence and apoptosis.³² BID, a pro-apoptotic member of the Bcl-2 family, mediates the activation of caspase-9 by caspase-8 through the mitochondrial pathway. The ability of DBID to activate BAX can be inhibited by the anti-apoptotic Bcl-2 proteins resulting in apoptosis inhibition by sequestering DBID, leading to reduced Bax activation.^{33,34}

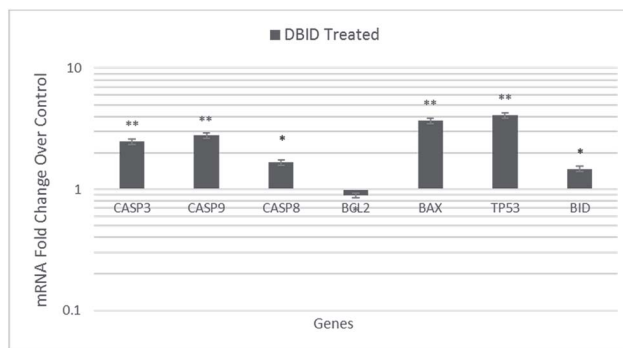


Fig. 7 Gene expression analysis of HT-29 cells treated with the IC_{50} concentration of DBID for 24 h. Results are expressed as mean \pm standard deviation of three replicates * $p < 0.05$, ** $p < 0.01$.



Moreover, the intrinsic or mitochondrial pathway is mediated by the alteration of the pro-apoptotic (Bax) and anti-apoptotic (Bcl2) ratio. Bcl2L (Bcl2-Like), BclXL (Bcl2 related protein long isoform) and other anti-apoptotic Bcl2 family members reside in the outer mitochondrial membrane and prevent cytochrome c release.³⁵ These results revealed that the treatment with DBID promotes apoptosis by alerting the Bax/Bcl2 ratio by increasing the expression of Bax protein and down-regulating the expression of Bcl-2 as well as upregulating the genes involved in apoptosis. These findings of the current study are consistent with previous studies reported by several researchers.^{28–31,36,37}

Conclusions

The present study demonstrates the antiproliferative activity of a new benzindole derivative (DBID) against colon cancer cell line, HT-29, without being toxic to the normal colon cell (CCD 841 CoN). The quantitatively and qualitatively investigation of caspase activities by using luminescence spectrophotometry, fluorescence microscopy (AO/PI), flow cytometric Annexin V and gene expression provided in the present study, proposed that the antiproliferative effect of DBID on colon cancer cell line could be due to the occurrence of apoptosis through the generation of intracellular ROS, induction of cellular caspase activities as demonstrated by, initiation of both the intrinsic and extrinsic apoptosis pathways and upregulation of apoptotic genes. In conclusion, DBID which is a novel synthetic compound shows potential and promising chemotherapeutic properties.

Acknowledgements

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