IN VITRO EVALUATION OF ANTI-CANCER ACTIVITY OF CURCUMINOIDS FROM TURMERIC (CURCUMA LONGA L.) AGAINST MULTIDRUG RESISTANT TUMOR CELL LINES.

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Abstract

Cervical cancer is the second most common cancers among women worldwide. Chemotherapy is the treatment of cancer with one or more cytotoxic drugs as part of a standard regimen. One of the major causes of chemotherapy failure in cancer treatment is multidrug resistance (MDR). It shows resistance to general anticancer drugs used and shows cross resistant to many different structurally unrelated drugs causing multidrug resistance. MDR is over expression of the ATP-binding cassette (ABC) transporters (efflux pump) such as p-glycoprotein (ABCB1 / p-gp). To overcome drug resistance most obvious response that widely employed is to use combination drug therapy. The association of phytochemicals with chemotherapy shows synergistic effects that target simultaneously multiple pathways and help to kill cancer cells and slowdown the onset of drug resistance. The present study was undertaken to assess the chemotherapeutic activity of curcuminoids against the cervical carcinoma cell line HeLa and drug resistant cell lines. Curcuminoids have broad spectrum of pharmacological properties. The IC₅₀ values of curcuminoids revealed that curcumin had a higher cytotoxicity against HeLa cell lines (IC₅₀= 4.30µg/ml) than DMC and BDMC are (IC₅₀= >100 and 10.43µM) respectively. The cytotoxicity of curcuminoids against drug resistant cell lines revealed that higher cytotoxicity at BDMC treatment at IC₅₀= 7.8 and 5.2µM on KB and KBChR8-5 respectively. Therefore BDMC act on MDR cell lines more effectively than other curcuminoids which revealed that it may have synergistic effect that target the pathway related to multidrug resistance and block the efflux pump to increase the accumulation of anticancer drugs and induce cytotoxicity.
treatment it is called cancer recurrence, in such cases patients should get regular treatments with chemotherapy, radiation therapy throughout their life time. One of the reasons for cancer recurrence may be development of drug resistance to common chemotherapeutic drugs used in cancer treatment. Human increasingly employs chemicals as chemotherapeutic agents to treat cancer; the repeated use of these chemicals often leads to becoming ineffective due to other onset of resistance by the target cells.

Tumor cells that are initially sensitive to a broad range of drugs can frequently develop resistance to a group of anticancer drugs. (Gottesman and Ling, 2006). The human multidrug-resistance (MDR1) gene encodes a membrane transporter p-glycoprotein (p-gp) a member of ATP binding cassette family (Borst et al., 2000). The expression of MDR gene plays important role in multidrug resistant phenotype and also confers to cross resistance to a large group of lipophilic cytotoxic compounds. Multidrug resistance in human and rodent cell lines results in decreased intracellular drug accumulation and correlates with the increased expression of MDR genes due to the presence of efflux pump (Gottesman and Pastan, 1988).

The basis of drug resistance have been studied in laboratory using drug resistance cell lines that have been isolated by exposing to increasing amounts of any one of the chemotherapeutic agents to various cancer cells in order to develop resistance to that particular drug (Ling and Thompson, 1974). These isolated resistant cells are frequently resistant to particular drug but also shows cross resistant to other drugs.

Akiyama et al., screened human cultured cancer cells for two properties: high efficiency of cloning and sensitivity to common MDR drugs such as colchicine, vinblastine and doxorubicin. (Akiyama et al., 1985). It was actually a subclone of HeLa, a cervical adenocarcinoma cell line that had contaminated and overgrown quite a few putative human cancer cell lines and had contaminated and distributed as KB (Nelson-Rees and Flandermeyer., 1976). The growth properties and drug sensitivity on HeLa (KB) has proved to be a very useful human cancer cell line for studying development of drug resistance (Boshart et al., 1984).

A large number of non cytotoxic compounds known as chemosensitizers or MDR modulators sensitize resistant cells for the action of cytotoxic drugs. Several generations of p-gp modulators are developed in reversing p-gp mediated multidrug resistance (MDR). The basic strategy to circumvent MDR is to co-administration of an anticancer drug with a chemosensitizer that impairs p-gp function resulting in enhanced intracellular anticancer drug accumulation (Leonard et al., 2002). It is necessary to identify natural compounds from plant origin that reverse the MDR phenotype, sensitize cancer cells to conventional chemotherapy without undesired toxicological effects. Dietary phytochemicals have been found to be very promising in reversing the resistance to anticancer.

Curcuma longa L. (turmeric) is a typical herbaceous plant that characterizes the family Zingiberaceae. Curcumin (C), main colouring substance in Curcuma longa L. and two related compounds, demethoxycurcumin (DMC) and bisdemethoxycurcumin (BDMC), were altogether known as curcuminoids (Govindarajan, 1980). The total of curcuminoids which are about 4-6%, turmeric also contains 2-4% essential oil and various volatile oils, including turmerone, atlantone, and zingiberone (Merina Benny Antony, 2003). Since the curcuminoid pigments vary in chemical structures, it is possible that the physico-chemical characteristics as well as the functional properties and biological role of curcuminoids would vary among them. The extraction of curcuminoids were done by soxhlet extraction of turmeric powder with acetone gave a yield of about 5.0% containing 43% curcuminoids in 4 to 6 hours. (Revathy et al., 2011). A number of studies are undertaken to separate curcuminoid pigments by thin layer chromatography (TLC), high-performance thin-layer chromatography (HPTLC), and column chromatography (CC) (Gupta et al., 1999). HPLC method was sensitive, precise, and accurate for detection and quantification of curcuminoids in the extract of rhizome Curcuma longa L. (Sompol et al., 2009). Characterization of compounds by GC-MS, FTIR, NMR confirms the identity of curcuminoids. The cytotoxicity of drugs were analysed by MTT Assay (Mosmann, 1983) is a sensitive, quantitative and reliable colorimetric assay that measures viability, proliferation and activation of cells.

In the present study we have demonstrated extraction, identification, purification and characterization of curcuminoids from turmeric rhizome. Further we have examined the cytotoxic properties of individual curcuminoids on drug resistance cell line.
Materials and Methods:-  
**Extraction of curcuminoids:**
Curcuma longa L. (Turmeric) rhizome was collected from Assam - Lakhadong variety. Fresh rhizomes were cleaned, washed, dried and blended to powder. About 20g of turmeric powder was taken into a thimble and placed in a soxhlet apparatus and extracted using acetone as solvent for 6 hours. The dark brown extract yielded was then cooled, filtered, concentrated using rotary evaporator. Extracts were analysed for curcuminoids by HPLC analysis. Extract were tested in TLC pre-coated silica gel (Merk-60 F254,0.25mm thick) plate developed using a camag twin trough glass tank with the mobile phase Chloroform:methanol (95:5) and the plate was developed to a height of about 10cm. After development, plates were removed and dried and spots were visualized in UV light.

**Column chromatography:**
About 5gm of crude Curcuminoids were mixed with 8gm of silica gel and loaded on to the glass column (60-120 mesh) and eluted with chloroform followed by chloroform:methanol with increasing polarity. All the collected fractions were subjected to TLC silica gel 60 F254 plate using chloroform:methanol (95:5) as the developing solvent system and detected as yellow spots. And fractions with similar Rf values were pooled and the organic solvent was removed by rotary evaporator.

**Purification of each curcuminoids:**
The individual Curcuminoids collected from the column chromatography was dissolved in methanol and heated. After complete dissolution chloroform was added to get the ratio methanol:Chloroform 5:2 and kept at 5°C for overnight. The crystals obtained were separated by filtration. The crystals were precipitated with petroleum ether. The purity of Individual crystals were analysed in HPLC.

**Characterization of Curcuminoids:**  
**Estimation of Curcuminoids by HPLC analysis:-**
Weighed accurately 25mg sample and dissolved in 25ml acetone Filtered through 0.2 m membrane filter before injection. Samples were analysed by HPLC in a Shimadazer LC 20A0 liquid chromatograph system with SPD-M20AuV detector in isocratic mode. 20μl of sample was injected and the elution was carried out with gradient solvent systems with a Column used was C18 (250X4.6mm), mobile phase 40% THF and 60% water containing 1% citric acid, pH adjusted to 3.0 using concentrated potassium hydroxide solution and measured in wavelength of 420nm. (Cooper et al., 1994).

**Gas chromatography – Mass Spectrometer (GC-MS):-**
Molecular weight of compounds was analysed by GC-MS using JOEL GCMATE II GC Mass spectrometer. Each curcuminoids were dissolved in methanol, the solution was then injected. Perfluorokerosene (PFK) is used as reference sample (Biemann, 1962).

**Fourier Transform - Infra Red Spectroscopy (FTIR):-**
Curcuminoids samples are dispersed in 100mg KBr and pressed to form a pellet and analysed on Perkin Elmer Spectrum One FT-IR instrument. (Colthup, 1990).

**Nuclear Magnetic Resonance Spectroscopy (NMR):-**
Spectra of 13C and 1H were determined in DMSO-d6, operating at 200 and 50 MHz respectively, using JEOL GSX-400 NMR spectrometer. Tetramethyl silane (TMS) was used as internal standard. (Slichter, 1990).

**Cell culture:-**
The KB, KB-ChR-8-5 human cervical carcinoma cell lines and HeLa cell lines were purchased from NCCS Pune, Maharashtra, India. The cell lines were cultured in Dulbecco ‘s modified Eagle s medium (DMEM) with 2mM L-glutamine, 10% fetal calf serum (FCS), 1.5g/L sodium bicarbonate, 1mM non-essential amino acid and 1.0mM sodium pyruvate. Only the KBChR8-5 cell line was routinely maintained in 10ng/ml colchicines and subsequently grown in colchicine-free medium for one week prior to drug treatment. The cell lines were maintained in a humidified incubator with an atmosphere of 5% air and 5% CO₂ at 37°C.
Real-time PCR:-
Total cellular RNA from KB and KBChR8-5 cells were extracted by TRIzol Reagent (Chomvzynski et al., 1987). (Invitrogen, Carlsbad, CA, USA) and treated with RQ1 RNase-free DNase. Total RNA was reverse-transcribed to generate cDNA using RT-PCR kit (Promega Corporation, Madison, USA). The cDNA pool was subjected to QRT-PCR by using SYBR green PCR Master Mix on the AB 7500 fast real time PCR system (Applied Biosystems), Primers for MDR1 gene forward primer sequence CAGAGCAAGAGGCCATCCT and the reverse primer sequence TGAAGTCTCAAACATGAT and GAPDH were used as internal control,

Table 1:- The reaction mixture was prepared as follow.

<table>
<thead>
<tr>
<th>Component</th>
<th>Volume</th>
</tr>
</thead>
<tbody>
<tr>
<td>10x PCR buffer with MgCl₂</td>
<td>5 µl</td>
</tr>
<tr>
<td>10mM dNTP mix</td>
<td>1 µl</td>
</tr>
<tr>
<td>10 µM forward primer</td>
<td>5 µl</td>
</tr>
<tr>
<td>10 µM reverse primer</td>
<td>5 µl</td>
</tr>
<tr>
<td>cDNA</td>
<td>5 µl</td>
</tr>
<tr>
<td>Taq polymerase (5U/µl)</td>
<td>0.2 µl</td>
</tr>
<tr>
<td>Nuclease free water</td>
<td>28.8 µl</td>
</tr>
<tr>
<td>Final reaction volume</td>
<td>50 µl</td>
</tr>
</tbody>
</table>

Polymerase chain reaction (PCR):-
40 cycles of amplification were done and water was amplified for negative control. The following conditions were used in QPCR: 2 min at 50°C, 10 min at 95°C, and 40 cycles of 15 s at 95°C and 1 min at 60°C. Total 10 µL of each PCR product as well as DNA Ladder was electrophoresed in 1X Tris/acetate/EDTA (TAE) electrophoresis buffer on a 1% agarose gel at 100 volts for 20-30 minutes, visualized using UV transilluminator and photographed after staining with Ethidium bromide (Huang et al., 2006).

MTT assay procedure:-
The MTT [3-(4,5-dimethyl-2-thiazoly)-2, 5-diphenyl-2H-tetrazolium bromide] assay was used to determine drug sensitivity. (James et al., 1987). KB, KBChR8-5 and HeLa cells in the exponential growth phase were harvested by trypsinization and seeded into 96-well plate at a concentration of 3500 cells per well. The cells were treated in triplicate with gradient concentration of curcuminoids ranging from 0.01μg, 0.1μg, 1.0, 10 and 100 μg and incubated at 37°C for 24 hours. The IC50 value defined as the drug concentration required to reduce cell survival to 50% as determined by the relative absorbance of MTT. IC50 value was calculated by (mean absorbance in test wells) / (mean absorbance in control wells) x 100.

Results:-
Characterization of curcuminoids High - Performance Liquid Chromatography (HPLC):-
In our study the total extract contained 40.1% of curcuminoids were found in Assam variety turmeric among that 23.9% curcumin, 10% DMC 6.1% BDNC by HPLC analysis. The purity profile of isolated individual curcuminoids were analysed by HPLC. C, DMC, BDNC showed single peaks at retention times of 10.81, 12.79 and 13.03 min respectively. The identity of each peak was confirmed by determination of retention times and by spiking with standards. Purity profile for each curcuminoids were calculated as 99%, 98%, 95% purity for C, DMC, BDNC respectively.

GC-MS:-
The full scan mass spectra of each curcuminoids isolated were detected shows molecular weight of each individual compounds of C, DMC, BDNC was found to be 368.31, 337.77, and 307.95 respectively.

FT-IR analysis:-
The presence of absorption bands of specific stretching regions obtained at their respective frequency confirms the structure of compounds. IR absorption bands (cm⁻¹)

For C showed 3413.76(O-H str); 1629.14(conjugated C=C str); 1585.60 (-unsaturated C=O str); 1455.64 (aromatic ring str); 1288.65, 1028.68 (=C-O-CH₃ str); 1140.52 (C-OH str).
For **DMC** showed 3326.98 (O-H str); 1627.30 (conjugated C=C str); 1575.24 (-unsaturated C=O str); 1437.33 (aromatic ring str); 1263.07, 1025.21 (=CO-CH3 str); 1134.67 (C-OH str).

For **BDMC** showed 3230.06 (O-H str); 1629.40 (conjugated C=C str); 1575.64 (-unsaturated C=O str); 1508.90 (aromatic ring str); 1165.85 (C-OH str).

**NMR analysis:**

**13C NMR** chemical shift (ppm): Curcumin 183.6, 149.8, 148.4, 141.1, 126.8, 123.5, 121.5, 116.1, 111.8, 101.2, 56.1.

DMC- 183.7, 183.5, 160.27, 149.82, 148.47, 141.16, 140.83, 130.78, 126.82, 126.29, 123.61, 121.27, 116.39, 116.18, 111.77, 101.40, 56.1.

BDMC - 183.66, 160.26, 140.82, 130.78, 126.30, 121.25, 116.38 and 101.41. The presence of signal at 56.1 in C and DMC denotes the presence of methoxyl group while there is no methoxyl group present in BDMC.

**1H NMR** chemical shift (ppm): Curcumin 9.65, 7.57, 7.32, 7.16, 6.84, 6.77, 6.06, 3.84.

DMC- 10.03, 9.68, 7.57, 7.55, 7.33, 7.16, 7.14, 6.84, 6.78, 6.74, 6.04, 3.84.

BDMC- 10.04, 7.57, 7.53, 6.83, 6.71, 6.04. Signal corresponding to the methoxyl group at 3.84 is found in C and DMC and absent in BDMC.

This demonstrates crude curcuminoids has three major compounds curcumin, demethoxycurcumin, bisdemethoxycurcumin and that the purity of each curcuminoids is in the range of 95-99%.

**Real time PCR:**

Total RNA was extracted from KB and KBChR8-5 cells by TRIzol Reagent and converted into cDNA using cDNA synthesis kit. Quantitative Real-Time PCR was set up with Power SYBR Green Master-mix for MDR1 gene expression using gene specific primers. Expression of Genes was normalized to GAPDH and control cell line relative expression was normalized with respect to KB cDNA shown in (Fig. 1)

![Fig 1: HumanMDR1 gene expression analysis in KB and KBChR8-5 cells: MDR1 gene expression were measured and normalized to GAPDH expression, double distilled water (DDH₂O) was used as a negative control. KB cell in lane 1, KBChR8-5 cell in lane 2, the data are means ± SEM of three independent experiments, and significantly different from the vehicle control (p<0.05).](image-url)
MDR1 gene expression in KB and KBChR8-5 cells showed significant difference in expression level. KBChR8-5 cells showed greater expression with 7.1 fold increase from KB cells, whereas very mild expressions were detected in KB cells. The MDR1 gene expression normalized with GAPDH were analyzed by gel run, the band formed shown in (Fig. 2).

**Real time PCR analysis of MDR-1:-**

![Marker: 100 bp DNA ladder](image1)

Lane 1: GAPDH – 294 bp

Lane 2: MDR - 1 - 202 bp

*Fig 2:* Real-time PCR analysis of MDR-1 gene. PCR product was run in 1% agarose gel and 100 bp DNA ladder was used as marker.

**MTT assay:-**

The dose response curve and the effect of three curcuminoids on KB, KBChR8-5 cell lines and HeLa were determined by cytotoxicity assay. Dose response cytotoxicity profiles for three curcuminoids C, DMC, and BDMC were established for KB, KBChR8-5 cell lines and HeLa were shown in figure 2 A, B, C. The percent of viable cells was calculated to determine the IC\(_{50}\). The IC\(_{50}\) value of C, DMC, and BDMC in KB, KBChR8-5 and HeLa cell lines is described in table 2.

**Table 2:** Cytotoxicity of three curcuminoids in KB, KBChR8-5 and HeLa cell lines.

<table>
<thead>
<tr>
<th>Drugs</th>
<th>IC(_{50}) µM</th>
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<tbody>
<tr>
<td></td>
<td>KB</td>
</tr>
<tr>
<td>C</td>
<td>18.3</td>
</tr>
<tr>
<td>DMC</td>
<td>22.8</td>
</tr>
<tr>
<td>BDMC</td>
<td>7.8</td>
</tr>
</tbody>
</table>
Fig 3: Cytotoxic activity of Curcuminoids on KB, KBChR8-5 and HeLa cell lines

![Graph of KB Cell Line Cytotoxic Activity](image)

Fig. 3a: Percentage growth against KB cells

![Graph of KB-CHR-8-5 Cytotoxic Activity](image)

Figure 3b: Percentage growth against KBChR8-5 cells.
Figure 3c: Percentage growth against HeLa cells.

Fig. 3: Cytotoxicity of C, DMC, BDMC in KB (3a), KBChR8-5 (3b), HeLa (3c) cells studied at various concentrations of curcuminoids were added and incubated for 24 hours in 37°C. The experiments were conducted in triplicates and the values represent mean ± S.D of three independent experiments. Curcumin ♦, DMC ■, BDMC▲. The data of KB, KBChR8-5 and HeLa cells as mean ± S.D (n=3) shown in 3a, 3b, 3c respectively.

Discussion:
Cancer is a major worldwide public health problem. Treatment for cervical cancer using chemotherapeutic drugs can be possible in early cancer stages, but the major obstacle in treatment failure is due to development of drug resistance. The major impediment in cancer chemotherapy is multidrug resistance due to overexpression of ABC transporters. The overexpression of ABCB1 occurs in 40-50% of cancer patients and is associated with poor clinical outcome (Jamroziak et al., 2004).

The mechanism of drug resistance is characterized by a decrease in drug accumulation resulting from overexpression of energy-dependent multidrug efflux pump known as multidrug transporter (ABC transporter). The major mechanism responsible to MDR in cancer cells is the overexpression of MDR1 gene product resulting increased production of a 170,000 dalton p-glycoprotein in the plasma membrane associated with multiple drug resistance in drug resistant CHO cells (Ling and Thompson, 1974).

Overexpression of p-gp has been well established as the cause of the MDR phenotype in many in vitro selected drug resistant cell lines. KB cell lines have been used to catalog the changes in specific protein synthesis associated with the development of multidrug resistance results in reduced drug accumulation which might be caused by activating efflux or decreasing uptake of anticancer drugs. KB-V1 cells were selected by subjecting KB cells in a step-wise fashion to increasing concentration of vinblastine. KB-V1 cells have been shown to express only p-gp at a higher level on their plasma membrane. The level of p-gp in KB-V1 cell membrane is about 1% of total plasma membrane of drug resistant cell lines (Ambudkar et al., 1992).

Likewise colchicine resistance clone KBChR8-5 with multidrug resistance is derived from the KB cell line (Fojo et al., 1985). KBChR8-5 cells were selected by subjecting KB cell lines in a stepwise fashion to increase the concentration of colchicine (Akiyama et al., 1985). The amount of purified p-glycoprotein thus obtained accounted for approximately 3 to 4% of the total plasma membrane protein in colchicine-resistant mutant cell line (Riordan and Ling, 1979). As KB and its stepwise derivative KBChR8-5 cell lines have been characterized extensively with respect to the phenomenon of drug resistance proving overexpression of MDRI gene, so we decided to use KB and
KBChR8-5 cell lines to assess the effect of curcuminoids on the cytotoxic activity and expression of MDR 1 gene, to see whether introduction of curcuminoids reduced the drug resistance of the cells.

We analysed the presence of MDR1 gene in KB and KBChR8-5 cells by real time PCR. KBChR8-5 cells showed higher level of MDR1 gene expression at 7.1 fold increase in MDR1 when compared to KB cells, but KB cells showed very mild expression of MDR1 gene.

There are vast number of heterogeneous class of molecules present in the diet they are generally called as phytochemicals includes vitamins like carotenoids and food polyphenols like, phytoalexins, phenolic acids indoles, flavonoids (Sporn and Suh, 2002). More than 10,000 phytochemicals have been described and among them flavanoids include about 6000 compounds (Hairborne, 1993). These phytochemicals play major role in chemotherapy for cancer patients. They are present in the herbal constituents fruits, berries and spices, In spices are mint, rosemary, garlic, piper nigrum, curcumin, ginseng, onion (Bansal et al., 2009). The traditional uses of turmeric or natural curcuminoids in folk medicine are multiple, including antioxidant, anti-cancer, anti-inflammatory, anti-fungal, anti-parasitic, anti-venom, anti-mutagenic activity in vitro, (Peret-Almeida, 2005).

Recently it was reported that the effect of curcuminoids was examined on the proliferation of MCF-7 human breast tumor cells that demethoxycurcumin was the best inhibition of MCF-7 cells followed by curcumin and bisdemethoxycurcumin (Simon, 1998). Bisdemethoxycurcumin (BDMC) is active for modulation of MDR-1 gene expression (Anuchapreeda, 2004).

Curcuminoids are curcumin and its derivatives varies in structure and pharmacological activity. Therefore it is important to obtain curcuminoid pigments in high purity for detailed study of their biological properties. Our present study demonstrates isolation and separation of curcuminoids in pure form, thin layer chromatography revealed that the Rf value of curcuminoids shown in TLC were 0.75, 0.55, and 0.27, for C, DMC, BDMC respectively. Better resolution of Rf value showed that chloroform and methanol can be suitable solvent for the separation of compounds in column chromatography. Further purification results in curcumin as bright yellow needle shaped crystals, DMC as light yellow crystals, BDMC as reddish orange color crystals. The structure and purity of isolated curcuminoids were determined by GCMS, FTIR, and NMR. The purity of curcuminoids were analysed by HPLC from the percentage calculation using peak area and retention time, it was found to be > 95% purity for each compound.

The curcuminoids were found more toxic for cancer cells than normal cell line. In this regards, therapeutic index’ is an important parameter to select samples for developing drugs. The result of our experiment revealed in MTT assay to determine the relative cytotoxicity of curcuminoids in KB and KBChR8-5 cell lines and HeLa cell lines. Each curcuminoids have different IC_{50} values in the range of 5-25μM. There was a decrease in the viability of individual cell line with increasing concentration of curcuminoids treated. As HeLa cells are cancer cells that are used generally for cancer studies, the reaction with curcuminoids showed increased Cytotoxicity on curcumin treatment than DMC, BDMC in HeLa cells. The cytotoxic activity of curcumin may be due to inducing apoptosis by inhibiting proliferation of cancer cells by cell cycle arrest. Chearwae et al., studied cytotoxic activity of three curcuminoids on KB-V1 and its parental cell lines results in IC_{50} range 25-90μM cytotoxicity of each compound on both cell lines are nearly equal. Curcumin showed greater cytotoxic effect (Chearwae et al., 2004).

KB and KBChR8-5 cell lines are derived from HeLa cell lines shows increased production of MDR-1 gene, BDMC act on MDR cell lines more effectively than other curcuminoids. The cytotoxicity of curcuminoids against drug resistant cell lines showed that higher cytotoxicity at BDMC treatment on KB and KBChR8-5. The IC_{50} value of drugs on MDR subline were nearly equivalent to that of parental cell line. Individual curcuminoids vary in chemical structure and the knowledge of structure-activity relationship in combination with conventional chemotherapy is useful for drug design. our result implies that BDMC act on MDR cell lines more effectively than other curcuminoids which revealed that it may have synergistic effect that target the site related to multidrug resistance and helps in accumulation of anticancer drugs by blocking the efflux pump to increase the cytotoxicity and induce apoptosis. BDMC can be used as effective chemosensitizer combination with conventional chemotherapy for circumventing MDR in cervical cancer.
Conclusion:
In summary, curcuminoids can inhibit tumor growth in drug resistant cell lines. BDMC in curcuminoids can synergistically induce the down regulation of MDR-1 gene and increase accumulation of anticancer drugs to produce typical apoptotic morphological changes. BDMC can be used as effective chemosensitizer combination with conventional chemotherapy for circumventing MDR in cervical cancer. Novelty in chemical structure further suggests its importance in future MDR tumor.

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References:


