

Comparison of Inhibitory Effect of Curcumin Nanoparticles and Free Curcumin in Human Telomerase Reverse Transcriptase Gene Expression in Breast Cancer

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ABSTRACT

Purpose: Telomerase is expressed in most cancers, including breast cancer. Curcumin, a polyphenolic compound that obtained from the herb of *Curcuma longa*, has many anticancer effects. But, its effect is low due to poor water solubility. In order to improve its solubility and drug delivery, we have utilized a β -cyclodextrin-curcumin inclusion complex. **Methods:** To evaluate cytotoxic effects of cyclodextrin-curcumin and free curcumin, MTT assay was done. Cells were treated with equal concentration of cyclodextrin-curcumin and free curcumin. Telomerase gene expression level in two groups was compared by Real-time PCR. **Results:** MTT assay demonstrated that β -cyclodextrin-curcumin enhanced curcumin delivery in T47D breast cancer cells. The level of telomerase gene expression in cells treated with cyclodextrin-curcumin was lower than that of cells treated with free curcumin ($P=0.001$). **Conclusion:** Results are suggesting that cyclodextrin-curcumin complex can be more effective than free curcumin in inhibition of telomerase expression.

Introduction

Telomerase activity is observed in more than 85% in the most cancer cells.¹ Telomerase is active in 74% of breast Carcinomas.^{2,3} Therefore, targeting the telomerase in this cancer could be promising step in its treatment.⁴ For this purpose it is better to use natural compounds such as Curcumin (CUR). Curcumin (commonly known as turmeric) is obtained from *Curcuma longa*.⁵ Curcumin has many pharmacological applications such as anticancer, with low or no intrinsic toxicity.⁶ Despite all these, curcumin suffers from low water solubility and bioavailability.⁷ To improve its solubility, β -Cyclodextrin was used for encapsulation of curcumin. β -Cyclodextrin is a semi-natural compound with low toxicity, which could enhance drug bioavailability.⁷ To the best of our knowledge, comparison of inhibitory effect of free CUR and CD-CUR on hTERT gene expression in T47D cell line has never been done so far. Anti-telomerase effect of curcumin has been studied previously in lung cancer cell line^{8,9} and preparation of β -cyclodextrin-curcumin inclusion complex for improvement of curcumin stability and solubility has been studied.⁷ In our previous study we showed the inhibitory effect of β -Cyclodextrin-curcumin in telomerase gene expression in breast cancer cell line.¹⁰ Rajeswari *et al* reported that cyclodextrin enhance the bioavailability of insoluble drugs by increasing the drug solubility.¹¹ Yadav and

coworkers showed that Cyclodextrin-complexed curcumin had superior attributes compared with free curcumin for cellular uptake.¹² Murali *et al* found that CD-CUR inhibit the growth of prostate cells higher than free curcumin.⁹ Therefore in this study we investigated the effect of β -Cyclodextrin-curcumin (CD-CUR) and free CUR (CUR) on human telomerase reverse transcriptase (hTERT) gene expression. Anticancer effect of free CUR and CD-CUR in breast cancer cell line, T47-D was compared. The level of telomerase gene expression after 24 h exposure was measured by Real-time PCR.

Materials and Methods

Cell culture and cell line

T47D cell line (breast cancer epithelial like cell line) was prepared from Pasteur Institute cell bank of Iran, code: C203. This cell line was cultured in RPMI1640 (Gibco, Invitrogen, UK) supplemented with 10% heat-inactivated fetal bovine serum (FBS) (Gibco, Invitrogen, UK), 2 mg/ml sodium bicarbonate, 0.05 mg/ml penicillin G (Serva co, Germany), 0.08 mg/ml streptomycin (Merck co, Germany) and incubated in 37°C with humidified air containing 5% CO₂.

In vitro cytotoxicity (MTT assay)

Cells in the exponential phase of growth were exposed to CD-CUR inclusion complex or free CUR. Cytotoxic

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effect of CD-CUR inclusion complex and free CUR was studied by 24, 48 and 72 h MTT assay. 2×10^3 cell/well was plated in a 96-well plate (Coastar from Corning, NY). After 24 h incubation, cells were treated with different concentrations (5-100 μ M) of CD-CUR and free CUR for 24, 48 and 72 h in the quadruplicate manner. Media containing equivalent amounts of CD in PBS or DMSO was used as control. After different exposure duration, medium was removed and then fed of the cells with 200 μ L of fresh medium. Cells were incubated for 24 h, then 50 μ L of 2 mg/ml MTT (Sigma co, Germany) dissolved in PBS was added to each well and plates were covered with aluminum foil and incubated for 4 h. In the next step, wells' content was removed and 200 μ L pure DMSO and 25 μ L Sorensen's glycine buffer were added. Finally amount of formazan was determined by measuring the absorbance at 570 nm using an ELISA plate reader (with a reference wavelength of 630 nm).

Cell treatment

After determining of IC_{50} , 2.5×10^5 cells in 25 cm^2 flasks were treated with 3 concentrations lower than IC_{50} of 24h CD-CUR (5, 10 and 15 μ M). Culture flasks were incubated for 24 h. For control cells, the same volume of 10% DMSO without CUR and CD-CUR was added to flask of control cells. An equivalent amount of CD in PBS was used as another control. Culture flasks were incubated in 37°C containing 5% CO₂ with humidified atmosphere incubator for 24 h exposure duration.

Real-Time PCR (qRT-PCR) Assay

After the RNA extraction (by the TRIZOL kit, Cinnagene, Iran), cDNA synthesized according the instructions (First Strand cDNA Synthesis Kit fermentase, K1622).

For real-time PCR, according to our previous study¹⁰, hTERT primers (Genbank accession: NM_198255, bp 2165-2362) and beta actin primers (Genbank accession: NM_001101, bp 787-917) were used. For hTERT, a 198 bp amplicon and for beta actin a 131 bp amplicon were generated in a 25 μ l reaction mixture that contained: 5 pmole of the forward and reverse PCR primers of hTERT (5'CCGCCTGAGCTGTACTTTGT3', 5' CAGGTGAGCCACGAAGTGT3' respectively) or for beta actin (5'TCCCTGGAGAAGAGCTACG3', 5'GTAGTTTCGTGGATGCCACA3' respectively), 2X PCR Master Mix Syber Green I, and 2 μ l of the cDNA was used. Each DNA sample was divided so that hTERT and beta actin could be amplified, in parallel, and in duplicated from equal amounts of starting cDNA separately. 25 μ l reactions contained the following final concentrations: 1X of MaximaTM SYBR Green/ROX qPCR Master Mix (including MaximaTM Hot start Taq Polymerase, MaximaTM SYBR Green qPCR Buffer, SYBR Green I and ROX Passive reference dye), 5pmole of each primer and 2 μ l of the cDNA. Negative controls were

prepared each time, consisting of 2 μ l ddH₂O instead of the cDNA template. The sample tubes were placed into the (Rotor-Gene 6000, Corbet) with the following settings as manufacture protocol.

Results and Discussion

Cell toxicity studies

In this study to evaluate the cytotoxic effect (MTT assay) of CD-CUR and free CUR, T47D breast cancer cell line were treated with different concentration (5-100 μ M) of CD-CUR and free CUR for 24, 48 and 72h. IC_{50} after 24 h treatment with CD-CUR and CUR was 18 μ M and 22 μ M respectively (Figure 1). Drug free CD as well as DMSO 2% showed an absorbance value equivalent of 90 and 92% of control respectively suggesting that CD and DMSO 2% have low effect on the cells. Cells treated with concentrations more than 60 μ M of CD-CUR and free CUR for 48 and 72h were died completely. IC_{50} after 48 h treatment with CD-CUR and CUR was 13 and 17 μ M respectively (Figure 2 and 3). Considering the fact that IC_{50} values for different treatment durations are not similar, it might be concluded that the effect of CD-CUR and CUR on T47D cell line is time-dependent. Moreover IC_{50} show that effect of CD-CUR on cells is more than free CUR, Demonstrating the enhanced uptake of CD-CUR with respect to CUR.

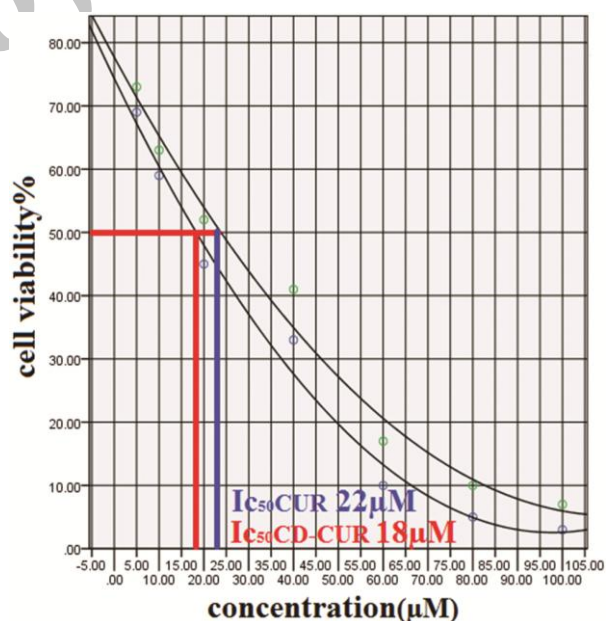


Figure 1. MTT assay graph of 24 h cell treatment with CD-CUR and CUR.

Real-time PCR

Figure 4 and 5 demonstrate the results of telomerase gene expression study at T47D breast cancer cell line after 24 h of CD-CUR and free CUR exposure. β -Cyclodextrin-curcumin suppresses cell proliferation in T47D breast cancer cells. Cells were treated with curcumin (CUR) or CD-CUR for 24 h. Cell proliferation was determined by MTT assay and

normalized to cells treated with equivalent amounts of respective controls (DMSO for curcumin and cyclodextrin for CD-CUR). The level of hTERT mRNA was normalized to mRNA levels of the housekeeping gene, Beta actin, within each sample. The differences of $2^{-\Delta\Delta Ct}$ values were calculated. Increasing $2^{-\Delta\Delta Ct}$ amount resulted in enhanced expression of mRNA levels. Data analyses of real-time PCR showed that with increasing the concentration of CD-CUR and free CUR, a decreasing trend is observed in mRNA levels of hTERT. Compared to CUR, in the same concentration, CD-CUR resulted in a lower mRNA level of hTERT and lower level of hTERT mRNA expression (Figure 5). The difference between two groups was statistically significant ($p=0.001$, $n=4$) (Figure 6).

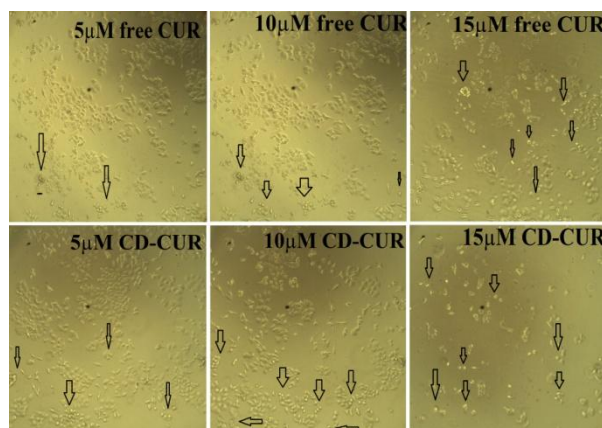


Figure 4. β -Cyclodextrin-curcumin and free curcumin treatment cells (Images were taken by Phase contrast microscope).

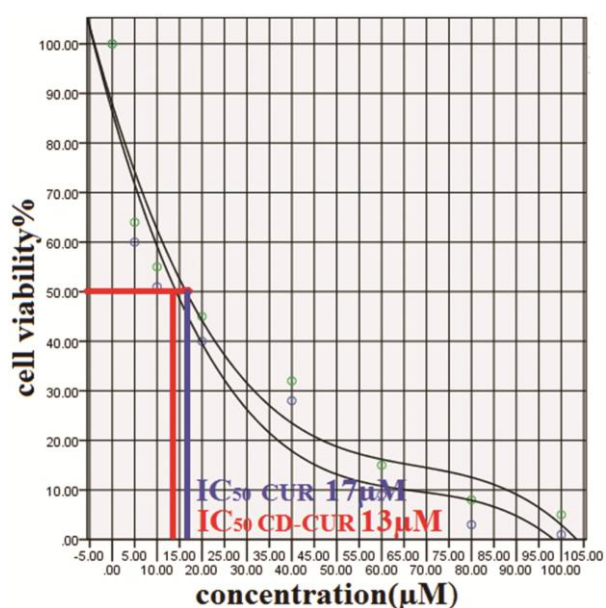


Figure 2. MTT assay graph of 48 h cell treatment with CD-CUR and CUR.

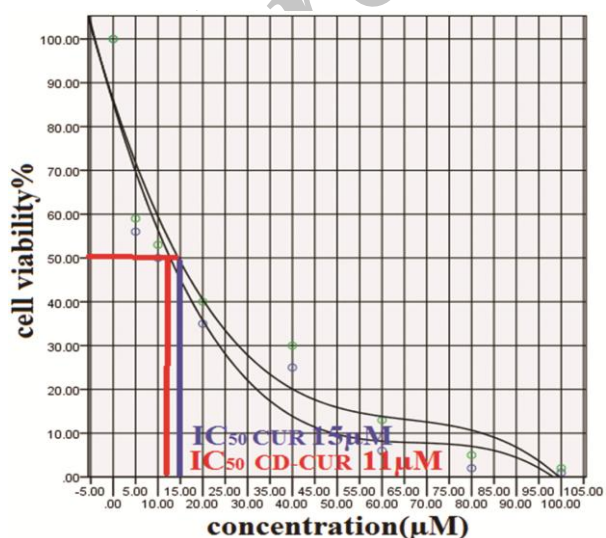


Figure 3. MTT assay graph of 72 h cell treatment with CD-CUR and CUR.

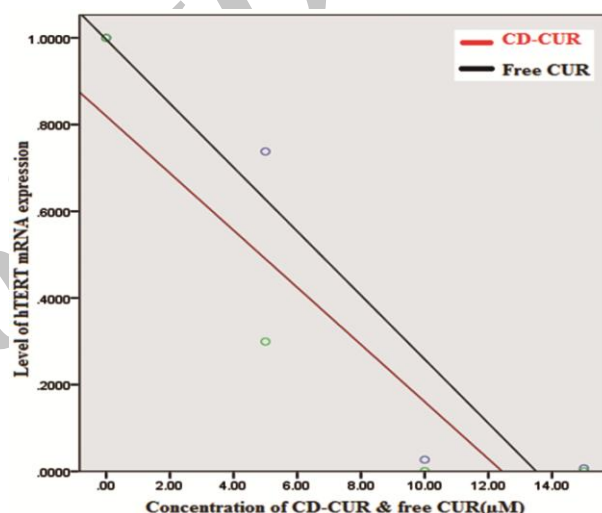


Figure 5. Level of hTERT mRNA expression in cells treated with CD-CUR or free CUR.

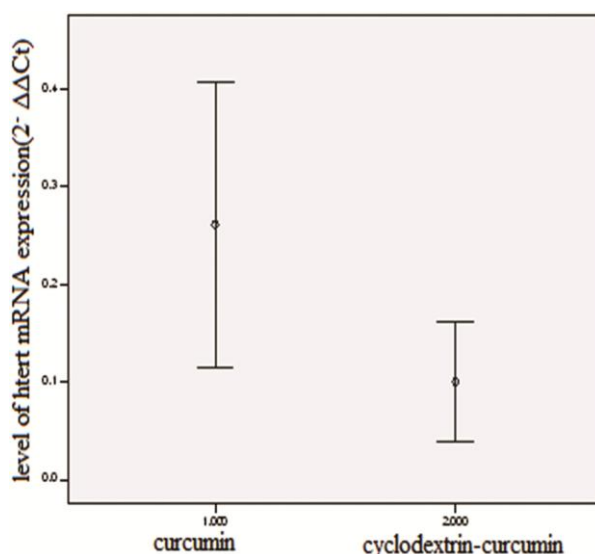


Figure 6. comparison of hTERT mRNA expression in curcumin as well as cyclodextrin-curcumin treated cells. The difference between two groups was statistically significant ($p=0.001$, $n=4$).

Conclusion

Our results showed that CD-CUR and free CUR had inhibitory effect on T47D breast cancer cell line. This inhibition was time and dose dependent. β -cyclodextrin-curcumin resulted in higher cell toxicity than free curcumin in breast cancer cell line. There was a significant difference between cells treated with CD-CUR and cells treated with free CUR in the levels of telomerase gene expression. The quantity of telomerase was decreased in the cells treated with CD-CUR in comparison to cells treated with free CUR ($P=0.001$). It may be concluded that relative to free CUR, CD-CUR inclusion complex inhibit the telomerase gene expression in T47D cell line more effectively.

Acknowledgements

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Conflict of interest

There is no conflict of interest in this study.

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