Curcumin inhibits the metastasis of K1 papillary thyroid cancer cells via modulating E-cadherin and matrix metalloproteinase-9 expression

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Abstract The anti-metastatic effect of curcumin on papillary thyroid cancer K1 cells and its underlying mechanisms were investigated. Curcumin at 12.5, 25 and 50 μM promoted mesenchymal–epithelial transition and decreased the migration rate of K1 cells by 24–87 %. Its mechanism may involve the up-regulation of E-cadherin expression levels and down-regulation of the activity and expression of matrix metalloproteinase-9.

Keywords Curcumin · E-cadherin · Matrix metalloproteinases · Metastasis · Thyroid cancer

Introduction

Papillary thyroid carcinoma (PTC) is the most frequently occurring form of differentiated thyroid cancer. Although the prognosis for PTC using traditional therapies is usually good, PTC can undergo de-differentiation and progression to become an anaplastic carcinoma, a formidable disease characterised by widespread invasion and early distant metastasis (Zuo et al. 2010). The loss of intercellular adhesion and acquisition of degradative properties are prerequisites for tumour cells to become fully invasive (Nawrocki-Raby et al. 2003), which is the beginning of metastasis. As a result, there is significant interest in developing an agent with the potential to restrain both of these prerequisites and thus inhibit metastasis.

An accumulating body of data supports the concept that phytochemicals in the human diet are devoid of toxicity and may have numerous long-lasting beneficial effects on human health (Shankar and Srivastava 2012). Considering its unique anti-metastatic property and low toxicity, curcumin (diferuloylmethane), the chief component of the spice turmeric and derived from the rhizome of the East Indian plant Curcuma longa, is one such potential candidate. Curcumin has been demonstrated to affect a number of the cellular adhesion molecules that are involved in the processes of metastasis (Ray et al. 2003), such as E-cadherin. E-cadherin expression is important in maintaining the integrity of intercellular adhesion through binding to various catenins (including β-catenin), and the loss of E-cadherin is associated with an increased tendency for tumour metastasis (Onder et al. 2008). In addition, the degradation of the basement membrane and extracellular matrix occurs primarily through matrix metalloproteinases (MMPs). Of these proteases, the...
expression of MMP-9 and 2 is high in various malignant tumours, and is closely related to the ability of these cells to invade and metastasise (Weng and Yen 2012). Many reports have shown that curcumin inhibits the expression of MMP-9 both in vitro and in vivo (Kunnumakkara et al. 2008).

As curcumin has an ill-defined effect on the metastasis of PTC and related mechanisms, in this paper we show how it affects the epithelial–mesenchymal transition (EMT) and cell migration involved in cell metastasis. We further discuss its impact on E-cadherin and MMPs.

Materials and methods

Reagents

Curcumin (purity ≥94 %), dimethyl sulfoxide, concanavalin A, propidium iodide and Hoechst 33342 were from Sigma. Dulbecco’s Modified Eagle Medium (DMEM) and foetal bovine serum (FBS) were purchased from Gibco (Logan, UT, USA) and Sijiqing Biological Engineering Materials Co Ltd. (Hangzhou, China), respectively. Bovine serum albumin (BSA) was purchased from Sangon (Shanghai, China). All other chemicals were analytical reagents, and purchased from Sinopharm Chemical Reagent Co Ltd. (Shanghai). Curcumin was dissolved in DMSO at 10 mg/ml, and stored at −20 °C until dilution before use.

Cell line and culture conditions

Papillary thyroid cancer cell line K1 was obtained from the European Collection of Cell Cultures and maintained in DMEM containing 10 % (v/v) FBS, 100 U penicillin/ml and 100 U streptomycin/ml in a humid atmosphere of 5 % (v/v) CO₂ and 95 % (v/v) air at 37 °C. K1 cells in the log phase were plated in cell culture plates (Corning, NY, USA). Control cells were treated with the same medium, but without curcumin or solvent. The solvent control contained an equivalent amount of DMSO corresponding to the highest used concentration of curcumin.

Cell viability assay

Cell viability was determined by propidium iodide and Hoechst 33342 staining (see Song et al. 2012).

Cell spreading assay

Ninety-six-well culture plates were coated with concanavalin A (Con A; 25 µg/ml) in phosphate-buffered saline (PBS) overnight at 4 °C, blocked with 0.2 % BSA for 2 h at room temperature and then washed twice with PBS. The cells were harvested with trypsin/
EDTA, washed twice and resuspended in DMEM at $8 \times 10^4$ cells/ml. Then, to 100 µl cells different concentrations of curcumin (12.5, 25 and 50 µM) were added. The plates were incubated for another 2 h at 37 °C, and then photographed.

Migration assay

The transwell migration assay was performed as described previously (Rajoria et al. 2011) with some modification.

Gelatin zymography

Zymography assay was performed as described previously (Forsyth et al. 1999) with some modification.

Western blot assay

K1 cells pretreated with curcumin (12.5, 25 or 50 µM) were incubated for 24 h. A western blot was then performed (see Song et al. 2012). The antibodies were anti-β-actin (1:1,000, v/v), anti-E-cadherin (1:1,000, v/v) or anti-MMP-9 (1:100, v/v) (Santa Cruz Biotechnology, CA, USA) and an appropriate horseradish peroxidase-conjugated secondary antibody (goat anti-mouse IgG, 1:500, 1:500 or 1:250, respectively) (Santa Cruz Biotechnology).

Statistical analysis

The results were expressed as mean ± standard errors (SE), and three group comparisons were evaluated using the Student’s t test. Differences were considered statistically significant at $P < 0.05$.

Results

Curcumin inhibits the cell viability of K1 cells

Figure 1a shows that the number of red-emitting fluorescent cells increased as the curcumin concentration increased. Compared to the control groups, treatment with curcumin at 12.5 and 25 µM increased the cell death rate to 2.6 ± 0.02 % ($P = 4.96 \times 10^{-1}$) and 5.5 ± 0.05 % ($P = 2.33 \times 10^{-1}$), respectively, whilst at 50 µM, it increased the death rate to 21.6 ± 0.04 % ($P = 6.17 \times 10^{-3} < 0.01$). Figure 1c shows that 50 µM curcumin also inhibited cell viability in a time-dependent manner after 24 h of treatment.

Curcumin inhibits K1 cell spreading

As illustrated in Fig. 2a, all cells treated with solvent alone spread out with large and flat lamellipodia. 12.5 µM curcumin had a modest effect on the

![Fig. 2](Effects of curcumin on K1 cell spreading. Cells with or without different concentrations of curcumin (12.5, 25 and 50 µM) were added to plates pre-coated with Con A (25 µg/ml), incubated for an additional 2 h at 37 °C and then photographed with a microscope. Arrows indicate non-spreading cells. SC solvent control.)
morphology of cell spreading (Fig. 2b). However, curcumin 25 μM partially inhibited cell spreading on Con A, as can be seen from the arrows in Fig. 2c. Moreover, in Fig. 2d, the cells pre-treated with 50 μM curcumin were small and round.

Curcumin inhibits K1 cell migration

Figure 3A, B shows that curcumin significantly inhibited cell migration in a dose-dependent manner in this study. After cells had been treated with 12.5, 25 or 50 μM curcumin, the migration rate of the K1 cells through the membrane decreased to 87.4 ± 12 % ($P = 3.55e-1$), 45 ± 1.3 % ($P = 1.68e-6 < 0.01$), 24.2 ± 5.8 % ($P = 2e-4 < 0.01$), respectively.

Curcumin up-regulates the expression of E-cadherin

Figure 4 shows that curcumin increased the expression of E-cadherin in a dose-dependent manner. In particular, the protein expression level of E-cadherin increased significantly when the cells were treated with 50 μM curcumin.

Curcumin down-regulates the activity and expression of MMP-9

Figure 5a shows that the activity of MMP-9 was notably inhibited by curcumin in a dose-dependent manner compared with the controls. In addition, we tested protein expression of MMP-9 through a western blot assay and found that the cytoplasmic expression level of MMP-9 was also down-regulated by curcumin (Fig. 5b). More importantly, at curcumin up to 50 μM, almost no obvious MMP-9 expression was observed.

Fig. 3 Effects of curcumin on cell migration. Cancer cell migration ability was examined using a transwell system. After being serum-starved for 24 h, cells with or without curcumin (12.5, 25 and 50 μM) were seeded onto the upper wells of transwells, while a conditional medium (DMEM with 10 % FBS) was added to the lower wells. Cells were incubated at 37 °C for 3 h. A Morphology of cells stained with Hoechst 33342 and inspected with a fluorescence microscope. B Quantification of cells attached to the lower surface of the polycarbonate filter. All data are represented as the mean ± SE of three independent experiments. SC solvent control. *$P < 0.05$ versus SC. **$P < 0.01$ versus SC (Student’s two tailed t test)

Fig. 4 Effects of curcumin on expression of E-cadherin. Western blot analysis of the expression level of E-cadherin of cells pretreated with or without curcumin (12.5, 25, 50 μM) for 24 h. Beta-actin was used as a loading control. All data are represented as the mean ± SE of three independent experiments. SC solvent control. *$P < 0.05$ versus control. **$P < 0.01$ versus control (Student’s two tailed t test)
Discussion

Over the past several years, concerted efforts have been made to define the pathways involved in invasion and metastasis, and it has been suggested that the reactivation of embryonic pathways could promote cell movement and induce epithelial to mesenchymal transition and collective cell migration (Vasko and Saji 2007). In a previous study, we found that curcumin could induce apoptosis in K1 papillary thyroid cancer cells (Song et al. 2012). In this paper, we further discuss how it affects EMT and the cell migration involved in cell metastasis.

We found treatment with curcumin induced dramatic morphological changes in K1 cells, which altered from a spindle-like, mesenchymal phenotype to become a more pronounced epithelial phenotype (Fig. 2). Moreover, the results of cell migration using the transwell assay showed curcumin to have a unique anti-migration property, as evidenced by the very high inhibition rate on K1 cell migration of 25–50 µM for 24 h. An equal amount of total proteins was loaded into each lane. All data are represented as the mean ± SE of three independent experiments. SC solvent control. *P < 0.05 versus control. **P < 0.01 versus control (Student’s two tailed t test)

Conclusion

We provide the first evidence of curcumin’s ability to promote mesenchymal–epithelial transition and decrease tumour cell migration that is involved in the invasion and metastasis of papillary thyroid cancer. Its anti-metastatic mechanism may be involved in up-regulating E-cadherin expression levels and down-regulating the activity and expression of MMP-9. These data suggest that curcumin is a potential preventive agent for the adjuvant treatment of thyroid cancers.

Acknowledgments This study was supported by grants from the National Natural Science Foundation of China (Nos. 31071491 and 31171627), Program for New Century Excellent Talents in University (NCET-09-0436), Natural Science Foundation of Jiangsu Province (No. BK2010156) and “Twelfth 5-year” National Key Technology R&D Program of China (2012BAD33B05).
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