Effects of curcumin on growth of human cervical cancer xenograft in nude mice and underlying mechanism

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Abstract
The present study investigated the effects of curcumin (Cur) on growth of human cervical cancer xenograft in nude mice and underlying mechanism. The nude mice modeled with human cervical cancer HeLa cell xenograft were treated with normal saline (control), 3 mg/kg Cisplatin, 50, 100 and 200 mg/kg Cur, respectively. The animal body weight and growth of tumor were measured. The expressions of Bax, Bcl-2, p53, p21, HIF-1α, VEGF and MIF protein in tumor tissue were determined. Results showed that, after treatment for 20 days, the tumor mass and tumor volume in 100 and 200 mg/kg Cur group were significantly lower than control group (P < 0.05). The expressions of Bax, p53 and p21 protein in tumor tissue in 200 mg/kg Cur group were significantly higher than control group (P < 0.05), and the expressions of Bcl-2, HIF-1α, VEGF and MIF protein in tumor tissue in 200 mg/kg Cur group were significantly lower than control group (P < 0.05). Cur can inhibit the growth of HeLa cell xenograft in nude mice. The possible mechanism may be related to its up-regulation of Bax, p53 and p21 protein expression in tumor tissue, and down-regulation of Bcl-2, HIF-1α, VEGF and MIF protein expression.

Keywords: curcumin; cervical cancer; xenograft; nude mice.

Practical Application: Curcumin may be applied to treatment of human cervical cancer.

1 Introduction
Curcumin (Cur) is a polyphenol extracted from the root of turmeric plants. It is used not only in food and condiment in daily life, but also as the anti-inflammatory (Motterlini et al., 2000), anticoagulant (Kim et al., 2012), anti-atherosclerosis (Olszanecki et al., 2005) medicine. In addition, Cur can be applied to the treatment of cancer (Anand et al., 2008), but the mechanism is complex. It is generally believed that, the anti-tumor mechanism of Cur is related to its induction of tumor cell apoptosis and inhibition of tumor cell growth and metastasis (Cheng et al., 2013). In addition, Cur has the function of reversing the drug-resistance of tumors (Li et al., 2011). Cervical cancer is one of the most common malignant tumors in women. The incidence of cervical cancer only seconds to that of the breast cancer (Adams et al., 2009). The chemotherapy is the main method for the treatment of advanced cervical cancer. However, as the currently commonly used chemotherapeutic agents cannot specifically act with the tumor cells, the therapy will result in a lot of side effects, such as myelosuppression, alopecia, nerve damage, gastrointestinal reactions, kidney toxicity, etc., which greatly limit its clinical application (Chung et al., 2007; Kodama et al., 2001; Ke et al., 2012). This study established the nude mice xenograft model of human cervical cancer HeLa cells, and investigated the effects of Cur on the tumor growth and the related mechanism. The objective was to provide the theoretical and experimental basis for the clinical application of Cur.

2 Materials and methods
2.1 Materials
Human cervical cancer HeLa4 cells were provided by the Institute of Oncology, Guangdong Medical University (Guangzhou, China). Cur (purity 98.5%) was purchased from Sigma-Aldrich Corp. (MO, USA). Cisplatin (DDP) injection was provided by Jiangsu Hansoh Pharmaceutical Co., Ltd. (Lianyungang, China). BALB/c nude mice (SPF grade, female, 4-5 weeks old, 16-20 g) were provided by Guangdong Medical Experimental Animal Center (Foshan, China). The nude mice were single-cage raised in the condition avoiding strong light and noise (temperature 21 ± 2 °C; relative humidity 30-70%; 12/12-h day-night cycle; free to feed and water). The western blot kits were provided by Shanghai Sangon Biological Engineering Technology And Service Co., Ltd. (Shanghai, China). Other reagents were provided by Sigma-Aldrich Corp. (MO, USA). Electronic balance (FA2004) was provided by Shanghai Shangping Scientific Instrument Co., Ltd. (Shanghai, China). Vernier caliper (8012) was purchased from Shanghai SANTO Hardware Tools Co., Ltd. (Shanghai, China).

2.2 Establishment of nude mice xenograft model of HeLa cells
HeLa cells in logarithmic growth phase were taken, and the single-cell suspension (3 × 10⁶ cells/L) was prepared. The right armpit was nude mice was disinfected using iodine, and then
0.2 ml HeLa single-cell suspension was subcutaneously inoculated at the right armpit. The growth of xenograft tumor was observed every day, and the diameter was measured. When the diameter of all xenograft tumors was more than 2 mm, the model was considered successfully established.

2.3 Animal grouping and treatment

After establishment of HeLa cell xenograft model, 40 nude mice were randomly divided into control, DDP, 50 mg/kg Cur, 100 mg/kg Cur and 200 mg/kg Cur group, 8 nude mice in each group. The nude mice in later 4 groups were intraperitoneally injected with 3 mg/kg DDP, and 50, 100 and 200 mg/kg Cur, respectively. The nude mice in control group were intraperitoneally injected with equal volume of normal saline. The administration was performed once per two days, and was last for 20 days. During the administration, the mind state, response, activity and diet were observed.

2.4 Determination of nude mice body weight and growth of tumor

At the treatment beginning (day 0) and on the day 5, 10, 15 and 20 after treatment beginning, the body weight of nude mice was measured using electronic balance. On the day 20, all nude mice were executed by cervical dislocation, and the tumor block was stripped, the maximal diameter of the tumor (a) and the short diameter (b) were measured using vernier caliper. Then, the tumor was weighted using electronic balance. The tumor volume was calculated as follows: tumor volume = (a × b²)/2. The tumor inhibition rate was calculated as follows: inhibition rate (%) = (1 - tumor mass in treatment group/tumor mass in control group) × 100%.

2.5 Determination of expressions of apoptosis-related proteins, hypoxia inducible factor-1α, vascular endothelial growth factor and macrophage migration inhibitory factor in tumor tissue

The tumor tissue was homogenized, and the protein was extracted. The expressions of apoptosis-related proteins including B-cell lymphoma-2 (Bcl-2), Bcl-2 associated X protein (Bax), p53 and p21, hypoxia inducible factor-1α (HIF-1α), vascular endothelial growth factor (VEGF) and macrophage migration inhibitory factor (MIF) in tumor tissue were determined using western blot assays.

2.6 Statistical analysis

All statistical analysis was carried out using SPSS13.0 software (SPSS Inc., Chicago, IL, USA). The data were presented as mean±SD. The difference between two groups was analyzed using t test. P < 0.05 was considered as statistically significant.

### Table 1. Effects of Cur on growth of tumor.

<table>
<thead>
<tr>
<th>Group</th>
<th>Tumor mass (g)</th>
<th>Tumor volume (cm³)</th>
<th>Inhibition rate (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>1.8 ± 0.2</td>
<td>1.0 ± 0.1</td>
<td>0.0</td>
</tr>
<tr>
<td>DDP</td>
<td>1.0 ± 0.2ab</td>
<td>0.6 ± 0.1ab</td>
<td>42.9</td>
</tr>
<tr>
<td>50 mg/kg Cur</td>
<td>1.7 ± 0.3</td>
<td>1.0 ± 0.2</td>
<td>1.7</td>
</tr>
<tr>
<td>100 mg/kg Cur</td>
<td>1.2 ± 0.3ab</td>
<td>0.7 ± 0.1ab</td>
<td>31.1</td>
</tr>
<tr>
<td>200 mg/kg Cur</td>
<td>1.1 ± 0.2ab</td>
<td>0.6 ± 0.1ab</td>
<td>39.6</td>
</tr>
</tbody>
</table>

*abP <0.05 compared with control group; P < 0.05 compared with 50 mg/kg Cur group. DDP: cisplatin; Cur: curcumin.

3 Results

3.1 Effects of Cur on body weight of nude mice

During the experimental period, the nude mice in DDP group presented decreased activity, feeding and drinking. The nude mice in other 4 groups presented good general condition, with no occurrence of obvious adverse reaction. Before treatment (day 0) and on day 5 after treatment beginning, there was no significant difference of body weight of nude mice among 5 groups (P > 0.05). On the day 10, 15 and 20, the body weight in DDP group was significantly decreased, compared with day 0 and day 5, respectively (P < 0.01). In addition, the body weight in DDP group was significantly lower than each other group at day 10, 15 and 20, respectively (P < 0.01) (Figure 1).

3.2 Effects of Cur on growth of tumor

After treatment for 20 days, the tumor mass and tumor volume in DDP, 100 mg/kg Cur and 200 mg/kg Cur group were significantly lower than those in control group and 50 mg/kg Cur group, respectively (P < 0.05). In addition, there was no significant difference of tumor mass or tumor volume among DDP, 100 mg/kg Cur and 200 mg/kg Cur group, respectively (P > 0.05). The tumor inhibition rate in DDP, 50 mg/kg Cur, 100 mg/kg Cur and 200 mg/kg Cur group were 42.9%, 1.7%, 31.1% and 39.6%, respectively (Table 1).

3.3 Effects of Cur on expression of apoptosis-related proteins in tumor tissue

As shown in Figure 2, the relative expression amounts of Bax, p53 and p21 protein in DDP group were significantly higher than that in control group, respectively (P < 0.05), and...
the expression of Bcl-2 protein in DDP group was significantly lower than that in control group (P < 0.05). The expression of Bax protein in 200 mg/kg Cur group was significantly higher than control group (P < 0.05). The expression of Bcl-2 protein in 100 mg/kg Cur and 200 mg/kg Cur group was significantly lower than control group, respectively (P < 0.05). The expression of p53 protein in 200 mg/kg Cur group and the expressions of p21 protein in 50 mg/kg Cur group and 100 mg/kg Cur group were significantly higher than control group, respectively (P < 0.05).

3.4 Effects of Cur on expression of HIF-1α and VEGF protein in tumor tissue

Figure 3 showed that, the relative expression amounts of HIF-1α and VEGF protein in DDP, 100 mg/kg Cur and 200 mg/kg Cur group were significantly decreased, respectively, compared with control group (P < 0.05). In addition, the expression of VEGF protein in 100 mg/kg Cur group and 200 mg/kg Cur group were significantly lower than that in 50 mg/kg Cur group (P < 0.05).

3.5 Effects of Cur on expression of MIF protein in tumor tissue

The relative expression amounts of MIF protein in DDP, 50 mg/kg Cur, 100 mg/kg Cur and 200 mg/kg Cur group were significantly decreased, respectively, compared with control group (P < 0.05). There was no significant difference among DDP, 50 mg/kg Cur, 100 mg/kg Cur and 200 mg/kg Cur group (P > 0.05) (Figure 4).

Figure 2. Effects of Cur on expression of apoptosis-related proteins in tumor tissue (times to control group). aP <0.05 compared with control group; bP <0.05 compared with 50 mg/kg Cur group. DDP, cisplatin; Cur, curcumin; Bax, Bcl-2 associated X protein; Bcl-2, B-cell lymphoma-2.

Figure 3. Effects of Cur on expression of HIF-1α and VEGF protein in tumor tissue (times to control group). aP <0.05 compared with control group; bP <0.05 compared with 50 mg/kg Cur group. DDP, cisplatin; Cur, curcumin; HIF-1α, hypoxia inducible factor-1α; VEGF, vascular endothelial growth factor.
Bcl-2 gene is the specific survival gene which inhibits the cell apoptosis. It plays an important role in the regulation of cell apoptosis. It can resist various forms of cell death, and prolong the life-span of cells, leading to the increase of cell number (Kim et al., 2006). Bax gene is the apoptosis-promotion gene. Bax gene and Bcl-2 gene belong to the same gene family. Bax can not only inhibit the apoptosis inhibition effect of Bcl-2, but also directly promote the cell apoptosis (Findlay et al., 1997). Results of this study showed that, the expression of Bax protein in 200 mg/kg Cur group was significantly higher than that in control group (P < 0.05), and the expression of Bcl-2 protein in 100 mg/kg Cur and 200 mg/kg Cur group was significantly lower than control group, respectively (P < 0.05). This indicates that, Cur can up-regulate the expression of Bax protein and down-regulate the expression of Bcl-2 protein in HeLa cell xenograft in nude mice, which may be related to its inhibitory effects on tumor growth.

p53 is a transcription factor composed of 393 amino acids. It interacts with other intracellular proteins, and is involved in cell cycle control, gene regulation, cell differentiation, apoptosis and tumor suppression (Takahashi, 1997). As a direct downstream product of p53, p21 plays an important role in cell cycle regulation by binding and inhibiting cyclin dependent kinase. In addition, p21 can inhibit the mediated and non-mediated cell apoptosis. In some specific cases, it can promote cell apoptosis (Bao et al., 2006). Results of this study showed that, the expression of p53 protein in 200 mg/kg Cur group and the expressions of p21 protein in 50 mg/kg Cur group and 100 mg/kg Cur group were significantly higher than control group, respectively (P < 0.05). This indicates that, the inhibitory effects of Cur on tumor growth may be related to its up-regulation on the expression of p53 and p21 protein in tumor.

HIF-1 is composed of α and β subunits. HIF-1α is the basic expression protein, and HIF-1α is an oxygen-regulated protein. It is found that, HIF-1α is closely related to the occurrence and development of many kinds of tumors, and it is highly expressed in prostate cancer, colon cancer, cervical cancer, lung cancer and so on (Hu et al., 2014). VEGF is an important vascular growth factor. It has stronger regulation effects on vascular endothelial cells proliferation, basement membrane degradation, endothelial cell migration and angiogenesis, with high specificity (Xiao et al., 2004). It is proved that VEGF is a key medium for tumor angiogenesis. The expression of VEGF is high in many animal and human malignant tumors, whether in mRNA level or protein level. Inhibition of VEGF level can inhibit the tumor growth (Ferrara, 2010). As found in this study, the relative expression amounts of HIF-1α and VEGF protein in DDP, 100 mg/kg Cur and 200 mg/kg Cur group were significantly decreased, respectively, compared with control group (P < 0.05). This indicates that, as the same with DDP, Cur can decrease the expressions of HIF-1α and VEGF protein in tumor tissue.

MIF is a pleiotropic cytokine produced by activated T cells. It does not belong to any currently known cytokine family. It is named due to that it can inhibit the random movement of macrophages in vitro. MIF is a classic proinflammatory cytokine, and can activate the inflammatory reaction through a variety of ways (Cooke et al., 2009). In addition, MIF is a unique cytokine
which can promote the occurrence of malignant tumors. It presents high expression in a variety of malignant tumors, and accelerates the tumor growth, invasion, and metastasis by promoting the malignant transformation of oncogene, inhibiting the function of tumor suppressor gene p53, and increasing the secretion of extracellular matrix degrading enzymes (Mitchell et al., 2002). In the present study, the relative expression amounts of MIF protein in DDP, 50 mg/kg Cur, 100 mg/kg Cur and 200 mg/kg Cur group were significantly decreased, respectively, compared with control group (P < 0.05). This indicates that, Cur can decrease the expressions of MIF protein in tumor tissue, and there is no dose difference.

5 Conclusions
Cur can inhibit the growth of HeLa cell xenograft in nude mice. The possible mechanism may be related to its up-regulation of Bax, p53 and p21 protein expression in tumor tissue, and down-regulation of Bcl-2, HIF-1α, VEGF and MIF protein expression. In addition, Cur has no obvious effect on the body weight of nude mice. This study has provided the theoretical and experimental basis for the clinical application of Cur. However, whether there are other mechanisms of Cur on the tumor inhibition should be further studied.

References
Motterlini, R., Foresti, R., Bassi, R., & Green, C. J. (2000). Curcumin, an antioxidant and anti-inflammatory agent, induces heme oxygenase-1 and protects endothelial cells against oxidative stress. Free Radical Biology & Medicine, 28(8), 1303-1312. PMid:10889462. http://dx.doi.org/10.1016/S0891-5849(00)00294-X.


