Honokiol induces apoptosis through p53-independent pathway in human colorectal cell line RKO

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Abstract

AIM: To investigate the signal pathway of honokiol-induced apoptosis on human colorectal carcinoma RKO cells and to evaluate whether p53 and p53-related genes were involved in honokiol-treated RKO cells.

METHODS: Cell cycle distribution and subdiploid peak were analyzed with a flow cytometer and DNA fragment with electrophoresis on agarose gels. Transcriptional level of Bax, Bcl-2, Bid and Bcl-xl was accessed by RT-PCR. Western blotting was used to measure p53 protein expression and other factors related to apoptosis. Proliferation inhibition of two cell lines (RKO, SW480) with high expression of p53 and one cell line with p53 negative expression (LS180) was monitored by MTT assay.

RESULTS: Honokiol induced RKO cell apoptosis in a dose-dependent manner. The mRNA expression level and protein level of Bid were up-regulated while that of Bcl-xl was down-regulated, but no changes in Bax and Bcl-2 were observed. Western blotting showed p53 expression had no remarkable changes in honokiol-induced RKO cell apoptosis. LS180 cells treated with honokiol exhibited apparent growth inhibition like RKO cells and SW480 cells.

CONCLUSION: Honokiol can induce RKO cells apoptosis through activating caspase cascade by p53-independent pathway.

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INTRODUCTION

Honokiol is a major active constituent extracted from the bark of Magnolia officinalis (Chinese name for Houpu) (Figure 1). It has a variety of pharmacological effects, such as anti-inflammatory[1], antithrombosis[2], anti-arrhythmic[3], antioxidant[4] and anxiolytic effects[5]. Recently, honokiol has been reported to exhibit a potent cytotoxic activity by inducing cell apoptosis in some cell lines[6-8]. Honokiol-triggered apoptotic process is accompanied with down-modulation of Bcl-XL, release of mitochondrial cytochrome C in CH27 cells[9].

Cells undergoing apoptosis are characterized by distinct biochemical and morphological changes. Many of the biochemical and morphological events of apoptosis are a direct result of caspase-mediated cleavage of specific substrates[10,11]. Together with caspase, Bcl-2 family is involved in inducing cell apoptosis and identified as essential components of the intracellular apoptotic signaling pathways. Cell viability versus death is determined by the relative abundance of various members of the Bcl-2 family, acting in concert with other proteins in the death pathway[12,13].

In many tumor cells, wild-type p53 is considered to participate in apoptosis in response to DNA damage[14]. P53 may transactivate apoptotic regulators, such as Bcl-2[15-17] and Bax[18,19]. Recent studies have shown that p53 plays a role in apoptosis by the mitochondrial-mediated apoptotic pathway[20]. Activation of p53 upregulates Bax[19] and increases the ratio of Bax:Bcl-2, releases cytochrome C and other polypeptides from the intermembrane space of mitochondria into cytoplasm. Cytochrome C activates caspase cascade[12,13].

It is well known that a broad range of agents can induce tumor cell apoptosis through p53-regulated manner by Bax/mitochondria/caspase-9 pathway. But up to now, it is not clear that whether wild-type p53 takes part in honokiol-induced apoptosis. In this present study we examined whether wild-type p53 and p53-related gene were involved in honokiol-treated RKO cells.

MATERIALS AND METHODS

Materials

RPMI 1640 medium was obtained from Gibco BRL. Newborn bovine serum was supplied by Sijiqing Biotechnology Co. (Hangzhou, China). Monoclonal antibodies to Bax, Bcl-xl, Bcl-2, β-actin and p53 were purchased from NeoMarkers, Fremont, CA, USA. Honokiol was got from the National Institute for Pharmaceutical and Biological Products, Beijing, China. The drug was dissolved in DMSO with the stock concentration of 10 mg/mL. It was further diluted in culture medium with the final DMSO concentration <1%. 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) and propidium iodide (PI) were purchased from Sigma Chemical Corporation, USA.

Cell culture

Human colorectal cell lines RKO, SW480 and LS180 were provided by Cancer Institute of Zhejiang University. Three cells were maintained in RPMI 1640 medium (Gibco BRL) supplemented with 100 mL/L heat-inactivated fetal bovine serum (Si-Ji-Qing Biotechnology Co, Hangzhou, China), 100 U/mL penicillin and 100 µg/mL streptomycin at 37 °C in a 50 mL/L CO₂ atmosphere.

Detection of DNA fragmentation

To detect DNA fragments, RKO cells were collected and lysed with lysis buffer containing 50 mmol/L Tris-HCl (pH 7.5),...
20 mmol/L EDTA, and 10 g/L NP-40. Then 10 g/L SDS and RNase (5 µg/mL) were added to the supernatants, and incubated at 56 °C for 2 h, followed by incubation with proteinase K (2.5 µg/mL) at 37 °C for 2 h. After DNA was precipitated by addition of both ammonium acetate (3.3 mol/L) and ethanol (995 mL/L) dissolved in a loading buffer, DNA fragmentation was detected by electrophoresis on 15 g/L agarose gels and visualized with ethidium bromide staining.

Cell cycle analysis by FCM
Honokiol-treated RKO cells (5, 10, 15 µg/mL) and vehicle were fixed with 700 mL/L alcohol for 15 min at 4 °C, then stained with 1.0 µg/mL propidium iodide (PI, Sigma, USA). The red fluorescence of DNA-bound PI in individual cells was measured at 488 nm with a FACSCalibur (Becton Dickinson, USA) and the results were analyzed using ModFit 3.0 software. Ten thousand events were analyzed for each sample.

RT-PCR assay
RKO cells 1×10⁶ were seeded on 24-well plate. After 24-h culture, cells were treated with 5, 10 µg/mL honokiol and vehicle for 48 h. Total RNA from RKO cells was extracted using Trizol (Invitrogen, USA). In RT-PCR, cDNA synthesis was performed using a RNA PCR kit (TaKaRa Biomedicals, Osaka, Japan) with the supplied oligo dT primer (Table 1). Reverse transcription was performed using thermal program of 92 °C for 2 min, then 38 cycles of annealing for 30 s at 55 °C, extension for 90 s at 72 °C. As an internal control, semiquantitative analysis of β-actin was also amplified. This primer pair of β-actin had an optimal annealing temperature of 55 °C with 20 cycles and yielded a 350-bp PCR product. Samples were separated on 20 g/L agarose gel and visualized with ethidium bromide staining under UV light.

Western blotting
RKO cells (5×10⁶) treated with 5, 10 µg/mL honokiol and vehicle respectively for 24 h were lysed by 4 g/L trypsin containing 0.2 g/L EDTA, then collected after washed twice with phosphate-buffered saline (PBS, pH 7.4). Total protein extract from RKO cells was prepared using cell lysis buffer [150 mmol/L NaCl, 0.5 mol/L Tris-HCl (pH 7.2), 0.25 mol/L EDTA (pH 8.0), 10 g/L Triton X-100, 50 mL/L glycerol, 12.5 g/L SDS]. The extract (30 µg) was electrophoresed on 12 g/L SDS-PAGE and electroblotted onto polyvinylidene difluoride membrane (PVDF, Millipore Corp., Bedford, MA) for 2 h in a buffer containing 25 mmol/L Tris-HCl (pH8.3), 192 mmol/L glycine and 200 mL/L methanol. The blots were blocked with 50 g/L nonfat milk in TBST washing buffer for 2 h at room temperature and then incubated at 4 °C overnight with antibodies. All antibodies were diluted in TBST according to the manufacturer’s instructions. After washed at room temperature with washing buffer, the blots were labeled with peroxidase-conjugated secondary antibodies.

Cell proliferation assay
RKO cells, SW480 and LS180 cells (1×10⁶ in 100 µL) were seeded on 96-well plates in triplicate respectively. Following a 24-h culture at 37 °C, the medium was replaced with fresh medium containing vehicle control or various concentrations of honokiol with peroxidase-conjugated secondary antibodies. The blots were blocked with 50 g/L nonfat milk in TBST washing buffer for 2 h at room temperature and then incubated at 4 °C overnight with antibodies. All antibodies were diluted in TBST according to the manufacturer’s instructions. After washed at room temperature with washing buffer, the blots were labeled with peroxidase-conjugated secondary antibodies.

Statistics
Statistical significance was determined by Student’s t-test. A P value of 0.05 or less was considered significant.

RESULTS
DNA fragmentation detection in RKO cells
In honokiol-treated RKO cells, a degradation of chromosomal DNA into small internucleosomal fragments was evidenced by the formation of 180-200 bp DNA ladder on agarose gels (Figure 2), hallmark of cells undergoing apoptosis. No DNA ladders were detected in the samples isolated from control cultures. These results indicated that honokiol induced an apoptotic cell death in RKO cells.

Effect of honokiol on cell cycle phase distribution
RKO cells were exposed to the increased concentrations of honokiol (5-15 µg/mL) for 48 h, and the cell growth was analyzed using flow cytometry. In the absence of honokiol, the cell populations were at G1, S, and G2/M phases (Figure 3), accompanied with increased concentrations of honokiol by a concomitant increase of the G1 phase (Table 2). From Figure 3, we considered the peak areas of subdiploid were up at the

Table 1 Expected size of PCR amplification product with each apoptosis modulator primer pair

| Apoptosis modulator | Upstream primer (5’-3’) | Downstream primer (5’-3’) | Size (bp) |
|---------------------|-------------------------|---------------------------|-----------|
| Bax                 | ACCAAGAAAGCTGAGCGAGGT   | ACAAACATGTGTCAGGTCTGC     | 332       |
| Bcl-xl              | GGAGCTGGGTGGTATCTTC     | CCAGAACGAGTCTCCAAT         | 379       |
| Bid                 | GAGC CCG GTG CCT CAG GA | ATG GTC ACG GTC TGC CA     | 586       |
| Bcl-2               | TTGTGCTTCTTGAGTTGCG     | TACTGCTTTAAGGACCTTTT       | 332       |
| β-actin             | AGGCCAAGCCGGGAGATGACC   | GAAGTCCAGGGCGACGTAGCAC     | 350       |
increased concentrations of honokiol. This observation led to a suggestion of G1 arrest. When the cells were exposed to honokiol at 10 µg/mL and above, subdiploid peak was significantly increased and the percentage of apoptotic cells in 10,000 cells was 14.1% and 20.31% respectively (Table 2).

Table 2 Effect of honokiol on cell cycle distribution and apoptosis in RKO cells

| Treatment (µg/mL) | Cell cycle distribution (%) | Apoptosis (%) |
|-------------------|-----------------------------|---------------|
|                   | G0/G1 | S   | G2/M |                  |
| Control           | 44.76 | 45.09 | 10.14 | 0.21            |
| 5                 | 48.03 | 42.37 | 9.60  | 4.51            |
| 10                | 54.04 | 40.30 | 5.66  | 14.10           |
| 15                | 58.94 | 39.36 | 1.70  | 20.31           |

Cell cycle distribution was determined after 48 h of treatment in each group. The tabulated percentages were an average calculated on the results of three separate experiments. *P<0.01, **P<0.01 vs corresponding control group.

Semi-quantification of Bid, Bax, Bcl-2 and Bcl-xl mRNA expression

mRNA expression of apoptosis-related genes in response to different levels of honokiol was assessed. The Bcl-xl mRNA expression was significantly reduced in RKO cells exposed to honokiol, while Bid mRNA expression was remarkably up-regulated. No apparent changes of Bax and Bcl-2 were observed (Figure 4).

Effect of honokiol on expression of p53-related gene and p53

p53 and p53-related gene were found to be importantly involved in apoptosis induced by many agents. To study their role in honokiol-treated RKO, we monitored Bax, Bcl-2, Bcl-xl, Bid and p53 protein levels. Results in Figure 5 shows that no significant changes of Bax, Bcl-2 and p53 were found compared with vehicle. The level of Bcl-xl was increased remarkably after treated with 5 and 10 µg/mL honokiol respectively. The protein levels were measured by quantitative Western blot analysis after normalized with β-actin content.

Inhibition of proliferation in RKO, Sw480 and LS180 cells

RKO cells had positively expressed high wild-type p53 and SW480 cells had positive expression of high mutant p53, while LS180 had negative p53 antigen expression[26-29]. Cells treated with various concentrations of honokiol resulted in a dose-dependent cytotoxicity in three cells. As shown in Figure 6, honokiol-treated LS180 exhibited apparent growth inhibition like RKO cells and Sw480. Honokiol-mediated cytotoxicity occurred at the concentration of 5 µg/mL and above. A significant decrease in cell number was seen at 10 µg/mL. The concentration leading to a 50% decrease in cell number (IC50) was approximate 10.33, 12.98, 11.16 μg/mL in RKO, SW480 and LS180 cells respectively.

DISCUSSION

p53 is a crucial protein in cellular stress response. p53-dependent arrest of cells in the G1 phase of the cell cycle was an important component of the cellular response to stress[23]. Wild-type p53 was considered to participate in apoptosis in response to DNA damage in many tumor cells[14,24]. When cells received UV or ionizing radiation and were exposed to anticancer drugs, p53 protein was accumulated[25,26]. The increased p53 could transactivate its downstream target genes to induce cell cycle arrest, DNA repair, and apoptosis[27,28]. p53-dependent apoptosis was also observed in vivo[29]. However, p53-independent apoptotic cascades after administration of anticancer drugs or r-irradiation have been described.

In the present study, honokiol-treated RKO cells could induce apoptosis in a concentration-dependent manner, with the ratio of G1 phase increased. The agent also caused an increase of the expression of caspase-3 and caspase-7 in a dose-dependent manner, did not cause the significant increase
of caspase-9 (data not shown). In CH27 cells it promoted to release cytochrome C and activate caspase-3[6]. p53-dependent apoptosis was activated by the Bax/ mitochondrial/caspase-9 pathway. Bcl-2 and Bax expression were regulated by p53 both in vitro and in vivo, and Bax was a direct target of p53 transcriptional activation[30]. Bax, a proapoptotic member of the Bcl-2 family, was located in the outer membrane of mitochondria[25]. Increased in the ratio of Bax/Bcl-2 could cause changes in the membrane potential of mitochondria, consequently, cytochrome C and other polypeptides were released from the intermembrane space of mitochondria into cytoplasm. Once released, cytochrome C could activate procaspase-9 by self-cleavage[12,13] and then other caspases[12].

According to previous data, honokiol led to cell G1 arrest and cytochrome C release like p53. Thus it is necessary to evaluate the mRNA and protein expression of Bax and Bcl-2. Our results showed that honokiol-induced apoptosis of RKO cells was accompanied with up-regulation of Bid and down-modulation of Bcl-xl, but did not involve the regulation of Bcl-2 or Bax protein expression. Since pro-apoptotic Bax is a p53 downstream target, the lack of change of Bax and Bcl-xl supports the view that honokiol probably does not trigger the p53/Bax-mediated apoptosis pathway. Next we examined the expression of p53 in honokiol-treated RKO cells. We found that no significant change of p53 was observed (Figure 5). Finally we analyzed honokiol-induced apoptosis of the three cells with different p53 expression. Our data showed that honokiol-treated LS180 and SW480 cells exhibited apparent growth inhibition like RKO cells. These results suggest that honokiol-induced caspase activation and cell apoptosis are entirely controlled by the p53-independent pathway.

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