RESEARCH PAPER

4-O-methylhonokiol, a PPAR γ agonist, inhibits prostate tumour growth: p21-mediated suppression of NF- κ B activity

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BACKGROUND AND PURPOSE

The effects of 4-O-methylhonokiol (MH), a constituent of *Magnolia officinalis*, were investigated on human prostate cancer cells and its mechanism of action elucidated.

EXPERIMENTAL APPROACH

The anti-cancer effects of MH were examined in prostate cancer and normal cells. The effects were validated *in vivo* using a mouse xenograft model.

KEY RESULTS

MH increased the expression of PPAR γ in prostate PC-3 and LNCap cells. The pull-down assay and molecular docking study indicated that MH directly binds to PPAR γ . MH also increased transcriptional activity of PPAR γ but decreased NF- κ B activity. MH inhibited the growth of human prostate cancer cells, an effect attenuated by the PPAR γ antagonist GW9662. MH induced apoptotic cell death and this was related to G₀-G₁ phase cell cycle arrest. MH increased the expression of the cell cycle regulator p21, and apoptotic proteins, whereas it decreased phosphorylation of Rb and anti-apoptotic proteins. Transfection of PC3 cells with p21 siRNA or a p21 mutant plasmid on the cyclin D1/ cyclin-dependent kinase 4 binding site abolished the effects of MH on cell growth, cell viability and related protein expression. In the animal studies, MH inhibited tumour growth, NF- κ B activity and expression of anti-apoptotic proteins, whereas it increased the transcriptional activity and expression of PPAR γ , and the expression of apoptotic proteins and p21 in tumour tissues.

CONCLUSIONS AND IMPLICATION

MH inhibits growth of human prostate cancer cells through activation of PPAR γ , suppression of NF- κ B and arrest of the cell cycle. Thus, MH might be a useful tool for treatment of prostate cancer.

Abbreviation

CDK, cyclin-dependent kinase; CNBr, cyanogen bromide; DAPI, 4',6-diamidino-2-phenylindole; GSK-3 β , glycogen synthase kinase 3 β ; IAP1, inhibitor of apoptosis 1; iNOS, inducible NOS; MH, 4-O-methylhonokiol

Introduction

The root and stem bark of *Magnolia officinalis* are used in oriental medicine and this medicinal herb and its bioactive constituents, such as honokiol, obovatol and magnolol, have been shown to exhibit a variety of biological actions, including anti-tumour, anti-microbial, anti-inflammatory, anti-thrombotic and anxiolytic effects (Lee *et al.*, 2011). Several studies have shown that honokiol has anti-angiogenic, anti-proliferative and anti-invasive activities in colon and prostate cancer cells as well as in lung, breast and ovarian cancer cells (Yang *et al.*, 2002; Wang *et al.*, 2004; Wolf *et al.*, 2007; Hahm *et al.*, 2008; Fried and Arbiser, 2009). Magnolol has also been shown to induce apoptosis and suppress the proliferation of cancer cells, and inhibit tumour metastasis in various cancer cells, including those of the colon and prostate (Lin *et al.*, 2002; Lee *et al.*, 2009). Further, we recently found that obovatol also inhibits colon and prostate cancer cell growth (Lee *et al.*, 2008).

PPARs are members of the nuclear hormone-receptor superfamily of ligand-dependent transcriptional factors and have three major subtypes (α , β and γ). PPARs have been demonstrated to have a number of pathophysiological roles and there is mounting evidence suggesting that PPAR γ is important in the control of cancer cell growth (Kim *et al.*, 2003). PPAR γ is up-regulated in many malignant tissues and PPAR γ ligands induce terminal differentiation, cell growth inhibition and apoptosis in a variety of cancer cells including those of the prostate (Haydon *et al.*, 2002; Yamakawa-Karakida *et al.*, 2002). PPAR γ negatively regulates gene expression of cell proliferation and pro-apoptotic genes in a ligand-dependent manner by antagonizing the activities of NF- κ B in prostate cancer cells (Hirsch *et al.*, 2011). Cell-cycle regulatory proteins, such as p21, have been shown to play a critical role in the PPAR γ -induced inhibition of prostate cancer cell growth (Radhakrishnan and Gartel, 2005). Moreover, the well-known PPAR γ agonist, 15-deoxy-PGJ₂ and other activators induce inhibitory effects on cancer cell growth in a PPAR γ -dependent manner through inactivation of NF- κ B (Kim *et al.*, 2003; Papineni *et al.*, 2008).

NF- κ B is a transcription factor regulating various genes involved in the production of inflammatory cytokines, chemokines, cell adhesion molecules and growth factors (Richmond, 2002). NF- κ B mediates tumour promotion and progression, angiogenesis, metastasis of cancer cells and promotes resistance to various chemotherapeutics by increasing the expression of genes participating in cancer development (Pikarsky *et al.*, 2004; Dolcet *et al.*, 2005). NF- κ B also regulates the expression of a number of anti-apoptotic proteins and apoptotic genes as well as proliferation genes (Wang *et al.*, 1998). Interestingly, NF- κ B is constitutively activated in various solid tumours including animal and human prostate tumours (Lind *et al.*, 2001; Suh and Rabson, 2004). Constitutive activation of NF- κ B has been found to up-regulate the expression of anti-apoptotic and proliferation genes, and therefore, disrupt the balance between apoptosis and proliferation leading to cancer cell growth (Garg and Aggarwal, 2002). Thus, inhibition of NF- κ B has been presented as a key regulator of cell growth of various cancer cells (Lee *et al.*, 2002; Allen *et al.*, 2007). We have also demonstrated that suppression of NF- κ B by naturally occurring compounds,

such as infliximol, thiacecremonone and cobra toxin, inhibits colon and prostate cancer cell growth (Son *et al.*, 2007; Ban *et al.*, 2009a; 2009b).

Functional inactivation of the p21 pathway has been observed in most human tumours (Biggs and Kraft, 1995) and p21 is an important cellular checkpoint protein for the inhibition of cell growth through the control of cyclin-cyclin-dependent kinases (CDKs) activities in human cancer cells (Ivanovska *et al.*, 2008). NF- κ B and PPAR γ are also important in p21-mediated cell cycle arrest of human cancer cells (Hellin *et al.*, 2000; Prakobwong *et al.*, 2011); the expression of p21 is increased in the resistant human cancer cells in a NF- κ B- and PPAR γ -dependent manner (Chang and Miyamoto, 2006; Jarvis *et al.*, 2005). Moreover, many investigators, using components and crude extracts isolated from the *Magnolia* family, have shown that p21-mediated cell cycle arrest leads to cell growth inhibition (Hahm and Singh, 2007; Lee *et al.*, 2007; Chen *et al.*, 2009). We recently isolated a bioactive compound, 4-O-methylhonokiol (MH) from *M. officinalis*, and found it suppressed NF- κ B activity in macrophages and colon cancer cells (Oh *et al.*, 2009; 2012). The growth inhibitory effects of the bioactive constituents of *M. officinalis*, obovatol, honokiol and magnolol, are thought to be associated with the inhibitory ability of these compounds on NF- κ B activity in cancer cells (Lee *et al.*, 2007; 2008). In addition, we recently found that MH-induced inhibition of colon tumour growth was accompanied by a suppression of NF- κ B activity (Oh *et al.*, 2012).

Thus, we hypothesized that MH inhibits the growth of human prostate cancer cells through activation of PPAR γ and inactivation of NF- κ B, and these effects are mediated by the induction of p21. Here, we showed that MH enhances PPAR γ activity but decreases NF- κ B activity, and thus inhibits prostate cancer cell growth through induction of p21.

Methods

Reagents

MH (Supporting Information Figure S1A) was isolated from the bark of *M. officinalis* Rehd. *et* Wils, by subsequently extracting with *n*-hexane, ethyl acetate and *n*-BuOH, and then identified by ¹H-NMR and ¹³C-NMR as described elsewhere (Oh *et al.*, 2009). MH was supplied by Bioland Ltd. (Chungnam, Korea) as a brown oilish solution of 99.6% purity. The stock solution of MH was diluted in 0.05% DMSO (final concentration, 50 mM).

Cell culture

PC-3, LNCaP human prostate cancer and PWR-1E human prostate normal cells were obtained from the American Type Culture Collection (ATCC, Manassas, VA, USA). The PC-3 (11 passages) and LNCaP (43 passages) prostate cancer cells were grown in RPMI1640, and the CCD 112-CoN cells were grown in Eagle's minimum essential medium with 10% FBS, 100 U·mL⁻¹ penicillin, and 100 µg·mL⁻¹ streptomycin. The PWR-1E cells were cultured in a keratinocyte growth medium supplemented with 5 ng·mL⁻¹ human recombinant EGF and 0.05 mg·mL⁻¹ bovine pituitary extract (Gibco/Invitrogen, Carlsbad, CA, USA).

Cell growth assay

Cells (5×10^4 cells per well) were plated onto 24-well plates. The cell growth inhibitory effect of MH was evaluated in cells treated with MH (0–30 μM) for 0–72 h, using an excluded trypan blue assay.

Transfection and assay of luciferase activity

Cells (1×10^5 cells per well) were plated in 24-well plates and transiently transfected with pNF- κB -Luc plasmid ($5 \times \text{NF-}\kappa\text{B}$; Stratagene, La Jolla, CA, USA) or plasmid pFA-GAL4-PPAR γ , using a mixture of plasmid and lipofectamine PLUS in OPTI-MEN according to manufacturer's specification (Invitrogen). The transfected cells were treated with MH in the absence (for assay of PPAR γ activity) or presence (for assay of NF- κB activity) of TNF- α (10 ng·mL $^{-1}$) for 8 h. To induce NF- κB luciferase activity, we co-treated the cells with TNF- α (10 ng·mL $^{-1}$). Luciferase activity was measured by using the luciferase assay kit (Promega, Madison, WI).

Electromobility shift assay

Electromobility shift assay was performed according to the manufacturer's recommendations (Promega) as described elsewhere (Oh *et al.*, 2009).

Western blot analysis

Protein (40 μg) from cultured cells was separated on a SDS/12%-polyacrylamide gel, and then transferred to a nitrocellulose membrane (Hybond ECL, Amersham Pharmacia Biotech Inc., Piscataway, NJ). The membranes were then immunoblotted with primary specific antibodies: rabbit polyclonal antibodies for PPAR γ , p65, bax, bcl-2, GSK-3 β (1:500 dilution, Santa Cruz Biotechnology Inc. Santa Cruz, CA, USA), caspase-3, caspase-9, pp53, survivin, c-IAP1 (1:1000 dilution, Cell Signaling Technology, Inc. Beverly, MA), and cIAP2 (1:1000 dilution, Abcam Ltd., Cambridge, UK) and mouse monoclonal antibodies for p50, p53, CDK6 (1:500 dilution, Santa Cruz Biotechnology Inc.), CDK4, p21, cyclin D, cyclin E (1:1000 dilution, Medical and Biology Laboratories Co. Ltd., Nagoya, Japan), iNOS and COX-2 (1:1000 dilution, Cayman Chemical Company, Ann Arbor, MI, USA). The blot was then incubated with the corresponding conjugated anti-rabbit IgG-HRP (1:2000 dilution, Santa Cruz Biotechnology Inc.).

Cell-cycle analysis by flow cytometry

Subconfluent cells were treated with MH (0–30 μM) in culture medium for 0–72 h. The analysis methods are as described elsewhere (Ban *et al.*, 2009a).

Transfection of siRNA and p21 mutant plasmid

Cells were transfected with 100 nM of CDKN1A (p21), NF- κB (p50 or 65) siRNA or non-specific siRNA (Bioneer, Co., Daejeon, Korea) and p21 mutant plasmid (CDKN1A p21, obtained from Dr Anindya Dutta, Department of Biochemistry and Molecular Genetics, University of Virginia School of Medicine, Charlottesville, VA, USA), which has mutated cyclin D1/CDK4 complex binding site (Cy1 site, $\Delta 17$ –24) using WelFect-EX plus transfection reagent (WelGENE, Seoul, Korea) prepared in a serum-free culture medium at 37°C (Hsu

et al., 2007). After 6 h, a complete medium was added and the cells were further cultured for 24 h.

Detection of apoptosis

Cells (1×10^4 cells per well) were cultured in 8-well chamber slides (BD Biosciences, Bedford, MA, USA) for 24 h. The method used to detect apoptotic cell death was as described elsewhere (Ban *et al.*, 2009b). The apoptotic index was determined as the number of DAPI-stained TUNEL-positive stained cells divided by the total number of cells counted $\times 100$.

Pull-down assays

MH bead conjugation was prepared and a pull-down assay conducted as described previously (Shim *et al.*, 2008). MH was conjugated with cyanogen bromide (CNBr)-activated Sepharose 4B (Sigma Chemical Co.). Briefly, MH (1 mg) was dissolved in 500 μL of coupling buffer (0.1 M NaHCO $_3$ and 0.5 M NaCl, pH 6.0). The CNBr-activated Sepharose 4B was swelled and washed with 1 mM HCl, then washed with the coupling buffer. CNBr-activated Sepharose 4B beads were added to the MH-containing coupling buffer and incubated at 4°C for 24 h. The MH-conjugated Sepharose 4B was washed with three cycles of alternating pH wash buffers (buffer 1, 0.1 M acetate and 0.5 M NaCl, pH 4.0; buffer 2, 0.1 M Tris-HCl and 0.5 M NaCl, pH 8.0). MH-conjugated beads were then equilibrated with binding buffer (0.05 M Tris-HCl and 0.15 M NaCl, pH 7.5). The control unconjugated CNBr-activated Sepharose 4B beads were prepared as described earlier in the absence of MH. For the pull-down assay, PPAR γ proteins (Abcam Ltd.) or cell lysate from PC-3 prostate cancer cells were incubated with MH-Sepharose 4B beads in reaction buffer (50 mM Tris, pH 7.5, 5 mM EDTA, 150 mM NaCl, 1 mM dithiothreitol, 0.01% Nonidet P-40, 2 $\mu\text{g}\cdot\text{mL}^{-1}$ BSA, 0.02 mM PMSF, 1 $\times$ proteinase inhibitor). The beads were washed five times with buffer (50 mM Tris, pH 7.5, 5 mM EDTA, 150 mM NaCl, 1 mM dithiothreitol, 0.01% Nonidet P-40, 0.02 mM PMSF), and proteins bound to the beads were analysed by immunoblotting with the PPAR γ antibodies or cell lysates.

Molecular modelling

The dimeric structure of PPAR γ complexed with nitrated fatty acids LNO $_2$ with 2.4 Å resolution (PDB ID: 3CWD) (Cheng *et al.*, 2009) was selected for docking with MH and only half of it was used for the study. For MH, rotational flexibility was allowed for all ligand backbone bonds. The initial search with the VINA docking calculation was done at the binding pocket where the LNO $_2$ ligand binds. Then, to get a better sampling of all possible binding sites on the PPAR γ , the search was expanded to include one monomer of the PPAR γ protein. The grid box was centred on the PPAR γ monomer and the size of the grid box was adjusted to completely include the whole PPAR γ monomer. All bonds able to be rotated on the MH were allowed to rotate during the molecular simulation. Docking experiments were performed at various exhaustive values of the default value, 25, 50 and 100. All these experiments identified the binding site with the highest affinity at the same binding pocket with essentially identical binding conformation on the MH backbone.

Antitumour activity study in in vivo xenograft animal model

Six-week-old male BALB/c athymic nude mice were purchased from Japan SLC (Hamamatsu, Japan). All experiments were approved and carried out according to the Guideline for 'Care and Use of Animals' of the Chungbuk National University Animal Care Committee. All studies involving animals are reported in accordance with the ARRIVE guidelines for reporting experiments involving animals (Kilkenny *et al.*, 2010; McGrath *et al.*, 2010). SW620 and PC3 cells were injected s.c. (1×10^7 cells in 0.1 mL PBS per animal) into the lower right flanks of mice. After 20 days, when the tumours had reached an average volume of 300–400 mm³ or about 50 mm³ (for prostate), the tumour-bearing nude mice were i.p. injected with MH (40 and 80 mg·kg⁻¹ dissolved in 0.1% DMSO) twice per week for 3 weeks. Cisplatin (10 mg·kg⁻¹, Choongwae Pharma Co., Seoul, Korea) was also i.p. injected once a week as a positive control. The group treated with 0.1% DMSO was designated as the control. The tumour volumes were measured with vernier calipers and calculated by the following formula: $(A \times B^2)/2$, where A is the larger and B is the smaller of the two dimensions.

Immunohistochemistry

Immunohistochemistry was performed as described elsewhere (Ban *et al.*, 2009a) with 5 µm thick tissue sections using primary mouse PCNA, PPAR γ , Ki-67 and p65 (1:200 dilution) antibodies or primary rabbit p50 and cleaved caspase-3 antibody (1:100 dilution) followed by a secondary biotinylated anti-mouse or rabbit antibody. For quantification, 200 cells at three randomly selected areas were assessed, and the positively stained cells were counted and are expressed as a percentage of stained cells.

Data analysis

Data were analysed using GraphPad Prism 4 software (Version 4.03, GraphPad Software, Inc., La Jolla, CA, USA). Data were assessed by one-way ANOVA. If the *P*-value in the ANOVA test was significant, the differences ($P < 0.05$) between the pair of means were assessed by Dunnett's test. Data are presented as mean $\pm$ SD from three independent experiments with triplicates.

Results

MH increased the expression, transcription and DNA binding activities, and nuclear translocation of PPAR γ

To investigate whether the inhibitory effect of MH on cell growth is dependent on PPAR γ , the expression of PPAR γ was first determined in two prostate cancer cells. The expression of PPAR γ was easily detected in the untreated cells was increased (3- or 3.5-fold increase) after MH treatment (20 µM) in both cells (Figure 1A). To investigate whether MH is able to activate PPAR γ -dependent transcription in these cells, the functional status of endogenous PPAR γ was determined using a luciferase reporter assay system. MH significantly increased PPAR γ transcription activity, in a concentration-dependent manner, in cells transiently transfected with the PPAR γ con-

struct (Figure 1B). The other related compounds isolated from *M. officinalis* had a similar effect on PPAR γ transcription activity that was about 90% of that induced by a well-known PPAR- γ agonist, 15-deoxy-PGJ₂, determined in PC3 cells (Supporting Information Fig. S1B). We also performed a gel mobility shift assay to assess whether MH increased the DNA binding activity of PPAR γ . The DNA binding activity of PPAR γ in untreated cells was not significant, but a substantial increase of binding activity was observed in the cells treated with MH and other compounds isolated from *M. officinalis* (Figure 1B). This binding activity was inhibited in the presence of cold PPAR response element (PPRE) oligonucleotides, but not in unrelated labelled PPRE oligonucleotides and the DNA-binding complex was supershifted by anti-PPAR γ (data not shown). MH caused nuclear translocation of PPAR γ , which is localized predominantly in the perinuclear region and cytoplasm in the untreated cells (Figure 1C).

In the molecular docking experiment, MH was predicted to bind at the identical ligand binding site as other PPAR γ agonists, such as 15-deoxy-PGJ₂ (Waku *et al.*, 2009) and the fatty acid ligand LNO₂ (Cheng *et al.*, 2009). This implies that MH may activate PPAR γ in the same manner as 15-deoxy-PGJ₂ and other PPAR γ agonists. According to the binding model, MH also forms hydrophobic interactions with Phe²⁸², Cys²⁸⁵, Ile³²⁶, Tyr³²⁷, Leu³³⁰, Phe³⁶³, Met³⁶⁴, Leu⁴⁵³, Leu⁴⁶⁵ and Leu⁴⁶⁹ (Figure 1D). One hydrogen bond was observed between the hydroxyl group of the MH and the hydroxyl side chain of Ser²⁸⁹ in PPAR γ (Figure 1E). The interaction was thus assessed in a pull-down assay using MH-sepharose 4B beads. The binding of MH to PPAR γ was then detected by immunoblotting with anti-PPAR γ in *in vitro* and *ex vivo* tests and MH was found to bind with recombinant PPAR γ protein (Figure 1F).

MH inhibited cancer cell growth, caused G₀/G₁ phase arrest and induced apoptotic cell death, and PPAR γ antagonist reversed the effects of MH

We then investigated whether PPAR γ activation resulted in cell growth inhibition. MH (0–30 µM) treatment resulted in a significant concentration- and time-dependent inhibition of prostate cancer cell growth (Figure 2A). However, MH was not cytotoxic in the normal prostate PWR-1E cells at the concentrations tested (Figure 1C). Even though it was clear that the LNCaP grew much faster than the PC-3 cells, in contrast to previous observations (Horoszewicz *et al.*, 1983; Lin *et al.*, 1998), it is noteworthy that the doubling times of these two cells are similar (Yu *et al.*, 2008), and the doubling time of LNCaP (androgen dependent cells) can be easily changed by the culture condition such as concentration of serum, and presence of other hormones and status of specific genes carrying as demonstrated by Horoszewicz *et al.* (1983) and Ghosh *et al.* (2006). We next measured apoptotic cell death, to determine whether MH-induced cell growth inhibition was due to apoptotic cell death. Apoptotic cell numbers were increased to 3 ± 5 , 10 ± 6 , 30 ± 20 and $75 \pm 2\%$ in PC3 cells, and 0 ± 6 , 6 ± 2 , 50 ± 6 and $63 \pm 2\%$ in LNCaP cells by 0–30 µM MH, respectively (Figure 2B). To further study the involvement of PPAR γ in the MH-induced cell growth inhibition and induction of apoptotic cell death, we employed the PPAR γ antagonist SW9662 (0–30 µM). Pretreating cells with SW9662 30 min

Figure 1

Expression of PPAR γ in prostate PC-3 and LNCap cells. (A) Cells were treated with 20 μ M MH for 24 h. Whole cell extracts were prepared and PPAR γ expression was determined by Western blot analysis as described in *Methods*. The value above the band indicates fold increase of PPAR γ expression. (B) Transcriptional activation determined by luciferase activity. Luciferase activity was determined in the cells transfected with PPAR γ plasmid construct after treatment with MH for 24 h as described in *Methods*. Values are mean $\pm$ SD of three experiments, with triplicate results for each experiment. (C) Immunoreactivity of PPAR γ in PC-3 cells and in LNCaP cells treated with 20 μ M MH. Untreated cells showed that PPAR γ was localized predominantly in the perinuclear region and cytoplasm. Cells treated with MH demonstrated nuclear translocation of PPAR γ . D-(a), Representations of the interaction between PPAR γ and MH. MH binds to a tight-binding cavity within the agonist binding pocket in the ligand-binding domain of PPAR γ shown in surface model. D-(b), The surrounding peptide residues that interact with MH are shown in the second picture. (E) Hydrogen-bonding interaction between hydroxyl group of MH and hydroxyl side chain of Ser²⁸⁹ of PPAR γ . (F) MH-Sepharose 4B was used to pull down with PPAR γ protein (Abcam Ltd.). WB, Western blot.

before incubating them with MH for 72 h reversed MH-induced cell growth inhibition in a dose-dependent manner (Figure 2C), and also reversed the induction of apoptotic cell death (Figure 2D). To determine whether the apoptotic cell death resulted from cell cycle arrest, we examined cell cycle arrest for 72 h. Exposure of cells to MH (0–30 μ M) increased the number of prostate cancer cells in the G₀/G₁ fraction in a concentration- and time-dependent manner (Figure 3A).

MH altered the expression of regulatory proteins involved in cell growth, cell cycle and apoptotic cell death

From the apoptotic cell death and cell cycle pattern, MH decreased the expression of anti-apoptotic proteins of bcl-2

and the inhibitor of apoptosis protein 1/2 (cIAP1/2) as well as GSK-3 β and survivin, but increased the expression of pro-apoptotic proteins of bax, cleaved caspase-3 and cleaved caspase-9 (Figure 3B). Moreover, MH significantly decreased the expression of COX-2 and iNOS, which are involved in the tumour promotion of prostate cancer (Figure 3B). MH also caused a rapid and marked decrease in the protein levels of CDK2, CDK4, cyclin D1 and cyclin E (Figure 3C). As p21 and p27 regulated the transition of G₀/G₁ by binding to CDK/cyclin complexes, which phosphorylate Rb, leading to cell-cycle progression (Hsu *et al.*, 2007), we determined the effect of MH on p21 and p27 protein levels and on the phosphorylation of Rb by immunoblotting. As shown in Figure 3C, MH markedly increased the protein expression of p21,

Figure 2

Effect of MH and WS9662 on cancer cell growth and apoptotic cell death. (A and C) Cells were treated with 0–30 μ M MH for different times (0–72 h) (A) or treated with WS9662 for 72 h (C). Cell counting was performed as described in *Methods*. (B and D) Cells were treated with 0–30 μ M MH for 24 h or the cells were pretreated with WS9662 for 24 h. DAPI and TUNEL staining were performed as described in *Methods*. Columns, mean of triplicate; bars, SD. * P < 0.05 indicates statistically significant differences from the untreated group. # P < 0.05 indicates statistically significant differences from the MH treated group.

slightly increased that of p27, but suppressed Rb phosphorylation (Figure 3C).

MH inhibited NF- κ B activity and cancer cell growth

Highly activated NF- κ B is associated with cell survival as well as with the resistance against therapeutics of prostate cancer cells. Activation of PPAR γ and inhibition of cancer cell growth could be coupled with inhibition of NF- κ B. Thus, we determined the ability of MH to inhibit the DNA binding activity of NF- κ B. In untreated human prostate cancer cells (PC3 and LNCaP), we found a high level of constitutive activation of the DNA binding activity of NF- κ B and MH (0–30 μ M) for 1 h inhibited this DNA binding activity of NF- κ B in a concentration-dependent manner (Figure 3D). In addition, MH (0–30 μ M) inhibited TNF- α -induced NF- κ B luciferase activity concentration-dependently (Figure 3E), and confocal

microscope analysis further demonstrated that the translocation of the NF- κ B subunits, p50 and p65, into the nucleus was also decreased by MH (Figure 3F). When we compared cell growth inhibition and the PPAR γ and NF- κ B DNA binding activities of honokiol, magnolol, obovatol and MH, the extent of cell growth inhibition was associated with the activation of PPAR γ (transcriptional and DNA binding activities); significant inhibition of NF- κ B was also observed in the cells treated with magnolol and obovatol (Figure 1B and D).

Knock-down of p21 abolished the inhibitory effects of MH on cancer cell growth and NF- κ B, and its activation of PPAR γ

p21 is a cell cycle inhibitory protein regulating cell cycle progression and is known to be a critical target of the *Magnolia* family of compounds. Thus, to further investigate the role of p21 in the MH-induced cell cycle arrest, we examined

Figure 3

Effect of MH on cell cycle, and expression of proteins regulating cell growth, cell cycle and apoptotic cell death, and NF- κ B DNA binding, transcriptional activity and translocation of p50 and p65. (A) Cells were treated with 0–30 μ M MH for 72 h. Cell cycle was performed as described in *Methods*. (B and C) Cells were treated with 0–30 μ M MH for 24 h, and used for Western blotting as described in *Methods*. The blots were representative of three experiments. (D and F) Cells were treated with 0–30 μ M MH for 1 h. Electrophoretic mobility shift assay (D) and immunofluorescence (F) were performed as described in *Methods*. Numbers, fold activation in relation to the control (D). (E) Cells were transfected with pNF- κ B-Luc plasmid (5 $\times$ NF- κ B) for 6 h and then incubated with complete media containing 0–30 μ M MH with TNF- α (10 ng mL⁻¹) for 8 h. Luciferase activity was assessed as described in *Methods*. Columns, mean of triplicate; bars, SD. #*P* < 0.05 indicates statistically significant differences from the untreated group. *P < 0.05 indicates statistically significant differences from the TNF- α -treated group.

the effect of siRNA p21 and a mutation of p21 on MH-induced cell growth inhibition. Knock-down of p21 with p21 siRNA abolished the inhibitory effect of MH on cancer cell growth (Figure 4A) and its induction of apoptotic cell death (Figure 4B). Moreover, in p21 siRNA-transfected cells MH-induced inhibition of the G₀/G₁ cell cycle-regulatory protein expression of CDK and CD4, translocation of p65 and p60, and expression of PPAR γ were all abolished (Figure 4C). Similarly, in cells transfected with mutant p21 (mutation of cyclin D1 and CDK4 complex binding site of p21) the inhibitory effects of MH on cell growth (Figure 4D), its induction of apoptotic cell death (Figure 4E), as well as cell cycle-regulatory protein expression and p50 and p65 translocation, and increased expression of PPAR γ were all abolished (Figure 4F). These results indicate that the PPAR γ and NF- κ B

signal-mediated inhibitory effects of MH on the growth of PC3 cells are associated with activation of the p21 signal pathway.

MH inhibited the growth of SW620 and PC3 tumours in an in vivo xenograft model

In PC3 xenograft studies, MH was administered i.p. daily for 4 weeks to mice with tumours ranging from 100 to 300 mm³ in volume. On day 28, the final tumour weight was recorded. Tumour volumes in mice treated with MH at 40 and 80 mg kg⁻¹, and cisplatin at 10 mg kg⁻¹ were 71.0, 57.7 and 46.6% of the control group in PC3 tumour xenografts, respectively. Tumour weight in mice treated with MH at 40 and 80 mg kg⁻¹ and cisplatin at 10 mg kg⁻¹ were 40.1, 30.9 and 22.1% of the control group in PC3 tumour xenografts, respec-

Figure 4

Transfection of PC3 cells with p21 siRNA or p21 mutant plasmid abolished the effect of MH on cell viability, apoptosis and related protein expression. (A–C) The cells were transfected with p21 siRNA for 6 h, grown for 24 h in complete media, and treated with 20 μ M MH for 24 h. (D–F) The cells were treated with p21 mutant plasmid which has mutated cyclin D1/cdk4 complex binding site (Cy1 site, 17–24) for 6 h, grown for 24 h in complete media, and treated with 20 μ M MH for 24 h. Cell counting (A,D), DAPI and TUNEL staining (B, E), and Western blotting (C, F) were performed as described in *Methods*. Columns, mean of triplicate; bars, SD. * $P < 0.05$ indicates statistically significant differences from the untreated group.

tively (Figure 5A). Immunohistochemical analysis of tumour sections by H&E, and proliferation antigens against PCNA and Ki67 staining revealed that both 40 and 80 mg·kg⁻¹ dose-dependently inhibited the growth of the tumour cells (Figure 5B). In addition, there was a trend towards decreased intensity of nuclear staining of p65 and p50 in MH-treated tumour tissue (Figure 5B). Moreover, PPAR γ immunoreactivity against anti-PPAR γ was also more intense in the tumours treated with MH than in untreated tumour tissues. Similar to its inhibitory effect *in vitro*, MH enhanced the DNA binding activity of PPAR γ , but inhibited NF- κ B activity in tumour tissue (Figure 5C). MH increased the expression of bax and cleaved caspases-3, but decreased the expression of bcl-2 in tumour tissue (Figure 5D). Immunohistochemical analysis also showed that the expression of cleaved caspases-3 positive cells was considerably increased in the MH-treated tumour

tissue compared with those of the control group. Apoptotic cell death was also significantly increased in the MH-treated tumour tissue (Figure 5B). Moreover, MH significantly increased the expression of p21 and PPAR γ in the tumour tissues (Figure 5B–D).

Discussion and conclusions

The present data show that MH inhibited the growth of prostate cancer cells through G₀/G₁ phase cell cycle arrest and induction of apoptotic cell death. These effects were associated with increased expression of PPAR γ and p21, and activation of PPAR γ but inhibition of NF- κ B activity. Knock-down of p21 with siRNA and mutation of p21 prevented this MH-induced increase in PPAR γ expression and activation,

Figure 5

Effect of 4-O-methylhonokiol (MH) on the tumour growth in PC3 xenografts, *in vivo* model. Mice with SW620 or PC3 xenografts were treated with MH (40 or 80 mg·kg⁻¹ everyday) or cisplatin (10 mg·kg⁻¹ once a week) for 4 weeks. (A) Tumour volume and weight were measured as described in *Methods*. Data represent mean of ten animals; bars show SD. *P < 0.05. (B) Immunohistochemistry; (C) electrophoretic mobility shift assay; and (D), Western blotting were performed as described in *Methods*. Columns, mean of three animals; bars, SD. *P < 0.05 indicates statistically significant differences from the untreated group.

NF- κ B inactivation, apoptotic cell death as well as the expression of CDK4/cyclin D1. These results suggest that MH inhibits the growth of prostate cancer cells by increasing the expression of p21 as a result of activation of PPAR γ , but inactivation of NF- κ B.

In the present study, MH activated PPAR γ and its nuclear translocation, but decreased NF- κ B activity and nuclear translocation of p65 and p50 in prostate cancer cells. MH also effectively and significantly induced apoptotic cell death in prostate cancer cells. These effects support the notion that activation of PPAR γ , but inactivation of NF- κ B, inhibit the growth of cancer cells by promoting apoptotic cell death (Garg and Aggarwal, 2002). Activation of PPAR γ and inactivation of NF- κ B are implicated in the regulation of growth arrest and/or apoptotic cell death by regulating expression of target genes such as bax, caspase-3, caspase-9, bcl-2, IAP,

survivin and GSK-3 β (Garg and Aggarwal, 2002). Our data clearly showed that MH inhibited the expression of genes involved in cell proliferation (cyclin D1 and COX-2) and of anti-apoptotic proteins (cIAP1/2 and bcl-2), which are regulated by PPAR γ and NF- κ B, but increased the levels of pro-apoptotic proteins (cleaved caspase-3 and 9, and bax) *in vitro* as well as *in vivo*. This study showed, for the first time, that MH increases PPAR γ activity and thus inhibited the growth of prostate cancer cells. These results corroborate previous findings where it was shown that PPAR γ agonists, such as troglitazone, curcumin and anti-tumoural pigment epithelium-derived factor, inhibit colon biliary and prostate cancer cell growth through activation of PPAR γ , but inactivation of NF- κ B (Yamakawa-Karakida *et al.*, 2002; Hirsch *et al.*, 2011; Prakobwong *et al.*, 2011). These effects of MH are similar to those obtained with honokiol and magnolol

isolated from *M. officinalis*; these compounds were found to have anti-tumour activity, both *in vitro* and *in vivo*, on a variety of cancer cells including prostate cancer cells through inactivation of NF- κ B, and inhibition of NF- κ B target gene expression (Wang *et al.*, 2004; Hahm *et al.*, 2008). In addition, we previously found that MH inhibited colon cancer cells through the inactivation of NF- κ B (Oh *et al.*, 2012). These data suggest that the ability of MH to enhance PPAR γ activation and its inhibitory action on the constitutive activation of NF- κ B are involved in its inhibitory effect on the growth of prostate cancer cells.

The p21 protein regulates G₁/S transition by inhibiting CDKs (Chen *et al.*, 1996). MH increased the induction of p21 protein and this was accompanied by a reduced expression of the G₀/G₁ cell cycle-regulatory proteins, cyclin D1 and CDK4. However, this reduction in expression of cyclin D1 and CDK4 was abolished in the cells after knock-down of p21 with siRNA. Likewise, cells transfected with p21, which had a cyclin/CDK binding site mutation, did not respond to MH. That is, the MH-induced cell growth inhibition, inhibition of the expression of cyclin D1 and CDK4, and phosphorylation of pRb were not observed in p21 mutant cells. These results suggest that the induction of p21 expression may be important in the MH-induced arrest of human prostate cancer cells in the G₀/G₁ phase of the cell cycle. Similar to our findings, honokiol and magnolol induce G₀/G₁ cell cycle arrest via induction of p21 in several cancer cells including prostate cancer cells (Hahm and Singh, 2007; Hsu *et al.*, 2007; Lee *et al.*, 2007; Chen *et al.*, 2009; Vaid *et al.*, 2010). We previously found that MH induced cell cycle arrest by induction of p21 in colon cancer cells (Oh *et al.*, 2012). PPAR γ and NF- κ B are also important for the p21-mediated cell cycle arrest of cancer cell growth (Hellin *et al.*, 2000). Increased NF- κ B-dependent p21 expression was observed in human cancer cells resistant to chemotherapeutics (Chang and Miyamoto, 2006). In the present study, the effects of MH on PPAR γ , and reciprocal converse effects of NF- κ B with p21 and p27 expression were found to correlate with its ability to inhibit cell growth and induce apoptotic cell death. The MH-induced PPAR γ activation and NF- κ B inactivation were abolished in the cells after knock-down of p21 with siRNA as well as in the cells transfected with mutant p21. We also found that MH, concomitant with the increased expression of p21 and p27, also enhanced the phosphorylation of Rb, the downstream target gene of p21. MH also increased the expression of p21 and p27 with decreased expression of the G₀/G₁ phase of cell cycle regulatory proteins of cyclin D1 and E, and CDK2 and CDK4 in prostate cancer xenograft nude mice tumour tissue. Similar to our results, a recent study demonstrated that an aqueous extract of *M. officinalis* increased p21 expression and decreased NF- κ B in human urinary bladder cancer cells (Lee *et al.*, 2007). Moreover, recent studies have shown that anti-cancer drugs are able to inhibit the growth of several human cancer cells through up-regulation of p21 accompanied by activation of PPAR γ , but inhibition of NF- κ B activation. For example, pioglitazone, a PPAR γ agonist inhibited p21-mediated prostate cancer cell growth (Hirsch *et al.*, 2011). PPAR γ stimulates sulindac sulfide-mediated p21 expression in prostate epithelial cells (Jarvis *et al.*, 2005). Soy peptide increased p21 in human breast MCF-7 tumour cells via

inactivation of NF- κ B, and thus inhibited their growth (Park *et al.*, 2009). Curcumin inhibited human breast cancer MDA-MB-231 cell growth via NF- κ B inactivation-induced up-regulation of p21 (Chiu and Su, 2009). Lycopene inhibited the growth of prostate cancer cells by inactivation of NF- κ B and up-regulation of p21 (Palozza *et al.*, 2010). We also found that MH inhibited colon cancer cell growth by p21-mediated NF- κ B inactivation (Oh *et al.*, 2012). These data and ours suggest that both the enhancing ability of MH on PPAR γ and its inhibitory effect on the constitutive activation of NF- κ B are related to the increased expression of p21, and the modification of these cooperative signals is involved in its inhibition of prostate cancer cell growth. However, it is not clear how changes in the expression of p21 alter PPAR γ and NF- κ B, or whether p21 is downstream of PPAR γ and/or NF- κ B or vice versa. It is noteworthy that there are PPRE as well as a NF- κ B-binding element on the p21 promoter (Han *et al.*, 2004; Sue *et al.*, 2009). Thus, increased activity of PPAR γ or reduced activity of NF- κ B induced by MH could enhance or inhibit the expression of p21. It is also interesting to note that the PPAR γ agonist thiazolidinedione inhibits cancer growth via inhibition of nucleophosmin (NPM) expression, which is known to induce p53 phosphorylation and p21 expression (Galli *et al.*, 2010). Thus, it is possible that the activating effect of MH on PPAR γ may affect the expression of proteins like NPM, which increase p21 expression. These data suggest that p21 acts downstream of PPAR γ and NF- κ B.

Pharmacokinetic data have shown that MH has good oral and intestinal absorption as determined by the Caco-2a with easy distribution into tissues (unpublished data). The dosage of MH used in the present *in vivo* study are within the range employed in other previous studies showing anti-tumour effects of the compounds isolated from *M. officinalis*; i.p. administration of honokiol at 80 mg·kg⁻¹ for 31 days displayed significant anti-cancer activity (Chen *et al.*, 2004). Treatment with liposomal honokiol at 25 mg·kg⁻¹ reduced the tumour sizes by 42% compared with the untreated tumour in Lewis lung carcinoma-bearing C57BL/6 mice (Hu *et al.*, 2008). In toxicology studies, we demonstrated that *M. officinalis* extract did not produce any toxic effects in the 21 and 90 day toxicity studies, and that up to 240 mg·kg⁻¹ is a no-observed-adverse-effects-level (Liu *et al.*, 2007), which is similar to the range of doses used in the present study if it is assumed the extract has about 15% of 4-O-methylenhonokiol, as described elsewhere (Oh *et al.*, 2009). We have also evaluated the long-term toxicity including carcinogenicity of MH using a prediction program (preADME version 1.0.2, BMDRC, Seoul, Korea), and found that it was predicted not to be toxic or carcinogenic to rodents (data not shown). Taken together, these results suggest that MH should be considered for further clinical investigation to determine its possible chemopreventive and/or therapeutic efficacy against human prostate cancer.

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Conflict of interest

None.

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Supporting information

Additional Supporting Information may be found in the online version of this article at the publisher’s web-site:

Figure S1 Chemical structure of MH, cell viability in normal prostate cells, and luciferase activity, DNA binding activity of

PPAR γ and NF- κ B by other components isolated from *Magnolia officinalis*. (A) Chemical structure of MH. (B) PWR-1E normal prostate cells were treated with various doses (0–30 μ M) MH for 72 h. Morphological changes were observed under a microscope. Cell viability was determined by direct cell counting using trypan blue as described in Methods. (C,D) Transcriptional and DNA binding activities were determined by luciferase activity and EMSA. Luciferase activity was determined in the cells transfected with PPAR- γ plasmid construct after treatment of MH and other several agents MH for 24 h as described in Methods. *Columns*, mean

of triplicate; *bars*, SD. Prostate cancer cells treated with MH and other agents were incubated for 1 h. Nuclear extract from cells treated with MH, honokiol, magnolol and obovotaol for 1 h was incubated in binding reactions of ^{32} P-end-labelled oligonucleotide containing the PPRE or κ B sequence (D). EMSA was performed as described in Methods. D, After Cells were treated with 0–20 μ M of MH, honokiol (H), magnolol (M) and obovatol (O) for 24 h, cell viability was determined by direct cell counting using trypan blue. *Points and Columns*, mean of triplicate; *bars*, SD. *, $P < 0.05$ indicates statistically significant differences from the untreated group.