Abstract. N-(4-hydroxyphenyl)retinamide (4-HPR or fenretinide), which is a synthetic analog of all-trans retinoic acid (ATRA), effectively inhibits the growth of several types of tumor cells; however, its molecular mechanism remains unclear. We found that 4-HPR altered the morphology of human liver cancer HepG2 cells and also inhibited their proliferation and suppressed the colony formation in a dose- and time-dependent manner. A wound healing assay revealed that 4-HPR significantly hindered HepG2 cell migration, and that this was accompanied by the phosphorylation of p38-MAPK (mitogen-activated protein kinase). Mechanistically, the MAPK-specific inhibitor SB203580 attenuated the inhibitory effects of 4-HPR on the migration of HepG2 cells. Moreover, we also observed that 4-HPR inhibited the activation and expression of myosin light chain kinase (MLCK) in HepG2 cells. Simultaneously, 4-HPR lowered the expression of F-actin and promoted the expression of E-cadherin. ML-7, a selective inhibitor of MLCK, significantly inhibited the migration of HepG2 cells while increasing the phosphorylation of p38-MAPK and the expression of E-cadherin, and decreasing the activation of MLCK and the expression of F-actin. In conclusion, 4-HPR inhibited the proliferation and migration of HepG2 cells, and p38-MAPK plays an important role in regulating these 4-HPR effects by reducing the activation of MLCK. The present study suggests that 4-HPR may be a potent antimetastatic agent.

Introduction

Human liver cancer, hepatoblastoma, is the most common malignant liver tumor in pediatrics (1). Although surgical excision of the tumor mass can be performed, challenges still remain for patients with vascular invasion and metastatic disease (2). Inhibition of metastasis is hence a key strategy for liver cancer treatment, and the discovery of potential inhibitors of metastasis could lead to improvements in therapy. Optional therapeutic schemes and auxiliary substances have been widely explored; however, only few are effective (3). Thus, the development of new drugs is required for optimal treatment with fewer complications.

All-trans retinoic acid (ATRA) has been effectively used for inducing the differentiation of acute promyelocytic leukemia cells (4); however, it does not cure patients with liver cancer, as liver cancer requires nearly a 10 times higher concentration than leukemic cells (5). Such high concentrations are not suitable for clinical use due to several side effects, including retinoic acid syndrome, skin dryness, and liver damage. N-(4-hydroxyphenyl)retinamide (4-HPR or fenretinide), an artificial variant of ATRA, exhibits markedly different effects to ATRA. Presently, 4-HPR is considered to be a drug with fewer side effects (6). Vaccari et al found that 4-HPR influences cell matrix interactions and blocks tumor progression to locally invasive malignancy (7). 4-HPR was found to reduce the incidence of breast cancer when used as a chemoprevention agent (8) and prevented secondary breast cancer in a phase III trial (9). Furthermore, 6 years later Kang et al demonstrated that 4-HPR inhibited the invasion of breast cancer cells (10). Similarly, Benelli et al demonstrated that 4-HPR hindered the migration and invasion of prostate cancer cells (11), and it has now entered clinical phase II trials (12). In the present study, we compared the antiproliferative effects of 4-HPR with ATRA on HepG2 cells, and
explored the functions and mechanisms of 4-HPR in modulating their migration capacity.

Mitogen-activated protein kinases (MAPKs) are highly conserved signaling pathway proteins, playing vital roles in deciding cell fate (13,14). p38-MAPK is activated by different stimuli, including chemical agents, cytokines and oxidative stress (13-15). Sustained activation of p38-MAPK induces cell death (14,16). 4-HPR-induced sustained activation of p38-MAPK, accompanied by cell apoptosis, has been reported in several types of tumors, including neuroblastoma, HeLa, T-cell leukemia/lymphoma cells (17-19); however, HepG2 cells were reported to be resistant to the apoptotic effect of 4-HPR (20). Whether 4-HPR influences the p38-MAPK pathway in HepG2 cells remains unclear.

Myosin light chain kinase (MLCK) is a crucial Ca²⁺/calmodulin-dependent effector, and controls the migration of smooth- and non-muscle cells through the phosphorylation of Ser19 and Thr18 on myosin light chains (MLC) (21). Previous studies have reported that the MLCK and activated myosin II are abundant in the lamellar protrusive structures of certain cell types during migration (22,23). Several studies have revealed p38-MAPK pathway links in the tumor cells treated with 4-HPR, and also some researches have found that MLCK is involved in the migration of tumor cells; however, the underlying mechanism that explains how these factors influence liver cancer is still unknown.

Therefore, we hypothesized that 4-HPR inhibits the proliferation and migration of liver cancer cells via MLCK and p38-MAPK signaling. This study was aimed to provide an experimental basis for the further application of 4-HPR in liver cancer therapy.

Materials and methods

Cell lines and major reagents. The human liver cancer cell line HepG2 (24) was obtained from the American Type Culture Collection (ATCC; Manassas, VA, USA). HyClone Dulbecco's modified Eagle's medium (DMEM; low glucose) was purchased from GE Healthcare Life Sciences (Logan, UT, USA). Fetal bovine serum (FBS) was purchased from Tianhang Biological Technology (Hanzhou, Zhejiang, China). Primary antibodies: Rabbit anti-human monoclonal MLCK (cat. no. ab92721) and E-cadherin (cat. no. ab40772), mouse anti-human monoclonal F-actin (cat. no. ab205) were obtained from Abcam (Cambridge, MA, USA); rabbit anti-human monoclonal phospho-p38 MAPK (cat. no. 4511), mouse anti-human monoclonal phospho-MLC (cat. no. 3675) were purchased from Cell Signaling Technology (Danvers, MA, USA); rabbit anti-human polyclonal MLC (cat. no. 15354-1-AP), mouse anti-human monoclonal GAPDH (cat. no. 60004-1-Ig) were obtained from Proteintech Group (Wuhan, Hubei, China); rabbit anti-human polyclonal p38 MAPK (cat. no. sc-7149) and mouse anti-human monoclonal β-actin (cat. no. sc-47778) were obtained from Santa Cruz Biotechnology (Santa Cruz, CA, USA). All secondary antibodies (cat. nos. AP124P and AP132P) were obtained from Millipore (Billerica, MA, USA). Western blot primary antibody diluent was obtained from Beyotime Institute of Biotechnology (Beijing, China). Enhanced chemiluminescence reagent Plus (ECL) reagents were purchased from Thermo Fisher Scientific (Waltham, MA, USA). 4-HPR was obtained from MedChem Express (Deer Park, NY, USA). ATRA and ML-7 was purchased from DC Chemicals (Shanghai, China). SB203580 was obtained from Selleck Chemicals (Houston, TX, USA). 4-HPR, ATRA, ML-7 and SB203580 were dissolved in a small amount of dimethyl sulfoxide (DMSO) before addition to the complete cell culture medium. MTS was purchased from Promega (Madison, WI, USA).

Cell culture and morphologic observation after drug treatment. Cells were seeded in 6-well plates, and cultured in DMEM supplemented with 10% FBS, penicillin (100 U/ml) and streptomycin (100 µg/ml). The plates were incubated at 37°C with 5% CO₂ in a humidified atmosphere. When the cell density reached 40-50% confluency, the cells were treated with 4-HPR or ATRA at 5 or 10 µM or with DMSO alone for 48 h. Cell morphology was imaged using a microscope (Leica DMI3000B; Leica Microsystems, Wetzlar, Germany).

Cell viability assay. Cells (5x10³ cells/well) were plated in 96-well plates, and then treated with 4-HPR or ATRA (5 or 15 µM) at different time intervals (24-72 h), or persistently treated with a dose range of 1-25 µM of 4-HPR or ATRA for 48 h at 37°C. Following treatment, MTS (20 µl/well) was added to each well and incubated for 1 h at 37°C. Optical density (OD) values were measured at 490 nm using a Microplate reader (ELX800; BioTek, Winooski, VT, USA) at 37°C. Cell inhibition rate=(OD490 of the cell control group-OD490 of the experimental group)/OD490 of the cell control group.

Plate colony formation assay. When the cells were in logarithmic growth, they were plated in 6-well plates. After culturing overnight, the cells were treated with 4-HPR or ATRA (5 or 10 µM) for 48 h, and were then collected as single cell suspensions. Approximately, 2,000 cells/wells were plated into a fresh 6-well plate, and the plate was incubated for approximately 10 days. When the colonies were clearly visible with the naked eye, the medium was discarded and the cells were washed twice with phosphate-buffered saline (PBS), fixed with 4% paraformaldehyde, and then stained with 1% crystal violet. In five random visual fields, the colonies containing ≥50 cells were counted by an inverted microscope (Leica DMI3000B; Leica Microsystems) for each well.

Wound healing assay. The migratory ability of the HepG2 cells was determined by the wound healing assay. Cells were seeded in 12-well plates. When the growth reached 95% confluency, the cell monolayer was scratched with a sterilized 200-µl pipette tip, and then the cells were washed thrice with PBS. Furthermore, two concentrations (5 and 10 µM) of 4-HPR or ATRA were added to the cell culture medium. The migration rate of cells was determined by observation under a microscope at different time intervals (0, 24, and 48 h). The nick distance of the wound was measured by Image-Pro Plus software 6.0 (Media Cybernetics, Inc., Rockville, MD, USA).

Western blot analysis. Cells were treated with 4-HPR or ATRA (5 or 10 µM) for 48 h. Total cellular proteins were extracted with lysis buffer (Tris-HCl, pH 7.14, 150 mM NaCl,
1 mM EDTA, 1% Triton X-100, 0.1% SDS, 1 mM leupeptin and 1 mM PMSF) on ice for 30 min. All the samples were mixed with loading buffer, and then boiled for 5 min. The proteins were separated by 8-12% sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE). Furthermore, the proteins were transferred to polyvinylidene difluoride (PVDF) membranes, and blocked with 5% nonfat dry milk in Tris-buffered saline Tween-20 (TBST) buffer (20 mM Tris-HCl, pH 7.6, 150 mM NaCl, and 0.05% Tween-20) for 2 h at room temperature. The membranes were then incubated in WB primary antibody diluent with the indicated primary antibodies: MLCK (1:8,000), phospho-MLC (1:1,000), MLC (1:1,000), phospho-p38 (1:1,000), p38 (1:1,000), F-actin (1:500), E-cadherin (1:4,000), GAPDH (1:50,000), β-actin (1:1,000), respectively overnight at 4˚C. Thereafter, the membranes were incubated with the corresponding horseradish peroxidase (HRP)-conjugated secondary antibodies (diluted with TBST containing 5% non-fat dry milk) for 2 h at room temperature, and detected by chemiluminescence using an ECL kit. Specific complexes were revealed by enhanced chemiluminescence (Clinx Science Instruments, Shanghai, China). The image data were quantified using Quantity One software 4.6.2 (Bio-Rad Laboratories, Inc., Hercules, CA, USA).

**HepG2 cell treatment with p38-MAPK and MLCK inhibitors.** The pharmacological inhibitors of the p38-MAPK signaling pathway (SB203580, 25 µM) and MLCK (ML-7, 20 µM) were used in combination with 4-HPR or ATRA (10 µM) to investigate the role of p38-MAPK and MLCK in 4-HPR/ATRA-induced inhibition of HepG2 cell migration. Thereafter, wound healing assay and western blot analysis were used to measure the migration distances and protein levels in the cells.

**Statistical analysis.** Results are expressed as means ± standard deviation (SD). The data were assessed by one-way ANOVA using SPSS 10.0 software (SPSS, Inc., Chicago, IL, USA). P-values <0.05 were considered as statistically significant.

**Results**

4-HPR alters the morphology of HepG2 cells. In a previous study, Yang et al (20) found that HepG2 cells were resistant to the apoptotic effect of 4-HPR after 24 h treatment; however, 4-amino-2-trifluoromethyl-phenyl retinate (ATPR), another ATRA derivative, has been observed to inhibit HepG2 cell proliferation after 48 h of culture in our laboratory (25). Based on this report, we chose two concentrations of 4-HPR (5 and 10 µM) to treat HepG2 cells for 48 h. Changes in the cell morphology and any inhibitory effects were examined by microscopy. The cells congregated neatly and closely in the solvent (DMSO) control (Fig. 1); however, after treatment with a low concentration of 4-HPR, the cell density and cell-to-cell contact was reduced. In the high concentration group, cell density was remarkably reduced, and the cell morphology was altered into a slender shape containing more filopodia when compared with the vehicle control (Fig. 1). These changes could be associated with cell migration after 4-HPR treatment. Cell density was reduced only slightly in the high concentration ATRA group, and was not accompanied by any noticeable change in the cell morphology.

4-HPR inhibits the proliferation of HepG2 cells. A cell viability assay was used to probe the effects of 4-HPR and ATRA on the proliferation of HepG2 cells. Cell viability was distinctly inhibited in a dose- and time-dependent manner by 4-HPR (Fig. 2A and B). At the same concentration, the inhibitory effect of 4-HPR was more intense than that of ATRA (P<0.05). In addition, there was no obvious difference between the cell control and the DMSO control group. Based on the aforementioned data, the DMSO control, and an incubation
4-HPR INHIBITS THE PROLIFERATION AND MIGRATION OF HepG2 CELLS

Figure 2. Effects of 4-HPR on the proliferation of HepG2 cells. Cells were incubated with the indicated concentrations of 4-HPR or ATRA for the indicated times. (A) The dose-effect of 4-HPR on HepG2 cell viability was measured by cell viability assay. (B) The time-effect of 4-HPR on HepG2 cell viability was determined using cell viability assay. (C) Colony formation map and mean numbers of colonies. All values are presented as mean ± SD. n=6. *P<0.05 compared with the control group; #P<0.05 compared with the ATRA group. 4-HPR, fenretinide; ATRA, all-trans retinoic acid.

Figure 3. Effect of 4-HPR on the migration ability and expression of p38-MAPK in HepG2 cells. HepG2 cells were treated with 4-HPR or ATRA at different concentrations for 48 h. (A) The treatment suppressed the migration of HepG2 cells. (a) Migration image. (b) Analysis of the migration rate. Calculation method: Relative migration ratio=(start distance-end distance)/start distance. All values are presented as mean ± SD. n=6. 24 h: #P<0.05 compared with the cell group; 48 h: *P<0.05 compared with the cell group. (B) (a) Western blot analyses revealed that the phosphorylation of p38 was significantly increased. Lane 1, cell control; lane 2, DMSO; lane 3, 5 µM ATRA; lane 4, 10 µM ATRA; lane 5, 5 µM 4-HPR; and lane 6, 10 µM 4-HPR. (b) Analysis of contrast gray value. All values are presented as mean ± SD. n=3. *P<0.05 compared with the DMSO group; #P<0.05 compared with the ATRA group. 4-HPR, fenretinide; ATRA, all-trans retinoic acid.

time of 48 h were chosen for further studies. A plate colony formation assay was used to assess the colonizing ability of HepG2 cells in vitro. After 10 days of culture, the density and size of the colonies were both reduced in a dose-dependent manner by 4-HPR (P<0.05); however, they were only changed slightly at the high concentration of ATRA compared with the control group (P<0.05). The mean numbers of colonies in the 4-HPR group were lower than those in the ATRA group.
(P<0.05), which was consistent with the morphological change and cell viability analyses (Fig. 2C).

4-HPR hinders the migration of HepG2 cells. We investigated the effect of 4-HPR and ATRA on the migration of HepG2 cells using a wound healing assay. 4-HPR inhibited the migration of HepG2 cells (Fig. 3A) in a dose-dependent manner (P<0.05, compared with the control), consistent with the data from the cell proliferation assay. 4-HPR inhibited cell migration at two concentrations of 5 and 10 µM (P<0.05, compared with the control), whereas ATRA only produced inhibition at 10 µM (P<0.05, compared with the control).

4-HPR increases the activation of p38-MAPK in HepG2 cells. Treatment of cells with 4-HPR increased the phosphorylation of p38 (p-p38) in a dose-dependent manner, and the expression of p38 was also increased in the high concentration group, as revealed by Western blot analysis (P<0.05, compared with the control). ATRA also slightly increased the phosphorylation of p38 (p-p38) (P<0.05, compared with the control); however, the increased levels elicited by 4-HPR were much higher than those produced by ATRA (P<0.05) (Fig. 3B). These results indicated that 4-HPR may inhibit the migration of HepG2 cells through the p38-MAPK signaling pathway.

p38-MAPK inhibitor reverses the inhibitory effect of 4-HPR on HepG2 cell migration. To determine whether 4-HPR inhibits HepG2 cell migration via p38-MAPK signaling, the cells were pretreated with the p38-MAPK inhibitor SB203580 for 1 h, and then exposed to 4-HPR (10 µM) for 48 h. 4-HPR-induced inhibition of migration was abrogated by the presence of the SB203580 at both 24 and 48 h (P<0.05, compared with 4-HPR alone) (Fig. 4). These data indicate that 4-HPR stimulates p38 activity leading to migration inhibition in the liver cancer cells.

4-HPR decreases the expression of MLCK and phosphorylation of MLC in HepG2 cells, and a p38-MAPK inhibitor had an inverse effect. Since myosin light-chain kinase (MLCK) plays a crucial role in cell migration and metastasis, we investigated the expression of this protein, as well as its substrate (MLC) and product (p-MLC). This revealed that 4-HPR markedly reduced the expression of MLCK and the phosphorylation of MLC (p-MLC), and also decreased the expression of MLC (P<0.05, compared with the control) (Fig. 5A). Concomitantly, 4-HPR inhibited the expression of F-actin and increased the expression of E-cadherin (P<0.05, compared with the control) (Fig. 5B). ATRA reduced the expression of p-MLC (P<0.05, compared with the control); however, it caused no obvious change in the MLCK levels. 4-HPR reduced the expression of MLCK and the phosphorylation of MLC to a much greater extent than ATRA (P<0.05). However, when the cells were pretreated with SB203580, p-MLC and F-actin were upregulated and E-cadherin was downregulated when compared to 4-HPR or ATRA treatment alone (P<0.05) (Fig. 5C and D). These observations fit well with the wound healing assay data presented above.

Reduction of MLCK activation inhibits the migration of HepG2 cells. To verify the association between HepG2 migration and the MLCK signaling pathway, a wound healing assay was performed using ML-7 (a specific inhibitor of MLCK) in the culture medium. After treatment with ML-7 for 24 or 48 h, the migration rate of HepG2 cells was suppressed when compared with the control (P<0.05) (Fig. 6). Moreover, the group treated with both 4-HPR and ATRA combined with ML-7 exhibited greater inhibition rates than the group treated with 4-HPR or ATRA alone (P<0.05) (Fig. 6).

ML-7 increases the phosphorylation of p38 and inhibits the activation of MLCK in HepG2 cells. The underlying mechanism, that is, the nature of the signaling pathway tied to MLCK repression and p38-MAPK activation after 4-HPR treatment, remained unclear. Therefore, we investigated the protein expression after ML-7 treatment (Fig. 7A-C). ML-7 not only inhibited the activity of MLCK by reducing the expression of p-MLC, but also activated p38-MAPK by enhancing the expression of p-p38 in HepG2 cells (P<0.05, compared with the control). ML-7 also altered the expression of F-actin and E-cadherin, and ML-7 combined with 4-HPR or ATRA further increased the levels of p-p38 compared to 4-HPR or ATRA alone (P<0.05).
Discussion

Patients with liver cancer often have unfavorable prognoses and short lifespans due to early metastasis (26). The process of metastasis involves tumor cell escape, migration, invasion of the basement membrane, and growth at new locations (27). Knowledge concerning the circumstances that favor liver cancer cell metastasis will aid in finding treatment options that can control the growth and metastasis of liver cancer.

4-HPR is a known retinoid analog that is active against several types of tumors that arise via different ontological mechanisms (28). A phase II clinical study in adults with prostate cancer revealed good compatibility with 4-HPR (29), and similar results were obtained in a neuroblastoma in a phase I study in children (30). Antitumor activity of 4-HPR was also observed in medulloblastoma (31), human pancreatic cancer (32), chronic myeloid leukemia (33), and in a lung cancer xenograft mouse model (34). Moreover, Sogno et al reported that 4-HPR is effective in inhibiting angiogenesis (35).

In our study, we compared the effect of 4-HPR with ATRA. 4-HPR potently inhibited the growth of and colony formation of HepG2 cells, and suppressed cell migration. Compared with ATRA, the inhibitory effect of 4-HPR on cell growth and colony formation was achieved at a lower concentration (5 µM). We observed that the IC50 of HepG2 cells was approximately 12.5 µM in the cell viability assay. Notably, pediatric neuroblastoma patients who received oral doses of 4-HPR achieved a blood serum concentration of 12.9 µM (30).
the effective concentrations of 4-HPR in HepG2 cells implies that 4-HPR may be a candidate for liver cancer therapy. When HepG2 cells were treated with 4-HPR for only 24 h, a significantly slower migration of the cells was observed; however, no effect could be detected using ATRA for the same amount of time. ATRA required much higher concentrations and longer incubation times to achieve the same inhibition rates as 4-HPR. These results indicate that much lower amounts of 4-HPR are required for growth suppression and migration of liver cancer cells, and hence, the drug may be applied clinically with less toxicity.

The main antitumor activity of 4-HPR is the induction of apoptosis by retinoic acid receptor-dependent or -independent mechanisms (8,36). 4-HPR also reduces the plasma concentrations of retinol and retinol binding protein (37). To determine whether the cytotoxic effect of 4-HPR in HepG2 cells is due to the induction of apoptosis, we analyzed the expression levels of proteins involved in this process; however, we found no marked changes in such proteins upon 4-HPR treatment (data not shown). A previous report is consistent with our results (20); hence, the mechanism of 4-HPR action might be different in HepG2 cells. The p38-MAPK pathway has been reported to mediate various cellular behaviors that are closely related to tumor initiation and progression (38). Nevertheless, the regulation of p38-MAPK in tumor development is complicated and controversial, involving responses of various cells and cancer types (39). In this study, we found that 4-HPR inhibited the migration of HepG2 cells by significantly inducing the activation of p38-MAPK. When the p38-MAPK inhibitor SB203580 was added to the culture system preceding 4-HPR treatment, the inhibitory effect on migration was ameliorated. This allows for the preliminarily conclusion that 4-HPR inhibits the migration of HepG2 cells via stimulation of the p38-MAPK pathway.

MLCK confers the rat pituitary adenoma cells with a slow and directional motility (40), and phosphorylation of MLC markedly improves the invasion and migration ability of gastric cancer cells (41). Leiomyosarcoma patients with high expression of MLCK or p-MLC have shorter life spans than the patients with low expression of these proteins (42). In our study, treatment of HepG2 cells with 4-HPR for 48 h resulted in the downregulated expression of MLCK. Simultaneously, the expression of MLCK and p-MLC were significantly suppressed. When the activation of MLCK was inhibited by ML-7, cell migration was retarded and accompanied by reduced p-MLC levels. In addition, ML-7 in combination with 4-HPR or ATRA enhanced the inhibitory effect on the migration of HepG2 cells, compared to 4-HPR or ATRA alone. Based on our results, 4-HPR decreased the proliferation and migration of HepG2 cells in association with activation of p38-MAPK and inhibition of MLCK. We measured the protein expression of MLCK and p-MLC after SB203580 pretreatment and found that p-MLC increased compared to 4-HPR treatment alone. Meanwhile, in cells treated with ML-7 and 4-HPR, p-p38 was upregulated compared to treatment with 4-HPR alone. These results provide evidence of a reciprocal cross-talk between MLCK and p38-MAPK.

**Figure 7.** Effect of 4-HPR and ML-7 on the expression of MLCK, E-cadherin, F-actin and phosphorylated MLC and p38. (A) ML-7 was used to treat HepG2 cells. The phosphorylation of MLC was evidently decreased. In addition, the protein expression of MLC was decreased by ML-7 combined with 4-HPR or ATRA. All values are presented as mean ± SD. n=3, *P<0.05, †P<0.05 compared with DMSO group. (B) Phosphorylated (p)-p38 was increased by ML-7. All values are presented as mean ± SD. n=3, *P<0.05 compared with DMSO group, †P<0.05 compared with ATRA group and ○P<0.05 compared with 4-HPR group. (C) The protein expression of E-cadherin was increased and F-actin was decreased by ML-7. All values are presented as mean ± SD. n=3, *P<0.05, †P<0.05 compared with DMSO group. For A-C: Lane 1, DMSO; lane 2, ML-7; lane 3, ATRA; lane 4, ATRA+ML-7; lane 5, 4-HPR; lane 6, 4-HPR+ML-7. 4-HPR, fenretinide; ATRA, all-trans retinoic acid; ML-7, a specific inhibitor of MLCK.
Erk-MAPK governs the cell movement via p-MLC (47). We also found an altered expression of p-p38 preceded that of p-MLC after 4-HPR treatment (data not shown). Thus, we presently believe that 4-HPR generates its effects on HepG2 cells via inhibiting MLCK activation through the p38-MAPK signaling pathway. The exact mechanism of HepG2 cell migration involving p38-MAPK via p-MLC, however, needs to be further investigated.

Collectively, the present study using the HepG2 cell line demonstrated a marked potential effects of 4-HPR on liver cancer. 4-HPR potentially inhibits the biological behaviors involved in liver cancer metastasis, and may be an alternative therapeutic agent for its prevention. Despite these findings, further studies on the specific targets of 4-HPR in these signaling pathways are required, as well as therapeutic experiments using in vivo models are warranted.

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Availability of data and materials
The datasets used during the present study are available from the corresponding author upon reasonable request.

Authors' contributions
YW, WW and LZ conceived and designed the study. LZ performed the experiments. HL provided the cell line. QZ and HL gave experimental guidance. LZ and DH analyzed the experimental data and wrote the paper. YW reviewed and edited the manuscript. All authors read and approved the manuscript and agree to be accountable for all aspects of the research in ensuring that the accuracy or integrity of any part of the work are appropriately investigated and resolved.

Ethics approval and consent to participate
All experimental protocols were approved by the Institutional Review Board of Anhui Medical University (Hefei, China).

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

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