A novel angiogenesis inhibitor impairs lovo cell survival via targeting against human VEGFR and its signaling pathway of phosphorylation

YM Zhang¹, BL Dai¹, L Zheng¹, YZ Zhan¹, J Zhang¹, WW Smith², XL Wang¹, YN Chen¹ and LC He*¹

Colorectal cancer represents the fourth commonest malignancy, and constitutes a major cause of significant morbidity and mortality among other diseases. However, the chemical therapy is still under development. Angiogenesis plays an important role in colon cancer development. We developed HMQ18–22 (a novel analog of taspine) with the aim to target angiogenesis. We found that HMQ18–22 significantly reduced angiogenesis of chicken chorioallantoic membrane (CAM) and mouse colon tissue, and inhibited cell migration and tube formation as well. Then, we verified the interaction between HMQ18–22 and VEGFR2 by AlphaScreen P-VEGFR assay, screened the targets on angiogenesis by VEGF Phospho Antibody Array, validated the target by western blot and RNAi assay in lovo cells. We found HMQ18–22 could decrease phosphorylation of VEGFR2 (Tyr1214), VEGFR1 (Tyr1333), Akt (Tyr326), protein kinase Cα (PKCα) (Tyr163) and phospholipase-Cγ1 (PLCγ1) (Tyr771). Most importantly, HMQ18–22 inhibited proliferation of lovo cell and tumor growth in a human colon tumor xenografted model of athymic mice. Compared with normal lovo cells proliferation, the inhibition on proliferation of knockdown cells (VEGFR2, VEGFR1, Akt, PKCα and PLCγ1) by HMQ18–22 decreased. These results suggested that HMQ18–22 is a novel angiogenesis inhibitor and can be a useful therapeutic candidate for colon cancer intervention.

**Cell Death and Disease** (2012) 3, e406; doi:10.1038/cddis.2012.145; published online 11 October 2012

**Subject Category:** Cancer

Angiogenesis is the process of new capillary formation from pre-existing blood vessels and plays an important role in the growth and spread of cancer. Previous studies suggest that angiogenesis mediates tumor growth and metastases, and that inhibition of new blood vessel formation by antiangiogenesis agents is an approach to fight cancer. As a target for cancer chemotherapy, blocking angiogenesis could be a strategy to arrest tumor growth. A number of antiangiogenic agents have been reported to inhibit tumor angiogenesis such as sorafenib and sunitinib. Sorafenib inhibits angiogenesis via inhibition of VEGFR2, VEGFR3 and/or platelet-derived growth factor receptor-β (PDGFR-β). Sunitinib is a tyrosine kinase (TK) inhibitor that targets VEGFR1 and VEGFR2, PDGFR-α and β, c-KIT, and the FLT-3 and RET kinases. Vascular endothelial growth factor receptor (VEGF) is one of the most important regulators of angiogenesis and a key drug target in anticancer treatment. VEGF binding to its receptor (VEGFR) leads to cell proliferation and new vascular formation by TK pathway. The VEGF/VEGFR pathway becomes an attractive target for anticancer drug design. Ligands binding VEGFRs in the cell membrane induces receptor dimerization and activation of the latter, and autophosphorylation of specific tyrosine residues within the dimeric complex. VEGF primarily binds to three transmembrane receptors with intracellular TK activity: VEGFR1 (Flt-1), VEGFR2 (Flk-1) and VEGFR3 (Flt-4). Although both VEGFR1 and VEGFR2 are expressed in the vascular endothelium, the angiogenic activities of VEGFs are transduced mainly through VEGFR2. VEGFR2 is the predominant receptor in angiogenic signaling. Its activation regulates endothelial cell migration, proliferation, differentiation, survival as well as vessel permeability and dilation. Activation of VEGFR1 mediates the growth and survival effects of VEGF, so VEGFR1 may function as a negative regulator of angiogenesis by binding VEGF and preventing its binding to VEGFR2. VEGFR3 is predominantly expressed on lymphatic endothelial cells. Recent research on the VEGF/VEGFR pathway has led to the development of novel antiangiogenic agents. Clinical trials have shown inhibitors to this pathway are effective in reducing tumor size, metastasis and blood vessel formation.

There are various molecular players and signaling cascades involved in the VEGF/VEGFR pathway, such as the phosphatidylinositol 3-kinase (PI3K)/AKT, Ras/Raf/mitogen-activated protein kinase (MAPK) and phospholipase-Cγ1/protein kinase C (PLCγ1/PKC) pathway. These signaling pathways regulate important cellular functions including cellular proliferation, migration, angiogenesis and apoptosis.

HMQ18–22 (Figure 1) was a novel derivative of taspine. Taspine was originally identified from a screen of *Radix et Rhizoma leonticis* (Hong Mao Qi in Chinese) using cell

1Institute of Materia Medica, School of Medicine, Xi’an Jiaotong University, Xi’an, China and ²Department of Pharmaceutical Sciences, School of Pharmacy, University of Maryland, Baltimore, MD, USA

*Corresponding author: LC He, School of Medicine, Xi’an Jiaotong University, No. 76, Yanta Weststreet, #54, Xi’an, Shaanxi Province 710061, People’s Republic of China. Tel: +86 29 82656264; Fax: +86 29 8265 5451; E-mail: helc@mail.xjtu.edu.cn

**Keywords:** HMQ18–22; angiogenesis; VEGFR; inhibition; signaling pathway

**Abbreviations:** VEGF, vascular endothelial growth factor; VEGFR, VEGF receptor; FBS, fetal bovine serum; ELISA, enzyme-linked immunosorbent assay; WB, western blot analysis; PCR, polymerase chain reaction; CAM, chicken chorioallantoic membrane

Received 15.5.12; revised 14.8.12; accepted 21.8.12; Edited by A Stephanou
membrane chromatography. We previously found that taspine could enter cells and had good affinity to overexpressed VEGFR cell membrane chromatography model and displayed anticancer and antiangiogenesis properties and therefore we used it as a leading compound for anticancer agents development with the aim to increase activity and solubility. A series of ring-opened and biphenyl derivatives have been designed and synthesized using dissection strategies. Among the derivatives, we found HMQ18–22 had good activity and inhibition on lovo cell. In the present study, we investigated the effects and mechanisms of HMQ18–22 on angiogenesis using tissue and cell model in vitro and mouse models in vivo.

Results

HMQ18–22 inhibited the angiogenesis of CAM and mouse colon tissue. To test the effects of HMQ18–22 on chorioallantoic membrane (CAM) blood vessels, we treated them with HMQ18–22 (0, 2.5, 5, 10 μg/per egg) for 72h, and found HMQ18–22 inhibited the CAM angiogenesis in a dose-dependent manner (Figures 2a and b). The mean IC50 for large vessels, small vessels and capillaries was 3.42 μg/egg. In the negative group, rich blood vessels grew and the density and area of the CAM blood vessels increased with incubation time increased. HMQ18–22 inhibited the new capillary vessels formation.

To assess whether HMQ18–22 alters microvessels growth, we used rat colon tissues as a model. We dissected rat colon tissues and subcultured the resultant cells. Microvessel outgrowths of colon tissue in matrigel were increased with the culture time increased (Figure 2ci–c5). Many new microvessels grew after 5 days, and the density and area of the blood vessels increased with incubation time increased. HMQ18–22 inhibited the new capillary vessels formation.

To assess whether HMQ18–22 alters microvessels growth, we used rat colon tissues as a model. We dissected rat colon tissues and subcultured the resultant cells. Microvessel outgrowths of colon tissue in matrigel were increased with the culture time increased (Figure 2ci–c5). Many new microvessels grew after 5 days, and the density and area of the blood vessels increased with incubation time increased. HMQ18–22 inhibited the new capillary vessels formation.

HMQ18–22 inhibited cell migration and tube formation. To test the effect of HMQ18–22 on cell migration and tube formation, HUVEC cells were induced by VEGF and treated with vehicle or HMQ18–22. HUVEC cells were incubated on matrigel for 48h with VEGF, forming an extensive and enclosed network of tubes. HMQ18–22 significantly decreased the number of the tube structure (Figure 3a) at the concentrations of 0.5–8.0 μmol/l, respectively. Furthermore, HMQ18–22 also inhibited the HUVEC cell migration (Figure 3b). In addition, tube formation of colon tissue in the fibrin matrices showed HMQ18–22 also inhibited the number of the tube structure compared with untreated control tissues (Figures 3c and d).

HMQ18–22 decreased phosphorylation of VEGFR2, VEGFR1, Akt, PKCα and PLCγ-1 involved in angiogenesis. HMQ18–22 decreased cell survival in lovo and HUVEC cells, and showed dose-dependent inhibition on cell growth (Figure 4a). It suggests that HMQ18–22 displays antiproliferative effect on the lovo and HUVEC cells. The lance assay was used to assess the effects of HMQ18–22 on VEGFR2 kinase activity. The optimized used concentrations of reaction system were as follows: VEGFR2 kinase 0.003767 ng/ml, ATP 1.332 μM and substrate 121.4 nM, respectively. The IC50 of HMQ18–22 on VEGFR2 kinase activity was over 5000 nM, suggesting that HMQ18–22 did not alter VEGFR2 kinase activity effectively. The AlphaScreen assay, an effective method for screening a broad range of targets, was used to determine the effect of HMQ18–22 on VEGFR. We found that HMQ18–22 acted on VEGFR by inhibiting its phosphorylation (Figure 4b).

To further assess whether HMQ18–22 alters the downstream signaling events of VEGFR, a phospho-specific antibody microarray (PVE185) targeting the VEGF Phospho signaling pathway was used. This antibody array included 190 VEGF-related proteins (85 pairs), each with six replicates (raw data in Supplementary Table 2). The paired antibodies for the same (but unphosphorylated) target sites were also included in the array to allow determination of the relative level of phosphorylation. Using a cutoff ratio of 0.88, we identified five pairs of phosphorylation sites of tyrosine, namely VEGFR2 (Tyr1244), VEGFR1(Tyr1335), Akt(Tyr326), PKCα (Tyr67) and PLCγ-1(Tyr71), were decreased by HMQ18–22 in lovo cell.
To validate the antibody array results, we tested the phosphorylation status of VEGFR2 (Tyr1214), VEGFR1 (Tyr1333), Akt (Tyr326), PKCζ (Tyr657) and PLC-γ1 (Tyr771) in Lovo cell by western blot analysis. Phosphorylation of Raf1 (Tyr331) was not affected from the results of antibody array, thus being used as the negative control protein.

HMQ18–22 significantly reduced the phosphorylation of VEGFR2 (Tyr1214), VEGFR1 (Tyr1333), Akt (Tyr326), PKCζ (Tyr657) and PLC-γ1 (Tyr771) (Figure 4c). Therefore, the results of microarray with western blotting verified that VEGFR2 (Tyr1214), VEGFR1 (Tyr1333), Akt (Tyr326), PKCζ (Tyr657) and PLC-γ1 (Tyr771) were the targets of HMQ18–22.
To further assess the roles of VEGFR2, VEGFR1, Akt, PKCα and PLCγ-1 phosphorylation in cancer cell growth, siRNA targeting these proteins were transfected into cells and followed by HMQ18–22 treatment for 48 h. IC50 of HMQ18–22 in survival of lovo cells transfected with siRNA targeting VEGFR2, VEGFR1, Akt, PKCα and PLCγ-1 was 59.66, 41.67, 88.79, 49.98 and 24.48 μM (Figure 4d), respectively, while IC50 in untransfected control cells was only 14.01 μM (Figure 4a).

HMQ18–22 inhibited tumor growth of xenografted model in athymic mice. To further assess the effects of HMQ18–22 on tumor growth in vivo, human tumor models xenografted in athymic mice were used (Figure 5a). HMQ18–22 significantly inhibited tumor growth in lovo xenografted athymic mice. Compared with the untreated group, HMQ18–22 significantly inhibited tumor growth at a rate of 21.25 and 44.84%, respectively. Furthermore, there was no change in athymic mice body weight during the experiment (the mean body weight at start was 21.14 ± 1.32g, 21.50 ± 0.98g and 21.71 ± 1.16g in the control group, 50.0 mg/kg and 100.0 mg/kg group, and at end was 25.51 ± 2.95g, 25.02 ± 3.26g and 25.92 ± 4.65g, respectively), which suggested that the dose of HMQ18–22 did not have a toxic effect on athymic mice. We further tested the phosphorylation status of VEGFR2(Tyr1214), VEGFR1 (Tyr1333), Akt(Tyr326), PKCα (Tyr657) and PLCγ-1(Tyr771) in lovo tumor tissue of athymic mice by western blot analysis. We found that HMQ18–22 significantly decreased the phosphorylation of VEGFR2(Tyr1214), VEGFR1 (Tyr1333), Akt(Tyr326), PKCα (Tyr657) and PLCγ-1(Tyr771) in tumor tissues compared with the untreated group (Figure 5b). Figure 5c indicated that the proteins phosphorylation of VEGFR2, VEGFR1, Akt, PKCα and PLCγ-1 in the HMQ18–22-treated group were significantly downregulated in a dose-dependent manner compared with the negative control (P<0.05). The results were in line with the corresponding data in lovo cells.
Colorectal cancer represents the fourth commonest malignancy, and constitutes a major cause of significant morbidity and mortality among other diseases. In this study, the novel compound, HMQ18–22 (a novel taspine analog) significantly reduced angiogenesis of CAM and mouse colon tissue, inhibited cell migration and tube formation and decreased...
phosphorylation of VEGFR2, VEGFR1, Akt, PKCa and PLCg1. Most importantly, HMQ18–22 inhibited tumor cell growth in xenografted model in athymic mice. These results suggest that HMQ18–22 can be a useful therapeutic candidate for colon cancer intervention.

HMQ18–22 was synthesized by our group for cancer drug development. Our data showed that HMQ18–22 significantly blocked angiogenesis using CAM model. Angiogenesis plays a vital role in growth, intravasation and metastatic spread of cancer. Inhibition of angiogenesis provides a good chance of preventing cancer from becoming malignant. Tumor angiogenesis shows a markedly increasing proliferation of endothelial cell and has significant functional and structural differences in the vascular plexus. In solid tumors, angiogenesis is well characterized as a critical step for growth, invasion and metastasis.\textsuperscript{3,22,23} To further validate our results of HMQ18–22, we established a new colon angiogenesis tissue model. In this model, the tissues cultured from colon were grown in a three-dimensional matrix (usually fibrinogen or matrigel). In this model, new capillary vessels erupted from the tissue in matrigel, and new erupted endothelial cells grew from tissues in fibrinogen proliferated and formed tube. These results generated by total nutritive index difference between fibrinogen and matrigel. The different type from the two culture mediums in the colon tissue model could be used to assess angiogenesis and tube formation at tissue and organ Level. HMQ18–22 inhibited the sprouting of new vessel from the tissue and tube formation by the erupted endothelial cells, further supporting the notion that HMQ18–22 suppressed the neovascularization.

The VEGF/VEGFR pathway was an important regulator in tumor angiogenesis. It has led to the development of novel antiangiogenic agents. The various downstream signaling molecular players, such as PI3K/AKT, MAPK and PLCg/ PKC, have the specific functions including cellular proliferation, migration, angiogenesis and apoptosis.\textsuperscript{16–18} Phospholipase Cg1 (PLCg1), a very important member of phospholipase C (PLC) families, is upregulated in many cancer tissues and cancer cell lines and has been found to participate in many physical processes such as cell proliferation, motility, differentiation, cytoskeleton rearrangement and so forth.\textsuperscript{24,25} There is an intimate relation between PLCg1 and PKCa. Activated PLCg1 can catalyze the hydrolysis of phosphatidylinositol 4,5-bisphosphate (PIP2) to inositol 1,4,5-triphosphate (IP3) and diacylglycerol (DAG). DAG and IP3 are second messengers to activate PKC. A decrease in the extent of tyrosine phosphorylation of PLCg1 has also been proved to be positively regulated by PKCa.\textsuperscript{26,27} So, downregulation of surface receptor expression represents a mechanism in which decreased PLC might block PKC activation. Furthermore, in the signaling pathway, PI3K/AKT mediated cell proliferation, migration, invasion, angiogenesis and metastasis.\textsuperscript{28} Akt is a kinase in the signal transduction, making it a potential target. The Akt/PKB kinase has been implicated in the genesis and/or progression of numerous human tumors, because AKT has a wide range of downstream targets that regulate endothelial cell functions such as migration, growth, proliferation and apoptosis.\textsuperscript{29} HMQ18–22 acted on VEGFR and inhibited the phosphorylation of VEGFR by p-VEGFR AlphaScreen assay. However, VEGFR kinase activity was not affected by HMQ18–22, indicating that the inhibition of VEGFR phosphorylation is not due to inhibition on kinase of VEGFR2 Kinase. Using a phospho-antibody array to screen the potential target in the VEGF/VEGFR pathway, we found that

**Figure 5** HMQ18–22 inhibited tumor growth in nude mice bearing human colon cancer xenografts. \( (a) \) The representative xenografts of human colon cancer in mice. \( (b) \) HMQ18–22 decreased the phosphorylation of VEGFR2 (Tyr\textsuperscript{1214}), VEGFR1 (Tyr\textsuperscript{1333}), Akt (Tyr\textsuperscript{326}), PKCa (Tyr\textsuperscript{353}) and PLCg1 (Tyr\textsuperscript{771}) in the tumor tissues by western blot analysis. \( (c) \) Quantitation data of \( (b) \). Data were expressed as mean values \( \pm \) S.D. \( (n = 3) \). **P < 0.01 versus the untreated control.**
the phosphorylation of VEGFR2, VEGFR1, Akt, PKCζ and PLCζ-1 were inhibited by HMQ18–22. We further confirmed the array data by western blots using antiphosphorylation antibodies against VEGFR2, VEGFR1, Akt, PKCζ and PLCζ-1 in culture cells and in vivo human colon cancer mouse models. Knockdown of VEGFR2, VEGFR1, Akt, PKCζ or PLCζ-1 by siRNA significantly attenuated tumor inhibitory effects of HMQ18–22. Taken together, these results suggest that HMQ18–22 inhibits tumor angiogenesis and tumor growth by downregulating the phosphorylation signaling of VEGFR2, VEGFR1, Akt, Raf1, PKCs and PLCζ-1. A compound exerts its action by entering the cell or acting on the cell membrane receptor with good affinity. In previous reports, taspine could enter the cell or act on the cell membrane receptor determined by HPLC-MS and cell membrane chromatography.30 HMG18–22 is a derivative of taspine, so HMG18–22 should exert its inhibition and downregulate the phosphorylation signaling by entering the lovo cells and being with good affinity with VEGFR-1 and VEGFR-2.

In conclusion, our results demonstrate that HMQ18–22 might be a novel angiogenesis inhibitor that reduces angiogenic responses in vivo and in vitro by blocking VEGFR signaling pathways. The effectiveness of HMQ18–22 in disrupting colorectal tumor growth provides a promising anticancer agent for further clinical trial for HMQ18–22 to colorectal cancer treatment.

Materials and Methods
Reagents. HMQ18–22 was from the Research and Engineering Center for Natural Medicine, Xi’an Jiaotong University. Trypsin and fibrinogen were from Sigma (St Louis, MO, USA). Human VEGF was from Peprotech Asia (Rehovot, Israel). WST, protease inhibitor cocktail and phosphatase inhibitor cocktail were from Roche (Roche Tech., Mannheim, Germany). Fibrin matrices were from Finch (San Jose, CA, USA). VEGFR2 kinase was from Genzyme Biosciences (Kobe, Japan). HTRF VEGFR2 kinase kit was purchased from Cobiol (Codiert, France). P-VEGFR2 AlphaScreen SureFire kit was purchased from Perkin-Elmer (Boston, MA, USA). Anti-phospho-VEGFR2 (Tyr1175), anti-phospho-VEGFR1 (Tyr1335), anti-phospho-Akt (Tyr422), anti-phospho-PKCζ (Tyr85), anti-phospho-PLCζ-1 (Tyr771) and anti-phospho-Raf1 (Tyr340/Tyr341) were purchased from Cell Signaling (Cell Signaling Tech., Danvers, MA, USA). Rabbit anti-60 kDa GAPDH was purchased from Santa Cruz Biotech (Santa Cruz, CA, USA). Rabbit anti-mouse IgG, goat anti-rabbit IgG, BCA protein assay reagent kit and enhanced chemiluminescent (ECL) plus reagent kit were obtained from Pierce (Pierce Biotech, Rockford, IL, USA). Total RNA extracted kit was from Fastagen (Fastagen, China). Revert AID First-strand cDNA synthesis kit was from Fermentas (Hanover, Lithuania). RNase A was from Fermentas. Other reagents used were analytical grades.

Animals and cell culture. In all, 6-8 weeks old male mice were used for all experiments and each procedure was approved by the regional authorities according to China animal-care regulations. Lovo cells from Shanghai Institute of Cell Biology in the Chinese Academy of Sciences were maintained in RPMI-1640 with 10% FBS. HUVEC from ATCC were cultured in F-12K media supplemented with 0.1 mg/ml heparin, 0.5 mg/ml endothelial cell growth supplement (ECGS) and 10% FCS.

CAM assay. CAM was prepared as described.31,32 Briefly, a circular window was opened aseptically on the egg shell, and the eggs were replaced into the incubator for 24 h at 10 μl HMQ18–22 was added to the CAM surface in every egg. Doses of 2.5, 5 and 10 μg/egg were used here. At least 10 eggs were used for each sample dose. The embryos were further incubated for 72 h after administration. The anti-angiogenic response was assessed by counting. Five randomly chosen fields were evaluated for each specimen. The total vessels number of large vessels, small vessels and capillaries in the fields were counted, and the mean values ± S.D. were calculated. The large, small vessels and capillaries meant the main blood vessels, vessels grown from main blood vessels and vessels grown from small vessels. The inhibitory effect on blood vessels could be observed by comparing the vascular change between the HMQ18–22 group and the negative control group.

Angiogenesis and tube formation of mouse colon tissue. An assay of mouse colon tissue co-cultured was used as the in vitro angiogenesis model. For preparation of fibrin matrices and matrigel, 3 mg/ml solution of fibrinogen containing 300 μg/ml ε-aminocaproic acid (in DMEM) and matrigel (in DMEM) was incubated on ice for 10 min. A sandwich structure was prepared in the 24-well plate (Costar, Corning, NY, USA). The colon tissues of mouse were cut into 0.5–1.0 mm3 tablets, placed into the sandwich structure, and covered with DMEM medium containing 20% bovine serum and HMQ18–22 (1.0, 4.0, 16.0 μmol/l) or vehicle alone. The sprouting vessels and tube formation were recorded and analyzed. The sprouting vessels and tube formation were assessed by counting. Five randomly chosen fields were evaluated. The total sprouting vessels and tube structures in the fields were counted, and the mean values ± S.D. were calculated.

Tube formation assay. A 48-well plate coated with 0.2 ml matrigel per well was added to solidify at 37°C for 1 h. Each well was seeded with 1 × 104 HUVEC and cultured in DMEM containing 50 ng/ml VEGF at various concentrations (1.0, 2.0, 4.0, 8.0 μmol/l) of HMQ18–22 or vehicle alone for 48 h. The enclosed networks of tubes were photographed from five randomly chosen fields under a microscope.

Migration assay. A cell migration assay was performed using a transwell system, which allows cells to migrate throughout an 8-μm pore size polycarbonate membrane of millicell.33,34 HUVEC cells were plated (104 cells well) in medium containing HMQ18–22 at concentrations of 0.5, 1.0, 2.0, 4.0, 8.0 μmol/l in the upper chamber of a 12-well plate. The lower chamber was filled with 1.5 ml medium containing 10% FBS. After 48 h, cells remaining on the upper surface of the membrane were scraped and the cells on the lower surface of the membrane were fixed with cold methanol for 15 min and stained with 0.2% crystal violet. Cells that had migrated to the bottom of the membrane were visualized and counted using an inverted microscope. For each repetition, cells in five randomly selected fields were counted and averaged. Data were expressed as a ratio to the untreated group.

Lance assay for VEGFR2 kinase activity. VEGFR2 kinase was determined by lance assay. In all, 2 μl kinase and 2 μl substrate were added to the 384-well plate, and HMQ18–22 at various concentrations was then added to the assay plate. In all, 2 μl ATP was added and the reaction was allowed to proceed at 37°C for 30 min. The TK-antibody labeled with Eu3+ -cryptate and streptavidin-XL665 was then added with EDTA to detect the phosphorylated product at room temperature for 1 h. Then, the fluorescence of the resulting solution was measured at 665 and 615 nm using the plate reader of Perkin-Elmer Victor 5. The kinase activity was expressed by the ratio of A665 × 10/7A651.

AlphaScreen P-VEGFR assay. AlphaScreen assays for p-VEGFR proteins were carried out using SureFire assay kits (Perkin-Elmer). Lovo cells were seeded at a density of 1 × 105 cells per well in 96-well plates. HMQ18–22 at various concentrations (2, 4, 8, 16 μmol/l) was added to the wells for 48 h. The cells were treated with 100 ng/ml VEGF 15 min after serum starving overnight, lysed and agitated. Then, the reactions were carried out according to the assay kit instructions. The plates were read on an Enspire by Reader (Perkin-Elmer) using standard AlphaScreen settings.35,36

VEGF phospho antibody microarray analysis. The expression profile of VEGF phospho-related proteins was detected and analyzed using a human VEGF Phospho Antibody Array kit (PVE185, Fullmoon Biosystems, Sunnyvale, CA, USA). Protein microarray analysis was carried out using the protocol provided. Briefly, protein was extracted, lysate was purified and then the protein was labeled by biotinylation. The resulting biotin-labeled proteins were diluted 1:20 in coupling solution before being applied to the array for conjugation. To prepare the antibody
microarray, it was blocked for 30 min, then dried and the array was incubated with the biotin-labeled cell lysates at 4 °C. After the array slide was washed three times, the conjugation-labeled protein was detected by Cy3-Streptavidin. The analyzed results were expressed by the ratio of phosphorylation/unphosphorylation.27,28

Cell viability assay. Lovo cells (1 × 10^5) and HUVEC (5 × 10^4) were cultured in 6-well microtiter plates, and fresh medium with or without HMQ18–22, was added for 48 h. Cell proliferation reagent WST-1 was added and incubated at 37 °C and 5% CO2 for 1 h. Absorbance was then measured at 440 nm with a microplate reader (Bio-Rad, Hercules, CA, USA).

Transient transfection of siRNA. For in vitro knockdown experiments, a smart pool of double-stranded siRNA against VEGFR2, VEGFR1, Raf, PLC, and PKCα as well as nonspecific siRNA was obtained from Shanghai GenePharma for transfection, siRNA was delivered at a final concentration of 50 nM using Lipofectamine 2000 reagent (Invitrogen, Carlsbad, CA, USA) according to the manufacturer’s instructions.29 The sense and antisense sequences were in Supplementary Table 1. We incubated the cells for 24 h to allow knockdown of VEGFR2, VEGFR1, Raf, PLC, and PKCα. These cells were used for proliferation assays.

Western blot analysis. The tissues isolated from nude mice treated with or without HMQ18–22 and lovo cells treated with or without HMQ18–22 for 48 h were prepared by extracting proteins with RIPA lysis buffer containing protease inhibitor cocktail and phosphatase inhibitor cocktail on ice. Cell lysates were analyzed for western blot analysis with primary antibodies [VEGFR2(Tyr1175)], VEGFR1 (Tyr1165), Akt (Tyr308), PKCα (Tyr657), PLCγ1 (Tyr1576) and Raf1(Tyr259)] followed by enhanced chemiluminescence.40 The protein expression was analyzed using Quantity one, 1-D analysis software (Version 4.4, Bio-Rad).

Acknowledgements. This work was supported by National Natural Science Foundation of China (Grant No. 81001447).

Conflict of Interest. The authors declare no conflict of interest.
40. Kortylewski M, Swiderski P, Hermann A, Wang L, Kowolik C, Kujawski M et al. In vivo delivery of siRNA to immune cells by conjugation to a TLR9 agonist enhances antitumor immune responses. Nat Biotechnol 2009; 27: 925–932.

41. Kim HJ, Oh GS, Lee JH, Lyu AR, Ji HM, Lee SH et al. Cisplatin ototoxicity involves cytokines and STAT6 signaling network. Cell Res 2011; 21: 944–956.

Supplementary Information accompanies the paper on Cell Death and Disease website (http://www.nature.com/cddis)