

HSP90 expression and its association with wighteone metabolite response in HER2-positive breast cancer cells

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Abstract. It is well known that heat shock protein 90 (HSP90) overexpression is correlated with poor prognosis and chemo-resistance in human malignant cancers. At the same time, wighteone, or 6-prenyl-5,7,4'-trihydroxyisoflavone, a major isoflavone component of the ornamental tall tree *Erythrina suberosa*, has been demonstrated to exhibit a potent anti-proliferative effect on human leukemia HL-60 cancer cell lines. In this study, the effects of wighteone on the proliferation of HER2-positive breast cancer cells were investigated, and the action mechanism was explored. MCF-7 HER2-positive breast cancer cells were treated with various concentrations of wighteone. The growth inhibitory rate of the cells was calculated by MTT assay, apoptosis was detected by flow cytometry, and the expression level of HSP90 was assessed by western blot analysis. The addition of wighteone at concentrations ranging from 1-10 g/ml in the medium for 48 h had a marked inhibition on the proliferation of HER2-positive cancer cell lines. The growth inhibitory rates with 0.5, 2 or 8 mM wighteone were significantly higher compared with the control group. Apoptosis in the wighteone-treated cells was also significantly higher compared with the control group. The expression level of HSP90 in the wighteone group was significantly lower than that in the control group. Our findings demonstrated that wighteone effectively inhibited the proliferation of HER2-positive cancer cell lines, and this is considered to be the result of downregulating HSP90 receptor and downstream signaling.

Introduction

Breast cancer is one of the most common and significant diseases affecting females. Breast tumors are comprised of phenotypically diverse populations of breast cancer cells. Current estimates

indicate that one in eight females in American who reach the age of 95 is likely to develop breast cancer (1). The treatment of advanced breast cancer is often unsuccessful and may cause disfigurement, making early detection a high priority in the medical management of the disease. The level of morbidity and high incidence of this disease is notable. Within the next two decades the incidence of cancer is expected to rise worldwide. It is noted that the management of cancer is still not under control and new effective drugs are lacking (2). Proteins associated with the human epidermal growth-factor receptor kinase (ERBB or HER) signaling network have proven to be useful targets for diagnostic imaging with radioimmunoconjugates due to their overexpression in various cancer phenotypes. In particular, the overexpression of the HER2/neu (also known as ERBB2) was revealed to be associated with increased tumor aggression, metastatic potential and poor prognosis for disease-free survival in patients with breast, colorectal, ovarian, lung, prostate and salivary gland tumors (3,4).

HER2/neu was revealed to be a key target for anticancer drugs due to its intrinsic involvement in the phosphatidylinositol-3-kinase-Akt/protein kinase B and the mitogen-activated protein kinase (MAPK) pathways. These pathways suppress apoptosis and promote tumor cell survival, gene transcription, angiogenesis, cellular proliferation, migration, mitosis and differentiation. There are three notable types of anti-HER2/neu therapeutics; namely, monoclonal antibodies (mAbs) directed against extracellular ligand-binding and dimerization epitopes, tyrosine-kinase (TK) inhibitors and heat shock protein 90 (HSP90) inhibitors. Examples of each therapeutic type include pertuzumab and trastuzumab (which block dimerization and suppress signaling by binding to extracellular domains II and IV, respectively), the HER2/neu TK inhibitor lapatinib, and HSP90 inhibitors including geldanamycin derivatives, SNX-5422, NVP-AUY922, BIIB021 and PU-H71 (5-10). HSP90 is a molecular chaperone protein essential for the function of the multiple growth and survival pathways required for the maintenance and progression of cancer (11). In breast cancer in particular, the overexpression of HSP90 is associated with poor outcome and decreased survival (12,13). Trastuzumab and related mAb fragments have been radiolabeled with a wide range of radionuclides, and quantitative immune positron emission tomography (PET) imaging has been employed to assess the effect of Hsp90 inhibitors on the expression levels of HER2/neu (14-18). The quantification of changes in HER2/neu

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expression in response to HSP90 treatment has the potential to facilitate patient-specific dose regimes.

During the last few years, the focus of drug development has shifted to natural chemotherapeutic agents from plants, which may be further modified to enhance their potential and reduce their side effects. Natural products have been particularly useful in cancer treatment, and if biologicals and vaccines are discounted, then 75% of approved chemotherapeutic agents are natural agents (19). *Erythrina* is a genus of flowering plants in the pea family, Fabaceae, with ~130 species reported. It is distributed worldwide in tropical and subtropical regions in addition to certain more temperate regions, and ~50% of its varieties have been studied. The alkaloids isolated from various species of *Erythrina* demonstrate hypotensive, anticonvulsant, hypnotic and analgesic properties, among others (20). *Erythrina* plant species are widely used in folk medicine to treat health conditions including agitation, insomnia, anxiety and inflammation (21,22). A previous study has revealed that an alcoholic extract of the stem bark of *E. suberosa* induces apoptosis in human promyelocytic leukemia HL-60 cells (23). The aim of the current study was to investigate the anticancer potential of the flavonoid wighteone isolated from the stem bark of *E. suberosa*, and to study HSP90 expression and its correlation with wighteone metabolite response in HER2-positive breast cancer cells.

Materials and methods

Reagents and cell lines. The breast cancer cell line MCF-7 was provided by the Breast Cancer Research Department of the Inner Mongolia Autonomous Region People's Hospital, Hohhot, China. Cells were cultured in Dulbecco's modified Eagle's medium (DMEM) with 10% fetal bovine serum (FBS) (Gibco®; Thermo Fisher Scientific, Inc., Waltham, MA, USA), 1% glutamine (Sigma-Aldrich, St. Louis, MO, USA), 100 IU/ml penicillin (Sigma-Aldrich) and 100 mg/l streptomycin (Sigma-Aldrich). The MTT kit was purchased from Sigma-Aldrich. Annexin V-fluorescein isothiocyanate (FITC) and the propidium iodide (PI) Annexin V-EGFP apoptosis detection kit were purchased from KGI Chemical Corporations (Nanjing, China), monoclonal mouse anti-rat HSP90 antibody (cat. no. ab13492; 1:1,000) from Abcam (Cambridge, UK) and monoclonal mouse anti-rat GAPDH antibody (cat. no. sc-32233; 1:1,000) from Santa Cruz Biotechnology, Inc. (Dallas, TX, USA). Wighteone was obtained as in previous studies (2).

3-(4,5-dimethylthiazolyl)-2,5-iphenyltetrazolium bromide (MTT) assay. Cell viability was determined using the MTT assay (Sigma-Aldrich). Firstly, cells were divided into four groups: group A: cells were treated with phosphate-buffered saline (PBS); group B: cells were treated with 0.5 mM/l wighteone; group C: cells were treated with 5.0 mM/l wighteone; group D: cells were treated with 10.0 mM/l wighteone. Cells from groups A, B, C and D were plated in 96-well plates (2x10⁴ cells/well) and cultured overnight in DMEM with 10% FBS. Following culture for 12, 24 and 48 h, the cells were treated with 0.5 mg/ml MTT for 4 h and lysed with dimethyl sulfoxide (Sigma-Aldrich). Absorbance rates were measured at 560 nm using a microplate reader (iMark™ Microplate Absorbance Reader; Bio-Rad Laboratories, Inc., Hercules, CA, USA).

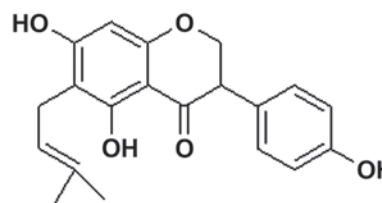


Figure 1. Structure of wighteone.

Annexin V-FITC/PI analysis. Apoptosis detection was conducted using the Annexin V-FITC/PI apoptosis detection kit, according to the manufacturer's instructions. Briefly, all four groups of cells were harvested by trypsinization, washed in PBS and stained with Annexin V-FITC conjugate and PI. Cells were then analyzed by flow cytometry (BD FACS-Calibur™, BD Biosciences, San Jose, NJ, USA) using BD CellQuest acquisition and analysis software.

Western blot analysis. The total volume of all formulations was 10 µl. The cells were diverted into a 1.5 ml Eppendorf tube (Eppendorf, Hamburg, Germany) which contained radioimmunoprecipitation assay buffer schizolysis liquid with protease inhibitor (Sigma-Aldrich). After being put on ice for 5-10 min, the mixture was agitated to make it fully dissolve, and put on ice for 30 min. Then the mixture was centrifuged (100 x g) at 4°C for 20 min, and the top clear liquid was collected and electrophoresed.

Statistical analysis. The data are presented as the mean ± standard deviation from at least three independent experiments. One way analysis of variance was used and all data analyses were performed using SPSS 13.0 software (SPSS, Inc., Chicago, IL, USA). P<0.05 was considered to indicate a statistically significance difference.

Results

Wighteone inhibits MCF-7 cell proliferation. The structure of wighteone is shown in Fig. 1. The results obtained from the MTT assay revealed that treatment with wighteone significantly inhibited cancer cell proliferation compared with the control PBS group (P=0.03). The results are shown in Table I and Figs. 2 and 3. Moreover, when increasing the concentrations of wighteone, the proliferation rates were gradually decreased.

Apoptosis. The experimental studies revealed that the apoptosis rates in groups A, B, C and D were 3.02, 4.22, 6.54 and 14.98%, respectively. However, only the apoptotic rate in group D differed significantly compared with that in group A (P=0.003). The apoptotic rate in group D also demonstrated a significant difference compared with that in group B (P=0.008; Fig. 4).

HSP90 protein expression. HSP90 protein expression of MCF-7 cells following treatment with wighteone is shown in Fig. 5. HSP90 expression was significantly lower (P=0.03) in groups B, C and D compared with that in group A. Moreover, the HSP90 expression at 48 h was significantly lower compared with that at 24 h (P=0.04).

Table I. Inhibition of MCF-7 cell proliferation following treatment with wighteone.

Group	Wighteone, mM/l	Inhibition rate of MCF-7 cancer cells (mean \pm SD)		
		12 h	24 h	48 h
A	Control	0	0	0
B	0.5	4.90 \pm 0.27	5.62 \pm 0.12	6.44 \pm 0.07
C	2.0	6.34 \pm 0.24	15.18 \pm 0.28	21.36 \pm 0.12
D	8.0	7.73 \pm 0.33	21.16 \pm 0.11	29.17 \pm 0.02

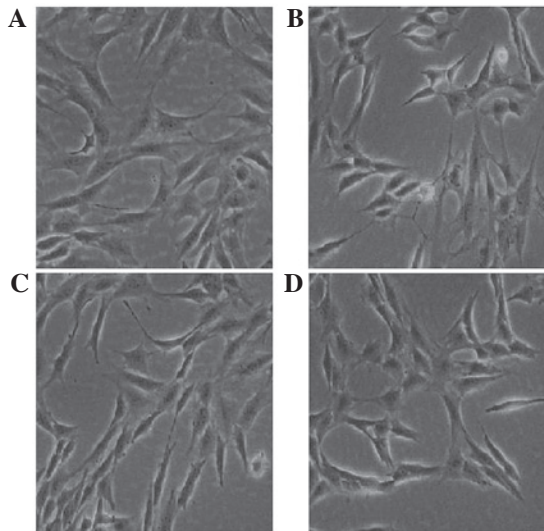


Figure 2. Inhibition of cell proliferation following treatment with various concentrations of wighteone at 48 h. (A) Phosphate-buffered saline; (B) 0.5 mM wighteone; (C) 2 mM wighteone; (D) 8 mM wighteone. Cells were visualized by phase contrast and fluorescence microscopy at a magnification of $\times 100$.

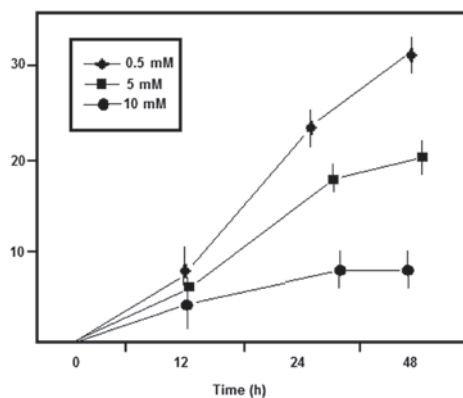


Figure 3. Inhibition of cell proliferation by wighteone *in vitro*. There was a statistically significant difference in the inhibition rate with 0.5, 5 and 10 mM wighteone vs. the control group ($P < 0.05$). Furthermore, with the increasing wighteone concentration the proliferation rates gradually decreased ($P < 0.05$).

Discussion

The use of plants as medicine dates back thousands of years, and they have become an essential source of the mainstream pharmacopoeia. Nature is an attractive source of new therapeutic candidate compounds as a vast chemical diversity

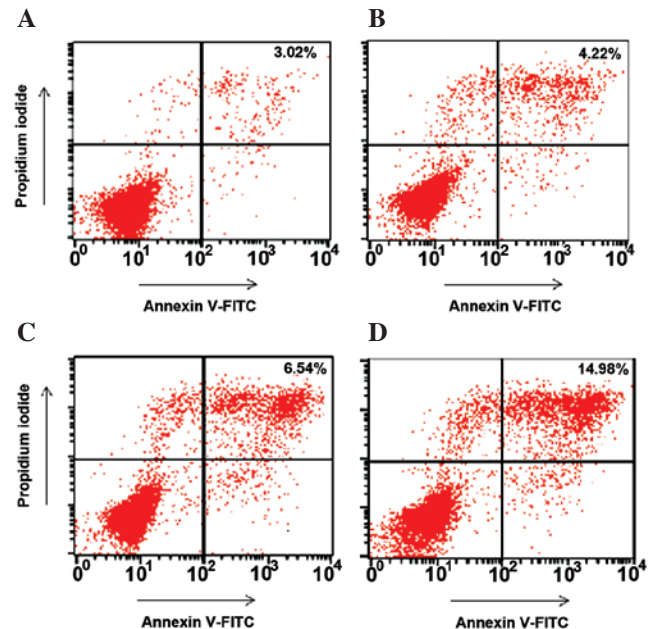


Figure 4. Wighteone-induced breast cancer cell apoptosis. The apoptotic rate in groups A, B, C and D was 3.02, 4.22, 6.54 and 14.98%, respectively. The apoptotic rate in group D differed significantly from that in group A ($P = 0.003$). The apoptotic rate in group D also demonstrated a significant difference to that in group B ($P = 0.008$). (A) Control group; (B) 0.5 mM group; (C) 5 mM group; (D) 10 mM group.

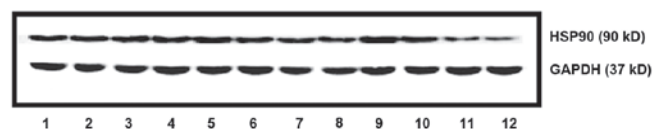


Figure 5. Heat shock protein 90 (HSP90) protein expression was assessed by western blot analysis at 0, 24 and 48 h. Channels 1, 5 and 9 correspond to group A; channels 2, 6 and 10 to group B; channels 3, 7 and 11 to group C; and channels 4, 8 and 12 to group D. HSP90 expression was significantly lower in groups B, C and D compared with group A. In addition, HSP90 expression at 72 h was significantly lower ($P < 0.05$) compared with that at 36 h.

has been observed in diverse plant species, animals, marine organisms and microorganisms. The advantage of botanical compounds is their ability to support the treatment of all phases of cancer. These compounds regularly interact with several targets simultaneously and often act synergistically. The *Erythrina* plant species is widely used in folk medicine to treat various health problems including agitation, insomnia, and anxiolytic and inflammatory processes. Recently, the

focus of drug development has shifted to the natural chemotherapeutic agents from plants, which may further be modified to enhance their potential and reduce their side effects. The identification of drugs from medicinal plants has played a significant role in the treatment of cancer; in fact, the majority of new clinical applications of plant secondary metabolites and derivatives over the last half century have been towards combating cancer. This forms the basis of the present study, in which our crucial aim was to develop a holistic molecular approach in consideration of the fact that several genes are mutated in cancer cells, which protects them from self-demise.

The predictive role of HSP90 tumor expression as a biomarker of activity of specific inhibitors remains unclear, although its overexpression is considered to confer a poor prognostic outcome in various tumors, including lung cancer, breast cancer and leukemia (24,25). With particular regard to breast cancer, the independent poor prognostic role of HSP90 overexpression in multivariate modeling was previously demonstrated in a large series of over 600 patients with a follow-up period of more than 10 years, together with large tumor size, nodal positivity, lower progesterone receptor level and high HER2 level (26).

All the experimental results revealed that with increasing concentrations of wigtheone the inhibition of proliferation of MCF-7 cells gradually increased. The apoptotic rate in the wigtheone-treated groups was higher compared with the control group ($P < 0.05$). From this, it is clear that wigtheone induces MCF-7 cell apoptosis *in vitro* and restrains cell proliferation. The activation of the AKT and MAPK pathways and the high expression of HSP90 protein are common in HER2-positive breast cancer (27) and there is a positive correlation between HSP90 and breast cancer pathological stage, local recurrence and distant metastasis, all of which result in poor patient prognosis (28). Our study reveals that wigtheone blocks the expression of HSP90 protein in MCF-7 cells, and that the inhibitory effect is enhanced with increasing drug concentrations. Wigtheone is able to control proliferation and promote apoptosis in cancer cells, and it is possible that wigtheone decreases the expression of HSP90 protein, which downregulates the key downstream molecular activation of the AKT and MAPK pathways.

In conclusion, the present study confirmed that wigtheone significantly inhibits proliferation and promotes apoptosis in HER2-positive breast cancer cells, and that this may be associated with the inhibition of HSP90 protein expression in cancer cells.

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