Increased Susceptibility to Apoptosis and Growth Arrest of Human Breast Cancer Cells Treated by a Snake Venom-Loaded Silica Nanoparticles

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**Key Words**

Apoptosis • Breast cancer • Nanoparticles • Proliferation • Snake venom

**Abstract**

**Background:** The development of effective treatments against metastatic cancers, including breast cancer, is among the most important challenges in current experimental and clinical cancer research. We recently demonstrated that *Walterinnesia aegyptia* venom (WEV), either alone or in combination with silica nanoparticles (WEV+NP), resulted in the growth arrest and apoptosis of different cancer cell lines. **Aims:** In the present study, we evaluated the impact of WEV alone and WEV+NP on human breast cancer cells isolated from cancer biopsies. **Methods:** The potential effects of WEV alone and WEV+NP on the proliferation, induction of apoptosis and generation of free radicals in breast cancer cells isolated from 80 patients clinically diagnosed with breast cancer were evaluated by flow cytometry and ELISA. **Results:** WEV alone and WEV+NP inhibited the proliferation, altered the cell cycle and enhanced the induction of apoptosis of the breast cancer cells by increasing the activities of caspase-3, caspase-8 and caspase-9. In addition, the combination of WEV and NP robustly sensitized the breast cancer cells to growth arrest and apoptosis by increasing the generation of free radicals, including reactive oxygen species (ROS), hydroperoxide and nitric oxide. The combination of WEV with NP significantly enhanced the anti-tumor effect of WEV in breast cancer cells. **Conclusion:** Our data indicate the therapeutic potential of the nanoparticle-sustained delivery of snake venom for the treatment of breast cancer.
Introduction

Despite major advances in the elucidation of the mechanisms of breast cancer progression and the development of novel therapeutic agents, breast cancer remains the second leading cause of mortality among women worldwide [1]. This mortality is almost invariably due to metastasis [1, 2]. Metastatic disease remains the most critical factor limiting patient survival, and the development of effective treatments against metastatic cancers, including breast cancer, is among the most important challenges in current experimental and clinical cancer research [3-5].

Natural toxins are recognized as sources for drugs against several human ailments, including cancers. In particular, a non-toxic dose of snake venom has been shown to both reduce the size of solid tumors and block the process of angiogenesis in ovarian cancer [6]. New anticancer agents must be identified to increase the number of available options and identify less toxic and more effective drugs. Snake venom is a complex mixture of many substances, including toxins, enzymes, growth factors, activators and inhibitors, with a wide spectrum of biological activities. Our recent studies have demonstrated the anti-tumor potential of snake venom from Walterinnesia aegyptia (WEV) on the human breast carcinoma cell line MDA-MB-231 and have shown its effect on normal murine peripheral blood mononuclear cells (PBMCs) [7]. In addition, other data have indicated that the snake venom toxin from Vipera lebetina turanica inhibits hormone-refractory human prostate cancer cell growth at nanogram concentrations; this effect is related to the NF-κB signal-mediated induction of apoptosis [8].

One mechanism by which chemotherapeutic agents kill tumor cells is by inducing apoptotic death pathways. The ability of cancer cells to escape from apoptosis and continue to proliferate is one of the fundamental hallmarks of cancer and is a major target of cancer treatment; therefore, the underlying mechanisms of apoptosis and cancer progression continue to be a focus of intense research. In a cell, apoptosis can be triggered through either the extrinsic pathway or the intrinsic pathway. In the extrinsic pathway, signal molecules, known as ligands, bind to transmembrane death receptors on the target cell to induce apoptosis through the activation of cellular caspases, while the intrinsic pathway is triggered by cellular stress and is mediated through a mitochondrial-dependent pathway. Caspase-dependent apoptosis includes the activation of caspase-3, caspase-8 and caspase-9, whereas the mitochondrial pathway involves the efflux of cytochrome C from the mitochondria to the cytosol to form apoptosomes with Apaf-1 and caspase-9, which lead to the activation of caspase-3 and subsequent apoptosis induction [9, 10]. In fact, both pathways are intricately related. Tumor cells contain fewer scavengers of free radicals than normal cells, and free radicals have been shown to participate in the mechanism of anticancer therapeutic agents; the production of large amounts of free radicals in tumor tissues may therefore have potential as a future anticancer therapy [11]. Consequently, modulation of the levels of reactive oxygen species (ROS) and other free radicals that induce oxidative stress has been proposed as a therapeutic approach to cancer [12]. Nitric oxide (NO) plays important physiological roles in vascular function and the inflammatory response. However, NO over-production induces DNA damage, mitochondrial uncoupling and increased ROS [12-14].

Nanoparticles loaded with chemical therapeutics have shown great promise for the treatment of cancer. When loaded with anticancer agents, nanoparticles can successfully increase drug concentrations in cancer tissues and act at the cellular level to enhance antitumor efficacy. The nanoparticles can be endocytosed and/or phagocytosed by cells, resulting in the internalization of the encapsulated drug [9]. Therefore, in the present study, we investigated the effects of WEV alone and in combination with silica nanoparticles (WEV+NP) on the proliferation and apoptosis of human breast cancer cells through monitoring the caspase activity and free radical levels.
Materials and Methods

Preparation of Walterinnesia aegyptia venom

Walterinnesia aegyptia snakes were collected from the central region of Saudi Arabia. No specific permits were required for the described field studies, and no specific permission was required for these locations/activities because the location was not privately owned or protected in any way and the field studies did not involve endangered or protected species. The snakes were kept in a serpentarium in the Zoology Department of the College of Science at King Saud University. The snakes were warmed daily for nine hours using a 100-watt lamp and were provided water *ad libitum*. The snakes were fed purpose-bred mice every 10 to 14 days. After the venom was milked from a single specimen of adult snake, the venom was lyophilized and reconstituted in 1X phosphate-buffered saline (PBS) prior to use.

Combination of snake venom with silica nanoparticles

As previously described [7, 15], a double mesoporous core-shell silica nanosphere was formed around a silica core by using an anionic surfactant to transform the solid silica core into a mesoporous core. To synthesize the solid silica core, 0.875 ml aqueous ammonia was added to a solution that contained 18 ml ethanol and 2.6 ml deionized water; then, 1.5 ml tetraethyl orthosilicate (TEOS) was added while the solution was vigorously stirred. The resulting mixture was heated to 30°C for 60 min, and the silica precipitate was then collected by centrifugation and washed three times with water. Second, to synthesize the mesoporous core-shell nanosphere, silica (SiO$_2$) particles were dispersed using an anionic surfactant in 15 ml H$_2$O and ultrasonicated for 10 min. To suppress the agglomeration of the silica cores, 1 g/l polyvinylpyrrolidone was added followed by constant stirring for 60 min. Next, 0.1 ml 3-aminopropyltrimethoxysilane (APMS), 0.2933 g (1 mmol) N-lauroylsarcosine sodium (Sar-Na) and 1.5 ml TEOS were added to the reaction mixture, which was stirred at 50°C for 2 h. The final solid was recovered by centrifugation, washed with deionized water and dried in an oven at 60°C for 12 h. Template removal was performed by heat treatment in an air stream at 550°C for 6 hours. After synthesizing the nanoparticles, 25 mg mesoporous silica nanoparticles was added to a solution of 50 mg/ml venom in 0.5 ml water. The suspension was stirred for 2 hours, and the evaporation of water was prevented. The mesoporous silica nanoparticles loaded with venom were recovered by high-speed centrifugation and then dried in a vacuum oven at 60°C. Transmission electron microscopy (TEM) was performed with a JECOL JSM-2100F electron microscope (Japan) operated at 200 kV. Nitrogen sorption isotherms were measured at 77 K with a Quantachrome NOVA 4200 analyzer (USA). Prior to taking measurements, the samples were degassed in a vacuum at 200°C for at least 18 hours. The Brunauer-Emmett-Teller (BET) method was utilized to calculate the specific surface areas (SBET) using adsorption data in a relative pressure range from 0.05 to 0.35. Using the Barrett-Joyner-Halenda (BJH) model, the pore volumes and pore size distributions were derived from the adsorption branches of the isotherms, and the total pore volumes (Vt) were estimated from the adsorbed amount at a relative pressure (P/P0) of 0.992.

Human breast cancer samples

Breast cancer tissue samples were obtained from 80 females with histologically proven breast cancer. The clinico-histopathological data of the patients are summarized in Table 1. All patients had been surgically treated at the South Egypt Cancer Institute of Assiut University in Egypt. Non-tumorigenic normal breast tissue samples were obtained from Assiut University Hospital. The tumor tissue specimens were taken at the time of surgery after informed written consent in accordance with South Egypt Cancer Institute ethical committee guidelines. This study was approved by the Ethical committee of South Egypt Cancer Institute, Assiut University, Egypt. This ethical committee is approved by U.S. Department of Health and Human Services (HHS) Institutional Review Board (IRB). IORG number: IORG0006563. OMB number: 0990-0279. The tumor and normal tissues were immediately disaggregated mechanically by passage through a 16-gauge stainless steel mesh, and the cells were either lysed in lysis buffer for the caspase activity test and the measurement of free radical levels or directly transferred to and maintained in a culture medium consisting of MEM supplemented with 10% heat-inactivated fetal bovine serum (FBS, EuroClone, Life Science Division, Milan, Italy). The anti-proliferative effects of WEV, NP and WEV+NP were determined on the non-tumorigenic and breast cancer cells isolated from human samples using the 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) uptake method. The cells were plated at 1x10^4 cells/ml in 2
ml culture medium in six-well Costar plates (Corning, Corning, NY). The cells were treated with different concentrations of WEV or WEV+NP for 1, 2, 6, 12, 24 or 48 h. Cytotoxicity was expressed as the relative percentage of the OD values measured in the control untreated (0), NP-, WEV- and WEV+NP-treated cells. Morphological changes following exposure to NP, WEV and WEV+NP were observed using a phase contrast inverted microscope (Olympus, Japan).

**Table 1.** Clinico-histopathological data for patients diagnosed with breast cancer

<table>
<thead>
<tr>
<th>Clinico-histopathological parameters</th>
<th>Patients (No.)</th>
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<tr>
<td>Histopathology</td>
<td></td>
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<tr>
<td>Infiltrating ductal carcinoma</td>
<td>61</td>
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<tr>
<td>Infiltrating lobular carcinoma</td>
<td>14</td>
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<tr>
<td>Other</td>
<td>5</td>
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<td>Grade I</td>
<td>53</td>
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<td>Grade III</td>
<td>27</td>
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<tr>
<td>Lymph node involvement</td>
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<td>N0</td>
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<td>N1</td>
<td>49</td>
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<td>N2</td>
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<td>Tumor burden</td>
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**CFSE proliferation assay and flow cytometry analysis**

Flow cytometry was performed with the FACSCalibur system (BD, San Jose, CA). All fluorocytometric data were subsequently analyzed and displayed with CELL QUEST software (BD, San Jose, CA). Each analysis included measurements for a minimum of 20,000 cells. The breast cancer cells were washed twice in PBS and stained with 0.63 µM carboxyfluorescein diacetate succinimidyl ester (CFSE) (Molecular Probes, Eugene, OR) for 8 min at room temperature. The residual CFSE was removed by washing 3 times in PBS. The CFSE-labeled cells were then seeded into 6-well plates, treated with NP, WEV or WEV+NP or left untreated (0) and grown for 4 days in RPMI cell culture medium. The cells were then stained with monoclonal antibodies. The CFSE fluorescence intensity was measured by flow cytometry. Isotype control antibodies were used to separate the positive and negative cells.

**Flow cytometry analysis of apoptosis and cell cycle analysis**

After treatment with NP, WEV or WEV+NP, the breast cancer cells were fixed and permeabilized with 70% ice-cold ethanol for at least 1 h and then washed twice with PBS. The DNA was stained by incubating the cells at 37°C for 1 h in 40 µg/ml propidium iodide and 100 µg/ml DNase-free RNase in PBS. The fluorescence area (FL2-A) is the main parameter in the cell cycle analysis; therefore, a histogram plot of FL2-A was used as a cell cycle graph. The cell cycle distributions were analyzed with Modfit LT 3.0 software (BD, San Jose, CA). Dead cells were identified using the Trypan blue dye exclusion test. The breast cancer cells were reanalyzed for the expression of annexin V and both annexin V and PI to identify early and late apoptosis, respectively.

**Measurements of caspase activity**

Caspase-3, caspase-8 and caspase-9 activities were evaluated using a fluorometric protease assay kit (MBL, Aichi, Japan) according to the manufacturer’s instructions.

**Measurements of ROS, hydroperoxide and nitric oxide levels**

The levels of ROS were determined using 2,7-dichlorodihydrofluorescein diacetate (H2DCF-DA) (Beyotime Institute of Biotechnology, Haimen, China). Tumor cells (1x10⁶) were directly treated with 10 µM H2DCF-DA dissolved in 1 ml PBS at 37°C for 20 min. The fluorescence intensity was monitored with an excitation wavelength of 488 nm and an emission wavelength of 530 nm. The levels of hydroperoxide were measured using the free radical analytical system (FRAS 2, IRAM, Parma, Italy). This test is a colorimetric assay that takes advantage of the ability of hydroperoxide to generate free radicals after reaction with specific transition metals. The concentrations of nitrite and nitrate were measured with a Griess assay reagent (NO₂⁻/NO₃⁻ detection kit; Dojindo, Kumamoto, Japan) according to the manufacturer’s instructions. In brief, the supernatant of the cell culture medium without phenol or serum was collected and then reacted with the Griess reagent. The azo coupling between the diazonium species, which are produced by the reaction of sulfanilamide with NO₂ and 1-naphthylethylenediamine was measured at 540 nm with an MRX microplate reader (Dynex Technologies, Inc., Chantilly, VA).
Statistical analysis

The data were first tested for normality using the Anderson–Darling test and then evaluated for variance homogeneity prior to further statistical analysis. The data were normally distributed and were expressed as the mean ± standard error of the mean (SEM). Significant differences among groups were analyzed by one- or two-way ANOVA followed by the Bonferroni multiple comparison test using PRISM statistical analysis software (GraphPad Software). The data were then reanalyzed by one- or two-way ANOVA followed by Tukey’s range test using SPSS software, version 17. Differences were considered significant at P < 0.05. *P < 0.05, WEV-treated vs. control; #P < 0.05, WEV+NP-treated vs. control; +P < 0.05, WEV+NP-treated vs. WEV-treated.

Results

**WEV and WEV+NP inhibit the growth of breast cancer cells**

Using a silica nanoparticles delivery system, we first investigated the ability of WEV and WEV+NP to induce growth arrest in breast cancer cells. The effects of WEV and WEV+NP on breast cancer cells and normal breast cells were examined at WEV concentrations of 0, 1, 5, 10, 20, 50, 100 and 1000 ng/ml and incubation times of 0, 1, 2, 6, 12, 24, 36 and 48 h. The resulting cytotoxic effects of WEV and WEV+NP were measured using the MTT uptake method. The results of five independent experiments (n=5) demonstrated that WEV and WEV+NP significantly inhibited the growth of breast cancer cells in a dose- and time-dependent manner (Fig. 1A & B). The IC_{50} values for WEV alone and WEV+NP were 50 ng/ml and 20 ng/ml, respectively. The effect was maximal at 12 h of incubation. Nevertheless, treatment with WEV or WEV+NP had no significant inhibitory effect on the viability of non-tumorigenic normal breast cells (Fig. 1C). The combination of WEV with NP (WEV+NP) significantly enhanced the inhibitory effect of WEV in breast cancer cells. The maximal inhibitory effects of WEV and WEV+NP on cell viability were observed 12 h after treatment with 50 ng/ml of WEV alone or 20 ng/ml of WEV+NP. Nevertheless, treatment with NP alone did not affect breast cancer cell viability.
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**Fig. 2.** WEV and WEV+NP decrease the proliferation of breast cancer cells. CFSE assays and flow cytometry were used to evaluate the ability of breast cancer cells to proliferate after treatment with vehicle, NP, WEV and WEV+NP. (A) One representative experiment showing the analysis of the CFSE staining of the breast cancer cells from one patient after gating on the viable cells. (B) The accumulated data for 80 patients are expressed as the mean percentage of proliferating cells ± SEM for vehicle-treated cancer cells (closed black bars), NP-treated cells (hatched bars), WEV-treated cells (closed gray bars) and WEV+NP-treated cells (dotted bars). P < 0.05, WEV-treated vs. control; #P < 0.05, WEV+NP-treated vs. control; +P < 0.05, WEV+NP-treated vs. WEV-treated.

**Treatment with WEV alone and WEV+NP inhibits the proliferation of breast cancer cells**

Because the proliferation process is crucial for the maintenance and progression of cancer cells, we monitored the effects of WEV alone or in combination with NP (WEV+NP) on the proliferation of breast cancer cells using a CFSE dilution assay followed by flow cytometry. The data from one representative experiment are shown as dot plots (Fig. 2A) and reveal that the percentage of proliferating cells was markedly decreased from 69% in NP-treated cells to 40.9% and 18.9% in WEV- and WEV+NP-treated cells, respectively. The pooled data for different patients diagnosed with breast cancer (n=80) demonstrated that treatment with WEV alone significantly reduced (P < 0.05) the proliferative capacity of breast cancer cells compared with vehicle-treated cells (Fig. 2B). Moreover, although NP had no effect on proliferation, the combination of NP with WEV significantly enhanced the inhibitory effect of WEV on breast cancer cells.

**Treatment with WEV and WEV+NP induces cell cycle arrest and apoptosis in breast cancer cells**

The inhibition of cancer cell proliferation, the cessation of cell-cycle progression and the induction of apoptosis have all been targeted in chemotherapeutic strategies for the treatment of cancer. Therefore, we used propidium iodide (PI) single staining and PI/annexin V double staining followed by flow cytometry analysis to determine if WEV and WEV+NP alter the cell cycle of breast cancer cells. The data from one representative experiment are presented as a histogram; the percentage of apoptotic cells was 11% in the NP-treated cells.
Treatment with WEV and WEV+NP markedly increased the percentage of apoptotic cells to 35% and 85%, respectively (Fig. 3A). The increased induction of apoptosis after treatment with WEV and WEV+NP was inversely correlated with a decrease in the percentage of cells in S phase to 59% and 25%, respectively. The breast cancer cells were also stained with PI/annexin V to discriminate between apoptotic, necrotic and viable cells, and the data are presented as dot plots. The data from one representative experiment indicate that the percentage of apoptotic cells clearly increased in WEV- and WEV+NP-treated cells compared with NP-treated cells (Fig. 3B). The accumulated data for cells isolated from different patients (n=80) demonstrated that treatment with WEV alone significantly (P < 0.05) potentiated apoptosis and diminished the percentage of cells in S phase in breast cancer cells compared with vehicle- and NP-treated cells (Fig. 3C). Although NP alone had no effect on apoptosis induction and the percentage of cells in S phase, the combination of NP and WEV significantly (P < 0.05) increased apoptosis induction and decreased the percentage of cells in S phase.
Treatment with WEV and WEV+NP induces apoptosis in breast cancer cells via direct activation of caspase activity

Because caspases are active mediators of apoptosis, the activities of caspase-3, caspase-8 and caspase-9 were monitored in the breast cancer cell lysates after treatment with vehicle, NP, WEV or WEV+NP. The accumulated data for 80 patients are expressed as the mean fold increase ± SEM for vehicle-treated cancer cells (closed black bars), NP-treated cells (hatched bars), WEV-treated cells (closed gray bars) and WEV+NP-treated cells (dotted bars). P < 0.05, WEV-treated vs. control; #P < 0.05, WEV+NP-treated vs. control; +P < 0.05, WEV+NP-treated vs. WEV-treated.

Treatment with WEV and WEV+NP induces the generation of ROS, hydroperoxide and nitric oxide

Tumor cells contain fewer radical scavengers than do normal cells, and free radicals have been shown to participate in the mechanism of anticancer therapeutic agents. Therefore, the generation of a large quantity of free radicals in tumor tissues may represent...
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Uncontrolled proliferation is significant in cell turnover and tumorigenesis. Therefore, we monitored the effects of WEV alone or in combination with NP on the proliferation of breast cancer cells with a CFSE dilution assay followed by flow cytometry analysis. Our data revealed that WEV alone and in combination with NP inhibited the proliferation of breast cancer cells. Uncontrolled proliferation is significant in cell turnover and tumorigenesis; thus agents that are able to inhibit proliferation may be useful as chemotherapeutic agents against breast cancer. Our data are consistent with Dowsett et al. who reported that increased apoptosis and decreased proliferation are common factors in the biological response of breast cancer to chemotherapy and endocrine therapy [25]. Positive clinical responses are associated with reduced proliferation during chemotherapy, endocrine therapy and chemoendocrine therapy. Anticancer agents may alter the regulation of the cell cycle machinery, resulting in cellular arrest at different phases of the cell cycle and thereby reducing the growth and proliferation of cancerous cells; cell cycle arrest may even induce apoptosis [26]. Several drugs have been designed to synthetically activate caspases, including peptides that contain the arginine-glycine-aspartate motif. These peptides are pro-apoptotic and have the ability to directly induce auto-activation of procaspase 3. They have also been shown to lower the activation threshold of caspase enzymes or activate caspases, contributing to an increase in the drug sensitivity of cancer cells [27]. Our data reveal that WEV alone and in combination with NP increased the activities of caspase-3, caspase-8 and caspase-9 in breast cancer cells. Tandon et al concluded that oxidative stress may have potential therapeutic applications in the development of anticancer drugs [28]. Potential anticancer drugs acting by this novel mechanism may prove to be useful in the future. Therefore, we measured the levels of ROS, NO and hydroperoxide, revealing an increase in the levels of free radicals after treatment with WEV and WEV+NP. Generally, chemotherapeutic drugs attack both normal and tumor cells non-specifically and may causing life-threatening side effects, necessitating the
targeted delivery of drugs to tumors [29]. Indeed, the therapeutic molecule must generally cross one or more biological membranes before diffusing through the plasma membrane to finally gain access to the appropriate organelle where the biological target is located. For a drug whose target is located intracellularly, deviating from this ideal path may not only decrease the drug’s efficiency but also entail side effects and toxicity. For these reasons, the design of carriers small enough to ferry the active substance to the target cell and the relevant subcellular compartment was proposed more than 30 years ago [30]. Various types of nanoparticles, such as liposomes, polymeric micelles, dendrimers, superparamagnetic iron oxide crystals and colloidal gold, have been employed in targeted therapies for cancer [31, 32]. Nanocarriers offer unique possibilities to overcome cellular barriers to improve the delivery of various drug candidates [26]. Recent studies have demonstrated efficient tumor targeting by nanoparticles through an enhanced permeability and retention effect [33-35]. The delivery of drug-loaded nanoparticles has achieved success in the treatment of advanced thyroid cancer and breast carcinoma [36, 37]. Here, while venom-free nanoparticles had no effect on breast cancer cells, the combination of WEV with nanoparticles increased the ability of WEV to kill breast cancer cells by 2-fold compared with WEV alone. Despite great interest in using nanoparticles in biomedical applications, a clear understanding of their cellular uptake and transport is still lacking. Nanoparticles appear to translocate across cells via clathrin- and macropinocytosis-mediated endocytosis [38]. The nanoparticles were also shown to be stable within the cytoplasm for at least 24 h and did not colocalize within the endosomal pathway. Furthermore, nanoparticle uptake was inhibited approximately 50% by genistein, an inhibitor of the caveolae-mediated pathway [39]. However, the clathrin-mediated endocytosis and macropinocytosis pathways were reduced by 17 and 24%, respectively, in the presence of the respective inhibitors. These findings suggest that PLL-g-PEG-DNA nanoparticles enter by several pathways and might therefore be an efficient and versatile tool to deliver therapeutic DNA [40]. Therefore, the combination of nanoparticles with WEV significantly enhanced the antitumor effects of WEV. Snake venom is a complex mixture of many substances, including toxins, enzymes, growth factors, activators and inhibitors, with a wide spectrum of biological activities. Furthermore, synergistic actions between the components of WEV likely contribute to the antitumor effects and the mechanisms that we observed. Subsequently, fractionation of the WEV components and studies of the combination of each fraction with NP are underway. This study revealed the unique biological effects of WEV and WEV+NP on breast cancer cells, which may permit these compounds to be utilized in treatments for breast cancer.

Disclosure Statement

The authors declare that they have no conflicts of interest, state that the manuscript has not been published or submitted elsewhere, state that the work complies with Ethical Policies of the Journal and the work has been conducted under internationally accepted ethical standards after relevant ethical review.

Abbreviations

Nanoparticles (NP); reactive oxygen species (ROS); Walterinnesia aegyptia venom (WEV); Walterinnesia aegyptia venom combined with nanoparticles (WEV+NP).

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References


