Antitumor effects of cecropin B-LHRH’ on drug-resistant ovarian and endometrial cancer cells

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Abstract

Background: Luteinizing hormone-releasing hormone receptor (LHRHr) represents a promising therapeutic target for treating sex hormone-dependent tumors. We coupled cecropin B, an antimicrobial peptide, to LHRH’, a form of LHRH modified at carboxyl-terminal residues 4–10, which binds to LHRHr without interfering with luteinizing hormone (LH) and follicle-stimulating hormone (FSH) secretion. This study aimed to assess the antitumor effects of cecropin B-LHRH’ (CB-LHRH) in drug-resistant ovarian and endometrial cancers.

Methods: To evaluate the antitumor effects of CB-LHRH’, three drug resistant ovarian cancer cell lines (SKOV-3, ES-2, NIH:OVCAR-3) and an endometrial cancer cell line (HEC-1A) were treated with CB-LHRH’. Cell morphology changes were assessed using inverted and electron microscopes. In addition, cell growth and cell cytotoxicity were measured by MTT assay and LDH release, respectively. In addition, hemolysis was measured. Furthermore, radioligand receptor binding, hypersensitization and minimal inhibitory concentrations (against Staphylococcus aureus, Klebsiella pneumoniae, Escherichia coli, Enterobacter cloacae, Pseudomonas aeruginosa, and Acinetobacter baumannii) were determined. Finally, the impact on tumor growth in BALB/c-nu mice was assessed in an ES-2 xenograft model.

Results: CB-LHRH’ bound LHRHr with high-affinity (dissociation constant, \( K_d = 0.252 ± 0.061\) nM). Interestingly, CB-LHRH’ significantly inhibited the cell viability of SKOV-3, ES-2, NIH:OVCAR-3 and HEC-1A, but not that of normal eukaryotic cells. CB-LHRH’ was active against bacteria at micromolar concentrations, and caused no hypersensitivity in guinea pigs. Furthermore, CB-LHRH’ inhibited tumor growth with a 23.8 and 20.4 % reduction in tumor weight at 50 and 25 mg/kg.d, respectively.

Conclusions: CB-LHRH’ is a candidate for targeted chemotherapy against ovarian and endometrial cancers.

Keywords: Luteinizing hormone releasing-hormone receptor, Cecropin B peptide, Ovarian cancer, Endometrial cancer

Background

Sex hormone-dependent tumors, including ovarian, endometrial, breast and prostate carcinomas, are the most common reproductive system tumors. Ovarian cancer is often detected at a late stage [1–3]; despite cytoreductive surgery and paclitaxel/platinum-based chemotherapy it frequently recurs, resulting in poor prognosis [4–6]. Although endometrial carcinoma presents at an early stage and responds well to surgery [7, 8], frequent recurrence also results in poor prognosis [8, 9]. Chemotherapy of sex hormone-dependent cancers is limited by the intrinsic or acquired drug resistance of tumor cells as well as the toxicity to normal cells of chemotherapeutic agents [5, 10, 11], whose side effect profiles limit the chemotherapeutic dosing [12, 13].

Targeted cytotoxic agents enable selective treatment of primary tumors and their metastases, reducing side effects and improving efficacy [2, 14–17]. Luteinizing hormone-releasing hormone receptor (LHRHr) may represent a useful target in sex hormone-dependent tumors as it is expressed in human breast (52 %), ovarian (80 %), endometrial (80 %), and prostate (86 %) carcinomas [18]. Potential drawbacks of targeted chemotherapy using natural LHRH as a binding partner include...
side effects resulting from interference with pituitary secretion of LH and FSH [19]. Previous LHRHr targeted cytotoxic therapies include cytotoxins, such as doxorubicin, which have non-specific cytotoxicity. Other LHRH targeted therapies used chemical approaches linking the cytotoxic to LHRH; however, such conjugates are readily hydrolyzed in the blood stream, releasing cytotoxic radicals before reaching their therapeutic targets, therefore inducing non-specific cytotoxicity [19]. Refined and improved strategies for LHRHr chemotherapy are necessary if these approaches are to be useful.

LHRH is a decapeptide that binds to receptors on pituitary gonadotropes, stimulating biosynthesis and secretion of FSH and LH, which regulate gonadal steroidsogenesis and gametogenesis in both sexes [20]. The carboxy-terminal residues 4–10 of LHRH are involved in receptor binding, while amino-terminal residues 1–3 activate the receptors [21]. As targeted chemotherapy using natural LHRH as a carrier may interfere with LH and FSH secretion, we constructed a modified peptide, LHRH, in which amino-terminal residues 1–3 are not included. Thus, LHRH is expected to target LHRHr positive carcinoma cells, while not interfering with LH and FSH secretion.

Cecropin B is an antimicrobial peptide (AMP) first characterized in 1980 [22, 23]; since then, thousands of similar molecules have been isolated from a wide range of organisms [24, 25]. AMPs are short peptides possessing net cationic charges, selective toxicity, rapid cytotoxic effects, broad antimicrobial spectra, and no documented resistance [24–26]. Some AMPs, including cecropins, are highly potent against cancer cells but not normal mammalian cells [27–31], making them attractive for the treatment of some cancers.

In this study, we developed a novel LHRHr cytotoxin by linking Cecropin B to the modified LHRHr receptor ligand, Cecropin B-LHRH (CB-LHRH) [32]. We hypothesized that the novel CB-LHRH may bind to LHRH receptors and deliver an effective broad-spectrum toxin specifically to tumor cells, without affecting healthy cells or readily inducing resistance. Therefore, we aimed in this study to assess the antitumor effects of CB-LHRH in the treatment of drug-resistant ovarian and endometrial cancers, using BALB/c-nu mice (a common model for cancer studies) harboring ES-2 xenografts.

Methods
The study was approved by the ethical committee of Women's Hospital, School of Medicine, Zhejiang University, Hangzhou, China. Informed consent was obtained from all individual participants included in the study.

Peptides and cytotoxic agents
CB-LHRH recombinant polypeptide (KWKFVKKIEKM GRNIRNGIVKAGPA1AVLGEAKALSYGLRPG) (Shanghai Sangon Biological Engineering Technology & Services Corporation, Shanghai, China) was synthesized by solid phase synthesis to a purity of 96 % as evaluated by high-performance liquid chromatography. It was identified by mass spectrographic analysis (MW 4,566.57D), dissolved in phosphate-buffered saline (PBS), and diluted to the desired concentration before use.

Drug-resistant tumor cell lines
Four drug-resistant cancer cell lines, including three ovarian cancer cell lines (SKOV-3, ES-2, NIH:OVCAR-3) and one endometrial cancer cell line (HEC-1A) (Cell Bank of Shanghai Biological Institute, Shanghai, China), were cultured in RPMI 1640 with 10 % fetal bovine serum, 1,000 IU/mL penicillin, and 100 mg/mL streptomycin at 37 °C and 5 % CO2. Cell cultures and subculture procedures were performed according to the recommendations of the ATCC Global Bioresource Center. ES-2, HEC-1A, and NIH:OVCAR-3 cells express LHRHr, while the SKOV-3 does not express LHRHr [33–35].

Clonogenic assay
A clonogenic assay was carried out as previously described [36]. 300 cells were seeded into 6-well dishes in 2 mL of medium. After overnight incubation, CB-LHRH was added at a final concentration of 12.5 μM and cells were further cultured for 10 d. Cells were then stained with crystal violet (0.5 % w/v). Colonies containing a minimum of 50 cells were counted using a dissection microscope. Each treatment was performed in triplicate and each experiment repeated three times.

Cytotoxicity assay
Lactate dehydrogenase (LDH) release was quantified using LDH Cytotoxicity Detection Kit (Roche Applied Science, Mannheim, Germany) according to the manufacturer’s instructions. Briefly, 100 μL of exponentially growing tumor cells (6 x 104 cells/mL) were seeded into 96-well plates, cultured overnight, and washed with RPMI 1640. Then, several CB-LHRH amounts (6.25, 12.5, 25, 50, and 100 μM) were completed to 100 μL with RPMI and added to cells, followed by 4 h incubation. LDH release was assessed by measuring absorbance at 490 and 630 nm, respectively, on a microplate reader (Bio-Tek Instruments Inc., Vermont, USA). Cytotoxicity was determined by the following formula: Cytotoxicity (%) = (exp. value - low control)/(high control - low control) x 100. All conditions were carried out in triplicate, and the experiments repeated three times.

Proliferation assay
The anti-proliferative activity of CB-LHRH was assessed by the 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl-tetrazolium
bromide (MTT) assay [37]. Briefly, 100 μL of exponentially growing tumor cells (6 × 10^4 cells/mL) were seeded in 96-well plates and cultured overnight. B-LHRH' was then added to each well at final concentrations of 6.25, 12.5, 25, 50, and 100 μM. After another 24 or 48 h, 10 μL of fresh MTT (Sigma–Aldrich, St. Louis, USA) at 5 mg/mL in Hank's salt solution was added for 4 h. After careful removal of the medium, 100 μL of dimethylsulfoxide (Sigma–Aldrich, St. Louis, USA) was added to wells followed by absorbance reading on a microplate reader (Bio-Tek Instruments Inc, Vermont, USA) at 490 and 630 nm, respectively. Cell viability was determined as follows: Cell viability (%) = (absorbance of treated wells - absorbance of blank control) / (absorbance of negative control - absorbance of blank control) x 100. The inhibition rate was derived as 1-Cell viability. Each experiment was performed in triplicate and repeated three times.

Cell morphology was monitored using inverted and electron microscopes after 1 h exposure to CB-LHRH' as previously described [38].

Electron microscopy
For transmission electron microscopy ES-2 cells grown on sterile slides were fixed with 2.5 % glutaraldehyde in pH 7.4 phosphate buffer for 1 h at 4 °C, followed by post-fixation in 1 % osmium tetroxide 1 h at 4 °C and dehydrated in an alcohol gradient (50 to 100 %). The attached cells were then dried by Lyophilization and coated with Au before examination with transmission electron microscopy (S-4800 SEM, Hitachi, Ltd. Tokyo, Japan).

For scanning electron microscopy ES-2 cells were grown and fixed under similar conditions, without slides. After post-fixation cells were infiltrated with Epon resin and sectioned with Leica UC6 ultramicrotome (Leica Microsystems Inc., LKB-II, Wetzlar, Germany). Sections of about 100 nm were post-stained with lead citrate and uranyl acetate. Grids were examined with scanning electron microscope (JEM-200CX TEM, JEOL, Ltd. Tokyo, Japan).

Cytotoxicity of CB-LHRH' in normal eukaryotic cells
The ability of CB-LHRH' to induce cytotoxicity in normal eukaryotic cells was assessed as previously described [39]. With informed consent and institutional review board approval, peripheral blood samples were obtained from healthy human donors. Red blood cells were separated by centrifugation at 1,000 g for 5 min followed by three washes with PBS. Washed red blood cells were resuspended in sterile Alsever’s solution (2.05 % dextrose, 0.8 % sodium citrate, 0.055 % citric acid, and 0.42 % sodium chloride, pH 6.1, hematocrit 5 %) in a plate and treated with CB-LHRH' at 100, 200, 300, 400, or 500 μM for either 30 min or 4 h at 37 °C with shaking. After centrifugation for 10 min at 600 g, the supernatant was collected for hemolysis measurement at 490 nm (hemoglobin absorbance). Adding equal volumes of water to red blood cells provided 100 % hemolysis (positive control); cell-free Alsever’s solution was used as negative control. Each experiment was performed in triplicate and repeated three times.

Radioligand receptor binding assays
Under approved animal research protocol standards, binding capacity to LHRHr was assessed in rat pituitary membranes by radioligand-binding assays as previously described [40]. The experiment was approved by the animal care and use committee of Fudan University.

Preparation of membrane fractions for primary pituicytes
To obtain primary pituicytes, SD male and female rats (180–240 g) were sacrificed and the pituitary glands dissected and washed twice with 10 mM Tris-HCl buffer, pH 7.4, containing 0.5 mM PMSF, 1.2 mM MgCl_2, 0.01 mM EDTA-Na_2. Tissue aliquots were homogenized using an automatic glass Potter homogenizer at 2000 rpm for 3 × 10 s. After nucleus and debris removal by centrifugation at 2000 g and 4 °C for 5 min, membrane preparation aliquots were collected by centrifugation at 2000 g and 4 °C for 20 min. Protein concentrations were determined by the Lowry method using bovine serum albumin (BSA) as a standard, diluted with 10 mM Tris-HCl buffer to 2 mg/mL stock at −70 °C.

Radioligand receptor binding assay
CB-LHRH' peptide was radioactively labeled with Iodine-125 (Perkin Elmer Life and Analytical Sciences, Waltham, USA) using Chloramine-T (Sigma–Aldrich, St. Louis, USA) and purified on Sephadex G15 chromatography (Sigma–Aldrich) using 0.1 mM aqueous acetic acid, containing 2.5 g/L BSA as eluent. The specific activity of 125I-CB-LHRH' was about 0.2 - 0.24 mCi/mmol with a purity greater than 95 %. In the binding assay, polypropylene tubes were pretreated overnight at 4 °C with 10 mM Tris-HCl, pH 7.4 containing 1 % BSA. Membrane fraction aliquots (100 μg protein in 50 μL per tube) were incubated with 200 nM gonadorelin (Tash Biotechnology, Shanghai, China) was used as competitor. Reactions were terminated by adding 1 mL of 10 mM Tris-HCl buffer (pH 7.4, 4 °C). The solutions were subsequently filtered onto Whatman filters, washed with 1 mL of 5 % trichloroacetic acid and transferred to a counting tube. The radioactivity was measured by a gamma counter (GC-300, AnHui uste ZonKia Scientific Instruments co. LTD, Hefei, China). Specific binding was determined by subtracting nonspecific from total binding. The equilibrium dissociation constant (Kd) and maximum binding capacity (Bmax) were derived by the Scatchard method.
Kd represents the concentration of drug inducing the half (50%) biggest effect. Bmax indicates the maximum binding volume combined per mg receptor proteins. Each experiment was performed in triplicate and repeated three times.

**Hypersensitization test**
Under approved animal research protocol standards, the hypersensitive reaction against CB-LHRH' (2 mg/mL, equal to 438 μM) was assessed in guinea pigs. According to the Chinese Pharmacopoeia, six animals with an average weight of 250 g (XiFengYang Special Economic Animal Farm, Huzhou, China) were administered an intraperitoneal injection of 0.5 mL of CB-LHRH' every other day three times, while three additional animals received PBS physiological saline injections. Of the six animals administered CB-LHRH', three received an intravenous injection of 1 mL CB-LHRH' at day 15, and the remaining three at day 22. Allergic reactions, such as piloerection, sneezing, lacrimation, dyspnoea, and hyperspasmsia, were then monitored for the next 30 min.

**Antibacterial assay**
The antibacterial activity of CB-LHRH' was evaluated by the agar dilution method [41]. Muller Hinton's agar medium plates containing serial dilutions of CB-LHRH' were inoculated with 2 μL of tenfold-diluted 0.5 McF bacteria and incubated for 22 h at 35°C. The lowest concentration of CB-LHRH' inhibiting bacterial growth was considered the minimal inhibitory concentration (MIC). Nine clinical strains of the following species were evaluated: *Staphylococcus aureus*, *Klebsiella pneumoniae*, *Escherichia coli*, *Enterobacter cloacae*, *Pseudomonas aeruginosa*, and *Acinetobacter baumannii*.

**Xenograft studies**
Five- to six-week-old female (15–20 g) athymic nude mice (BALB/c-nu) were obtained from Shanghai SLAC Laboratory Animal Corporation, China. The animals were housed in sterile cages under laminar flow hoods in a temperature controlled room with a 12-hour light/12-hour dark cycle, and fed autoclaved chow and water *ad libitum*. All experiments were carried out in accordance with the guidelines for the welfare ethics of experimental animals [42]. All the animal-related procedures were approved by the Animal Ethical Committee of Zhejiang University.

**Experimental protocol**
Cultured ES-2 ovarian cancer cells were resuspended at $1 \times 10^7$/ml and inoculated subcutaneously into the nude mouse armpit. When tumors reached 60 mm$^3$, they were extracted, cut into pieces of about 2 mm$^3$, and inoculated subcutaneously into the nude mouse armpit. When the tumors reached an average volume of about 100 mm$^3$, the mice were randomly divided into five groups ($n = 10$). Groups 1–3 received intra-tumor injection of CB-LHRH' twice daily at 12.5, 25 or 50 mg/kg/d, respectively. Group 4 received 2 mg/kg cisplatin, once every 3 days [43], and group 5 the same volume of PBS, administered twice daily.

Tumor volumes were assessed every 3 d as 0.5 × length × width$^2$ (where length is the longest diameter across the tumor and width the corresponding perpendicular diameter as measured by calipers). On day 13, 4 h after the last dose, mice were sacrificed, and tumors excised and weighed; tumor growth inhibition rate was derived as $(1 - \text{tumor weight treated}/\text{tumor weight control}) \times 100\%$.

**Response criteria**
Anti-tumor activity was evaluated as relative growth of tumor volume (T/C, value), the mean relative tumor volume (RTV) for the treatment group divided by the mean RTV for the control group, as follows: $T/C(%) = \text{RTV}_t/\text{RTV}_c \times 100\%$. RTV = $V_t/V_0$, where $V_t$ is the volume on any given day, and $V_0$ the volume at treatment start. Agents producing a T/C of >60% were considered to be inactive, those with a mean T/C of ≤60% considered to be active.

General toxicity was evaluated based on body weight, measured weekly.

**Statistical analysis**
Data are mean ± standard deviation (SD). Normality and homogeneity of variance assumptions were assessed. One way analysis of variance (ANOVA) was used for multiple group comparison, as required. SPSS 16.0 (SPSS, USA) was used for the statistical analysis. $p < 0.05$ was considered statistically significant.

**Results**
**CB-LHRH’ inhibited proliferation of cancer cell lines**
Three drug-resistant ovarian cancer cell lines (SKOV-3, ES-2, NIH:OVCAR-3) and one drug-resistant endometrial cancer cell line (HEC-1A) were incubated with 12.5 μM CB-LHRH’. After 10 d there were almost no colonies after CB-LHRH’ treatment, while control wells contained approximately 30–50 colonies, indicating that 12.5 μM CB-LHRH’ significantly inhibited the growth of drug-resistant cancer cells. Damage to all four cell lines was observed within 24 h of exposure to 12.5 μM CB-LHRH’ by light microscopy (data not shown). And the effect of CB-LHRH’ on ES-2 cell morphology was shown in Fig. 1. Typical cell damage included aggregation, detachment from the plate, inflation, cytoplasm leak, disintegration into granules, and decreased cell number (Fig. 1).
Electron microscopy revealed cell membrane blebbing, the disappearance or reduction of microvilli, cell shrinkage, increased cellular granularity, the formation and separation of apoptotic bodies, and cytoplasm leak in response to 25 μM CB-LHRH in ES-2 cell (Fig. 2).

All four cell lines incubated for 4 h with CB-LHRH exhibited concentration-dependent cytotoxicity ($P < 0.001$) as shown in Fig. 3. And there were no differences among the four cell lines (Fig. 3). In contrast, CB-LHRH caused no hemolysis of red blood cells from healthy human donors (data not shown).

Proliferation of all four drug-resistant cancer cell lines was inhibited by CB-LHRH in a concentration-dependent manner ($P < 0.001$) (Fig. 4a–d). And the greatest inhibition...
was obtained for ES-2 cells, and CB-LHRH' inhibited cell proliferation in ES-2 cell in a time-dependent manner ($P < 0.05$) (Fig. 4c). Inhibition of SKOV-3 and HEC-1A cell proliferation was weaker at the dose of lower than 25 μM and 50 μM, respectively (Fig. 4a and 4b). And inhibition of NIH:OVCAR-3 cell proliferation was up to 40 % even at the high dose of 100 μM (Fig. 4d).

**CB-LHRH' is a high-affinity ligand for LHRHr**

The binding affinity of CB-LHRH' to the LHRH receptor was assessed in rat pituitary membranes by radioligand-binding assays. Kd and Bmax values were 0.2526 ± 0.061 nM and 110.4 ± 17.68 fmol/mg protein, respectively (Fig. 5), indicating that $^{125}$I-CB-LHRH' is a high-affinity ligand for LHRHr.

**CB-LHRH' does not induce hypersensitivity**

The potential of CB-LHRH' to induce a hypersensitive reaction in mammals was assessed in the guinea pig model. Guinea pigs administered 0.5 mL 2 mg/ml CB-LHRH' intraperitoneally every other day three times ($n = 6$) survived the 15 days of treatment and their weights did not differ significantly from animals administered PBS ($n = 3$). Afterwards, three animals were administrated an additional 1 mL of 2 mg/ml CB-LHRH'. No anaphylaxis occurred in any of the six guinea pigs treated with CB-LHRH' (data not shown).

**CB-LHRH' inhibits Gram-negative bacteria**

The antibacterial activity of CB-LHRH' was evaluated by the agar dilution method. CB-LHRH' killed or inhibited the growth of all tested gram-negative bacteria at micromolar concentrations. The minimal inhibitory concentrations (MIC) were 1.93–7.72 μM, 3.86–7.72 μM, 3.86–
7.72 μM, 3.86–7.72 μM, ≥15.44 μM, and >15.44 μM for
Acinetobacter baumannii, Escherichia coli, Enterobacter cloacae, Klebsiella pneumoniae, Pseudomonas aeruginosa,
and Staphylococcus aureus, respectively (Table 1).

CB-LHRH’ inhibits ES-2 Xenograft growth in vivo
Nude mice were implanted with ES-2 and adminis-
tered intra-tumoral injection of CB-LHRH’ twice daily
at 0, 12.5, 25 or 50 mg/kg, or 2 mg/kg cisplatin,
one day every 3 days (n=10); thirteen days later 9, 8, 9, 7
and 8 surviving mice were obtained, respectively
(Tables 2 and 3).

However, administration of CB-LHRH’ at 50 and
25 mg/kg significantly inhibited the growth of ES-2 Xe-
 nografts by day 13, resulting in tumor sizes of 50 %
and 59 % the control values (P < 0.01), and 23.8 %
and 20.4 % decrease in tumor weights (P < 0.05), respec-
tively; no differences were found in animal weights.

Discussion
In this study, we developed a novel LHRHr cytotoxin
strategy by linking Cecropin B to the modified LHRH’
receptor ligand in order to deliver an effective broad-
spectrum toxin specifically to tumor cell targets, without
affecting gonadotrophin secretion in healthy cells or in-
ducing resistance.

Table 1 The MIC of the Cecropin B-LHRH’ against the six kinds
of clinically isolated bacteria

| Bacteria                  | MIC (μM)       |
|---------------------------|----------------|
| Acinetobacter baumannii   | 1.93–7.72      |
| Escherichia coli          | 3.86–7.72      |
| Enterobacter cloacae      | 3.86–7.72      |
| Klebsiella pneumoniae     | 3.86–7.72      |
| Pseudomonas aeruginosa    | > = 15.44      |
| Staphylococcus aureus     | >15.44         |

CB-LHRH’ binds to LHRHr with a low dissociation
constant of 0.252 ± 0.061 nM, indicating the high-affinity
of the modified peptide, which should have broad-
spectrum toxicity against cancer cells expressing LHRHr.
Indeed, CB-LHRH’ was cytotoxic to ES-2 and NIH:Ov-
CAR-3 (ovarian cancer cells) and HEC-1A (endometrial
cancer cells) which express LHRHr, while displaying
the weakest cytotoxicity in the LHRHr negative ovarian can-
cer cell line SKOV-3. Of note, the four cancer cell lines
sensitive to CB-LHRH’ are known to be resistant to
several cytotoxic drugs, including diphtheria toxin,
doxorubicin, cisplatin, and adriamycin, among others,
according to the ATCC Global Bioresource Center
(USA). For instance, ES-2 cells were derived from an
ovarian clear cell carcinoma characterized by unique
clinical features such as high incidence in stage I of the
disease, relatively strong resistance to conventional plat-
inum or taxane-based chemotherapies, and poor prog-
noses [44–46]. The overt cytotoxicity demonstrated for
CB-LHRH’ against several multidrug-resistant cancer
cell-lines, particularly ES-2 cells, indicates that the new
peptide might help in the treatment of some cancers re-
sistant to currently available chemotherapeutics; indeed,
CB-LHRH’ is a promising candidate for the treatment of
LHRHr dependent cancers.

Table 2 Therapeutic effect of Cecropin B-LHRH’ on the volume
of ES-2 tumors xenografted into nude mice (Mean ± SD)

| Group   | Dose (mg/kg,d) | Tumor Volume (mm³) | RTV | T/C (%) |
|---------|----------------|--------------------|-----|---------|
|         | d 1            | d 13               |     |         |
| CB-LHRH’| 50             | 308.91 + 110.19    | 965.92 + 212.31** | 3.13 | 50     |
|         | 25             | 364.18 + 112.16    | 1334.78 + 179.25** | 3.67 | 59     |
|         | 12.5           | 298.23 + 107.54    | 1416.71 + 353.71** | 4.75 | 76     |
| Cisplatin| 0.67           | 343.91 + 123.06    | 913.16 + 142.64** | 2.66 | 42     |
| PBS     | 326.56 + 85.00 | 2042.62 + 239.07   | 6.5 | –       |

In comparison to the PBS control, ** P < 0.01
patients with its antibacterial function, since bacterial infections are a major cause of morbidity and mortality in neutropenic patients following chemotherapy for malignancy [49].

Conclusion

LHRH' may represent a more promising therapeutic than LHRH, and cecropin B appears to be a potentially valuable component of targeted therapies. The peptide drug CB-LHRH’ described here is a potent candidate for cancer therapy, with broad anti-cancer spectrum and minimal toxicity to normal eukaryotic cells.

Ethics approval and consent to participate

The study was approved by the ethical committee of Women’s Hospital, School of Medicine, Zhejiang University, Hangzhou, China. Informed consent was obtained from all individual participants included in the study.

Consent for publication

Not applicable.

Availability of data and materials

The datasets supporting the conclusions of this article are included within the article.

Abbreviations

AMP: antimicrobial peptide; Bmax: maximum binding capacity; BSA: bovine serum albumin; FSH: follicle-stimulating hormone; Kd: equilibrium dissociation constant; LDH: lactate dehydrogenase; LH: luteinizing hormone; LHRH: luteinizing hormone-releasing hormone; LHRHr: luteinizing hormone-releasing hormone receptor; MIC: minimal inhibitory concentration; MITT: 3-(4, 5-dimethylthiazol-2-yl)-2,5-diphenyl-tetrazolium bromide; PBS: phosphate-buffered salt solution; RTV: relative tumor volume; T/C value: RTV of the treatment group/RTV of the control group.

Competing interests

The authors declare that they have no competing interests.

Authors’ contributions

XYL conceived and designed the experiments and write most of the manuscript. BS carried out Radioligand Receptor Binding Assays and analyzed the data. QC performed experiments and analyzed the data. XHZ participated in study design and performed the statistical analysis. YQY, FMW carried out Antibacterial Assays. XMZ conceived of the study, and participated in its design and coordination, and helped draft the manuscript. All authors read and approved the final manuscript.

Acknowledgments

This study was supported by grant from the National Natural Science Foundation of China (No. 30700896 and No. 81270672).

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Received: 13 October 2014 Accepted: 22 March 2016

Published online: 28 March 2016

Table 3 Therapeutic effects of Cecropin B-LHRH' on the weight of ES-2 tumors xenografted into nude mice (Mean ± SD)

| Group          | Tumor Weight (g) | Tumor growth inhibition rate (%) |
|----------------|------------------|----------------------------------|
| 50 mg/kgd CB-LHRH' | 1.31 ± 0.31*     | 23.8                             |
| 25 mg/kgd CB-LHRH'  | 1.37 ± 0.15*     | 20.4                             |
| 12.5 mg/kgd CB-LHRH' | 1.40 ± 0.81      | 18.4                             |
| Cisplatin            | 1.09 ± 0.25 **   | 36.4                             |
| PBS                    | 1.72 ± 0.24      | -                                |

In comparison to the PBS control, *P < 0.05; **P < 0.01
Each experiment was performed in triplicate and repeated three times.

The mechanisms underlying the antimicrobial and antitumor effects of cecropin include plasma membrane disruption via micellization, or pore formation by peptide-lipid interactions [47, 48]. Cationic peptides target characteristic, cell-surface, anionic lipids by electrostatic attraction that are ubiquitous and exclusive to micro-organisms, followed by membrane permeation and disruption. This simple electrostatic recognition makes the development of resistance difficult. Eukaryotic cell membranes have important amounts of sterols, and the membrane-stabilizing cholesterol protects normal eukaryotic cells from attacks by therapeutic peptides [47, 48]. Changes in lipid compositions of cancer cell membranes may be a reason why these peptides show specific toxicity to cancer cells [30]. In this study, loss of cytoplasm and disintegration of cancer cells, with no hemolytic activity in normal red blood cells, supports similar anticancer mechanisms of plasma membrane disruption.

CB-LHRH’ also induced no hypersensitive reactions in guinea pigs, suggesting that CB-LHRH’ may not cause severe allergy, a common problem of peptide drugs. This benign profile distinguishes the newly developed polypeptide from other chemotherapeutics, as the safety of chemotherapeutics is one of the most important features of contemporary drug treatment. However, these results need to be replicated in human subjects in order to rule out potential relevant inter-species differences. In addition, the anticancer effects of CB-LHRH’ was assessed after intra-tumoral administration, which does not fully recapitulate the targeting of tumors in the peritoneal space.

CB-LHRH’ also inhibited the growth of clinically isolated bacteria at micromolar concentrations, indicating a potent antibacterial effect. The antibacterial and anticancer functions of CB-LHRH’ suggest that the carboxyl-terminal extension of CB doesn’t alter its antibacterial properties. In addition to inhibiting growth of cancer cells, CB-LHRH’ may also benefit cancer
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