Antitumor Activity of Antimicrobial Peptides Containing CisoDGRC in CD13 Negative Breast Cancer Cells

Lei Hou¹, Xinhan Zhao¹*, Pei Wang¹, Qian Ning¹, Min Meng¹, Caigang Liu²

¹ Department of Oncology, The First Affiliated Hospital of Medical School of Xi’an Jiaotong University, Xi’an, Shaanxi Province, China, ²Department of Breast Surgery, General Surgery, the First Hospital of China Medical University, Shenyang, Liaoning Province, China

Abstract

Background: isoAsp-Gly-Arg (isoDGR) is a derivative of the Asn-Gly-Arg (NGR) motif, which is used as a targeted delivery tool to aminopeptidase N (CD13) positive cells. Recent studies have shown that cyclic isoDGR (CisoDGRC) has a more efficient affinity with $\alpha_v\beta_3$, a type of integrin that overexpresses in tumor cells. Antimicrobial peptides (AMPs) are an efficient antitumor peptide that specifically kills tumor cells. In the present study, we designed antimicrobial peptides containing the CisoDGRC motif (CDAK) and assessed its antitumor activity for CD13+/alpha_v/beta_3+ breast cancer cells (MCF-7 and MDA-MB-231) in vitro and in vivo.

Methods: In vitro: We assessed the cytotoxicity of CDAK for MCF-7 and MDA-MB-231 breast cancer cells, the human umbilical vein endothelial cell (HUVEC), and human foreskin fibroblasts (HFF). We performed an apoptosis assay using Annexin-V/PI, DNA ladder, mitochondrial membrane potential, and Caspase-3 and Bcl-2. The effect on cell cycles and affinity with cell were tested using flow cytometry and fluorescent microscopy and the effect on invasion was analyzed using an invasion assay. CDAK was injected intravenously into tumor-bearing athymic nude mice in vivo experiment.

Results: CDAK showed cytotoxic activity in MCF-7 and MDA-MB-231 cells, whereas HUVEC and HFF were less sensitive to the peptides. CDAK induced apoptosis, reduced mitochondrial membrane potential, promoted Caspase-3, and inhibited Bcl-2 expression in the two breast cancer cell lines. In addition, CDAK inhibited proliferation of cancer cells through S phase arrest, and own selective affinity with MCF-7 and MDA-MB-231 cells, inhibited the invasion of MDA-MB-231 cells. In vivo, CDAK significant inhabited the progression of the tumor and the generation of neovascularization.

Conclusion: Antimicrobial peptides containing the CisoDGRC (CDAK) motif could efficiently exhibit the antitumor activity for CD13+/alpha_v/beta_3+ breast cancer cells.

Introduction

Various compounds and particles have been coupled or added synthetically to the Asn-Gly-Arg (NGR) motif to target recognized aminopeptidase N (CD13), which are over-expressed in angiogenic blood vessels and some tumors cells, which increases antitumor activity. This includes chemotherapeutic drugs, gene molecular, and pro-apoptosis peptides [1,2,3]. Some tumors cells, however, express little or do not express CD13, which reduced the efficiency of NGR. Recent research has shown that the Asn residue of NGR can deamidate and generate Asp-Gly-Arg (DGR) not only has the canonical RGD/alpha_v/beta_3 contacts, but also establishes additional polar interactions [12]. In addition, studies have shown that the cyclic isoDGR (CisoDGRC) motif with a disulfide bridge constraint can increase the tumor targeting efficiency and stability of bent conformation [13]. CisoDGRC peptides as a targeted delivery of tumor necrosis factor alpha (TNF) has been explored and shown promising results. CisoDGRC thus appears to be a promising candidate as a tool for targeted delivery of anti-agents for CD13+alpha_v/beta_3+ tumor cells, such as MCF-7 and MDA-MB-231 cell lines [14,15].

Antimicrobial peptides (AMPs) have drawn attention as a promising alternative to current antitumor agents. Antimicrobial peptides are linear, cationic and alpha-helix-forming. Cationic amino acids electrostatically attract the anionic of phospholipids to disrupt negatively charged cell membranes or form non-permeabilized pore-like structures that permit translocation of the AMPs into the cytoplasm without cell lysis [16]. The outer leaflet of the cell membrane owns the negatively charged phosphatidylyserine (PS) in tumor cells. The negatively charged PS, however, is localized exclusively to the inner leaflet of the membrane in
normal cells [17]. The difference in the composition of the cell membrane between tumor cells and normal cells leads tumor cells to be more sensitive to AMPs. In addition, another possibility for tumor cells damage is the induction of apoptosis in cancer cells via mitochondrial membrane disruption after AMPs enter the cytoplasm [18].

In the present study, we conjugate CisoDGRC and AMPs to develop a novel tumor targeting AMP to exploit its antitumor properties for CD13-/-β5 tumor cells.

Materials and Methods

Design of the peptide

The sequence of antimicrobial peptides containing the CisoDGRC motif is shown in the following: (1) CisoDGRCLLIK-LAKLAKKLAKL (CDAK), (2) CRDGCAKALIKKLK-LILALK (CRLK). KLAKLAKKLAKLAK is a type of sequence from antimicrobial peptides [19]. As a linker, LLII fuses the two functional domains. CRLK was a random sequence of CDAK. Peptides were synthesized and then purified by high-performance liquid chromatography to 95% (Peptide Synthesis Facility, Guangzhou Bio Integration Biotechnology Company, Guangzhou, China), dissolved in deionized water to 1 mg/ml, and saved in −20°C.

Cell lines

Four cell lines as follows were purchased from the cell bank of the Chinese Academy of Sciences (Shanghai, China): (1) human breast cancer cell lines MCF-7 and MDA-MB-231; (2) human foreskin fibroblasts (HFF), and (3) human umbilical vein endothelial cells (HUVEC). Cells had a monolayer long-term culture in a flask of RPMI1640 medium supplemented with 10% fetal bovine serum (FBS; Hyclone, Logan, Utah, USA).

Cell viability assay

The MCF-7 and MDA-MB-231 cells were cultured in 96-well plates at a density of 5×10^4 cells/well. CDAK and CRLK were added to different wells, ranging from 10 μg/ml to 200 μg/ml, respectively. After 24 h, 48 h and 72 h, 3-[4,5-dimethylthiazol-2-yl]-2,5-diphenyltetrazolium bromide (MTT; Sigma-Aldrich, St. Louis, MO, USA) was added to each well and incubated for 4 h. The MTT was dissolved in 200 μL of Dimethyl sulfoxide (DMSO). Absorbance was detected using a microplate reader at 550 nm and 620 nm, respectively, with exposure at 400 ms. Images were recorded using a CDD camera (ECLIPSE TE2000-U, Nikon, Tokyo, Japan).

Fluorescence microscopy detections

Cells were incubated with CDAK and CRLK (10 μg/ml; Rhodamine B was connected to the K of the amino acid residue’s end) for 12 h and washed with PBS three times. They were detected by a fluorescence microscope (Ex and Em wavelengths were set at 550 nm and 620 nm, respectively, with exposure at 400 ms). Images were recorded using a CDD camera (ECLIPSE TE2000-U, Nikon, Tokyo, Japan).

Flow cytometry

Annexin-V/PI double notation detects apoptosis. Cells were harvested by trypsinization followed by washing with phosphate buffered saline (PBS) two times after being treated with CDAK using the dose of IC50 (MCF-7 were 190 μg/ml, MDA-MB-231 were 212 μg/ml; CRLK/200 μg/ml and control (PBS) for 24 h. They were then incubated with 500 μL Binding Buffer and Annexin-V-FITC (20 μg/ml) 10 ul at room temperature for 30 minutes. We then added propidium iodide (PI) (50 μg/ml), 5 ul for five minutes. The samples were then analyzed using flow cytometry (BD Biosciences, San Diego, CA, USA).

Mitochondrial membrane potential (Ψm) was analyzed using flow cytometry through the JC-1 mitochondrial membrane potential assay kit (Cell Technology Company, Minneapolis, MN, USA). Cells were harvested by trypsinization followed by washing with PBS two times after being treated with CDAK (the dose were 190 μg/ml and 212 μg/ml respectively in MCF-7 and MDA-MB-231), CRLK/200 μg/ml and control (PBS) for 24 h, then incubated with 500 μL JC-1 working fluid for 20 minutes, washed with 500 μL staining solution twice, and then suspension with an incubation buffer. The cells were analyzed using flow cytometry.

The affinity rate of peptides was analyzed using flow cytometry. Cells were seeded in 6-well plates at a density of 5×10^4 cells/well. After 24 h, the cells were treated with CDAK and CRLK (10 μg/ml; Rhodamine B was connected to the K of the amino acid residue’s end) for 12 h. At the end of the incubation, the cells were harvested by trypsin and washed with PBS three times. We then proceeded with the fluorescence-activated cell sorting (FACS).

Analysis of cell cycle dynamics and DNA fragmentation

After being incubated with CDAK (190 μg/ml and 212 μg/ml), CRLK (200 μg/ml) and control (PBS) for 24 h, 20 μM BrdU (Sigma-Aldrich, St. Louis, MO, USA) was added to the media for 30 min. Cells were harvested and fixed in ice-cold 75% ethanol for 24 h. Then, the cells were treated with 4N HCL, neutralized by 0.1 M borax, and washed with PBS containing 0.05% bovine serum albumin (BSA). The cells were incubated sequentially with anti-BrdU antibody (santa cruz biotechnology, California, US) and FITC-conjugate anti-mouse secondary antibody in the presence of 0.5% BSA and 0.5% Tween-20. Cells were added PI (50 μg/ml) and RNase A (50 μg/ml) for 20 minutes and analyzed by flow cytometry. Select a forward scatter (FSC) versus side scatter (SSC) window defined single cells and exclude cell doublets.

For DNA fragmentation analysis, DAN of cells which were treated with CDAK (190 μg/ml and 212 μg/ml), CRLK (200 μg/ml) and control (PBS) for 24 h were extracted using a selective apoptotic DNA ladder extraction KIT (Biomed Company, Beijing, China) and were analyzed using 1% agarose gels. DNA was visualized with ethidium bromide staining.

Western-blot

Cells were treated with CDAK (190 μg/ml and 212 μg/ml), CRLK (200 μg/ml) and control (PBS) for 24 h and washed with cooled PBS three times. They were then added to RIPA 140 ul and the protease inhibitors (2 mg/ml) for 30 minutes. The lysates were then centrifuged at 12,000 g at 4°C for 20 minutes and boiled in a loading buffer for 5 minutes. The total protein was normalized using a Bradford protein assay, and 150-microgram protein samples were loaded on sodium dodecyl sulfate-polyacrylamide gels and transferred to polyvinylidene difluoride membranes. The membrane was probed with rabbit monoclonal caspase-3, bel-2, αv, β3, and the CD13 antibody purchased from Abcam, (Abcam, Cambridge, UK) at 4°C overnight. After washing three times in Tris hydrochloride buffer (TBS) containing 0.1% Tween-20 (TBS-T), the second antibody was incubated with the membranes for 2 h at room temperature. The membranes were then washed with TBS-T three times for 10 minutes and exposed to X-ray film. Autoradiogram signals were quantified using a gel densitometric scanning program. The relative expression of protein was determined from the optical density ratio of the corresponding protein bands.
Invasion assay

Invasion assay was performed in a Matrigel-coated invasion chamber (Matrigel, BD Biosciences, San Diego, CA; Transwell Chamber, Millipore, Massachusetts, USA). MDA-MB-231 cells were plated in the upper chamber at a density of 1 x 10^5 cells/100 μl in serum-free RPMI-1640 medium and were then treated with CDAK and CRLK (10 μg/ml). The lower chamber was filled with RPMI-1640 medium containing 20% FBS. After incubating at 37°C for 24 h, the nonmigrated cells were scraped in the upper chamber with a cotton swab, and fixed the migrated cell on the lower surface of the porous membrane with methanol. The cells were then stained with crystal violet and counted by a light microscope.

In vivo efficacy in a xenograft model

The experiment was approved by the Animal Care and Use Committee of Xi’an Jiaotong University. MDA-MB-231 cells (2 x 10^6) were injected subcutaneously into the right flank of 6- to 9-week-old female BALB/cnu-nu athymic nude mice (Shanghai Siliäke Laboratory Animal Co., Ltd, Shanghai, China) weighing 18 to 20 g. When the tumor reached 60 mm³ in size, the mice were randomized into three groups: (1) CDAK (4 mg/kg); (2) CRLK (4 mg/kg); and (3) saline (control). They were then injected intravenously (50 μL/injection) three times a week for three weeks. Tumor volume was measured three times a week using calipers to calculate the tumor size using the following formula: length x width² x 0.5. All values are expressed as the mean ± SD.

Tumor-bearing athymic nude mice were sacrificed and the weights of the tumors were recorded. The tumor tissue, liver as well as lung tissues of mice were paraffin-embedded. The tumor paraffin sections were incubated for 10 minutes with 3% H₂O₂ deionized water to eliminate the endogenous peroxidase activity, washed in PBS three times for five minutes, 5% goat serum was added for 15 minutes, then incubated with mouse monoclonal CD105 antibody (Abcam, Cambridge, UK) at 4°C overnight and washed with PBS three times for five minutes. The biotin-labeled goat anti-mouse IgG was incubated with sections at 37°C for 15 minutes, and then the sections were washed by PBS three times for five minutes. Horseradish peroxidase-avidin enzyme working solution was added at 37°C for 15 minutes and washed with PBS. DAB was added to develop the color, and the nuclei were counterstained mildly with hematoxylin. The liver and lung paraffin sections were stained with hematoxylin and eosin (HE) and were independently evaluated by two pathologists.

Terminal deoxynucleotidyl transferase-mediated dUTP nick end-labeling (TUNEL) were examined in the lung and liver using TdT In Situ Apoptosis Detection Kit ( Trevigen, Gaithersburg, Maryland, USA) following the manufacturer’s protocols. Apoptosis cells were identified as having brown nuclei under a light microscope. The number of apoptosis cells was counted in five random fields (x400) in a blinded manner.

Statistical analysis

The experiments with more than two treatment groups and various treatment concentrations were tested by univariate ANOVA, followed by Bonferroni or Dunnett’s for multiple comparisons. All values are presented as the mean ± SD. An alpha level of <0.05 was used as the criterion of significance. Results were reproduced in three independent experiments.

Results

Test of cytotoxicity on CDAK for CD13 negative breast cancer cell

We firstly examined the expression of CD13 and α,β on the MCF-7, MDA-MB-231, HUVEC, and Fibroblast cells using Western-blot. As shown in Figure 1A, we did not detect the expression of CD13 in MCF-7 or MDA-MB-231 cells. In contrast, the two cell lines all expressed α,β. HUVEC and HFF cells showed a double positive expression on the protein of CD13 and α,β. The MCF-7 and HUVEC cells expressed more α,β compared to MDA-MB-231 cells. CD13 was expressed higher in HUVEC than HFF. We then assessed the cytotoxic activity of CDAK on the two kinds of CD13⁺/α,β⁺ breast cancer cells and the two normal cells through MTT. The results indicated that the treatment with CDAK induced dose-dependent cytotoxicity in the MCF-7 and MDA-MB-231 cells (Figure 1B). We found significant cytotoxicity at the >40 μg/ml concentration (P<0.05).

However, the cytotoxicity decreased after 48 h. The IC50 of CDAK (the peptides’ concentration that induces 50% inhibition of cell growth) were 190 μg/ml and 212 μg/ml for MCF-7 and MDA-MB-231 cells respectively at 24 h (Table 1), which were 1/4 to 1/5 compared to HUVEC and HFF cells, respectively. These data suggest that MCF-7 and MDA-MB-231 cells were susceptible to CDAK more than normal cells. There was a correlation between the expression of α,β and the cytotoxicity of CDAK in cancer cells.

Characterization of the breast cancer cell death mechanism by CDAK

To explore whether apoptosis played important role in the cytotoxicity of CDAK, MCF-7 and MDA-MB-231 cells were treated with CDAK for 24 h. As shown in Figure 2A and 2B, the outcome of the Annexin V/PI detection showed that CDAK increased the percentage of apoptosis cells in both the MCF-7 and MDA-MB-231 cells (P<0.05). The DNA ladder assay showed that no DNA ladder was detected in the MCF-7 and MDA-MB-231 cells treated with CRLK. However, the formation of DNA nucleosome ladders was clearly detected in the MCF-7 and MDA-MB-231 cells treated with CDAK. In addition, the expression of Caspase-3 and Bcl-2 were detected by Western-blot (Fig. 2C). Caspase-3 increased 8.5 times and 2.8 times in MCF-7 and MDA-MB-231 cells treated with CDAK compared to control (P<0.05), and Bcl-2 decreased 96% and 92% in MCF-7 and MDA-MB-231 cells treated with CDAK compared with control (P<0.05).

Previous experiments have shown that AMPs induce tumor cell apoptosis by disrupting the mitochondria. To confirm this mechanism further, mitochondria membrane potential (Ψm) was assessed using JC-1 staining. The disrupting of the mitochondria along with dropping of Ψm, and the variance of fluorescence ratio (red/green) indicate the variance of Ψm. After treated with CDAK, the mitochondria membrane potential decreased 33% and 28% respectively in MCF-7 and MDA-MB-231 cells compared to CRLK (P<0.05) (Figure 2D).

Affinity of CDAK in CD13 negative breast cancer cells

Rhodamine B was connected to the K of the amino acid residues’ end. MCF-7, MDA-MB-231, HUVEC, and Fibroblast cells were treated with CDAK and CRLK (10 μg/ml) for 12 h. The affinity rate of CDAK was then assessed by flow cytometry (Figure 2E). The result showed that CDAK had significant higher affinity rate with both MCF-7 and MDA-MB-231 cells compared to HUVEC and HFF cells (P<0.05). At the same time, we further assessed the affinity using fluorescence microscopy. As shown in...
Figure 3A, CDAK affinity was evident in the MCF-7, MDA-MB-231 and HUVEC cells, but not in the Fibroblast cells. These data further verified that CisoDGRC motif has the ability to specifically distinguish and bind tumor cells.

The effect of CDAK on cell cycle in CD13 negative breast cancer cells
We analyzed the effect of CDAK on the cell cycle. The results showed that the cells were significantly arrested in the S phase after incubated with CDAK for 24 h ($P < 0.05$) (Fig. 3B). The change between the control and CRLK did not show a significant difference ($P > 0.05$).

CDAK inhibits invasion of CD13 negative breast cancer cell
A transwell chamber invasion assay was performed using the MDA-MB-231 cells, which are high invasion cancer cells, to determine whether CDAK is capable of inhibiting the invasion of the MDA-MB-231 cells. As shown in Figure 3C, invasion of the MDA-MB-231 cells were significantly inhibited in the CDAK compared with the control group and CRLK ($P < 0.05$).

The effect of CDAK’s antitumor activity on breast cancer xenograft growth
Finally, we tested activity of CDAK in female athymic nude mice implanted with MDA-MB-231 cells. The CDAK and CRLK concentration used in the present study refers to a previous animal study [20]. As shown in Figure 4A and 4B, the tumors treated with CDAK were significant smaller than the control group ($P < 0.05$). The CRLK treated tumors showed a similar trend, but the differences did not reach a statistically significance compared with control group. For example, after being injected for 25 d, the average volume of tumor in control mice was about two- to three fold higher compared with mice treated with CDAK. The average weights of tumors were also lower in the CDAK group compared to the CRLK and control mice ($P < 0.05$).

As shown in Figure 4C, CDAK significantly inhibited the angiogenesis while not CRLK ($P < 0.05$). The data also indicated that CDAK inhibited angiogenesis in vivo. No abnormalities were observed in the lung and liver of mice in the histological examination (Fig. 4D). The TUNEL assay showed that apoptosis was not significant induced by CDAK in the lung and liver compared with control ($P > 0.05$) (Fig. 4E, table 2).
Therapy peptides are gaining increased popularity to treat cancerous tumors. This might be the case because we know that peptides are generally less expensive than antibody-based therapy due to the development of recombinant or solid-phase chemical synthesis techniques. And, compared with proteins, peptides have higher solubility in water, and lower immunogenicity [21,22]. In the present study, we linked two functional domains to construct a novel antimicrobial peptide containing isoDGR motif to selectively kill CD13/αvβ3+ breast cancer cells. The results showed that the

**Discussion**

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synthesized peptides have an efficient antitumor effect for CD13/αvβ3+ breast cancer cells in both vitro and vivo experiments.

CD13 were up-regulated in angiogenic blood vessels, tumor cells, pericytes, fibroblasts, and smooth muscle cells. NGR, therefore, is an efficient tool to deliver targeted treatment to these cells [23]. CD13 is not present in all tumors, in fact, it was found in only 36.2% of breast cancer patients [24]. NRG motif only indirectly killed tumor cells by disrupting the angiogenic blood vessel in CD13 negative tumor cells. Current studies, however, have discovered the isoDGR, a derivative of NGR, has a high affinity for αvβ3, which is expressed at low levels in epithelial cells and mature endothelial cells, but is up-regulated on tumor cells and tumor endothelial cells [25]. The isoDGR motif is therefore a proper candidate to be used as an antitumor agent, which can be targeted to CD13/αvβ3+ tumor cells. In the present study, we connected two functional domains, antimicrobial peptides and isoDGR, to conjugate novel antitumor peptides that selectively bind integrin αvβ3 to kill tumor cells. We used cyclic isoDGR, which could bind αvβ3 with an affinity >100-fold higher than that of linear isoDGR, to target kill αvβ3+ tumor cells. The experiment results demonstrate that the CisoDGRC motif has a higher affinity for MCF-7 compared to MDA-MB-231 cells, and the result of Western-blot assay shown MCF-7 have more expression of αvβ3 compared to MDA-MB-231. This further verified the relationship between CisoDGRC motif and αvβ3-positive tumor cells. Furthermore, the cytotoxicity assay demonstrated that synthesized peptides owned selective cytotoxicity for MCF-7 and MDA-MB-231 cells, especially MCF-7, but left normal cells unaffected. Prior studies have shown that the antimicrobial peptides sequence used in the present study is non-toxic outside of the cell, but toxic when internalized into the cell by disrupting the mitochondrial membrane [26]. The apoptosis studies showed synthesized peptides induced the apoptosis of MCF-7 and MDA-MB-231 cells, and disrupted mitochondrial membrane potential promotes the expression of Caspase-3 and inhibits Bcl-2. At the same time, the assay of the peptide...
cell cycle showed that the cells were arrested in the S phase, which demonstrates that CDAK not only promoted apoptosis, but also inhibited the proliferation of MCF-7 and MDA-MB-231 cells. In vivo experiment, the synthesized peptides significantly inhibited the xenograft growth. Other studies have shown that some antitumor agents adding the ligand of $\alpha_v\beta_3$, such as RGD [27], which could inhibit tumor angiogenesis that have a high expression of $\alpha_v\beta_3$. We used CD105 to stain tumor angiogenesis, and the result verified that CDAK significantly inhibited tumor angiogenesis. Regarding how the mechanism inhibited angiogenesis, $\alpha_v\beta_3$ is expressed on endothelial cell during neovascularization, but is not strongly expressed on quiescent endothelial cell [28]. We speculated, therefore, that AMPs containing $\text{CisoDGR}$ could target recognized endothelial cell of angiogenesis to induce the apoptosis. We will verify this in our next experiment. At the same time, the histochemical and TUNEL assays showed the peptides had little effect on normal tissue.

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AMPs containing $\text{CisoDGR}$, like all targeting antitumor drugs, offers the possibility of targeting the $\text{CD13}^+\alpha_v\beta_3^+$ breast cancer cells. The results of the present experiment showed that the synthesized AMPs owned a high selectivity for $\text{CD13}^+\alpha_v\beta_3^+$ breast cancer cell and less cytotoxicity toward normal cells. The

**Figure 4. Antitumor activity of CDAK in breast cancer xenograft in vivo.** (A) Tumor–growth curve. MDA-MB-231 cell lines were implanted into the right flank of female athymic mice. When the tumor size reached about 60 mm$^3$, mice received an intravenous injection of CDAK (4 mg/kg) and CRLK (4 mg/kg), or saline used as control (three times a week). The tumor volumes of the tumors were measure three times a week. The tumor volume of CDAK group was smaller than control and CRLK, $P < 0.01$. (B) The athymic mice were sacrificed as 25 d after being injected with CDAK and CRLK and the weight of each tumor was measured. The CDAK group had low weight compared to the control and the CRLK groups, $P = 0.003$. (C) The inhibition of angiogenesis was evaluated in tumor sections using immunohistochemistry with anti-CD105 antibody. Angiogenesis was quantified by image analysis of Microvessel density (MVD) (400× magnification). CDAK caused significant inhibition of angiogenesis, $P = 0.003$. (D) Images (200× magnification) of the lung and liver from athymic mice were obtained by staining with hematoxylin and eosin (HE). No significant damage was observed. (E) Representative photographs of the tumor section examined by TUNEL assay, light microscope (×400). The number of apoptotic cells was counter 5 random field in blinded manner. There differences of apoptotic cell were not statistic significant in the three groups (table 2) (ANOVA assay). The images were analyzed by pro plus 5.0. Each group had five nude mice. Data are shown as mean ± SD. *$P < 0.05$ versus control, **$P < 0.05$ versus CRLK.

**Table 2.** Comparison of apoptosis ratio in liver and lung of athymic mice (mean ± SD, n = 5).

| tissue | group | apoptosis rate (%) | P value (versus control) |
|--------|-------|--------------------|--------------------------|
| lung   | control | 0.14±0.01          |                          |
|        | CRLK   | 0.17±0.01          | 0.64                     |
|        | CDAK   | 0.16±0.02          | 0.48                     |
| liver  | control | 0.33±0.01          |                          |
|        | CRLK   | 0.35±0.23          | 1                        |
|        | CDAK   | 0.38±0.14          | 0.25                     |

The images were analyzed by pro plus 5.0. Each group had five nude mice. Data are shown as mean ± SD. *$P < 0.05$ versus control, **$P < 0.05$ versus CRLK.

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selective property provides the efficient antitumor activity. Further studies to examine the mechanisms of antitumors, safety, immunogenicity, and antitumor spectrum will broaden the indication of AMPs containing CisDGRC for the future.

**Conclusion**

The present study found that AMPs containing CisDGRC selectively kill CD13⁺/αvβ3⁺ breast cancer cells by inducing apoptosis and inhibiting proliferation. Furthermore, the vivo experiments showed that AMPs containing CisDGRC could significantly inhibited CD13⁺/αvβ3⁺ tumor progression and angiogenesis.

**Author Contributions**

Conceived and designed the experiments: XZ. Performed the experiments: LH XZ PW QN MM CL. Contributed reagents/materials/analysis tools: LH XZ PW QN MM CL. Wrote the paper: LH.

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