Structure and function of antimicrobial peptide penaeidin-5 from the black tiger shrimp Penaeus monodon

Shao-Yang Hu a,1, Jan-Hsiung Huang b,1, Wei-Ting Huang a, Yang-Hui Yeh a, Mark Hung-Chih Chen c, Hong-Yi Gong a, Tze-Ting Chio d, Tzu-Hsuan Yang b, Thomas T. Chen e, Jenn-Khan Lu d,⁎, Jen-Leih Wu a,⁎

a Institute of Cellular and Organismic Biology, Academia Sinica, Nankang, Taipei 115, Taiwan
b Institute of Microbiology and Biochemistry, National Taiwan University, Taipei 106, Taiwan
c Department of Biotechnology, Hung Kuang University, Taichung, Taiwan
d Department of Aquaculture, National Taiwan Ocean University, Keelung 202, Taiwan
e Department of Molecular and Cell Biology, University of Connecticut, Storrs, CT 06269, U.S.A.

Abstract

The gene for penaeidin-5, an antimicrobial peptide comprising 55 amino acids, was isolated from the hemocyte of black tiger shrimp (Penaeus monodon). RT-PCR expression tests revealed that penaeidin-5 was produced in hemocytes, gills, the intestine and muscle. Western blot analysis confirmed the penaeidin-5 was abundant in hemocytes, the intestine and hemolymph. Immunohistochemistry revealed penaeidin-5 in the cuticle and gills that are considered primary defense barriers. The deduced amino acid sequence of penaeidin-5 included a proline-rich N-terminal domain and a carboxyl-domain that contained six cysteine residues. Circular dichroism analysis revealed an α-helix in its secondary structure and the predicted 3D structure indicated two-disulfide bridges in the α-helix. Based on the sequence of penaeidin-5 peptide cDNA, synthetic penaeidin-5 was prepared to carry out functional tests. The synthetic peptide had efficient bacteriostatic and bactericidal activity against Aerococcus viridans, and also inhibited the growth of two filamentous fungi, Fusarium pisi and Fusarium oxysporum. To measure penaeidin-5 in vivo, black tiger shrimp were challenged with Vibrio alginolyticus and A. viridans. At 3 h post-challenge, penaeidin-5 was induced and bacterial numbers decreased significantly by 12 h and 24 h.

Keywords: Penaeus monodon; Antimicrobial peptides; Penseidins; Innate immunity

1. Introduction

Antimicrobial peptides (AMPs) are considered to be key component of the innate immune system and they are widespread throughout the animal and plant kingdoms. Since the discovery of the first AMP cecropin from diapausing pupae of the lepidopteran Hyalophora cecropia (Boman and Hultmark, 1987), many AMPs have been identified and characterized from a wide
variety of vertebrate and invertebrate species (Hertru et al., 1994). Based on amino acid sequences, structural features and functional features, AMPs have been divided into distinct subgroups with antimicrobial activities predominantly against particular pathogens (Bulet et al., 1999). Thus, more and more AMPs are being studied as potential therapeutic agents for aquaculture, agriculture and human medicine.

Penaeid shrimp are economically important aquaculture species. However, the production of shrimp was threatened by disease problems that may result in huge economic losses. Bacteria and fungi diseases may cause severe mortalities during the larval stages of shrimps and other crustaceans. For example, septicemia by the Gram-positive bacterium Aerococcus viridans can cause mortality in lobsters and crabs (Newman and Feng, 1982), and Gram-negative bacteria in the family Vibrionaceae, and fungi Fusarium spp. can cause mortality in shrimp (Song et al., 1993; Burns et al., 1979). To overcome such problems while also avoiding use of antibiotics, natural AMPs from shrimp have been considered. Penaeidins are AMPs that were first isolated from the plasma and hemocytes of Penaeus vannamei. The mature peptide contains a conserved leader sequence of 19 amino acids and a putative mature peptide of 55 amino acids (Chen et al., 2004). The deduced peptide of 74 amino acids. This peptide was worthwhile, especially with respect to pathogens of shrimp and other crustaceans.

2. Materials and methods

2.1. Animals

Juveniles from the black tiger shrimp, P. monodon, were obtained from different hatcheries of the Ilan, Taiwan. The average of body length and body weight of the adult shrimps were approximately 18.1±2.2 cm and 43.3±3.4 g, respectively. These shrimps were maintained in one ton tanks equipped with circulating seawater at 28 °C.

2.2. Semi-quantitative RT-PCR and Western blotting

Total RNA was isolated from the various tissues using TRIZOL reagent (Boehringer Mannheim), according to the manufacturer’s instructions. First-strand cDNA was synthesized in a 20 μl RT reaction from 250 ng of total RNA, using SuperScriptIII one-step RT-PCR (Invitrogen). The mRNA for the ubiquitous expression β-actin was used for normalization. The primers used in this study were as follows: penaeidin-5 forward primer 5′-ACC AGT CGG TGC TTG GCT CT, penaeidin-5 reverse primer 5′-GAG TAC TAC AAT TCC GAA TGT CC, β-actin forward primer 5′-CTT GTG GTT GAC AAT GGC TCC G and β-actin reverse primer 5′-TGG TGA AGG AGT AGC CAC GCT C. The first PCR cycles were: 94 °C, for 2 min flowed by 35 cycles at 94 °C for 30 s, 55 °C for 30 s, 72 °C for 30 s and a final cycle at 72 °C for 7 min. To perform the Western blot analysis, proteins from various tissues were extracted in homogenizing buffer (Tris–HCl 13.7 mM, Na2EDTA 20 mM, sucrose 25 mM, 9.6 mM, pH 7.4). The soluble proteins were separated by 10% Tricine SDS-PAGE and transferred to a nitrocellulose membrane using a Trans-Blot SD Semi-Dry Transfer Cell (Bio-Rad). The membrane was blocked by the NET buffer (0.25% gelatin, 50 mM NaCl, 0.05% Tween, 50 mM Tris–HCl, 5 mM EDTA, pH 7.5) at 4 °C overnight, and then incubated for 1 h with the penaeidin-5 polyclonal antibody (Genesis Biotech Inc.). After a 1 h incubation, the membrane was washed three times using PBS buffer containing 0.05% Tween 20 and then incubated with 1:7500 dilution of alkaline phosphatase conjugated anti-rat antibody. The membrane was washed and the location of the penaeidin-5 was detected using an ECL kit, according to the manufacturer’s protocol (Amersham). The protein for the ubiquitous translation β-actin was used for internal normalization.

2.3. Immunohistochemistry (IHC)

Gill and cuticular tissues from black tiger shrimp were washed in the PBS buffer (140 mM NaCl, 2.7 mM...
KCl, 10 mM Na₂HPO₄·12H₂O, 1.8 mM KH₂PO₄, pH 7.3) and then fixed in a solution containing 4% paraformaldehyde (PFA). After dehydration, the tissues were embedded in agarose and 8 μm sections were cut, mounted on poly-L-lysine coated slides, and stored at 4 °C until used. Immunohistochemistry of penaeidin-5 in gill and cuticular tissues was performed according to the previously developed protocol. Immunoreactions of the sections were carried out using polyclonal antibodies specific *P. monodon* penaeidin-5. This product was washed and incubated with 1:1000 dilution of horse-radish peroxidase (HRP)-conjugated anti-rat antibody (Bethyl). DAB substrate solution was used in the immunodetection according to manufacturer’s protocol (Amersco). Controls consisted in pre-absorbing the antibody overnight at 4 °C with synthetic penaeidin-5 (10 μg peptide/μg purified IgG).

### 2.4. Circular dichroism analysis and 3D modeling

Penaeidin-5 molecular ellipticity was measured under far-UV using a Jasco J-720 spectropolarimeter. The peptide was dissolved in distilled water to a final concentration of 0.2 μg/μl, and then analyzed at a resolution of 0.05 nm, from 260 nm to 280 nm, by ultra-violet spectroscopy using a cell of 0.1 cm path length. The three-dimensional structure of penaeidin-5 was predicted by using the MODELLER program and drawn with the program RasMol version 2.6.

### 2.5. In vitro susceptibility test

Antimicrobial activity assays were modified from the bacteriostatic and bactericidal assays previously described (Chio et al., 2005). The pathogens, growth medium and antimicrobial peptides used are shown in Table 1 using published peptide sequences, the antimicrobial peptides, cecropin A and B, magainin-II and penaeidin-5 were chemically synthesized by Genemed Synthesis Inc. (South San Francisco, CA). The purity of these antimicrobial peptides was >90% by HPLC analysis. The antibiotic tetracycline was used as positive control. They were dissolved in a PBS buffer for activity assays. Three different bacteria, *Aerococcus viridans*, *Fusarium pisi* and *Fusarium oxysporum* were used as models for the bacteriostatic and bactericidal/fungicidal assays. A single colony was picked up and inoculated into a 250 ml flask containing 50 ml of appropriate culture medium in a shaking incubator at 30 °C at 150 rpm. After 24 h cultivation, cell pellets were isolated by centrifugation at 3000×*g* for 15 min at 4 °C and washed twice in the PBS buffer. To obtain uniform (standard) cell concentrations, the washed cell pellets were re-suspended thoroughly in PBS buffer and the suspension was adjusted to an optical density (OD) of 1 at 540 nm. Subsequently, 10 ml of OD 1 suspension was diluted with 90 ml of appropriate culture medium (total 100 ml) to prepare the stock susceptibility test broth that was dispensed in 1.9 ml aliquots into 10 ml sterile assay tubes for activity assays.

### Table 1

<table>
<thead>
<tr>
<th>Peptide</th>
<th>Amino acid sequences</th>
<th>Organism source</th>
</tr>
</thead>
<tbody>
<tr>
<td>Penaeidin-5</td>
<td>QGYQGGYTRPFRPPYGGYHPVPVTCHRLSPLQARACCRLQGRCCDAKQTYG</td>
<td><em>Penaeus monodon</em></td>
</tr>
<tr>
<td>Cecropin A</td>
<td>KWKLFKKIENVGQRDGIKAAPAVVGVQATQIAK</td>
<td><em>Hyalophora cecropia</em></td>
</tr>
<tr>
<td>Cecropin B</td>
<td>KWKVFKKIEMGRNRINGIVAKAGPIAVLGEAKALG</td>
<td><em>Hyalophora cecropia</em></td>
</tr>
<tr>
<td>Magainin</td>
<td>GIGKFLHSAKFGKAVGEIMNS</td>
<td><em>Xenopus laevis</em></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Strain</th>
<th>Culture medium</th>
<th>Source</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Aerococcus viridans</em></td>
<td>Blood TBS (tryptic soy broth + 5% fetal bovine serum)</td>
<td>BCRC 14925*</td>
</tr>
<tr>
<td><em>Fusarium pisi</em></td>
<td>Potato dextrose broth</td>
<td>BCRC 35290*</td>
</tr>
<tr>
<td><em>Fusarium oxysporum</em></td>
<td>Potato dextrose broth</td>
<td>BCRC 35100*</td>
</tr>
</tbody>
</table>

* BCRC: Bioresources Collection and Research Center, Hsinshu, Taiwan.
To these assay tubes, 0.1 ml of serial dilutions of synthetic antimicrobial peptide solutions were added prior to cultivation on a shaking incubator at 30 °C at 150 rpm for 16 h. The minimal inhibitory concentration (MIC) was expressed as the lowest concentration that inhibited bacterial growth. To determine bactericidal activity of antimicrobial peptides, 100 μl aliquots of test broths from the MIC test were spread on either Blood TSB (for bacteria) or potato dextrose agar plate (for filamentous fungi) and incubated at 30 °C for 24 h and 48 h, respectively. The minimum bactericidal concentration (MBC) and minimum fungicidal concentration (MFC) were expressed as the minimum concentration at which there was no colony formation.

2.6. In vivo pathogen challenge tests

The marine bacterium *Vibrio alginolyticus* (SY4), a well-known shrimp pathogen was purchased from the...
American Type Culture Collection (ATCC, Manassas, Va.). A single colony of either *V. alginolyticus* or *A. viridans* was pick up and inoculated into a 250 ml flask containing 50 ml TSB broth (tryptic soy broth 30 g l$^{-1}$, NaCl 15 g l$^{-1}$) or Blood TSB broth, respectively, in a shaking incubator at 30 °C at 150 rpm. After overnight cultivation, the culture broth was centrifuged at 3000×g for 15 min at 4 °C to isolated the cell pellets. The pellets were washed twice and re-suspended thoroughly in sterile PBS buffer before adjustment to cell concentrations of 1.2×10$^5$ CFU/ml of *V. alginolyticus* and 10$^8$ CFU/ml of *A. viridans*. The juveniles black tiger shrimp was used for these challenge tests. Thirty black tiger shrimps were divided into two groups and each group was injected 100 $\mu$l of pathogen solution (104 CFU of *V. alginolyticus* per shrimp; 10$^7$ CFU of *A. viridans* per shrimp) into the muscle between third and fourth somite. The shrimp were then replaced in the cultural tanks for continued incubation. At various intervals thereafter, 5 $\mu$l samples of hemolymph were withdrawn from individual shrimp and mixed with an equal volume of anti-coagulant (27 mM sodium citrate, 336 mM NaCl, 115 mM glucose, 9 mM EDTA, pH 7.0) to prevent the hemocytes from coagulated. A 5 $\mu$l of mixture hemocytes solution from *V. alginolyticus* and *A. viridans* injected shrimps were inoculated into 3 ml of TSB and Blood TSB broth, respectively, and cultured at 30 °C at 150 rpm for 16 h. A 100 $\mu$l of cultured broth was spread on the TSB and Blood TSB agar plate and incubated at 30 °C for 24 h, respectively. The experiment was performed triplicate. The variation of bacterial colony-forming units was used to evaluate the response of endogenous penaeidin-5.

3. Results

3.1. Tissue distribution of penaeidin-5 mRNA

To investigate the tissue expression of penaeidin-5, the penaeidin-5 transcripts with a size of 400 bp was carried out with semi-quantitative RT-PCR in different tissues. The expression of penaeidin-5 was abundant in hemocytes and the intestine and intermediate in the gill and muscle. There was no signal in the eyes (Fig. 1A). The expression pattern of penaeidin-5 was also investigated by Western blotting using the penaeidin-5 polyclonal antibody. Mature penaeidin-5 could be recognized at 5.8 kDa, which agree with the expected molecular weight of the penaeidin-5 cDNA. Corresponding with the mRNA expression level, penaeidin-5 was found abundantly in hemocytes, the intestine and hemolymph (Fig. 1B). Based on the above results, penaeidin-5 existed mainly in the hemocytes. As shrimp gills are well known to be intensely vascularised, penaeidin-5 in gills was further confirmed by immunohistochemistry. The hemocytes only reacted with penaeidin-5 antibody and were located exclusively in the gill vessels (Fig. 2B); no positive reaction could be seen in the control (Fig. 2A). Furthermore, the subcuticular tissue was also investigated since it is involved in the first line of defense against pathogen. In comparison with the control, significant penaeidin-5

![Fig. 4. Predicted three-dimensional structure of penaeidin-5. (A) The structure of the full-length peptide was prepared using MODELLER and shows a CSαβ-type α-helix structure in the carboxy-terminal region. (B) Clustering of cationic and hydrophobic amino acids. Red, basic (positively charged) amino acids; blue, hydrophobic amino acids; yellow, α-helix. Other amino acids are shown in gray.](image-url)
labeling could be detected in sub-cuticular epithelium of the shrimp (Fig. 2C,D).

3.2. Circular dichroism and three-dimensional structure of the synthetic penaeidin-5

The proline-rich domain and cysteine-rich cyclic domain of penaeidins are unique features among antimicrobial peptides. In penaeidin-5, the linear proline-rich domain, which contains the triple Pro–Arg–Pro motif, consists of six proline residues at the N-terminal region while the cysteine-rich cyclic domain is located at the carboxy-terminal region. Circular dichroism analysis revealed a minimum signal of around 200 nm is indicative of an α-helix structure while a lower wavelength of 190 nm implied a strong contribution of poorly ordered structures that could possibly be random coils or loops (Fig. 3). A three-dimensional helical structure that contained two disulfide bridges was predicated by MODELLER. The helix structure was similar to the CSαβ-type antimicrobial peptides that contain a single α-helix and a pair of antiparallel β-sheets stabilized by multi-intramolecular disulfide bridges (Fig. 4). Because the CSαβ-type peptides possess antimicrobial activity against a wide spectrum of pathogens, the synthetic penaeidin-5 peptides was tested for bacteriostatic and bactericidal or/and fungicidal activity.

3.3. In vitro susceptibility test

The bacteriostatic and bactericidal or fungicidal activity of the antimicrobial peptides is shown in Table 2. Penaeidin-5 had strong antimicrobial activity against A. viridans. The minimum inhibitory and bactericidal concentrations were μM and 150 μM, respectively. In comparison to antibiotics and antimicrobial peptides in other species, penaeidin-5 had the lowest minimum bactericidal concentrations. In the fungi, the penaeidin-5 also showed a significant antimicrobial activity against F. pisi and F. oxysporum. Only 20 μM was required for fungicidal activity.

3.4. In vivo pathogen challenge test

In general, no large variation occurred in the bacterial number 6 h after V. alginolyticus or A. viridans were injected (Fig. 5). However, the bacterial number was decreased significantly form 1.22 × 10⁹ CFU/ml at 6 h to 4.37 × 10⁸ CFU/ml at 12 h and 3.21 × 10⁸ CFU/ml at 24 h in the V. alginolyticus injected samples (Fig. 5A). In the A. viridans injected samples, similar results were obtained. The bacterial number was decreased significantly from 1.23 × 10⁹ CFU/ml at 6 h to 4.9 × 10⁸ CFU/ml at 12 h and 3.21 × 10⁸ CFU/ml at 24 h in the V. alginolyticus injected samples (Fig. 5A).

Table 2

<table>
<thead>
<tr>
<th></th>
<th>Aerococcus viridans</th>
<th>Fusarium pisi</th>
<th>Fusarium oxysporum</th>
</tr>
</thead>
<tbody>
<tr>
<td>MIC (μM)</td>
<td>20</td>
<td>5</td>
<td>10</td>
</tr>
<tr>
<td>MBC (μM)</td>
<td>150</td>
<td>20</td>
<td>20</td>
</tr>
<tr>
<td>Penaeidin-5</td>
<td>20</td>
<td>5</td>
<td>10</td>
</tr>
<tr>
<td>Cecropin A</td>
<td>200</td>
<td>&gt;500</td>
<td>100</td>
</tr>
<tr>
<td>Cecropin B</td>
<td>30</td>
<td>400</td>
<td>100</td>
</tr>
<tr>
<td>Magainin II</td>
<td>1</td>
<td>5</td>
<td>1</td>
</tr>
<tr>
<td>Tetracycline</td>
<td>1</td>
<td>300</td>
<td>&gt;500</td>
</tr>
</tbody>
</table>

Values represent means of three experiments.
ml at 12 h and 1.1×10^8 CFU/ml at 24 h (Fig. 5B). We assumed that the decrease in bacteria in the pathogen injected shrimp was due, at least in part, to the induction of penaeidin-5. Therefore, we evaluated the expression of penaeidin-5 in the hemocytes of pathogen injected shrimp by Western blotting using ubiquitously expressed β-actin as an internal control. We found that expression level of penaeidin-5 was induced within 3 h after V. alginolyticus or A. viridans injection and that a higher protein level of penaeidin-5 was maintained up to 24 h (Fig. 6). Hence, we proposed that injection of the pathogens triggers production of penaeidin-5.

4. Discussion

Antimicrobial peptides have been established as key players in animal defense systems. The mature penaeidin-5 cDNA, which encode a 5.8 kDa antimicrobial peptide, was first cloned from the black tiger shrimp P. monodon. In a previous report, penaeidins from L. vannamei were demonstrated to be constitutively synthesized and stored in shrimp hemocytes where their mRNA expression level was highest (Destoumieux et al., 2000a,b). A similar result was seen in our experiments with high levels of expression in hemocytes and the intestine. The high level of penaeidin-5 expression gills probably resulted because they are intensively vascularised.

Some antimicrobial peptides from the crustacean have chitin-binding activity because of the cysteine-rich region in their COOH-terminal domain. This property is considered essential not only for antimicrobial activity, but also for chitin assembly in wound healing. Thus, it may also play an important role in preventing invasive infections (Destoumieux et al., 2000a,b). Penaeidin-5 displayed a similar motif in its COOH-terminal domain. Thus, we suppose that the penaeidin-5 also has chitin-binding properties that are involved in resistance to infections. Many reports indicate that specific structures of antimicrobial peptides have bactericidal activity against specific pathogens. A structural feature of the penaeidin family is that the mature peptides including an N-terminal cyclisation (Bachère et al., 2000). In our circular dichroism analysis, the lower wavelength of 190 nm implies a coil or loop structure. Thus, we assumed that it contains an N-terminal cyclisation as do other penaeidins. Comparing the results from circular dichroism with those from 3D structural modeling, we can expect that that an α-helix structure existed in the C-terminus of penaeidin-5. It contains six cysteine residues that are probably engaged in the formation of three intramolecular disulfide bridges that contribute to the formation of the α-helix structure. The helix structure was similar to that of CSαβ-type antimicrobial peptides such as drosomycin. It is noteworthy that the CSαβ motif antimicrobial peptides are effective against a wide variety of microbes such as Gram-negative bacteria, Gram-positive bacteria and yeasts. Therefore, some of the peptides from this family are being developed as drugs base on their specific antimicrobial features (Landon et al., 1997).

Comparison of penaeidin-5 with other penaeidins, the MIC concentration of penaeidin-2 and penaeidin-3 for Fusarium spp. was around 5 μM-10 μM; thus, the bacteriostatic activity of penaeidin-5 was similar to penaeidin-2 and penaeidin-3 (Destoumieux et al., 1999). In addition to other penaeidins, the antimicrobial peptides from other species also used to compare the antimicrobial activity. Cecropin A and B are a well-characterized family of antimicrobial peptides from the Hyalophora cecropia. They show a very broad spectrum of activities against Gram-positive and Gram-negative bacteria, also having antifungal activities such as Fusarium spp. and Aspergillus spp. (Boman, 1995; De Luca et al., 1998). Magainin-II is a natural antibiotic from frog skins with significant activity against the bacteria, fungi, protozoa, parasites and even viruses (Guerrero et al., 2004; Chinchar et al., 2004). We found that penaeidin-5, as well as cecropin A, cecropin B and magainin-II, were efficient bacteriostatic and bactericidal or/and fungicidal agents. The antimicrobial activity of penaeidin-5 was more efficient when compared with cecropin A and cecropin B. Since tetracycline is an antibacterial antibiotic, a high amount is necessary to act against fungi such as Fusarium spp. By contrast antimicrobial peptides like penaeidin-5 are much more effective against fungi. Based on the characteristic of penaeidin-5, we recently developed a recombinant penaeidin-5 as a fodder additive to enhance the immunity of marine animals. We are also developing transgenic shrimp that over-express penaeidin-5 in the hope that can be used as broodstock for the shrimp aquaculture industry.

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