In vitro activity of cationic peptides against *Neisseria gonorrhoeae* and vaginal *Lactobacillus* species: The effect of divalent cations

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Received 21 March 2012; revised 30 April 2012; accepted 5 May 2012

**ABSTRACT**

One of the new strategies for the prevention of HIV acquisition is the use of microbicides such as topical microbicides including antimicrobial and antiviral peptides. Ideally, new drug candidates should kill pathogens without determent to the normal bacterial flora considered important in health; such as hydrogen peroxide producing *Lactobacillus* species. The antimicrobial peptides LL-37 and LSA-5 were studied to determine their spectrum of activity against bacterial pathogens and normal flora organisms. The effects of divalent cations at biologically relevant concentrations were determined. We show the synthetic lytic peptide LSA-5 and the naturally occurring peptides LL-37 inactivate *Neisseria gonorrhoeae* but are less active against many normal flora members such as *Lactobacillus* species. Biologically relevant concentrations of calcium and magnesium prevented killing of sensitive strains. LSA-5 is more potent than LL-37, both are inhibited from killing sensitive strains by calcium and magnesium. Strains of *Lactobacillus iners* were killed by both microbicides even in the presence of the divalent cations. Antimicrobial peptides, such as LSA-5, have good potential for use in prevention of sexually transmitted disease, if formulated to sequester calcium and magnesium present in biological fluids.

**Keywords:** Microbicides; HIV; *Lactobacillus*; Antimicrobial Peptides; Cations; *Neisseria gonorrhoeae*

1. INTRODUCTION

Antimicrobial peptides, both host-derived and engineered, have great potential for the treatment and prevention of human diseases [1-3]. The rapid development of antibiotic resistance among many of the important pathogens extends the potential application of peptides to include topical and systemic uses. Internal uses such as intravenous or as a topical vaginal microbicides introduce the possibility of interference or inactivation by endogenous factors present in plasma or vaginal fluid. Microbicides are being developed for intra-vaginal use in the prevention of the acquisition of sexually transmitted infections (STIs) such as HIV, *Neisseria gonorrhoeae* and *Trichomonas vaginalis*. A microbicide must prevent infection by either rapidly killing or reducing the number of pathogens to below the infectious dose in a reasonably short time. Microbicides are evaluated by their killing efficacy, which describes the drug concentration required to reduce viability by 99.99% (a four log reduction) in a given time. This differs from conventional antibiotics where inhibition of growth is used to determine activity [4].

Lentivirus Lytic Peptides (LLPs) are highly conserved among the HIV isolates and are structurally similar to the magainins and human cathelicidin [5-8]. These peptides are based on the 28-residues of the C-terminus of the HIV-1 transmembrane protein gp41 [8]. Cellular toxicity results from the perturbation of the cell membranes [9-14]. By making specific amino acid substitutions different antimicrobial peptide derivatives have been devised with increased potency and specificity [6,9,10].

The normal healthy vagina has a predominance of hydrogen peroxide (H\(_2\)O\(_2\)) producing *Lactobacillus* species that are important in the maintenance of health and prevention of the STI acquisition [15,16]. Disease states such as bacterial vaginosis are characterized by a decrease or loss of H\(_2\)O\(_2\) producing lactobacilli [17]. Candidate microbicides are screened for their ability to specifically kill pathogens but not beneficial members of the normal flora such as *L. crispatus* and *L. jensenii*. 
The vaginal ecosystem is complex and dynamic. Vaginal fluid contains serum transudate, mucins and other proteins. Vaginal fluid has higher concentrations of calcium (8.5 mM) and magnesium (2.5 mM) than are found in plasma or interstitial tissues [18,19]. The pH of the vagina is usually around 4.0 - 4.5 but does increase to around pH 6.0 - 7.0 during menses or following intercourse. Therefore, testing at both low and high pH is advisable. We have examined several antimicrobial peptides to determine their efficacy in killing sexually transmitted pathogens and normal flora using conditions that mimic those found in the vagina.

We report here that the engineered peptide LSA-5 and the naturally occurring peptide LL-37 demonstrated the appropriate selectivity to kill pathogens such as Neisseria gonorrhoeae and non-beneficial bacteria such as Lactobacillus iners while not killing the beneficial Lactobacillus species. The activity of both peptides against N. gonorrhoeae and sensitive Lactobacillus isolates was inhibited by calcium and magnesium ions. These ions also inhibited other antimicrobial peptides.

2. MATERIAL AND METHODS

2.1. Antimicrobial Peptides


2.2. Microorganisms and Culture Conditions

Reference strains were obtained from the American Type Culture Collection, ATCC (Manassas, VA). Clinical isolates were obtained from swab specimens from women enrolled in clinical studies from Allegheny County Health Department and Magee-Womens Hospital. N. gonorrhoeae was identified using Gonocheck II® (PML Microbiologicals) and confirmed using an amplified nucleic acid test with the BDPProbe Tec ET instrument (strand displacement amplification assay [SDA]; Becton Dickens-on, Sparks, MD) [22]. Lactobacillus species were identified to the species level based on DNA-DNA homology to the type strains as previously described [17]. Organisms were stored frozen at −70°C in litmus milk until needed. Stock cultures were revived by inoculation onto either 5% sheep blood agar plates (Columbia blood agar base for Lactobacillus, PML Microbiologicals, Wilsonville, OR.) or chocolate agar (PML Microbiologicals, or prepared in house) for N. gonorrhoeae. Cultures were incubated at 37°C in air enriched to 6% CO2 overnight and evaluated for growth.

ACES buffer (N-(2-acetamido)-2-aminoethanesulfonic acid, Sigma Chemical Co. St. Louis, MO) was used at a pH of 7.0 and lactate buffer was used at pH 5.0. The isotonic strength of each lot of buffer was determined in a Wescor Vapro Pressure Osmometer 5520 (Logan, UT) and adjusted to 200 - 300 mosm/kg prior to use.

2.3. Antimicrobial Assays

In preliminary experiments, the activity and final peptide concentration of the peptides were compared when solid agents were first dissolved in distilled water or distilled water containing 0.01% glacial acetic acid. For both conditions comparable concentrations and experimental results were observed; therefore, in all subsequent work peptides were dissolved in distilled water to a concentration of 1.1 mM. Peptide concentrations of the working stock solutions were determined using a ninhydrin reaction [23]. Minimum cidal concentrations (MCCs) were determined in lactate (pH 5.0) and ACES (pH 7.0) buffers as previously described [4,24]. Initial concentrations used ranged from 40 µM down to as low as 0.01 µM. In subsequent work, more focused concentrations were used. Assays were performed in Costar (Corning, NY) polypropylene 96 well plates Briefly, isolated colonies were selected from fresh overnight culture plates and suspended in saline to a density of a 0.5 McFarland standard and diluting 1:10 in sterile saline. A 10 µl volume of the bacterial suspension was added to 90 µl of test solution. After incubation at 35°C for 30 min, 25 µl samples were taken and plated on to the appropriate medium, allowed to absorb and dry for 10 - 15 min then spread over the surface of the agar plate. This allows for an initial separation of organisms from the antimicrobial agent and facilitates the observation of inhibitory activity since organisms that are inhibited, but not killed, are observed as growth away from the point of sample application. Plates were incubated as described above for 24 h and evaluated for killing of the test microorganisms by counting colony forming units. Samples yielding 10 or fewer colony forming units represent a four log kill and were considered sensitive to killing.

The effects of divalent cations were determined by the addition of CaCl2 or MgCl2 prepared as 1 M stock solutions and filter sterilized. The salt solutions were added to the assay buffers prior to the addition of peptides or bacteria.
3. RESULTS

The minimum cidal concentrations (MCC) of LSA-5 and LL-37 for *N. gonorrhoeae*, *L. crispatus*, *L. iners* and *P. aeruginosa* are presented in Tables 1 and 2. For the strains where killing was observed, the general trend was that LSA-5 was more active as evidenced by the lower MCC values compared with LL-37.

The differences in the MCCs were five-fold higher for LL-37 at pH 7.0 than at pH 5.0. Differences in MCCs for the two peptides for *L. iners* and *P. aeruginosa* ranged from 2 to greater than 20 fold, respectively. The other microorganisms were consistently more sensitive to LSA-5 than LL-37. *N. gonorrhoeae* strains DOD 431 and UPS 2015 required higher concentrations than the other isolates tested >40 µM and 20 µM respectively.

Calcium and magnesium effects on antimicrobial activity. In early experiments we attempted to determine bacterial killing in rich media such as minimal essential medium (MEM) and trypticase soy broth, but killing was never observed. Because the concentration of divalent cations in MEM is about 26 mM, an equal concentration of EDTA was incorporated into MEM. EDTA binds calcium and magnesium ions effectively removing them from solution and therefore unavailable. This restored antimicrobial activity for sensitive bacteria; therefore, we systematically investigated the role of calcium and magnesium ions on killing by the panel of antimicrobial peptides in buffers appropriate for use with these ions. The results are presented in Tables 1 and 2. *N. gonorrhoeae* were protected by both ions, but calcium and magnesium afforded protection to more strains at pH 5.0 than 7.0, and calcium was more effective than magnesium.

*L. crispatus* was not killed by the LSA-5 at any concentration tested (up to 40 µM) and the divalent cations had no influence. *L. jensenii* demonstrated protection by calcium and magnesium ions among the strains with MCCs less than 10 µM (Table 1). *L. iners* was uniformly sensitive to killing by LSA-5 and was not protected from killing by the addition of the ions (Table 2). Experiments using LL-37 demonstrated the same trends as observed with the LSA-5, except 5 of 11 *L. iners* strains demonstrated MCCs to LL-37 greater than 20 µM, Table 2. Unlike LSA-5 there were no differences in MCCs for the individual *L. iners* strains with the LL-37 at the two different pHs (Table 2). Experiments with the structurally similar synthetic lytic peptides LBU-2, WLSA-5 and WLBU-2, that are under development for other therapies demonstrated protection by calcium and magnesium (data not shown) [6,9,12].

4. DISCUSSION

Condom use by males significantly reduces the risk of STD acquisition [25] but often women are coerced into unsafe sex practices by the unwillingness of their male partners to use condoms. The use of topical microbicides would allow protection in the absence of condom use. Antimicrobial peptides occur naturally, and are believed to play an important role in the host defenses [26,27].

<table>
<thead>
<tr>
<th>pH</th>
<th>Cations used and concentration</th>
<th><em>N. gonorrhoeae</em> n = 14</th>
<th><em>L. crispatus</em> n = 8</th>
<th><em>L. iners</em> n = 11</th>
<th><em>L. jensenii</em> n = 15</th>
<th><em>P. aeruginosa</em> n = 3</th>
</tr>
</thead>
<tbody>
<tr>
<td>pH 7.0</td>
<td>0</td>
<td>12 (86)</td>
<td>0 (0)</td>
<td>11 (100)</td>
<td>15 (100)</td>
<td>3 (100)</td>
</tr>
<tr>
<td>pH 5.0</td>
<td>0</td>
<td>12 (86)</td>
<td>0 (0)</td>
<td>11 (100)</td>
<td>ND*</td>
<td>0 (0)</td>
</tr>
<tr>
<td>pH 7.0</td>
<td>10 mM calcium</td>
<td>9 (62)</td>
<td>0 (0)</td>
<td>11 (100)</td>
<td>ND</td>
<td>0 (0)</td>
</tr>
<tr>
<td>pH 5.0</td>
<td>10 mM calcium</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>11 (100)</td>
<td>ND</td>
<td>0 (0)</td>
</tr>
<tr>
<td>pH 7.0</td>
<td>20 mM calcium</td>
<td>5 (33)</td>
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<td>11 (100)</td>
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<tr>
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<td>0 (0)</td>
<td>11 (100)</td>
<td>0 (0)</td>
<td>0 (0)</td>
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<tr>
<td>pH 7.0</td>
<td>10 mM magnesium</td>
<td>9 (62)</td>
<td>0 (0)</td>
<td>11 (100)</td>
<td>ND</td>
<td>0 (0)</td>
</tr>
<tr>
<td>pH 5.0</td>
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<tr>
<td>pH 7.0</td>
<td>20 mM magnesium</td>
<td>5 (33)</td>
<td>0 (0)</td>
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<td>1 (7)</td>
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<td>0 (0)</td>
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<td>1 (7)</td>
<td>0 (0)</td>
</tr>
<tr>
<td>MCC pH 7.0</td>
<td>0</td>
<td>0.5 - 1.0 µM*</td>
<td>&gt;40 µM</td>
<td>1.0 - 2.5 µM</td>
<td>&lt;1.0 - 2.5 µM</td>
<td>&lt;0.1 - 2.5 µM</td>
</tr>
<tr>
<td>MCC pH 5.0</td>
<td>0</td>
<td>0.05 - 0.5 µM</td>
<td>&gt;40 µM</td>
<td>0.5 - 1 µM</td>
<td>&lt;1 - 20 µM</td>
<td>0.5 - 2.5 µM</td>
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</tbody>
</table>

*N. gonorrhoeae* isolates DOD-431 and UPS 2015 had MCCs of 20 µM and >40 µM respectively. *Not determined.*
Table 2. Minimal cidal concentrations and the effects of calcium and magnesium ions and pH on killing of test organism by the antimicrobial peptide LL-37.

<table>
<thead>
<tr>
<th>pH</th>
<th>Cations present</th>
<th>N. gonorrhoeae</th>
<th>L. crispatus</th>
<th>L. iners</th>
<th>L. jensenii</th>
<th>P. aeruginosa</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>n = 14</td>
<td>n = 8</td>
<td>n = 11</td>
<td>n = 15</td>
<td>n = 3</td>
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<tr>
<td>pH 7.0</td>
<td>0</td>
<td>12 (86)</td>
<td>0 (0)</td>
<td>6 (55)</td>
<td>0 (0)</td>
<td>3 (100)</td>
</tr>
<tr>
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<td>0</td>
<td>12 (86)</td>
<td>0 (0)</td>
<td>6 (55)</td>
<td>0 (0)</td>
<td>3 (100)</td>
</tr>
<tr>
<td>pH 7.0</td>
<td>20 mM calcium</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>6 (55)</td>
<td>0 (0)</td>
<td>0 (0)</td>
</tr>
<tr>
<td>pH 5.0</td>
<td>20 mM calcium</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>6 (55)</td>
<td>0 (0)</td>
<td>0 (0)</td>
</tr>
<tr>
<td>pH 7.0</td>
<td>20 mM magnesium</td>
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<td>0 (0)</td>
<td>6 (55)</td>
<td>0 (0)</td>
<td>0 (0)</td>
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<tr>
<td>pH 5.0</td>
<td>20 mM magnesium</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>6 (55)</td>
<td>0 (0)</td>
<td>0 (0)</td>
</tr>
<tr>
<td>MCC pH 7.0</td>
<td>0</td>
<td>2.5 - 5.0 µM*</td>
<td>&gt;40 µM</td>
<td>5 - &gt; 20 µM</td>
<td>&gt;40 µM</td>
<td>4 µM</td>
</tr>
<tr>
<td>MCC pH 5.0</td>
<td>0</td>
<td>0.05 - 1.0 µM</td>
<td>&gt;40 µM</td>
<td>5 - &gt; 20 µM</td>
<td>&gt;40 µM</td>
<td>4 - 10 µM</td>
</tr>
</tbody>
</table>

* N. gonorrhoeae isolates DOD 431 and UPS 2015 had MCCs > 40 µM.

We have surveyed a number of antimicrobial peptides whose structures are based on the lytic portion of the HIV gp41 transmembrane glycoprotein, and found that LSA-5 appears to meet our requirements of killing pathogens such as N. gonorrhoeae at low concentrations, and not active against the protective/healthy microbiota of the vagina such as the hydrogen peroxide producing lactobacilli L. crispatus and L. jensenii.

The coating and duration of the epithelial and vaginal surface by products will vary with product formulation [28]. The standard for microbicides is a four log reduction in the viability within 30 min. Another useful measure a bactericidal activity is the decimal reduction value (D-value) which describes the time required by a given agent to reduce the viability of the test cultures by 90% or one log. The MCC simply describes the concentration at which organisms have a decimal reduction value (D value) of 6 minutes or less. Decimal reduction times for the test organisms in the buffers used (at or above the MCC) ranged from 1 - 1.5 min for N. gonorrhoeae and from 2 - 3 min for the lactobacillus (data not shown). Therefore the viability of the test organisms should have been reduced to less than 1/ml⁻¹ in 5 minutes. In theory, after 30 minutes a 20 log reduction in viability could be expected but the cultures were still viable in the presence of calcium or magnesium and peptide. This and the fact that all experiments used controls that omitted peptide for comparison, suggests the protection from the divalent cations is not an artifact of the system.

Comparing the activity of LL-37 (cathelicidin) with that of LSA-5 for the lactobacilli, the LSA-5 was much more active against the L. iners than was LL-37, ranging from 5 to greater than 20 times, at both pH 5.0 and 7.0. Most of the L. jensenii were more resistant to killing by LSA-5 at pH 5.0 than pH 7.0 which may reflect the natural habitat of this species; the vagina where the pH is about 4.0. Interestingly, the hydrogen peroxide producing species (L. jensenii and L. crispatus) were more resistant to the antimicrobial peptides than L. iners, which does not produce H₂O₂. The presence of L. iners has been associated with increased risk of HIV acquisition [29], suggesting decreasing its' numbers would be beneficial.

Caution should be used in the interpretation of the data obtained with N. gonorrhoeae at pH 5.0, since the organisms are sensitive to acidic pH. However, we have compared viability of strains of N. gonorrhoeae in the buffers used and GC broth buffer to pH 5.0 with lactate buffer, pH 7.0 with ACES buffer or phosphate buffer. At pH 7.0 there was no difference in viability, however, at pH 5.0 cells suspended in lactate buffer had a loss in viability which was about 20% after 30 minutes of incubation. The presence of added calcium to the lactate buffer did not increase survival. Therefore, the differences observed in the MCCs at pH 5.0 and pH 7.0 may reflect increased sensitivity to killing brought on by the acidic conditions. The same does not apply to Lactobacillus species tested whose natural habitat is the vagina where the pH is usually low (pH 4 - 5), suggesting the observations at acidic pH are not an artifact of the system.

Most of the N. gonorrhoeae isolates tested, 12 of 14, had low MCCs for LSA-5 (<1.0 µM) and LL-37 (5 µM). Isolates UPS 2015 and DOD 431 have proven to be resistant to killing by all the lytic peptides we have tested to date as well as nonoxynol-9 (29). We have observed no other phenotypic difference with these isolates. Strain DOD 431 demonstrated intermediate sensitivity to LSA-5, with an MCC some forty fold greater than most other strains tested (at pH 5.0). There are reports in the literature to indicate that Neisseria resistance may develop...

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through modification of the structure of the cell wall lipooligosaccharide and from acquisition of an efflux pump [30-32]. From these data, it is not possible to discriminate the underlying cause for the resistance. Perhaps of more significance is the observation of protection from killing by calcium and magnesium ions. Turner et al. reported difficulty in determining MICs for LL-37 when microbiological culture media was used [26]. When the medium was passed over an ion-exchange column they were able to observe activity. They also noted that the addition of 1 mM calcium resulted in an increase in the MIC for *Escherichia coli* but not *Listeria monocytogenes*. Their observations suggest a difference in the other most portions of the cells since *E. coli* is gram negative while *L. monocytogenes* is gram positive. The authors suggested that there are peptide fragments which may bind to or possibly precipitate the LL-37 to explain the reduced activity in complex microbiological media.

Although we cannot dismiss this possibility, our data suggest that divalent cations are primarily responsible for the reduced cidal activity of these lytic peptides since the addition of EDTA resulted in higher antimicrobial activity. Further, their observations with the addition of 1 mM calcium to the assays, is consistent with ours and supports our hypothesis that the ions have their effects on the bacteria, not the peptide.

The concentration of calcium and magnesium ions in vaginal fluid is about 8 mM and 2.5 mM respectively [18,19]. Many of the *N. gonorrhoeae* isolates were protected by as little as 1 mM calcium. Turner et al. using an MIC assay reported the protection from LL-37 of *Escherichia coli* but not *Listeria monocytogenes* by 1.25 mM calcium [26]. The mechanism for ion induced resistance probably relates to the lipophilic and cationic nature of the peptides. It is believed that the cationic portions of the peptide interact with the negative charge on the surface of the bacteria facilitating the lipophilic region of the peptide’s interaction with the membrane leading to disruption, and ultimately cell death. It is unlikely calcium and magnesium act to alter the structure of these peptides since all these antimicrobial peptides studied demonstrated the same effect. It is well established that the lytic effects of such peptides are directly related to the propensity of a peptide to form an α-helical structure when they interact with a membrane [6]. This property is influenced by the concentration of the peptide and the anions present [5,11]. Finally, if the peptides conformation was changed by the addition of cations the *L. iners* strains examined should have been protected as well, but were not. Calcium and magnesium cations are known to interact with the bacterial membranes, cytoplasmic and or outer membranes and prevent the binding of the peptide to target or stabilize the membrane [5,10,12-14]. It is significant that the inclusion of EDTA to complex media containing calcium and magnesium was adequate to bind sufficient divalent cations to allow the microbicidal properties of the peptides to function since many over the counter vaginal products contain EDTA, it would seem reasonable to assume that the topical administration of peptides in vivo would be effective. The inclusion of EDTA in the final formulated product may offer many additional benefits such as preservation, antioxidation and it may destabilize the membranes of target bacteria especially gram negative organisms.

The existence of such a resistant strain of *N. gonorrhoeae* as UPS 2015 would seem more problematical as it demonstrates the existence of microbicide resistance prior to any in vivo use of such synthetic lytic peptides as LSA-5.

It is difficult to describe or predict the availability of free calcium and magnesium ions in vivo. Large proportions are bound to protein or are complexed in other forms. The presence of similar naturally occurring lytic peptides such as LL-37, suggest they have some role and are active in vivo, even if these ions are present. However, when heat inactivated human or horse serum was added to assays in concentration of 30% or greater; inhibition of cidal activity was observed. These results confirm those of Johansson et al. with LL-37 who reported increases in MICs when serum was added. They also noted that there was complete inactivation of the LL-37 when it was first dissolved in serum rather than buffer [26]. The peptides were active when EDTA was added to the serum or when the serum was dialyzed against EDTA (data not shown). There was still a small inhibition of killing following dialysis of the sera. Additional studies are required to define other inhibitory components of serum. Preliminary data suggest that human serum albumin as well as α and β globulins bind the peptides and are therefore protective. If the observation is confirmed it would demonstrate that microbes have two separate means of using host factors to prevent killing by the microbicides. One is the presence of divalent cations present in secretions binds the bacteria resulting in their resistance to the drug. The other mechanism is the binding of the drugs to serum proteins making them ineffec-

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use in the prevention of acquisition of sexually transmitted infections.

5. ACKNOWLEDGEMENTS

This work was supported by grant NIAID 6PO1 AI39061 from the National Institutes of Health.

REFERENCES


